

Fourth Edition

THE CENTRAL NERVOUS SYSTEM

Structure and Function

Per Brodal



OXFORD

The Central Nervous System

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Structure and Function

Fourth Edition

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OXFORD

UNIVERSITY PRESS

2010

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Oxford University Press, Inc., publishes works that further
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Oxford New York
Auckland Cape Town Dar es Salaam Hong Kong Karachi
Kuala Lumpur Madrid Melbourne Mexico City Nairobi
New Delhi Shanghai Taipei Toronto

With offices in
Argentina Austria Brazil Chile Czech Republic France Greece
Guatemala Hungary Italy Japan Poland Portugal Singapore
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Published by Oxford University Press, Inc.
198 Madison Avenue, New York, New York 10016
www.oup.com

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Library of Congress Cataloging-in-Publication Data
Brodal, Per.

The central nervous system : structure and function / Per Brodal. — 4th ed.
p. ; cm.

Includes bibliographical references and index.

ISBN-13: 978-0-19-538115-3 (alk. paper)

ISBN-10: 0-19-538115-7 (alk. paper)

1. Central nervous system. I. Title.

[DNLM: 1. Central Nervous System—physiology. WL 300 B8645c 2010]

QP370.B76 2010

612.8'2—dc22

2009028992

1 3 5 7 9 8 6 4 2

Printed in the United States of America
on acid-free paper

Preface

This book is intended primarily for use by students of medicine, physical therapy, and psychology—that is, for use in neuroscience or neuroanatomy courses by students who need knowledge of the nervous system as a basis for later clinical study and practice. This fourth edition has been thoroughly revised and renewed. In addition to the updated and rewritten text, all figures have been redrawn and printed in full color to improve their impact, and many new ones have been added. The number of chapters has been increased to facilitate reading and grasp of the material. Further, each chapter begins with a short overview, setting the stage for what to come and emphasizing salient points.

My intentions remain the same as those of my father, Alf Brodal, when he wrote the Norwegian forerunner of this book more than 60 years ago: to stimulate understanding rather than memorization of isolated facts, while at the same time fostering a realistic attitude toward our still-limited ability to explain the marvels of the human brain.

The book aims to present the difficult subject of neuroscience so that those approaching it for the first time can understand it. Therefore, many details are left out that might be of great interest to the specialist but would merely obscure the essentials for the beginner. Everyday experiences and clinical examples are integrated throughout the text to help students link the new material with their prior knowledge and future profession. The nervous system, however, is exceedingly complex, both structurally and functionally, and much remains to be learned before we can answer many fundamental questions. Thus, while an undergraduate course can provide only partial insights, no one is served by a presentation that avoids controversial issues and areas of ignorance. Indeed, pointing out what we do not know is sometimes better than presenting an oversimplified version. For this reason I have also discussed how the data were obtained and the limitations inherent in the various methods.

The main challenge—for both the student and the scientist—is to understand how the nervous system solves its multifarious tasks. This requires an integrated approach, drawing on data from all fields of neurobiology, as well as from psychology and clinical research.

Textbooks sharing this goal nevertheless differ markedly in how they present the material and where they put the emphasis. Perhaps because my own field of research is the wiring patterns of the brain, I strongly feel that knowledge of how the nervous system is built—in particular, how the various parts are interconnected to form functional systems—is a prerequisite for proper understanding of data from other fields. A fair knowledge of brain anatomy is especially important for sound interpretations of the symptoms of brain disease. Textbooks of neuroanatomy often overwhelm the reader with details that are not strictly relevant for either functional analysis or clinical thinking. Neither does a strong emphasis on cellular mechanisms at the expense of the properties of neural systems seem the right choice if the aim is to help readers understand how the brain performs its tasks and how the site of a disease process relates to a patient's symptoms. Therefore, neither anatomical nor cellular and molecular details are included in this book if they cannot in some way be related to function. My hope is that the book presents a balance of cellular and neural systems material that is right for students.

In-depth sections and more advanced clinical material are clearly marked so that they should not disturb reading of the main text. Because the needs of readers differ, however, they are encouraged to read selectively and pick the material they find most relevant and interesting from their perspective, regardless of whether it is placed in the main text or in boxes. The frequent subheadings should facilitate such selective reading.

During the preparation of the former and the present editions, I have received help from several colleagues, for which I am truly grateful. Jan Bjaalie, Niels Christian Danbolt, Paul Heggelund, Jan Jansen, Harald Kryvi, Kirsten Osen, Ole Petter Ottersen, Eric Rinvik, and Jon Storm-Mathisen have all provided constructive criticism and advice. I also gratefully acknowledge the expert help of Gunnar Lothe and Carina Knudsen, who produced the photographic work.

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Oslo, Norway

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Introduction

A BIRD'S EYE VIEW OF THE NERVOUS SYSTEM

What are the main tasks of the nervous system? This question is not easily answered—our brains represent most of what we associate with being a human. At a superior level, the brain creates our reality: it selects, sorts, and interprets the overwhelming amount of information we receive from our bodies and the environment, and it controls behavior in accordance with its interpretations of reality. This control concerns behavior in a wide sense: one aspect is control and maintenance of the body and its inner milieu; another is our interaction with our surroundings and other human beings through actions and speech. A third aspect is our inner, subjective, mental reality that others can only partially know. In early childhood, the brain must create order and predictability so that we learn to relate successfully to ourselves and our environment.

The essential building block of the nervous system is the **neuron** (nerve cell), specialized for rapid conveyance of signals over long distances and in a very precise manner. Together, billions of neurons in the brain form complicated and highly organized networks for **communication** and **information processing**.

The nervous system receives a wealth of information from an individual's surroundings and body. From all this information, it extracts the essentials, stores what may be needed later, and emits a command to muscles or glands if an answer is appropriate. Sometimes the answer comes within milliseconds, as a **reflex** or automatic response. At other times it may take considerably longer, requiring cooperation among many parts of the brain and involving **conscious processes**. In any case, the main task of the nervous system is to ensure that the organism adapts optimally to the environment.

The nervous system is equipped with sense organs, **receptors**, that react to various forms of sensory information or stimuli. Regardless of the mode of stimulation (the form of energy), the receptors “translate” the energy of the stimulus to the language spoken by the nervous system, that is, **nerve impulses**. These are tiny electric discharges rapidly conducted along the nerve processes. In this way signals are conveyed from the receptors to the regions of the nervous system where information processing takes place.

The nervous system can elicit an external response only by acting on **effectors**, which are either muscles or glands. The response is either **movement** or **secretion**. Obviously, muscle contraction can have various expressions, from communication through speech, facial expression, and bodily posture to walking and running, respiratory movements, and changes of blood pressure. But one should bear in mind that the nervous system can only act on muscles and glands to express its “will.” Conversely, if we are to judge the activity going on in the brain of another being, we have only the expressions produced by muscle contraction and secretion to go by.

On an anatomic basis we can divide the nervous system into the **central nervous system** (CNS), consisting of the brain and the spinal cord, and the **peripheral nervous system** (PNS), which connects the CNS with the receptors and the effectors. Although without sharp transitions, the PNS and the CNS can be subdivided into parts that are concerned primarily with the regulation of visceral organs and the internal milieu, and parts that are concerned mainly with the more or less conscious adaptation to the external world. The first division is called the **autonomic** or **visceral nervous system**; the second is usually called the **somatic nervous system**. The second division, also called the cerebrospinal nervous system, receives information from sense organs capturing events in our surroundings (vision, hearing, receptors in the skin) and controls the activity of voluntary muscles (made up of cross-striated skeletal muscle cells). In contrast, the autonomic nervous system controls the activity of involuntary muscles (smooth muscle and heart muscle cells) and gland cells. The autonomic system may be further subdivided into the **sympathetic system**, which is mainly concerned with mobilizing the resources of the body when demands are increased (as in emergencies), and the **parasympathetic system**, which is devoted more to the daily maintenance of the body.

The **behavior** of a vertebrate with a small and—comparatively speaking—simple brain (such as a frog) is dominated by fairly fixed relationships between stimuli and their response. Thus, a stimulus, produced for example by a small object in the visual field, elicits a stereotyped pattern of goal-directed movements. Few neurons are intercalated between the sense organ and

the effector, with correspondingly limited scope of response adaptation. Much of the behavior of the animal is therefore instinctive and automatic, and not subject to significant change by learning. In mammals with relatively small brains compared with their body weights (such as rodents) a large part of their brain is devoted to fairly direct sensorimotor transformations. In primates, the relative brain weight has increased dramatically during some million years of evolution. This increase is most marked in humans with relative brain weight double that of the chimpanzee. In humans, there are few fixed relationships between sensations and behavior (apart from a number of vital reflexes). Thus, a certain stimulus may cause different responses depending on its context and the antecedents. Consequently, we often can choose among several responses, and the response can be changed on the basis of experience. Such flexibility requires, however, increased “computational power” in terms of number of neurons available for specific tasks. The more an animal organizes its activities on the basis of previous experience, and the more it is freed from the dominance of immediate sensations, the more complex are the processes required of the central nervous system. The behavior of humans cannot be understood merely on the basis of what happened immediately before. The British neuropsychologist Larry Weiskrantz (1992) puts it this way: “We are controlled by predicted consequences of our behavior as much as by the immediate antecedents. We are goal-directed creatures.”

The higher processes of integration and association—that is, what we call **mental processes**—are first and foremost a function of the **cerebral cortex**. It is primarily the vast number of neurons in this part of the brain that explains the unique adaptability and learning capacity of human beings. Indeed, the human brain not only permits adaptation to extremely varied environments, it also enables us to change our environment to suite our needs. This entails enormous possibilities but also dangers, because we produce changes that are favorable in the short run but in the long run might threaten the existence of our species.

STUDYING THE STRUCTURE AND FUNCTION OF THE NERVOUS SYSTEM

Some of the many methods used for the study of the nervous system are described in the following chapters—that is, in conjunction with discussion of results produced by the methods. Here we limit ourselves to some general features of neurobiological research.

Many approaches have been used to study the structure and function of the nervous system, from straightforward observations of its macroscopic appearance to determination of the function of single molecules. In recent years we have witnessed a tremendous development of methods,

so that today problems can be approached that were formerly only a matter of speculation. The number of neuroscientists has also increased almost exponentially, and they are engaged in problems ranging from molecular genetics to behavior. Although the mass of knowledge in the field of neurobiology has increased accordingly, more importantly, the understanding of how our brains work has improved considerably. Nevertheless, the steadily expanding amount of information makes it difficult for the scientist to have a fair knowledge outside his or her specialty. It follows that the scientist may not be able to put findings into the proper context, with danger of drawing erroneous conclusions.

Traditionally, methods used for neurobiological research were grouped into those dealing with **structure** (neuroanatomy) and those aiming at disclosing the **function** of the structures (neurophysiology, neuropsychology). The borders are far from sharp, however, and it is typical of modern neuroscience that anatomic, physiological, biochemical, pharmacological, psychological, and other methods are combined. Especially, cell biological methods are being applied with great success. Furthermore, the introduction of modern computer-based imaging techniques has opened exciting possibilities for studying the relation between structure and function in the living human brain. More and more of the methods originally developed in cell biology and immunology are being applied to the nervous system, and we now realize that neurons are not so different from other cells as was once assumed.

Animal Experiments Are Crucial for Progress

Only a minor part of our present knowledge of the nervous system is based on observations in humans; most has been obtained in experimental animals. In humans we are usually limited to a comparison of symptoms that are caused by naturally occurring diseases, with the findings made at postmortem examination of the brain. Two cases are seldom identical, and the structural derangement of the brain is often too extensive to enable unequivocal conclusions.

In animals, in contrast, the experimental conditions can be controlled, and the experiments may be repeated, to reach reliable conclusions. The properties of the elements of neural tissue can be examined directly—for example, the activity of single neurons can be correlated with the behavior of the animal. Parts of the nervous system can also be studied in isolation—for example, by using tissue slices that can be kept viable in a dish (*in vitro*) for hours. This enables recordings and experimental manipulations to be done, with subsequent structural analysis of the tissue. Studies in invertebrates with a simple nervous system have made it possible to discover the fundamental mechanisms that underlie synaptic function and the functioning of simple neuronal networks.

When addressing questions about functions specific to the most highly developed nervous systems, however, experiments must be performed in higher mammals, such as cats and monkeys, with a well-developed cerebral cortex. Even from such experiments, inferences about the human nervous system must be drawn with great caution. Thus, even though the nervous systems in all higher mammals show striking similarities with regard to their basic principles of organization, there are important differences in the relative development of the various parts. Such anatomic differences indicate that there are functional differences as well. Thus, results based on the study of humans, as in clinical neurology, psychiatry, and psychology, must have the final word when it comes to functions of the human brain. But because clinicians seldom can experiment, they must often build their conclusions on observations made in experimental animals and then decide whether findings from patients or normal volunteers can be explained on such a basis. If this is not possible, the clinical findings may raise new problems that require studies in experimental animals to be solved. Basically, however, the methods used to study the human brain are the same as those used in the study of experimental animals.

Ethics and Animal Experiments

Experiments on animals are often criticized from an ethical point of view. But the question of whether such experiments are acceptable cannot be entirely separated from the broader question of whether mankind has the right to determine the lives of animals by using them for food, by taking over their territories, and so forth. With regard to using animals for scientific purposes, one has to realize that a better understanding of human beings as thinkers, feelers, and actors requires, among other things, further animal experiments. Even though cell cultures and computer models may replace some of them, in the foreseeable future we will still need animal experiments. Computer-based models of the neuronal interactions taking place in the cerebral cortex, for example, usually require further animal experiments to test their tenability.

Improved knowledge and understanding of the human brain is also mandatory if we want to improve the prospects for treatment of the many diseases that affect the nervous system. Until today, these diseases—most often leading to severe suffering and disability—have only occasionally been amenable to effective treatment. Modern neurobiological research nevertheless gives hope, and many promising results have appeared in the last few years. Again, this would not have been possible without animal experiments.

Yet there are obviously limits to what can be defended ethically, even when the purpose is to alleviate human suffering. Strict rules have been made by government

authorities and by the scientific community itself to ensure that only properly trained persons perform animal experiments and that the experiments are conducted so that discomfort and pain are kept at a minimum. Most international neuroscience journals require that the experiments they publish have been conducted in accordance with such rules.

Sources of Error in All Methods

Even though we will not treat systematically the sources of error inherent in the various methods discussed in this book, certainly all methods have their limitations. One source of error when doing animal experiments is to draw premature conclusions about conditions in humans. In general, all experiments aim at isolating structures and processes so that they can be observed more clearly. However necessary this may be, it also means that many phenomena are studied out of their natural context. Conclusions with regard to how the parts function in conjunction with all of the others must therefore be speculative.

Purely anatomic methods also have their sources of error and have led to many erroneous conclusions in the past about connections between neuronal groups. In turn, such errors may lead to misinterpretations of physiological and psychological data. The study of humans also entails sources of error—for example, of a psychological nature. Thus, the answers and information given by a patient or a volunteer are not always reliable; for example, the patient may want to please the doctor and answer accordingly.

Revising Scientific “Truths” from Time to Time

That our methods have sources of error and that our interpretations of data are not always tenable are witnessed by the fact that our concepts of the nervous system must be revised regularly. Reinterpretations of old data and changing concepts are often made necessary by the introduction of new methods. As in all areas of science, conclusions based on the available data should not be regarded as final truths but as more or less probable and preliminary interpretations. Natural science is basically concerned with posing questions to nature. How understandable and unequivocal the answers are depends on the precision of our questions and how relevant they are to the problem we are studying: stupid questions receive stupid answers. It is furthermore fundamental to science—although not always easy for the individual scientist to live up to—that conclusions and interpretations be made without any bias and solely on the strength of the facts and the arguments. It should be irrelevant whether the scientist is a young student or a Nobel laureate.

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The Central Nervous System

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I

MAIN FEATURES OF STRUCTURE AND FUNCTION

GENERAL information about the structure and function of the nervous system forms a necessary basis for treatment of the specific systems described in subsequent parts of this book. **Chapters 1 and 2** describe the structure of nervous tissue and some basic features of how neurons are interconnected, while **Chapters 3, 4, and 5** deal with the functional properties of neurons as a basis for understanding communication between

nerve cells. **Chapter 6** provides an overview of the macroscopic (and, to some extent, the microscopic) structure of the nervous system with brief descriptions of functions. **Chapter 7** treats the membranes covering the central nervous system, the cavities within the brain, and the cerebrospinal fluid produced in these. Finally, **Chapter 8** describes the blood supply of the brain and the spinal cord.

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Structure of the Neuron and Organization of Nervous Tissue

OVERVIEW

The nervous system is built up of nerve cells, **neurons**, and special kinds of supporting cells, **glial cells** (discussed in Chapter 2). The nerve cells are responsible for the functions that are unique to the nervous system, whereas the glial cells are non-neuronal cells that primarily support and protect the neurons. Neurons are composed of a cell body called the **soma** (plural somata) and several processes. Multiple short **dendrites** extend the receiving surface of the neuron, while a single **axon** conducts nerve impulses to other neurons or to muscle cells. Neurons are characterized by their ability to respond to stimuli with an electrical discharge, a **nerve impulse**, and, further, by their fast **conduction** of the nerve impulse over long distances. In this way, signals can be transmitted in milliseconds from one place to another, either within the central nervous system (CNS) or between it and organs in other systems of the body. When the nerve impulse reaches the **synapse**, which is the site of contact between the axon and the next neuron, a substance called a **neurotransmitter** is released from the **axon terminal** that conveys a chemical signal from one neuron to the next.

Neurons are classified into two broad groups: **projection neurons** that transmit signals over long distances and **interneurons** that mediate cooperation among neurons that lie grouped together. Many axons are surrounded with a **myelin sheath** to increase the speed of impulse propagation. Nervous tissue contains some areas that look gray—**gray matter**—and others that look whitish—**white matter**. White matter consists of axons and no neuronal somata, and the color is due to the whitish color of myelin. Gray matter consists mainly of somata and dendrites, which have a gray color. Neuronal somata are collected in groups sharing connections and functional characteristics. In the CNS, such a group is called a **nucleus** and in the peripheral nervous system (PNS), a **ganglion**. A bundle of axons that interconnect nuclei is called a **tract**. A nerve connects the CNS with peripheral organs. Groups of neurons that are interconnected form complex **neuronal networks** that are responsible for performing the tasks of the CNS. A fundamental principle of the CNS is that each neuron influences many others (**divergence**) and receives

synaptic contacts from many others (**convergence**). A neuron contains a cytoskeleton consisting of various kinds of neurofibrils. They are instrumental in forming the neuronal processes and in transport of substances along them. By **axonal transport**, building materials and signal substances can be brought from the cell soma to the nerve terminals (anterograde transport), and signal substances are carried from the nerve terminal to the soma (retrograde transport).

NEURONS AND THEIR PROCESSES

Neurons Have Long Processes

Like other cells, a neuron has a **cell body** with a nucleus surrounded by cytoplasm containing various organelles. The nerve cell body is also called the perikaryon or **soma** (Figs. 1.1, 1.2, and 1.3). Long processes extend from the cell body. The numbers and lengths of the processes can vary, but they are of two main kinds: **dendrites** and **axons** (Fig. 1.1). The dendrites usually branch and form dendritic “trees” with large surfaces that receive signals from other nerve cells. Each neuron may have multiple dendrites, but has only one axon, which is specially built to conduct the nerve impulse from the cell body to other cells. The axon may have many ramifications, enabling its parent cell to influence many other cells. Side branches sent off from the parent axon are termed **axon collaterals** (Fig. 1.1). The term **nerve fiber** is used synonymously with “axon.”

Neurons Are Rich in Organelles for Oxidative Metabolism and Protein Synthesis

When seen in a microscopic section, the nucleus of a neuron is characterized by its large size and light staining (i.e., the chromatin is extended, indicating that much of the genome is in use). There is also a prominent nucleolus (Figs. 1.2 and 1.3). These features make it easy to distinguish a neuron from other cells (such as glial cells), even in sections in which only the nuclei are clearly stained. The many **mitochondria** in the neuronal cytoplasm are an indication of the high metabolic activity of nerve cells. The mitochondria depend entirely on

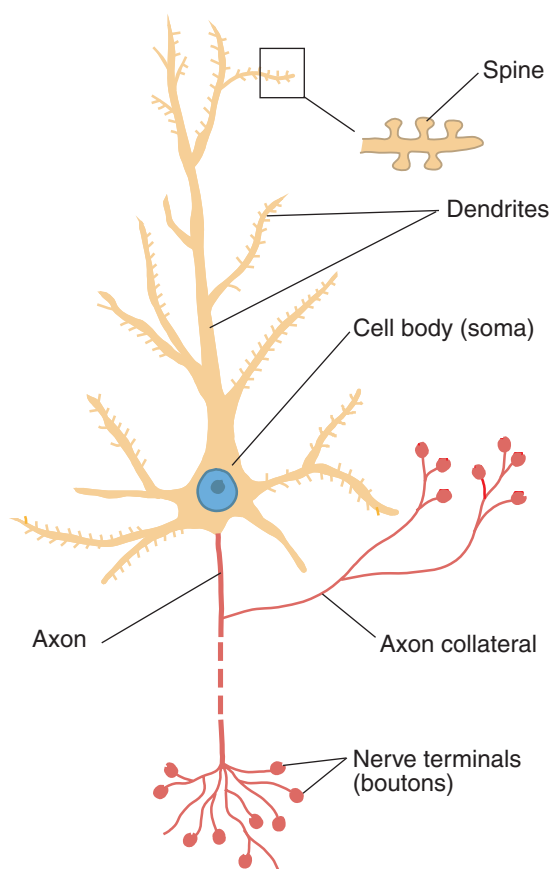


FIGURE 1.1 A neuron. Half-schematic to illustrate the neuron's main parts. The axon is red.

aerobic adenosine triphosphate (ATP) production and, unlike those in most other cell types, cannot utilize anaerobic ATP synthesis. **Glucose** is the substrate for ATP production in the mitochondria of nerve cells, which cannot, unlike in muscle cells, for example, use fat.

Neuronal somata also contain conspicuous amounts of free ribosomes and **rough endoplasmic reticulum** (rER) for synthesis of proteins. Large clumps of rER are seen via light microscopy in the cytoplasm of neurons greater than a certain size (Figs. 1.2 and 1.3). These were called tigroid granules or Nissl bodies long before their true nature was known. There are also as a rule several Golgi complexes, which modify proteins before they are exported or inserted in membranes. The large neuronal production of proteins probably reflects the enormous neuronal surface membrane, which contains many protein molecules that must be constantly renewed. Membrane proteins, for example, form ion channels and receptors (binding sites) for neurotransmitters, are constantly being recycled.

Dendrites Are Equipped with Spines

To study the elements of nervous tissue, it is necessary to use thin sections that can be examined microscopically.

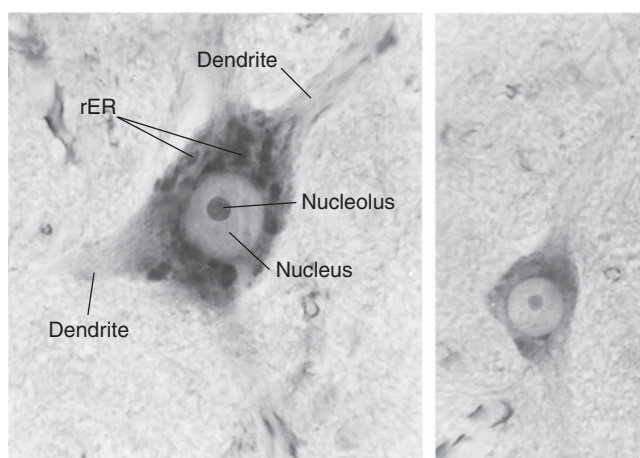


FIGURE 1.2 Neuronal somata (cell bodies). Two motor neurons, one small and one large, are shown. The large, pale nucleus has a distinct nucleolus. Only the cell body and the proximal parts of the dendrites are visible with the staining method used here. The stain (thionine) binds primarily to nucleic acids (DNA in the nucleus and RNA in the cytoplasm and nucleolus). The deeply stained clumps in the cytoplasm represent aggregates of rough endoplasmic reticulum (rER). Photomicrographs taken with a light microscope of a 20 μm thick section of the spinal cord. Magnification, $\times 800$.

Different staining methods make it possible to distinguish the whole neuron or parts of it from the surrounding elements (Figs. 1.2 and 1.4). It then becomes evident that the morphology of neurons may vary, with regard to both the size of the cell body and the number, length, and branching of the dendrites (Fig. 1.2; see also Figs. 33.5–6). The size of the dendritic tree is related to the number of contacts the cell can receive from other nerve cells. Dendrites often have small spikes, **spinae** (sing. spina) or **spines**, which are sites of contact with other neurons (Figs. 1.1, 1.7, and 1.8). The axons also vary, from those that ramify and end close to the cell body to those that extend for more than 1 meter (see Fig. 1.10; see also Figs. 21.3, 33.5, and 33.6). These structural differences are closely connected to functional differences.

Most Neurons Are Multipolar

Most neurons have several processes and are therefore called **multipolar** (Fig. 1.5). Special kinds of neurons, however, may have a different structure. Thus, neurons that conduct sensory signals from the receptors to the CNS have only one process that divides close to the cell body. One branch conducts impulses from the receptor toward the cell soma; the other conducts impulses toward and into the CNS. Such neurons are called **pseudounipolar** (Fig. 1.5). In accordance with the usual definition, the process conducting signals toward the cell body should be termed a dendrite. In terms of both structure and function, however, this process must be regarded as an axon. Some neurons have two processes, one conducting

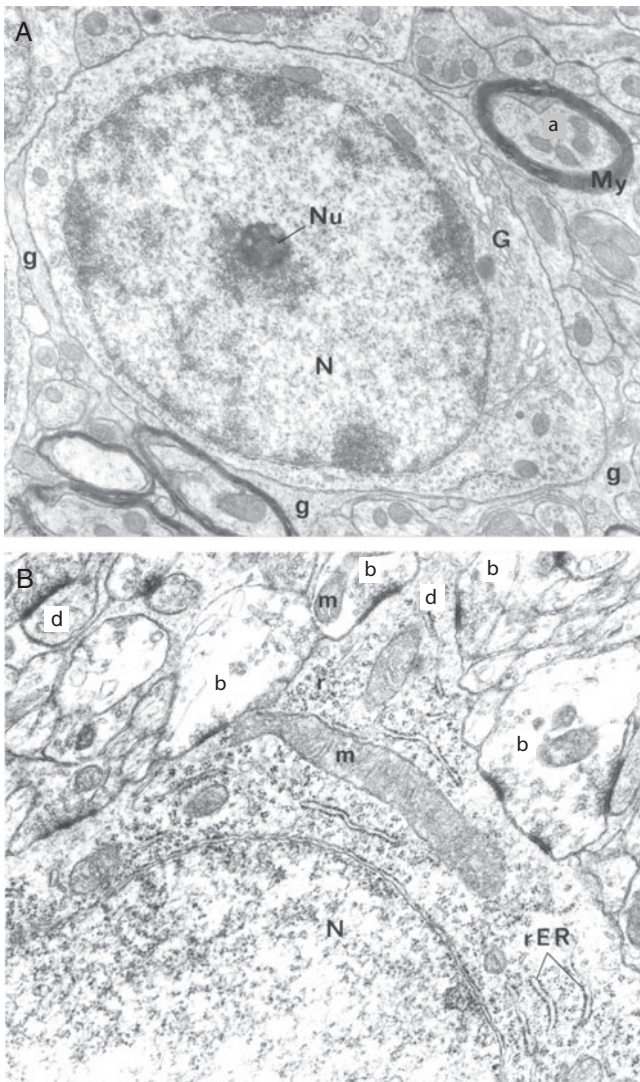


FIGURE 1.3 *Ultrastructure of the neuron.* Electron micrograph showing the cell body of a small neuron (A) and parts of a larger neuron (B). The nucleus (N) is light, due to extended chromatin, and contains a nucleolus (Nu). The cytoplasm contains rough endoplasmic reticulum (rER) and a Golgi complex (G)—that is, organelles involved in protein synthesis. The presence of many mitochondria (m) reflects the high oxidative metabolism of neurons. Nerve terminals, or boutons (b), forming axosomatic and axodendritic synapses are also seen. Glial processes (g) follow closely the surface of the cell body and the dendrites (d). a, axon; My, myelin. Magnifications, $\times 9000$ (top) and $\times 15,000$ (bottom).

toward the cell body, the other away from it (Fig. 1.5). Such neurons, present in the retina (see Fig. 16.7) and the inner ear (see Fig. 17.5B), are called **bipolar**. Also in these neurons both processes function as axons.

Communication between Nerve Cells Occurs at Synapses

The terminal branches of an axon have club-shaped enlargements called **boutons** (Figs. 1.1 and 1.6).

The term **terminal bouton** is used when the bouton sits at the end of an axon branch, and we also use the term **nerve terminal**. In other instances, the bouton is only a thickening along the course of the axon, with several such **en passage boutons** along one terminal branch (Fig. 1.8). In any case, the bouton lies close to the surface membrane of another cell, usually on the dendrites or the cell body. Such a site of close contact between a bouton and another cell is called a **synapse**. In the PNS, synapses are formed between boutons and muscle cells. The synapse is where information is transmitted from one neuron to another. This transmission does not occur by direct propagation of the nerve impulse from one cell (neuron) to another, but by liberation of signal molecules that subsequently influence the other cell. Such a signal molecule is called a **neurotransmitter** or, for short, a **transmitter** (the term “transmitter substance” is also used). The neurotransmitter is at least partly located in small vesicles in the bouton called **synaptic vesicles** (Figs. 1.6 and 1.7). How the synapse and the transmitters work is treated in Chapters 4 and 5. Here we restrict ourselves to the structure of the synapse.

The membrane of the nerve terminal is separated from the membrane of the other nerve cell by a narrow cleft approximately 20 nm wide (i.e., $2/100,000$ mm). This **synaptic cleft** cannot be observed under a light microscope. Only when electron microscopy of nervous tissue became feasible in the 1950s could it be demonstrated that neurons are indeed anatomically separate entities. In the electron microscope, one can observe that the membranes facing the synaptic cleft are thickened (Figs. 1.6 and 1.7), due to accumulation of specific proteins that are of crucial importance for transmission of the synaptic signal. Many of these protein molecules are receptors for neurotransmitters; others form channels for passage of charged particles. The membrane of the bouton facing the cleft is called the **presynaptic membrane**, and the membrane of the cell that is contacted is called the **postsynaptic membrane** (Fig. 1.6). We also use the terms pre- and postsynaptic neurons.

The **postsynaptic density** (Fig. 1.6; see also Fig. 4.17) connects to the cytoskeleton with actin filaments and other proteins. This connection probably anchors the postsynaptic receptors to the site of neurotransmitter release. In addition, certain proteins in the postsynaptic density, such as **cadherins**, bind to corresponding proteins in the presynaptic membrane to keep the nerve terminal in place (cadherins are present also in many other cell-to-cell contacts, e.g., in adherence contacts between epithelial cells). Other proteins in the postsynaptic density have modulatory actions on synaptic function, for example, by changing receptor properties. Synaptic modifications associated with learning involve structural and functional changes of the postsynaptic density.

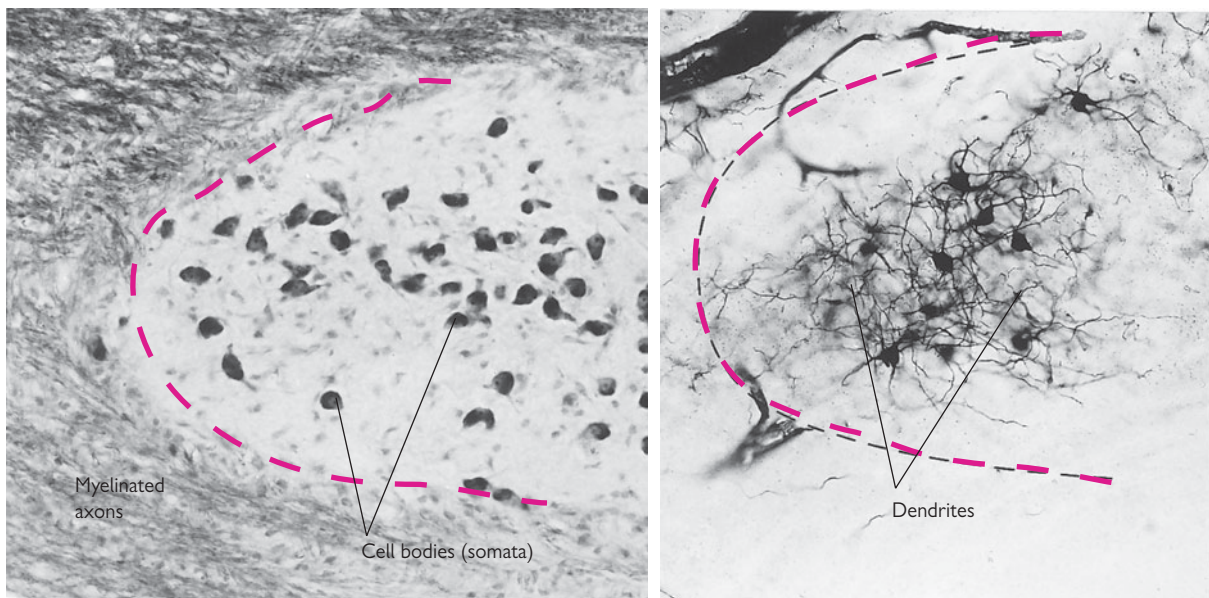


FIGURE 1.4 *Neurons.* Photomicrographs of sections stained with two different methods. **Left:** Only the cell bodies (somata) of a group of neurons are stained and visible in the section. The dark region surrounding the group of neurons contains myelinated fibers that are

also stained. **Right:** The same cell group, but treated via the Golgi method so that the dendrites and the cell bodies are visualized. Magnification, $\times 150$.

Synapses formed on the cell soma are called **axosomatic**, while synapses on dendrites are called **axodendritic** (Figs. 1.3 and 1.8). Where dendrites are equipped with spines, one or two **axospinous** synapses are always

formed with the spine head (Figs. 1.7B, 1.8, and 1.9). The functional role of spines is not fully understood (see Chapter 4). Boutons may also form a synapse with an axon (usually close to a terminal bouton of that axon), and such synapses are called **axoaxonic** (Fig. 1.8 and 1.9B). This enables selective control of one terminal only without influencing the other terminals of the parent axon. Axoaxonic synapses thus increase the precision of the signal transmission.

There are many more axodendritic than axosomatic synapses because the dendritic surface is so much larger. Every neuron has many thousands of synapses on its surface, and the sum of their influences determines how active the postsynaptic neuron will be at any moment.

Two Main Kinds of Nerve Cell: Projection Neurons and Interneurons

Some neurons influence cells that are at a great distance, and their axons are correspondingly long (more than a meter for the longest). They are called **projection neurons**, or Golgi type 1 (Fig. 1.10). Neurons that convey signals from the spinal cord to the muscles are examples of projection neurons; other examples are neurons in the cerebral cortex with axons that contact cells in the brain stem and the spinal cord (see Fig. 33.5). As a rule, the axons of projection neurons send out branches, or **collaterals**, in their course (Figs. 1.1 and 1.11; see also Fig. 33.5). Thus, one projection neuron may send signals to neurons in various other parts of the nervous system.

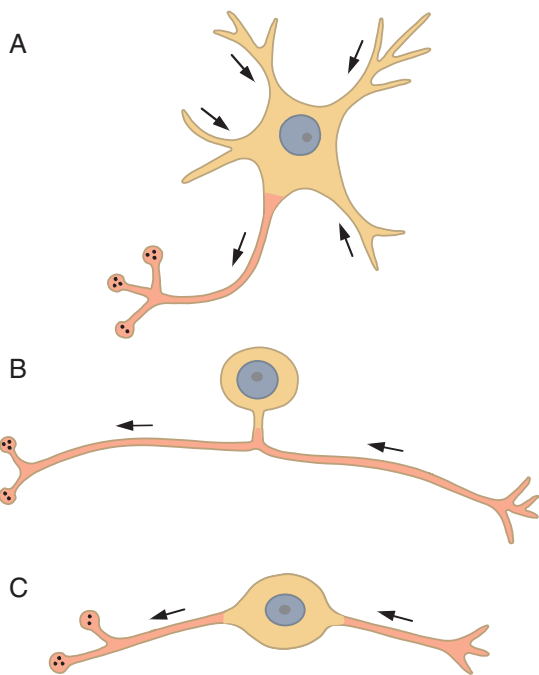


FIGURE 1.5 *Neurons exemplifying three different arrangements of processes.* Multipolar (A), pseudounipolar (B), bipolar (C). Arrows show the direction of impulse conduction.

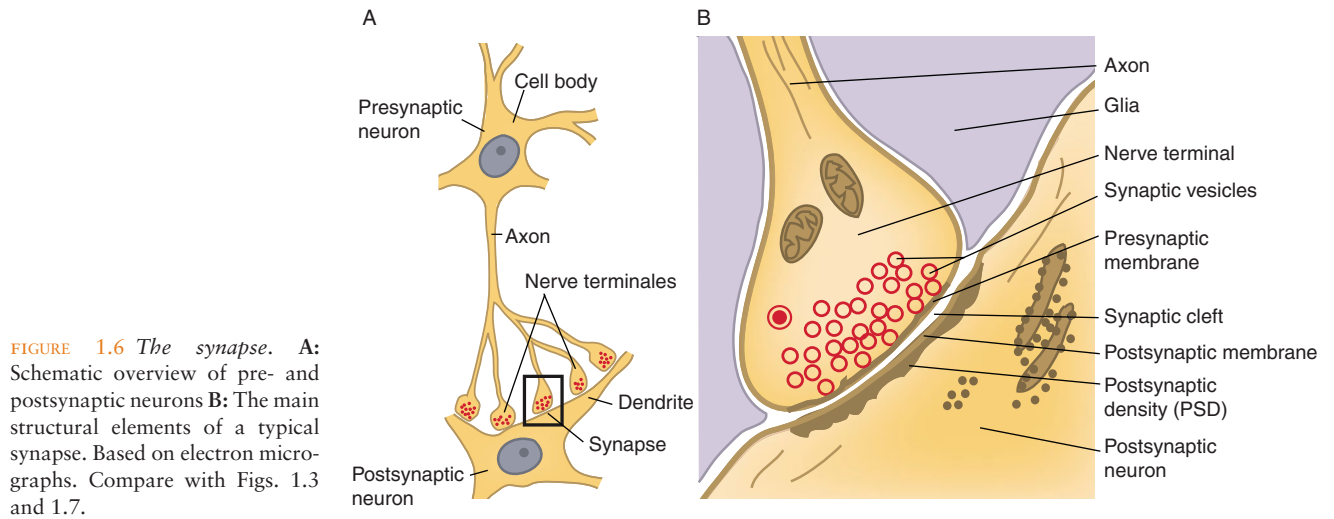


FIGURE 1.6 *The synapse.* **A:** Schematic overview of pre- and postsynaptic neurons **B:** The main structural elements of a typical synapse. Based on electron micrographs. Compare with Figs. 1.3 and 1.7.

The other main type of neuron is the **interneuron**, or Golgi type 2 (Fig. 1.10, see also Fig. 33.6), characterized by a short axon that branches extensively in the vicinity of the cell body. Its name implies that an interneuron is intercalated between two other neurons (Fig. 1.12). Even though, strictly speaking, all neurons with axons that do not leave the CNS are thus interneurons, the term is usually restricted to neurons with short axons that do not leave one particular neuronal group. The interneurons thus mediate communication between neurons within one group. Because interneurons may be switched on and off, the possible number of interrelations among the neurons within one group increases dramatically. The number of interneurons is particularly high in the cerebral cortex, and it is the number of interneurons that is so much higher in the human brain than in that of any other animal. The number of typical projection neurons interconnecting the various parts of the nervous system, and linking the nervous system

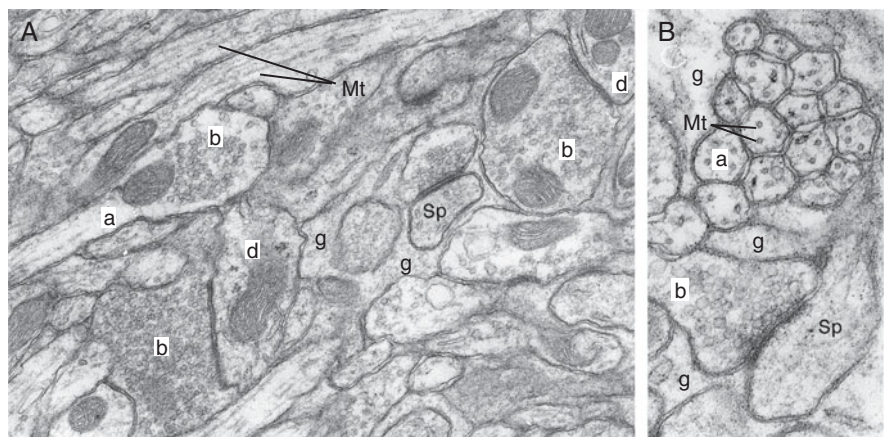
with the rest of the body, as a rule varies more with the size of the body than with the stage of development.

The distinction between projection neurons and interneurons is not always very clear, however. Many neurons previously regarded as giving off only local branches have been shown via modern methods also to give off long axonal branches to more distant cell groups. Thus, they function as both projection neurons and interneurons. In contrast, many of the “classical” projection neurons, for example, in the cerebral cortex (Fig. 33.5), give off collaterals that end within the cell group in which the cell body is located.

Tasks of Interneurons

Figure 1.12 shows how an interneuron (b) is intercalated in an impulse pathway. One might perhaps think that the simpler direct pathway shown below from neuron A to neuron C would be preferable. After all, the

FIGURE 1.7 *Synapses.* **A, B:** Electron micrographs showing boutons (b) in synaptic contacts with dendrites (d), and dendritic spines (Sp). Note how processes of glia (g) cover the dendrites and nerve terminals except at the site of synaptic contact. Note bundle of unmyelinated axons (a) in **B**. Microtubules (Mt) are responsible for axonal transport. Magnifications, $\times 20,000$ (**A**) and $\times 40,000$ (**B**).



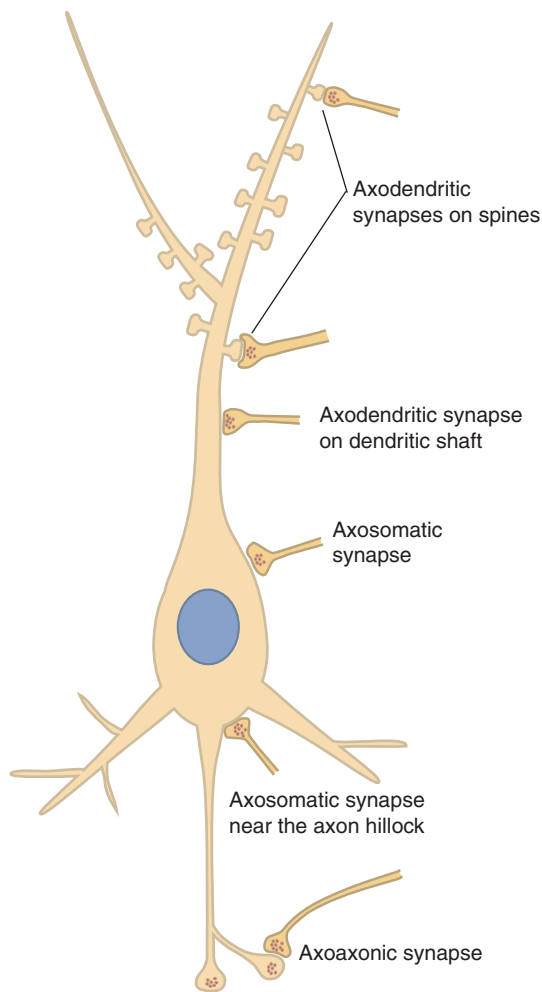


FIGURE 1.8 *The placement of synapses.* The position of a synapse determines (together with other factors) its effect on the postsynaptic neuron. A synapse close to the exit of the axon, for example, has much greater impact than a synapse located on a distal dendrite.

interneuron leads to a delay in the propagation of the signal from A to C, and this would be a disadvantage. Most important, however, is that the interneuron provides added **flexibility**. Thus, whether the signal is transmitted from a to c can be controlled by other synaptic inputs to interneuron b. Identical synaptic inputs to a neuron a may in one situation be propagated further by neuron c but in another situation not, depending on the state of interneuron b. This kind of arrangement may partly explain why, for example, identical stimuli may cause pain of very different intensity: interneurons along the pathways conveying sensory signals are under the influence of other parts of the brain (e.g., neurons analyzing the meaning of the sensory stimulus).

Figure 1.13 illustrates another important task performed by interneurons. Interneuron B enables neuron A to act back on itself and reduce its own firing of impulses. The arrangement acts to prevent neuron A from becoming excessively active. Thus, the **negative feedback** provided by the interneuron would stop the firing of neuron A. Such an arrangement is present, for example, among motor neurons that control striated muscle contraction (see Fig. 21.14).

Many Axons Are Isolated to Increase the Speed of Impulse Propagation

The velocity with which the nerve impulse travels depends on the diameter of the axon, among other factors. In addition, how well the axon is insulated is of crucial importance. Many axons have an extra layer of insulation (in addition to the axonal membrane) called a **myelin sheath**. Such axons are therefore called **myelinated**, to distinguish them from those without a myelin sheath, which are called **unmyelinated** (see Figs. 2.6 and 2.7).

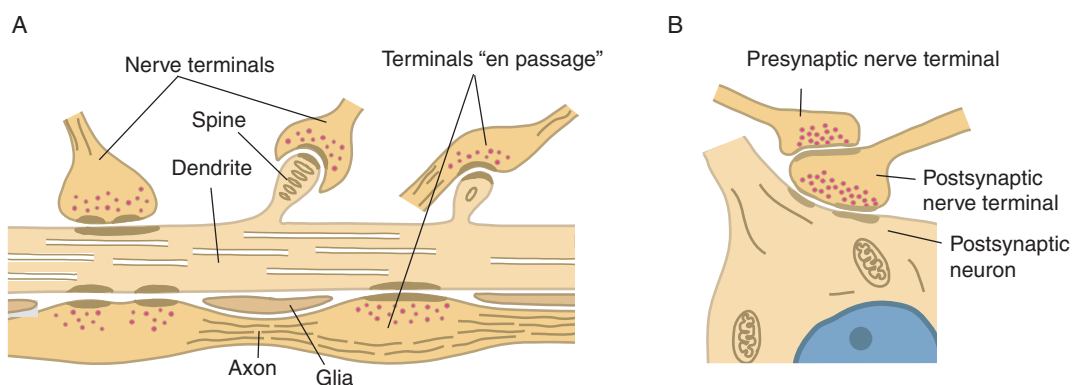


FIGURE 1.9 **A:** *Axodendritic synapses.* A nerve terminal (bouton) may form a synapse directly on the shaft of the dendrite or on a spine. The axon may also have several boutons en passage. **B:** *Axoaxonic*

synapse. The presynaptic nerve terminal influences—by usually inhibiting—the release of neurotransmitter from the postsynaptic nerve terminal.

FIGURE 1.10 *Projection neuron and interneuron.* A projection neuron sends its axon to neurons in other nuclei (cell groups), often at a long distance. The axon of an interneuron ramifies and makes synaptic contacts in its vicinity (within the same nucleus). Examples from the brain stem of a monkey, based on sections treated via the Golgi method, which impregnate the whole neuron with silver salts. The photomicrograph to the left is from the Golgi-stained section containing the drawn projection neuron. The depth of field is only a fraction of the thickness of the section (100 μm). Therefore, only part of the projection neuron is clearly visible in the photomicrograph.

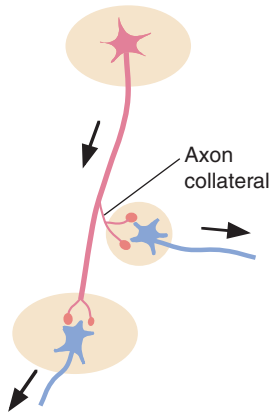
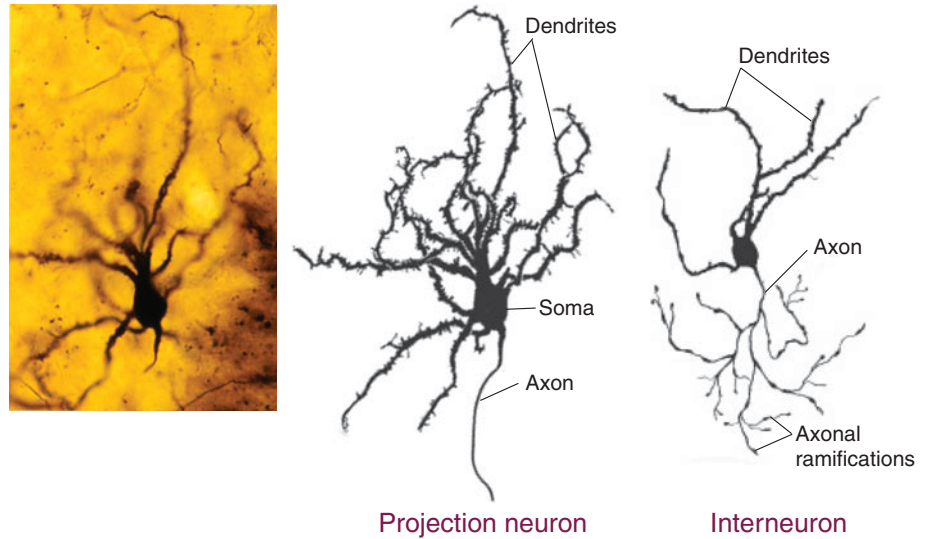


FIGURE 1.11 *Collateral of a projection neuron.* By sending off collaterals, a projection neuron may establish synapses in different cell groups (nuclei). Arrows show the direction of impulse conduction.

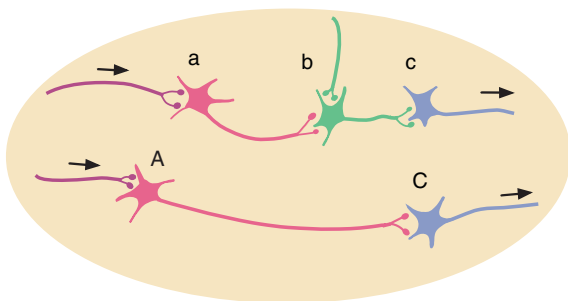


FIGURE 1.12 *An interneuron (b) intercalated in a pathway from neuron a to neuron c.* This arrangement increases the flexibility, as compared with the direct pathway from neuron A to C shown below. Arrows show the direction of impulse conduction.

Many of the tasks performed by the nervous system require very rapid conduction of signals. If unmyelinated axons were to do this, they would have to be extremely thick. Nerves bringing signals to the muscles of the hand, for example, would be impossibly thick, and the brain would also have to be much larger. Insulation is thus a very efficient way of saving space and expensive building materials. Efficient insulation of axons is, in fact, a prerequisite for the dramatic development of the nervous system that has taken place in vertebrates as compared with invertebrates.

Myelin and how it is formed is treated in Chapter 2, while the conduction of nerve impulses is discussed Chapter 3.

White and Gray Matter

The surfaces made by cutting nervous tissue contain some areas that are whitish and others that have a gray color. The whitish areas consist mainly of myelinated axons, and the myelin is responsible for the color; such regions are called **white matter**. The gray regions, called **gray matter**, contain mainly cell bodies and dendrites (and, of course, axons passing to and from the neurons). The neurons themselves are grayish in color. Owing to this difference in color, one can macroscopically identify regions containing cell bodies and regions that contain only nerve fibers in brain specimens (Fig. 1.14).

Neurons Are Collected in Nuclei and Ganglia

When examining sections from the CNS under the microscope, one sees that the neuronal cell bodies are not diffusely spread out but are collected in groups.

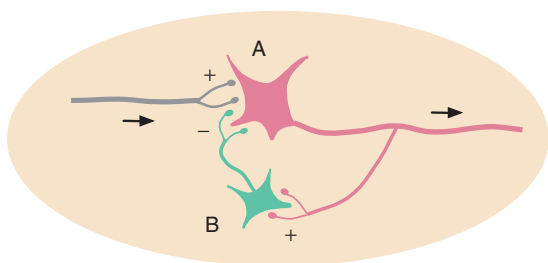


FIGURE 1.13 An interneuron (B) mediates negative feedback to the projection neuron (A). Arrows show the direction of impulse conduction.

Such a group is called a **nucleus** (Figs. 1.14, 1.15, and 1.16). Neurons collected in this manner share connections with other nuclei and constitute in certain respects a **functional unit**; thus, the neurons in a nucleus receive the same kind of information and act on the same (or similar) target. In the PNS, a corresponding collection of cell bodies is called a **ganglion**.

Axons that end in a nucleus are termed afferent, whereas axons that leave the nucleus are efferent. We also use the terms afferent and efferent for axons conducting toward and away from the CNS, respectively. Thus, sensory axons conveying information from sense organs are afferent, while the motor axons innervating muscles are efferent.

Axons Form Tracts and Nerves

Axons from the neurons of one nucleus usually have common targets and therefore run together, forming bundles. Such a bundle of axons connecting one nucleus

with another is called a **tract** (tractus; Figs. 1.15 and 1.16). In the PNS, a collection of axons is called a **nerve** (nervus; Fig. 1.16, see also Figs. 11.1 and 28.3). We also use the term **peripheral nerve** to emphasize that a nerve is part of the PNS. Tracts form white matter of the CNS, and likewise, peripheral nerves containing myelinated axons are whitish.

Schematically, the large tracts of the nervous system are the main routes for nerve impulses—to some extent, they are comparable to highways connecting big cities. In addition, there are numerous smaller pathways often running parallel to the highways, and many smaller bundles of axons leave the big tracts to terminate in nuclei along the course. The number of smaller “foot-paths” interconnecting nuclei is enormous, making possible, at least theoretically, the spread of impulses from one nucleus to almost any part of the nervous system. Normally, the spread of impulses is far from random but, rather, is highly ordered and patterned. As a rule, the larger tracts have more significant roles than the smaller ones in the main tasks of the nervous system. Consequently, diseases affecting such tracts usually produce marked symptoms that can be understood only if one has a fair knowledge of the main features of the wiring patterns of the brain.

COUPLING OF NEURONS: PATHWAYS FOR SIGNALS

In addition to the properties of synapses, which determine the transfer of signals among neurons, the function of the nervous system depends on how the various neuronal groups (nuclei) are interconnected (often called

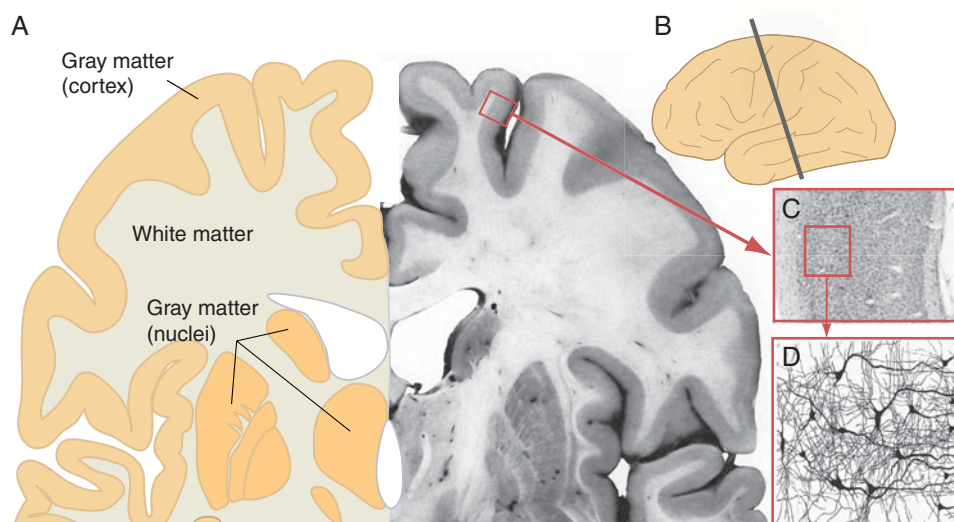


FIGURE 1.14 Gray and white matter. **A:** Drawing and photograph of an unstained frontal section through the human brain. **B:** The white matter consists only of axons and glial cells, whereas the gray matter contains the cell bodies, dendrites, and nerve terminals. **C:** Low-power photomicrograph of a section through the cerebral cortex (frame in A) stained so that only neuronal somata are visible (as small dots). **D:** Drawing of neurons in a section through the cerebral cortex (Golgi method). Only a small fraction of the neurons present in the section are shown.

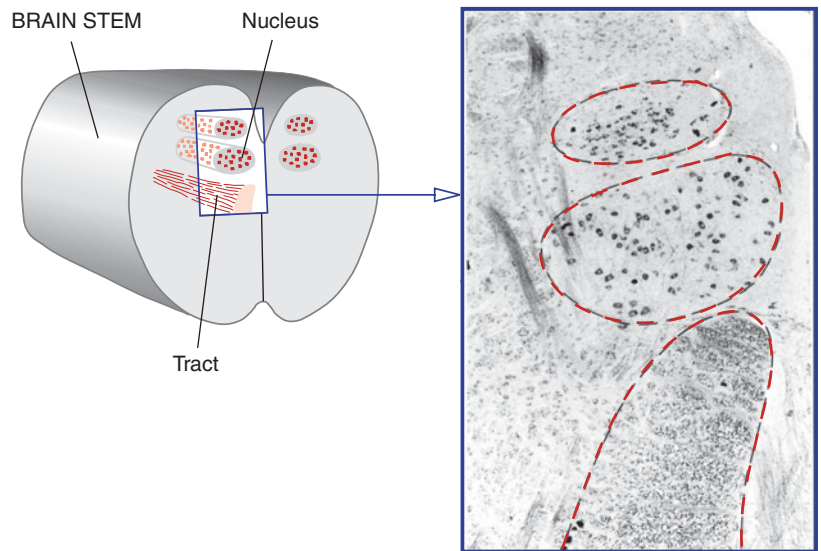


FIGURE 1.15 *Nucleus and tract.* Left: Schematic of part of the brain stem, showing the three-dimensional shape of two nuclei and a tract. Right: Photomicrograph showing the same structures in a section stained to visualize somata and myelinated axons. Magnification, $\times 75$.

the **wiring pattern** of the brain). This pattern determines the pathways that signals may take and the possibilities for cooperation among neuronal groups. Thus, although each neuron is to some extent a functional unit, it is only by proper cooperation that neurons can fulfill their tasks. We will describe here some typical examples of how neurons are interconnected, as such general

knowledge is important for understanding the specific examples of connections dealt with in later chapters.

Divergence and Convergence

A fundamental feature of the CNS is that each neuron influences many—perhaps thousands—of others; that is, information from one source is spread out. This phenomenon is called **divergence** of connections. Figure 1.17 shows schematically how a sensory signal (e.g., from a fingertip) is conducted by a sensory neuron to the spinal cord and there diverges to many spinal neurons. Each of the spinal neurons acts on many neurons at higher levels.

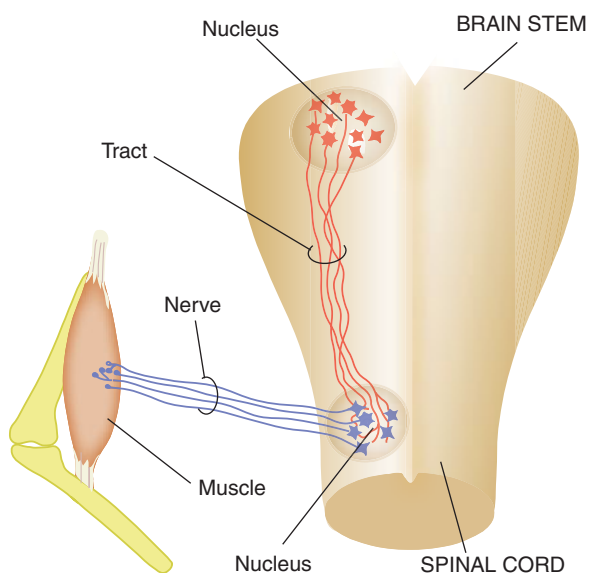


FIGURE 1.16 *Nucleus, tract, and nerve.* Three-dimensional schematic of parts of the brain stem, spinal cord, and a muscle in the upper arm. Axons from a nucleus in the brain stem form a tract destined for a nucleus in the spinal cord. The axons of the neurons in the spinal nucleus leave the CNS and form a nerve passing to the muscle.

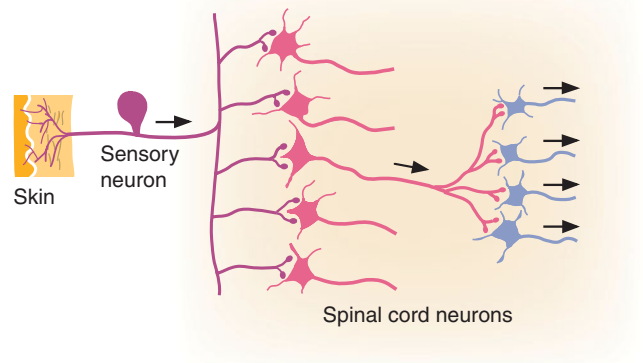


FIGURE 1.17 *Divergence of neural connections.* Highly simplified diagram. The axon collaterals of one sensory neuron contact many neurons in the spinal cord (red). Each of the spinal neurons contacts many other neurons (blue) in the cord or in the brain stem. In this way, the signal spreads from one neuron to many others. Arrows show the direction of impulse conduction.

Another equally ubiquitous feature, **convergence** of connections, is shown schematically in Fig. 1.18. It means that each neuron receives synaptic contacts from many other neurons. The motor neuron in Fig. 1.18 controls the contraction of a number of striated muscle cells (but could have been almost any neuron in the CNS). The motor neuron receives synaptic contacts from many sources (peripheral sense organs, motor neurons in the cerebral cortex that initiate voluntary movements, and so forth). In this case, the motor neuron represents the **final common pathway** of all the neurons acting on it.

The nerve impulses may not necessarily follow all the available pathways shown in Figs. 1.17 and 1.18 because, as a rule, many synapses must be active almost simultaneously to make a neuron fire impulses. Thus, more than one of the blue neurons in Fig. 1.18 must be active at the same time to bring the motor neuron to fire impulses and make the muscle contract. This phenomenon is termed **summation** and is exemplified further in Fig. 1.19. The many synapses axon a makes on neuron A brings the latter to fire a series of nerve impulses whenever axon a is active. But because of fewer synapses, the impact of axon a on neurons B and C is too weak to make them fire impulses. If, however, axons b and c are active simultaneously with a, their effects are summated so that neurons B and C may fire impulses. Summation is discussed further in Chapter 4.

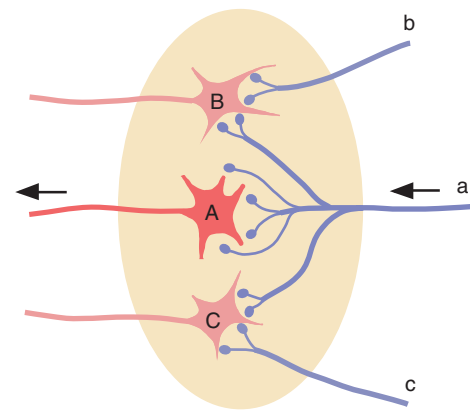


FIGURE 1.19 *Summation.* Many synapses must act on a neuron at the same time to make it fire impulses. Axon a makes many synaptic contacts with neuron A, and their effects summate so that the neuron A fires impulses. In contrast, axon a forms only few synapses with neurons B and C and is not able on its own to fire these neurons. If, however, also axons b and c send impulses at the same time as a, summation ensures that neurons B and C fire impulses. Arrows show the direction of impulse conduction.

Parallel Pathways and Reciprocal Connections

Figure 1.20 illustrates common types of connections among neuronal groups (nuclei). Figure 1.20A shows the principle of **parallel pathways**. There is one direct pathway from nucleus N1 to N2, and one indirect pathway that is synaptically interrupted in other nuclei (n1 and n2). Thus, some of the information reaching N2 is a direct consequence of the activity in N1, whereas information passing through n1 and n2 is modified by other connections acting on these nuclei (not shown). The abundance of such parallel pathways in the human cerebral cortex is one of the factors that explain its enormous flexibility and capacity for information processing (Fig. 1.23). Parallel pathways may, further, be of practical importance after partial **brain injury**. If, for example, the direct pathway between N1 and N2 is interrupted, the indirect one may at least partly take over the tasks formerly performed by the direct one (examples of this are discussed in Chapter 11, under “Restitution of Function”).

Reciprocal connections represent another common arrangement, in which a nucleus receives connections from the nuclei to which it sends axons (Fig. 1.20B). In many cases, such back-projections serve as **feedback**, whereby the first nucleus is informed of the outcome of the impulses emitted to the second one. If the influence was too strong, the feedback may serve to reduce activity, and vice versa if the influence was too weak. Among other actions, such feedback connections serve to stabilize the functioning of the nervous system. Thus, many of the symptoms appearing in neurological diseases are due to the failure of feedback mechanisms. Often, however, it is not obvious which one should be regarded as

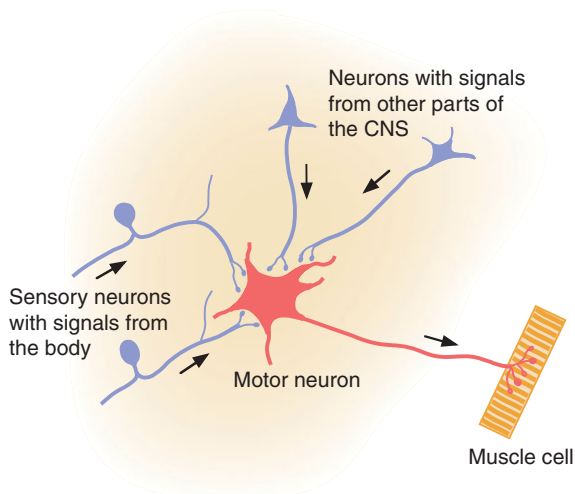


FIGURE 1.18 *Convergence of neural connections.* Synaptic inputs from many neurons (blue) converge onto one neuron (red). In this example, the red neuron is motor and sends its axon to striated muscle cells. The sum of all converging synaptic inputs determines the frequency of impulses sent from the motor neuron to the muscle cells—and thus their strength of contraction. Arrows show the direction of impulse conduction.

a feedback connection and which one as a **feed-forward** connection. Presumably, a single pathway may serve both purposes.

Couplings Contributing to Continuous Neuronal Activity

There is always electric activity in the CNS, because numerous neurons are firing impulses at any given time. In the cerebral cortex, for example, even during sleep there is considerable neuronal activity. How is this activity sustained, even in the absence of sensory inputs? In early embryonic life, groups of neurons become spontaneously active—that is, they fire impulses without any external influence (this is caused by development of special membrane properties). As the nervous system matures, neuronal behavior is governed more and more by synaptic connections with other neurons; nevertheless, some neurons remain spontaneously active. Another feature contributing to continuous activity is that, when activated, most neurons fire a train of impulses, not just one. Further, interneurons contribute to prolongation of activity, as schematically exemplified in Fig. 1.21. Impulses in axon *a* make neuron A fire impulses, propagated along its axon. At the same time, axon *a* makes interneuron 1 fire impulses, which act on neuron A and interneuron 2. The latter acts on neuron A to produce impulses. Owing to a delay of a few milliseconds at each synapse and the time for conducting the impulse in the axons, neuron A receives synaptic inputs over a prolonged period. This kind of coupling (in reality far more elaborate than shown in Fig. 1.21) can translate a brief synaptic input to long-lasting neuronal firing in a neuronal network. Working memory, that is, the ability to keep task-relevant information in mind for a while, depends on neurons that continue firing after a stimulus has stopped.

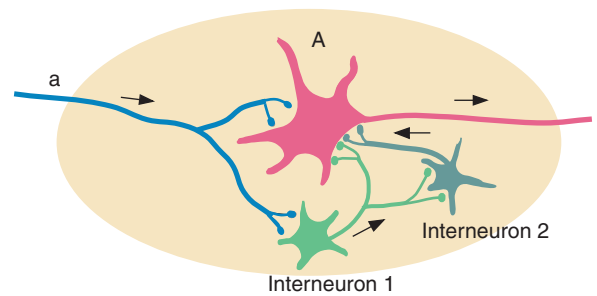


FIGURE 1.21 Interneurons that prolong the activity of a projection neuron (A) when activated by impulses in axon *a*. Arrows show the direction of impulse conduction.

Connections between the Two Halves of the Central Nervous System

Another important general feature of the CNS is that many nuclei have connections with both sides of the brain—so-called **bilateral connections** (Fig. 1.22A). Some tracts supply both sides with approximately the same number of axons (i.e., equal numbers of crossed and uncrossed axons), whereas other tracts are predominantly crossed (contralateral), with only a few axons supplying the same (ipsilateral) side. Although the functional significance of such bilateral connections may not always be clear, they can contribute to recovery of function after partial brain damage.

That the two sides of the CNS cooperate extensively is witnessed by the vast number of **commissural connections**—that is, direct connections between corresponding parts in the two brain halves (Fig. 1.22B). Such connections occur at all levels of the CNS, but the most prominent one connects the two halves of the cerebral hemispheres (corpus callosum; see Figs. 3.26 and 3.27). In humans, this pathway contains approximately 200 million axons.

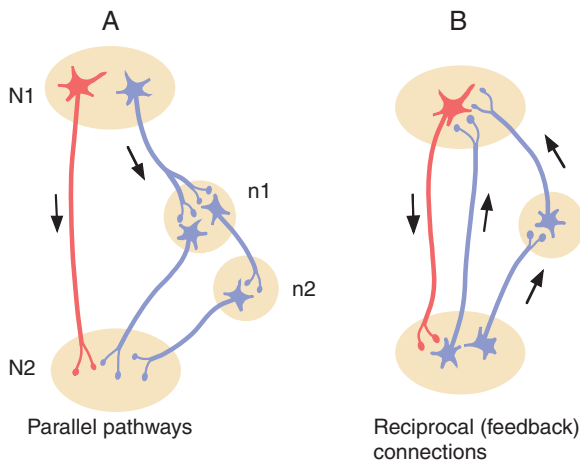


FIGURE 1.20 Examples of organization of neuronal pathways. Arrows show the direction of impulse conduction; N1, N2, n1, and n2 are nuclei in different parts of the CNS.

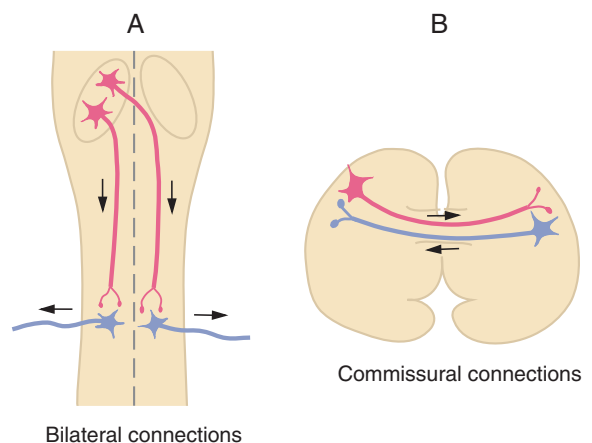


FIGURE 1.22 Examples of organization of neuronal pathways. Arrows show the direction of impulse conduction.

Single Neurons Are Parts of Neural Networks

The tasks of a neuron can be understood only in conjunction with the thousands of neurons with which it is synaptically interconnected. Further, functions of the brain are very seldom the responsibility of one neuronal group or “center” but, rather, the result of cooperation among many neuronal groups. Such cooperating groups or nuclei often lie far apart. For example, proper voluntary movements require cooperation among specific neuronal groups in the cerebral cortex, the cerebellum, and the basal ganglia deep in the cerebral hemispheres. Today we use the term **distributed system** rather than **center** when referring to the parts of the brain that are responsible for a specific function. Such a distributed system is a complicated **neural network** of spatially separate but densely interconnected neuronal groups. Figure 1.23 gives a very simplified example of such a network that could be dedicated to, for example, the subjective sensation of pain. Owing to the abundance of reciprocal connections, the signal traffic can take various routes within the network, and each neuronal group has connections outside the network. This means that a variety of inputs can activate the network—all presumably giving the same functional result (the sensation of pain, a specific memory, an emotion, and so forth). Nevertheless, it should not be surprising that each group, or **node**, might participate in several different, function-specific networks. Thus, as a rule one neuronal group participates in several tasks.

The organization of the brain in distributed systems is particularly clear with regard to **higher mental functions**. Language is a good example: there is not one center for language but specific neuronal groups in many parts of the cerebral cortex that cooperate. Other networks are responsible for attention, spatial orientation, object identification, short-term memory, and so forth.

Data-based models of neural networks have provided new insight into the workings of the cerebral cortex and how symptoms arise from partial destruction of networks.

Injuries of Neural Networks

An important feature of distributed systems is that **partial damage** can degrade their performance but seldom eliminate it. Sometimes partial damage may become evident only in situations with very high demands, for example, with regard to the speed and accuracy of movements, the capacity of short-term memory, and so forth. If the number of neurons participating in the network undergoes further reduction, however, performance may deteriorate severely. In such cases, symptoms may occur rather abruptly, even though the disease process responsible for the cell loss may have been progressing slowly for years. This is typical of degenerative brain diseases such as Parkinson’s disease and Alzheimer’s disease.

THE CYTOSKELETON AND AXONAL TRANSPORT

The cell bodies and processes of neurons contain thin threads called **neurofibrils**, which can be observed in specially stained microscopic sections (Fig. 1.24). The neurofibrils are of different kinds, but together they form the **cytoskeleton**—the name refers to its importance for development and maintenance of **neuronal shape**. The fact that neurons have very different shapes—with regard to dendrites, cell bodies, and axons—is due to cytoskeletal specializations. For example, the neurofibrils have a decisive role when axons grow for long distances, and the cytoskeleton serves to anchor synaptic elements at the postsynaptic density (see Fig. 4.6).

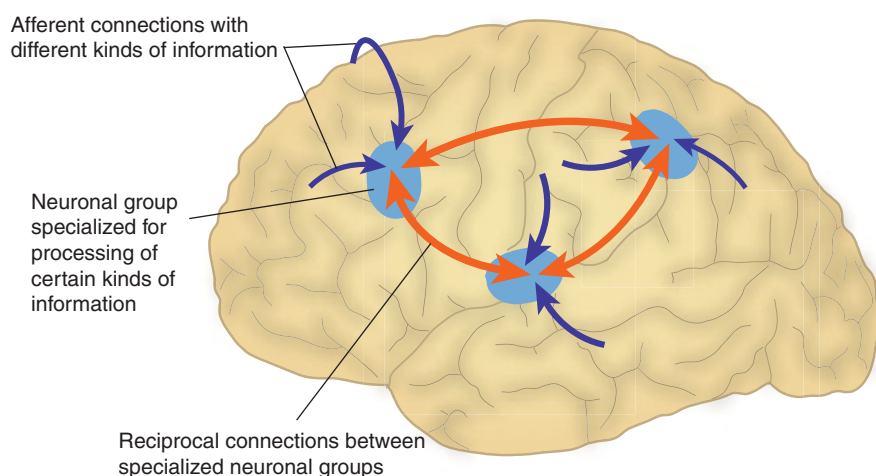


FIGURE 1.23 *Distributed neural networks.* Simplified. Three regions (groups of neurons) in the cerebral cortex are interconnected by reciprocal connections (red arrows). The collective activity of all parts of the network is responsible for its “product”—for example, the sensation of pain.

The neurofibrils of the cytoskeleton are also responsible for another important cellular function: the transport of **organelles** and **particles** in the neuronal processes. Although such transport takes place in both dendrites and axons, **axonal transport** (Fig. 1.25) has been most studied (mainly because, for technical reasons, transport in dendrites is much harder to study). It is obvious that neurons need direction-specific transport mechanisms. Thus, the organelles necessary for protein synthesis and degradation of particles are present almost exclusively in the cell body. Nevertheless, dendrites contain small amounts of mRNA located at the base of dendritic spines, which may enable a limited amount of protein synthesis important for synaptic changes related to learning and memory.

Transport from the cell body toward the nerve terminals is called **anterograde** axonal transport (Fig. 1.25). Examples of particles transported anterogradely are mitochondria, synaptic vesicles, proteins to be inserted in the axonal membrane, and enzymes for transmitter synthesis and degradation in the nerve terminals. Growth factors, synthesized in the cell body but liberated far away at the synapses, also require efficient anterograde axonal transport. Transport toward the cell body from the nerve terminals is called **retrograde** axonal transport. Retrograde transport brings signal

molecules of various kinds that are taken up by the nerve terminals to the cell body (Fig. 1.25). Often such molecules are produced by postsynaptic cells and released to the extracellular space. In the cell body (nucleus) the signal molecules can influence genetic expressions—that is, they can change protein synthesis. In this way, the properties of the neuron can be changed transiently or in some instances permanently. This is a form of **feedback**: ensuring that the neuron is informed of its effects on other cells and of the state of its target cells. In some instances, neurons even require this kind of feedback to survive. Retrograde transport also moves “worn-out” organelles to the cell body for degradation in lysosomes.

Components of the Cytoskeleton

Electron microscopic and biochemical analyses have shown that the cytoskeleton consists of various kinds of fibrillary proteins, making threads of three main kinds:

1. **Actin filaments** (microfilaments) and associated protein molecules (approximately 5 nm thick)
2. **Microtubules** (narrow tubes) and associated proteins (approximately 20 nm thick)
3. **Intermediary filaments** or neurofilaments (approximately 10 nm thick)

Actin (microfilaments) is present in the axon, among other places. There it has an important role during development. When the axon elongates, actin together with microtubules serves to produce movements of the **growth cone** (Fig. 9.16) at the tip of the axon (in general, actin is present in cells capable of movement, such as muscle cells). The growth cone continuously sends out thin fingerlike extensions (filopodia) in various directions. These probably explore the environment for specific molecules that mark the correct direction of growth. In addition, actin is probably important in maintaining the shape of the fully grown axon.

Microtubules and **microtubule-associated proteins** (MAPs) are present in all kinds of neuronal processes and are most likely important for their shape (Figs. 1.7 and 2.7). Of special interest is the relation of microtubules to the transport of substances in the neuronal processes. As mentioned, there is a continuous movement of organelles, proteins, and other particles in the axons and dendrites. Destruction of microtubules by drugs (such as colchicine) stops axonal transport.

The functional role of the **intermediary filaments** (neurofilaments) is uncertain, although they make up about 10% of axonal proteins. One function might be to maintain the diameter of thick myelinated axons, as internal scaffolding. Whatever their normal role is, it is noteworthy that neurofilaments are altered in several degenerative neurological diseases. In Alzheimer’s disease, for example, a characteristic feature is disorganized



FIGURE 1.24 *The cytoskeleton in neurons.* Drawing of neurons from the cerebral cortex, as appearing in sections stained with heavy metals to visualize neurofibrils. Both dendrites and axons (a) contain numerous neurofibrils. (From Cajal 1952.)

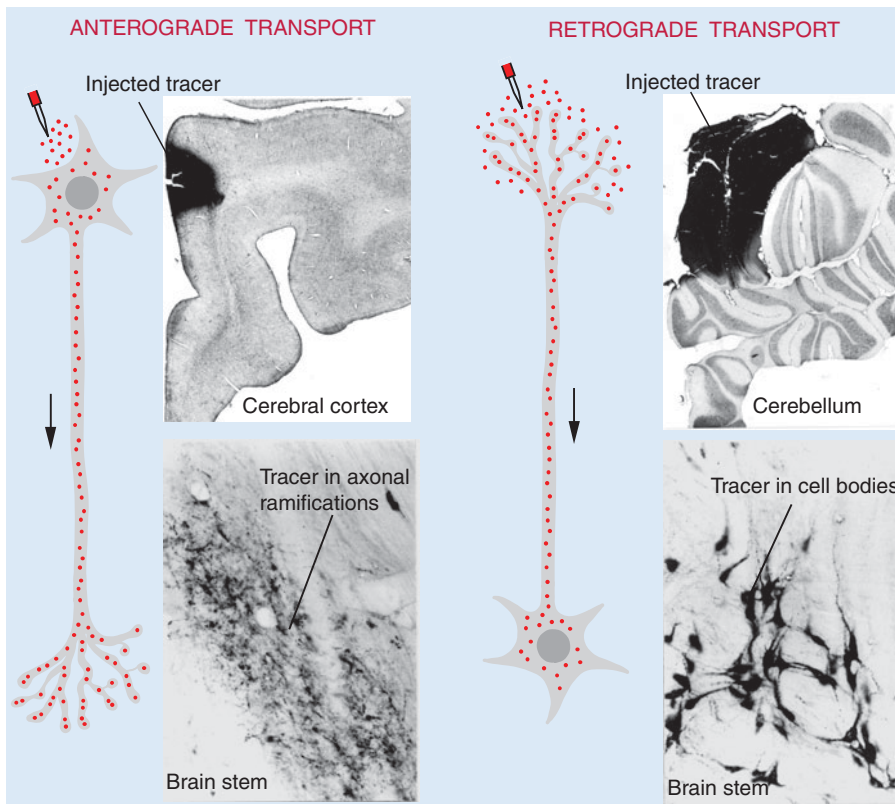


FIGURE 1.25 *Axonal transport.* The photomicrographs illustrate the use of axonal transport for tract tracing, that is, to map the connections in the CNS. A cat received injections of an enzyme (horse-radish peroxidase, HRP) in the cerebellum and in the cerebral cortex ($0.2 \mu\text{L}$ in each). The enzyme was taken up by endocytosis of neuronal cell bodies and terminals. Vesicles with enzyme were then transported anterogradely from the cerebral cortex to the brain stem (left) and retrogradely from the cerebellum to the brain stem (right). A black reaction product in the upper photomicrographs shows the extension of the tracer at the injection site. The anterogradely labeled terminal ramifications of the axons appear as black dust in the left lower photomicrograph, while retrogradely labeled cell bodies are seen in the right lower photomicrograph. Magnifications, $\times 8$ (upper) and $\times 150$ (lower)

tangles of intermediary filaments in the cerebral cortex (neurofibrillary tangles).

More about Axonal Transport and Its Machinery

The injection of radioactively labeled substances taken up by neurons has shown that axonally transported material moves in at least **two phases**. One phase is **rapid**, with particles moving up to half a meter per day; the other is **slow**, with movement of between 1 and 3 mm per day. The rapid phase carries mainly organelles and vesicles, that is, membrane-bound structures. The slow phase carries primarily enzymes and components of the cytoskeleton. As mentioned, microtubules are of particular importance for axonal transport. Each microtubule is composed of smaller building blocks of the protein **tubulin**. MAPs help the formation of tubes from many tubulin molecules. MAPs also anchor the microtubules to the cell membrane and to other parts of the cytoskeleton, such as neurofilaments. Two kinds of MAPs found only in neurons—**MAP2** and **tau**—stiffen the microtubules. Specific kinds of MAPs perform

anterograde and retrograde transport, respectively, serving as the “motors” of axonal transport. These MAPs are ATPases (enzymes that split ATP), and the released energy alters their form, thus producing movement. The transported particles, such as vesicles and mitochondria, move by temporarily binding to MAPs protruding from the microtubule, so that they appear to “walk” along the microtubule. One microtubule can transport in both directions, depending on the kind of motor to which a particle binds. Proteins belonging to the **kinesin** family are responsible for anterograde movement. Different varieties of kinesin appear to transport different “cargo”; for example, one variety transports mitochondria and another transports precursors of synaptic vesicles. **Dynein**, which is a more complex protein than kinesin, is responsible for the bulk of retrograde transport, although certain kinesins probably also contribute.

Injections into nervous tissue of substances that are transported axonally and later can be detected in tissue sections are widely used for tract tracing, that is, to reveal the “wiring pattern” of the brain (Fig. 1.25).

2 | Glia

OVERVIEW

Glial cells are the most numerous cells in the brain and are indispensable for neuronal functioning. Glial cells are of three kinds that differ structurally and functionally. **Astrocytes** have numerous processes that contact capillaries and the lining of the cerebral ventricles. They serve important **homeostatic functions** by controlling the concentrations of ions and the osmotic pressure of the extracellular fluid (water balance), thereby helping to keep the neuronal environment optimal. Astrocytes also take part in **repair processes**. **Oligodendrocytes** insulate axons by producing **myelin sheaths** in the central nervous system (CNS). **Microglial** cells are the **macrophages** of nervous tissue. **Schwann cells** are a specialized form of glial cells that form myelin sheaths in the peripheral nervous system (PNS). Apart from these specific functions, glial cells are involved in the **prenatal development** of the nervous system, for example, by providing surfaces and scaffoldings for migrating neurons and outgrowing axons.

TYPES OF GLIAL CELLS

Although they do not take part in the fast and precise information processing in the brain, glial cells are nevertheless of crucial importance to proper functioning of neurons. In fact, the number of glial cells in the brain is much higher than the number of neurons. The name *glia* derives from the older notion that glial cells served as a kind of glue, keeping the neurons together. Although improved methods have revealed hitherto unknown properties of glial cells, much still remains to be understood about their functional roles in the nervous system.

It is customary to group glial cells into three categories: **astrocytes**, or astroglia; **oligodendrocytes**, or oligodendroglia; and **microglial cells**, or microglia. Each is structurally and functionally different from the others. Astrocytes have numerous processes of various shapes whereas oligodendrocytes have relatively few and short processes (*oligo* means few, little). In routinely stained sections, glial cells can be distinguished from neurons by their much smaller nuclei. The identification of the various types, however, requires immunocytochemical methods to identify proteins that are specific to each type.

Specialized Forms of Glial Cells

In addition to the three main kinds, there are other, specialized forms of glial cells. The surface of the cavities inside the CNS is lined with a layer of cylindrical cells called **ependyma** (Fig. 2.3; see also Fig. 9.6). There are also special types of glial cells in the retina (**Müller cells**), the cerebellum (**Bergman cells**), and the posterior pituitary gland (**pituicytes**).

GLIAL CELLS AND HOMEOSTASIS

Astrocytes Contact Capillaries, Cerebrospinal Fluid, and Neurons

Astrocytes have structural features that make them well suited to control the extracellular environment of the neurons:

1. They have numerous short or long processes that extend in all directions (Figs. 2.1 and 2.2). Thus, the astrocytes have a very **large surface area** that enables efficient exchange of ions and molecules with the extracellular fluid (ECF).
2. Some processes contact the surface of **capillaries** with expanded **end “feet”** and cover most of the capillary surface (Figs. 2.3 and 2.4).
3. Some processes form a continuous, thin sheet (membrana limitans, also called glia limitans) where nervous tissue borders the **cerebrospinal fluid (CSF)**, that is, in the cavities inside the CNS and against the connective tissue membranes on its exterior (Fig. 2.3).
4. Other processes contact **neuronal surfaces**; in this manner, parts not contacted by boutons are covered by glia (Figs. 2.3 and 2.5). Glial processes usually enclose the nerve terminal (see Figs. 1.6 and 1.7).
5. Numerous **gap junctions** (nexus) couple astrocytes, allowing free passage of ions and other small particles among them (Fig. 2.4). Thus, apart from allowing electric currents to spread, astrocytes form continuous, large fluid volumes for distribution of substances removed from the ECF.

Glial Cells Communicate with Electric Signals and Influence Cerebral Blood Flow

Although glial cells do not send precise signals over long distances, they can produce brief electric impulses

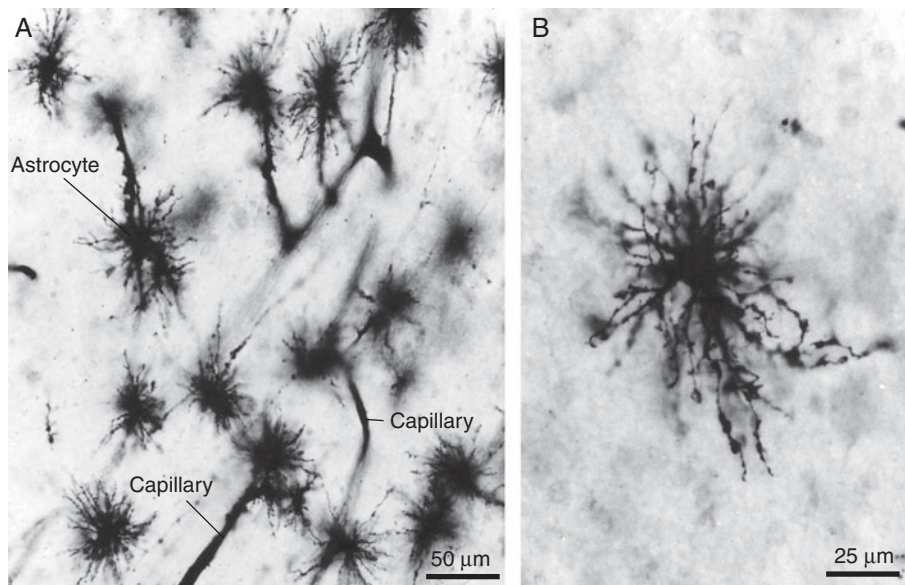


FIGURE 2.1 *Astrocytes*. Photomicrographs of Golgi-stained sections from the cerebral cortex. No neurons are visible. Note the close relationship between astrocytic processes and capillaries.

(currents) by opening of membrane **channels** for Ca^{2+} . Such an opening can be evoked by binding of neurotransmitters (e.g., glutamate) to G-protein-coupled receptors in the glial cell membrane. Thus, neuronal activity can directly influence the astrocytes, whereas the latter affects neuronal activity. Owing to the electric coupling (nexus) of the astrocytes, the “calcium signal” can presumably spread rapidly in networks of astroglial cells, and thus influence many neurons almost simultaneously, which, among other roles, can help synchronize the activity of neurons in a group. In light of the electric coupling among astrocytes, one might expect the population of neurons influenced by an astrocytic network to be quite large. Recent data, however, indicate

that the population can be surprisingly small, enabling spatially precise interactions among neurons and astrocytes. Thus, although it is well known that a specific sensory input (e.g., from a small spot in the visual field) activates neurons in a precisely defined, small part of the cortex, recent experiments (Schummers et al. 2008) suggest that astroglial cells are activated in a similarly precise manner (although a few seconds later than the neurons). Presumably, inputs from the periphery activate neurons that in turn activate astrocytes in their immediate vicinity. When activated, the astrocytes increase local **blood flow** (see Chapter 8, under “Regional Cerebral Blood Flow and Neuronal Activity”).

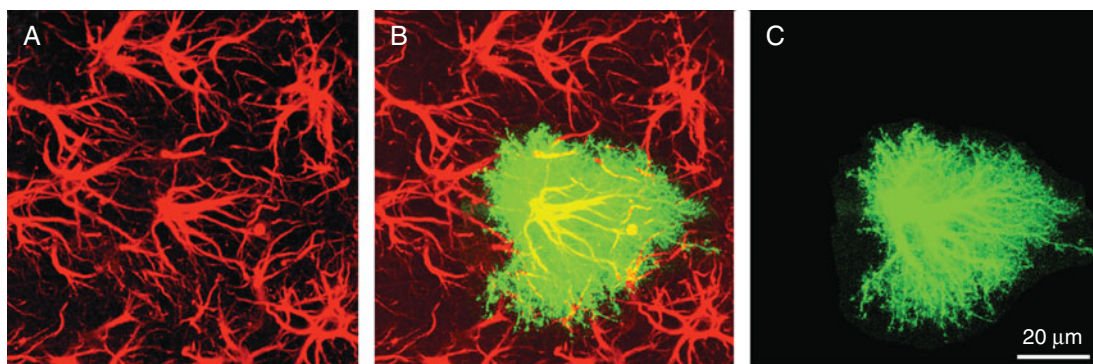


FIGURE 2.2 *Astrocytes*. **A:** Astrocytic processes visualized using an antibody against glial fibrillary acidic protein (GFAP) present in intermediary filaments. The antibody was labeled with a substance with red fluorescence. **B:** One of the astrocytes in **A** has been filled completely with intracellular injection of a substance with green fluorescence (Lucifer yellow), and reconstructed three-dimensionally. It is

obvious that the astrocytic processes are much more abundant and of finer caliber than in **A**. **C:** View of the injected astrocyte in **B** in isolation, showing to advantage its dense and bushy halo of processes. (Reproduced with permission from Wilhelmsson et al. (2004) and *The Journal of Neuroscience*.)

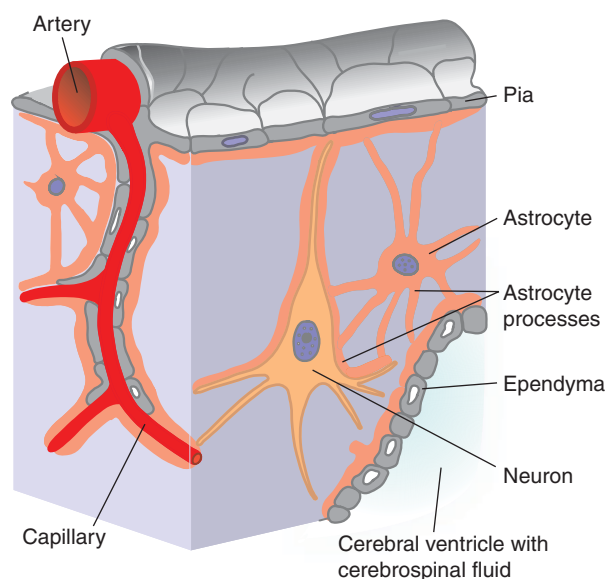


FIGURE 2.3 *The relationship between astroglia and neurons, blood vessels and the CSF.* The astrocytes cover the surface of the neurons and are also closely related to vessels, ependymal cells, and the innermost part of the cerebral meninges (pia).

Astroglia and the Control of Neuronal Homeostasis

Their intimate contact with neurons, capillaries, and the CSF places astroglial cells in a unique position to control the environment of the neurons, that is, the extracellular (interstitial) fluid of the brain (see Fig. 1.28). Such control is vitally important for three main reasons. First, neurons are exquisitely sensitive to changes in extracellular concentrations of ions and neurotransmitters. Second, the osmotic pressure (the water concentration) must be tightly controlled because the brain cannot expand in the skull. Third, adding even minute amounts of a substance may produce a substantial increase in its extracellular concentration, owing to the very limited extracellular space in the brain (less than 20% of total volume), as illustrated in electron micrographs showing only narrow slits between the cellular elements (see Figs. 1.3 and 1.7). Further, the tortuous shape of the extracellular space hampers free diffusion of particles.

With regard to extracellular **ions**, the control of K^+ (potassium ions) is particularly important. Thus, neuronal excitability is strongly influenced even by small changes in the amount of K^+ ions extracellularly, and as neurons fire impulses, K^+ ions pass out of the cell (Fig. 2.4). Prolonged or intense neuronal activity would therefore easily produce dangerously high extracellular levels of K^+ ions were it not for their efficient removal by glia. Further, astrocytes contribute to **extracellular pH** control by removing CO_2 .

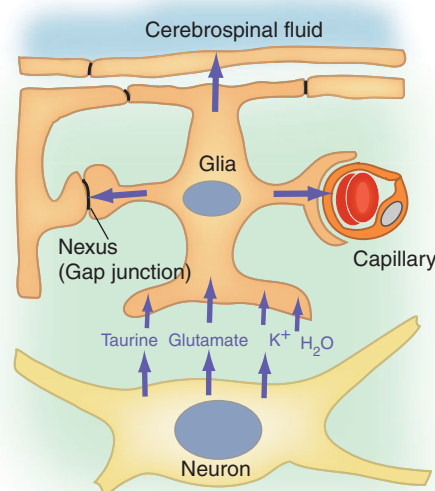


FIGURE 2.4 *Astroglia and the homeostasis of nervous tissue.* Schematic shows the close contacts between astroglial cells on the one hand and neurons, capillaries, and the CSF on the other. The astroglial cells are coupled by nexus (gap junctions), and thus form a large fluid volume for distribution of substances. Some important substances handled by astroglia are indicated (the transport is not always in the direction of the arrows). Surplus of water, K^+ ions, and the amino acid taurine can be transported to the blood and the CSF, thereby preventing their accumulation in the ECF. Next to glutamate, taurine is the amino acid with the highest concentration in the CNS and therefore significantly contributes to the osmolarity of the ECF. Taurine does not appear to function as a neurotransmitter, but its transport in and out of astroglia may be a mechanism for controlling the volume of the neurons. The neurotransmitter glutamate is treated differently, however. Glutamate is transformed to glutamine after uptake in glia, and thus loses its transmitter actions and becomes neutral to the neurons. Glutamine can therefore be returned to the ECF for subsequent uptake into neurons where it is used for resynthesis of glutamate. Because the neurons need large amounts of glutamate, this is an economic means to ensure a sufficient supply. (Based on Nagelhus 1998.)

Extracellular **neurotransmitter concentration** must be tightly controlled, because proper synaptic functioning requires that their extracellular concentrations be very low, except during the brief moments of synaptic release. Most neurotransmitters are indeed removed from the ECF near the synapses by **transporter proteins** in the membranes of neurons and astrocytes. Specific transporters have been identified for several neurotransmitters (discussed further in Chapter 5). Figure 2.5 gives an impression of the abundance of a specific kind of transporter proteins (for the ubiquitous neurotransmitter glutamate, which is neurotoxic in abnormally high concentrations).

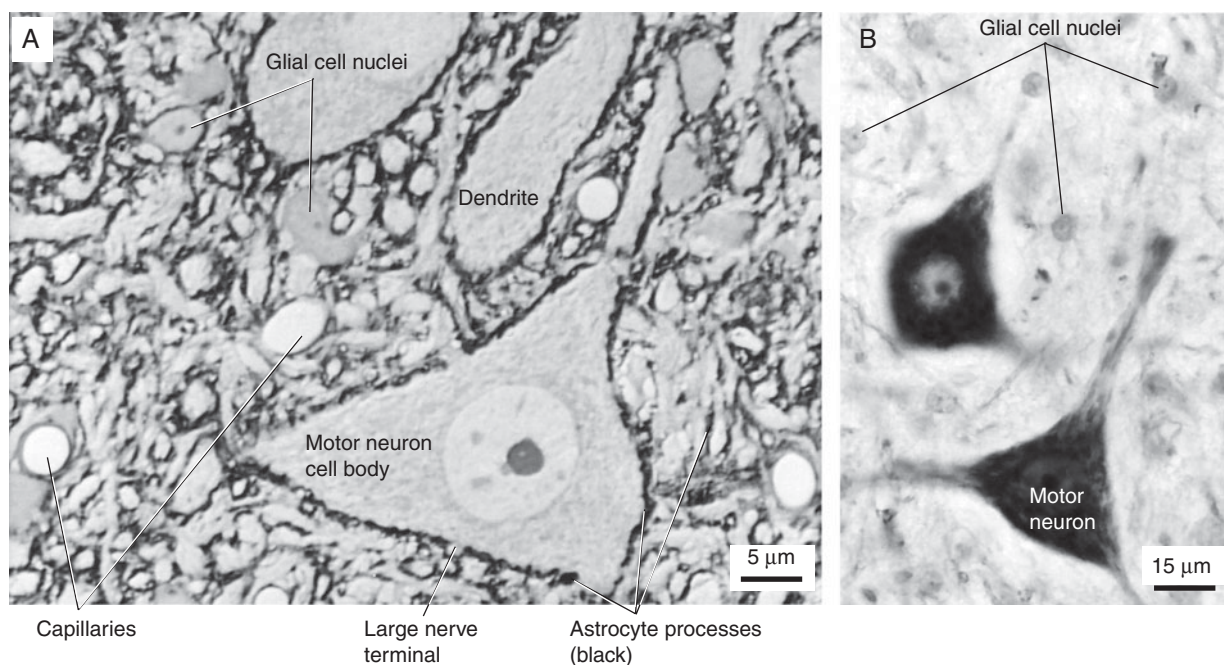


FIGURE 2.5 *Astroglial processes in nervous tissue.* **A:** Photomicrograph showing the distribution of a glutamate-transporter protein, as visualized via an immunocytochemical technique. In this 1 µm thick section from the spinal cord, the dark spots and bands are astrocytic processes expressing glutamate transporters. They outline the somata, dendrites, and capillaries. The picture illustrates both the capacity of astroglia to take up glutamate from the ECF and the enormous astroglial surface facing neurons and capillaries. The contours of dendrites

and neuronal somata are uneven because of synaptic contacts (thin arrow) breaking the otherwise continuous layer of astroglia. Capillaries are marked with asterisks. The cell body of an astroglial cell is marked with a thick arrow. (Courtesy of Drs. J. Storm-Mathisen and N.C. Danbolt, Department of Anatomy, University of Oslo.) **B:** For comparison, a photomicrograph of a thionine-stained section from the same part of the spinal cord as in A.

As mentioned, astrocytes are also involved in the control of the extracellular osmotic pressure, that is, in controlling the **water balance** of the brain (Fig. 2.4). Of particular interest in this respect are channels for transport of water—**aquaporins**—that are present in the membranes of astrocytes. Aquaporins were first described in kidney tubular systems, where they were shown to increase significantly the capacity for water passage. Interestingly, in the brain they are most abundant on the glial processes that are in close contact with capillaries and the CSF, that is, where one would expect them to be if they were involved in brain water balance. Exchange by astroglial cells of small neutral molecules, such as the amino acid **taurine**, may be another mechanism to control extracellular osmolarity.

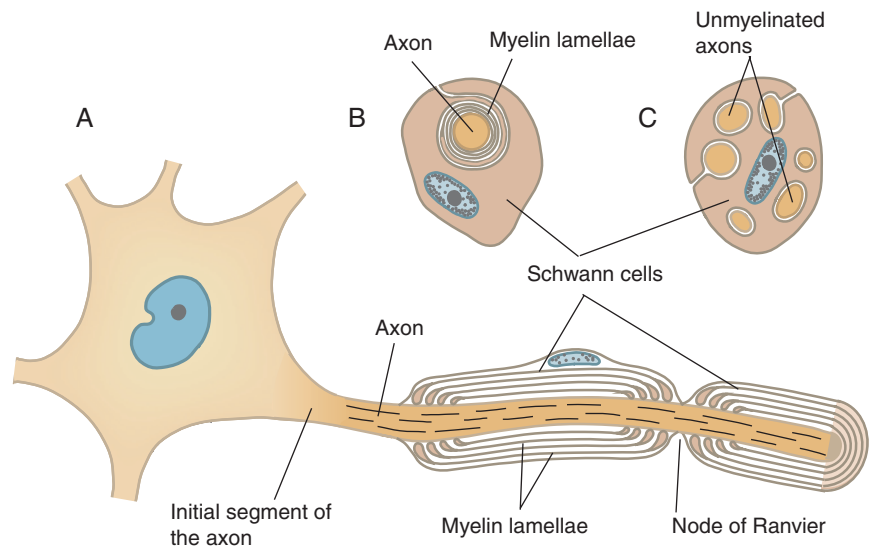
Finally, the layer of astrocytic processes surrounding brain capillaries helps to prevent many potentially harmful substances from entering the brain (see Chapter 7, under “The Blood–Brain Barrier”).

Aquaporins in Health and Disease

Two varieties of aquaporin are present in the brain. **AQP4** is located in the astrocyte membrane, and particularly concentrated in the end-feet region close to

capillaries and in glial processes bordering the CSF. **AQP1** is present in epithelial cells of the choroid plexus (which produces the CSF; see Chapter 7). In general, aquaporins increase water permeability of the cell membrane, thus allowing water to follow active ion transport. A function of AQP4 in the normal brain is probably to export water. Thus, AQP4-deficient mice have increased ECF volume compared to normal mice. Further, in so-called **vasogenic brain edema**, wherein water accumulates extracellularly, AQP4 contributes to removal of excess water. This kind of edema arises when the brain capillaries become leaky due to, for example, traumatic brain injury. On the other hand, when water accumulates intracellularly, as typically occurs in cerebral ischemia or hypoxia (e.g., in stroke), the presence of AQP4 seems to *increase* the edema by allowing more water to enter the astrocytes. Such **cytotoxic brain edema** is caused by failure of energy-dependent ion pumping, which reduces the ability of the cells to maintain osmotic stability. Brain edema is a serious and often life-threatening complication in many brain disorders, such as stroke and traumatic brain injuries. Therefore, the discovery of a relationship between aquaporins and brain edema led to an intensive search for drugs that can modulate the activity of aquaporins.

FIGURE 2.6 *Myelin sheath, myelination, and unmyelinated axons.* Schematics based on electron microscopic observations. **A:** Cell body with proximal parts of the dendrites and myelinated axon. The myelin sheath consists of lamellae formed by the membrane of glial cells (oligodendroglia, or Schwann cells). Each cell produces one segment of myelin. The node of Ranvier is the site of contact between two segments of myelin. The nerve impulse usually starts in the initial segment of the axon, and then “jumps” from one node of Ranvier to the next. **B:** Cross section of an axon in the process of becoming myelinated. The myelin sheath is formed when a glial cell wraps itself around the axon. **C:** Unmyelinated axons in the peripheral nervous system are surrounded by Schwann cell cytoplasm.



In animal experiments, inhibitors of AQP4 can reduce cytotoxic edema whereas they seem to worsen vasogenic edema. This complicates the search for the ideal drug because in human brain disorders the two kinds of brain edema usually coexist (although one may dominate depending on the specific disorder).

INSULATION AND PROTECTION OF AXONS

Oligodendrocytes and Schwann Cells

The myelin sheaths, which insulate axons, are formed by oligodendrocytes¹ in the CNS and by Schwann cells in the PNS. Although the structure and function of the myelin sheaths they produce are the same,² oligodendrocytes and Schwann cells are not identical. One difference is that a single oligodendrocyte usually sends out processes to produce myelin segments for several axons (up to 40), whereas each Schwann cell forms a myelin segment for only one axon (Fig. 2.6). A particularly interesting difference concerns their differential influence on **regeneration** of damaged axons. In the PNS, a cut axon can regenerate under favorable conditions, provided that viable Schwann cells are present. In

the CNS, however, such regeneration of axons does not normally occur, mainly because of inhibiting factors produced by oligodendrocytes.

In addition to forming myelin sheaths, oligodendrocytes and Schwann cells are important for survival of the axons. Thus, diseases affecting oligodendrocytes or Schwann cells produce axonal loss in addition to loss of myelin. In addition, oligodendrocytes and Schwann cells influence axonal thickness and axonal transport.

The Myelin Sheath

The **myelin sheath** forms an insulating cylinder around the axons (Fig. 2.6), reducing the loss of current from the axon to the surrounding tissue fluid during impulse conduction. This contributes to the much higher conduction velocity in myelinated axons than in unmyelinated axons (discussed further in Chapter 3 under “Impulse Conduction in Axons”). The thickest myelinated axons conduct at approximately 120 m/sec (versus less than 1 m/sec in unmyelinated axons).

The myelin sheath consists almost exclusively of numerous layers of cell membrane, as evident from electron micrographs (Figs. 2.6 and 2.7). The layers, or **lamellae**, are formed when a glial cell wraps itself around the axon (Fig. 2.6B). During this process, the cytoplasm of the glial cell is squeezed away so that the layers of cell membrane lie closely apposed. The composite of material ensheathing the axons is called **myelin**. Myelin is whitish in color because of its high lipid content.

The cell membrane forming the myelin has a unique lipid and protein composition. Among other components, myelin has a high content of cholesterol and various **glycolipids**. The glycolipids appear to be crucial

¹ We do not know whether *all* oligodendrocytes form myelin. Thus, their cell bodies are often closely apposed to neuronal cell bodies, suggesting that they may have other tasks in addition to myelination.

² Even though the myelin sheaths produced by oligodendroglial cells and by Schwann cells look the same, they differ significantly in their lipid and protein composition. For example, myelin basic protein (MBP) makes up a much larger fraction of the total myelin protein in the CNS than in the PNS, whereas **peripheral myelin protein-22 (PMP-22)** is absent in the CNS. Another example is **myelin-oligodendrocyte glycoprotein (MOG)**, which is expressed in the CNS only. Such differences may help explain why some diseases affect only myelinated axons in the CNS (e.g., multiple sclerosis), whereas others are restricted to peripheral axons.

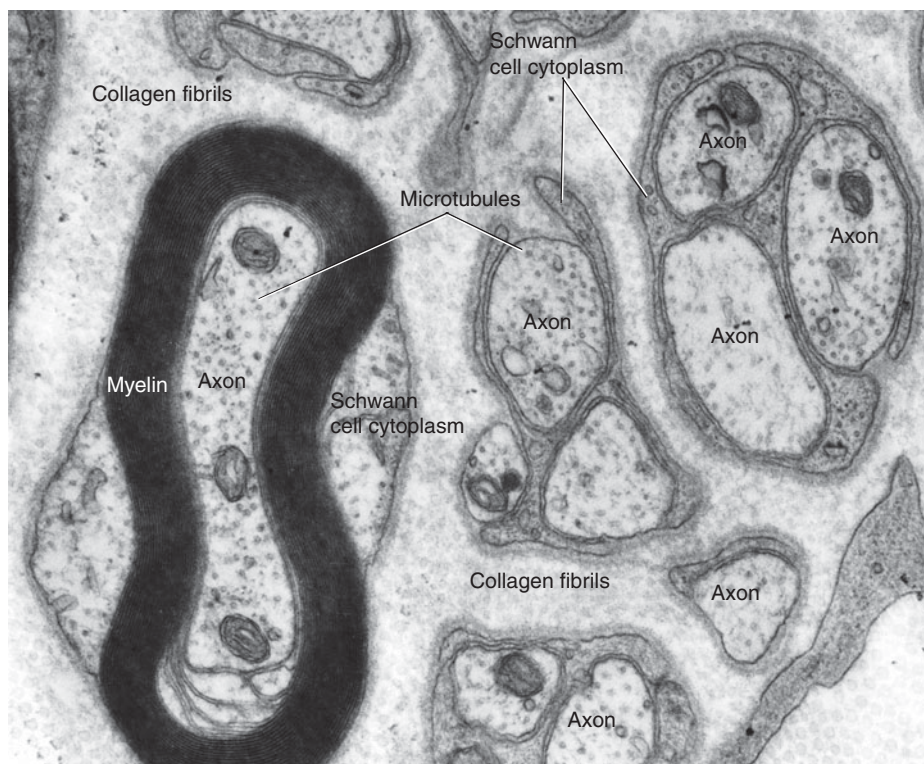


FIGURE 2.7 Myelinated and unmyelinated axons. (Detail from Fig. 2.8.) An axon is surrounded by myelin. The myelin lamellae are seen as dark stripes, arranged concentrically. The cytoplasm of the Schwann cell that is responsible for producing the myelin is seen externally. The unmyelinated axons are completely surrounded by Schwann cells. Between the axons are numerous collagen fibrils. Magnification, $\times 30,000$.

for the insulating properties of myelin. Certain **membrane proteins** related to the immunoglobulins bind the external (apposing) sides of the membranes tightly together. Another membrane protein, **myelin basic protein (MBP)**, seals the cytoplasmic sides of the membranes in the myelin lamellae so that very little cytoplasm (with poor insulating properties) takes up space in the myelin sheath. Mice with a mutation of the *MBP* gene make abnormal myelin and develop serious movement disorders.

Myelination of the axons starts prenatally, but many neural pathways in the human are not fully myelinated until 2 years after birth (see Chapter 9, under “Myelination”). The process of myelination is closely related to functional maturation of the brain.

Nodes of Ranvier

Longitudinal views of axons show that the myelin sheath is interrupted at intervals, forming the **nodes of Ranvier** (Fig. 2.6A). The nodes of Ranvier exist because the glial cells forming myelin lie in a row along the axon, each cell making myelin only for a restricted length, or segment, of the axon. When viewed in the electron microscope, the axolemma (the axonal membrane) is “naked” at the node; that is, it is exposed to the ECF. Thus, only at the node of Ranvier can current in the form of ions pass from the axon to the ECF (and

in the opposite direction). This arrangement makes it possible for the nerve impulse to “jump” from node to node, thus increasing the speed of impulse propagation (discussed further in Chapter 3). The distance between two nodes of Ranvier in the PNS may be 0.5 mm or greater.

Multiple Sclerosis

In demyelinating diseases of the nervous system, the myelin sheaths degenerate. The most common of these diseases is **multiple sclerosis (MS)**, which typically manifests in young adults and usually has a long course of increasing disability. Its cause is still unknown, but most likely environmental factors precipitate an inflammatory process in individuals with a certain inherited susceptibility. Histopathologically, isolated and apparently randomly distributed regions of inflammation and demyelination are characteristic. In these regions, called **plaques**, impulse conduction in the axons is severely slowed or halted, and usually the symptoms are ascribed to the loss of myelin. For some reason, the optic nerve is often the first to be affected, resulting in reduced vision. Later symptoms that usually occur in varying proportions are muscle weakness, incoordination, and sensory disturbances. In most patients, exacerbations of the symptoms occur episodically in the beginning, associated with fluctuation in the inflammatory process.

Thus, periods of marked symptoms (such as paresis of extremities) are followed by periods of partial recovery. The improvement of symptoms is ascribed to partial remyelination of the affected regions. After a variable time (often many years), the disease becomes progressive, with a steady deterioration of the patient's condition.

There is not always a clear relationship between degree of demyelination and symptoms, suggesting that the disease process also directly harms axonal conductance and axonal viability. Indeed, it is now well established that in MS not only myelin sheets but also axons degenerate from the beginning of the disease. Presumably, the number of axons lost at early stages is modest and brain plasticity may compensate for their loss. As the disease progresses, however, the axon loss becomes so large that permanent and steadily progressing disability ensues.

Intense research activity is devoted to clarifying the etiology and pathogenesis of MS. Although clearly the disease process includes both inflammation and degeneration, it was long held that inflammation was the primary phenomenon (perhaps evoked by autoimmunity), and that loss of nervous tissue was secondary. This is now being questioned, however. Thus, it seems possible that "... people who develop multiple sclerosis

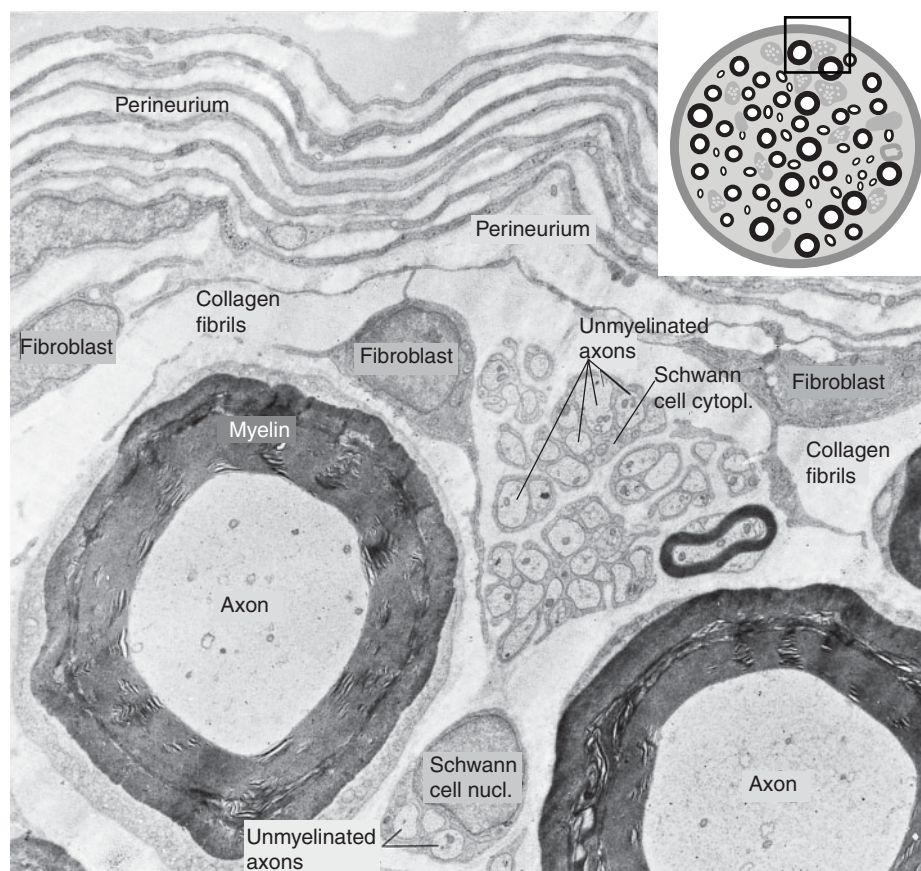
will be shown to have a (genetically determined) diathesis [disease disposition] that does indeed predispose to neurodegeneration ... but the exposure of that vulnerability requires an inflammatory insult without which the degenerative component does not manifest" (Compston, 2006, p. 563).

With regard to the inflammatory process, T lymphocytes, microglial cells, brain endothelial cells, and numerous immune mediators are involved, but their relative contributions are not fully understood. The role of microglia illustrates the complexity: they may contribute both to destruction of myelin and axons and to regenerative processes (such as remyelination), presumably depending on the local situation.

Unmyelinated Axons

As mentioned, unmyelinated axons conduct much more slowly (at less than 1 m/sec) than myelinated ones, because they are thinner and lack the extra insulation provided by the myelin sheath. In the CNS, unmyelinated axons often lie in closely packed bundles without any glial cells separating them (see Fig. 1.7). In the PNS, however, unmyelinated axons are always ensheathed in Schwann cells that do not make layers of myelin

FIGURE 2.8 *Peripheral nerve.* Electron micrograph of cross section of the sciatic nerve. The picture shows a small, peripheral part of a nerve fascicle. The perineurium surrounding the fascicle, is formed by several lamellae of flattened cells. Note the large difference in diameter among various myelinated axons. The thickness of the myelin sheath increases apace with the increase in axonal diameter. Between the myelinated axons are numerous unmyelinated ones. Collagen fibrils, produced by fibroblasts, fill most of the space between the axons. Magnification, $\times 4000$.



(Figs. 2.6–2.8). During early development, several axons become embedded in the cytoplasm of the Schwann cells by invagination of the Schwann cell membrane. This arrangement probably serves to protect the axon from harmful substances in the interstitial fluid. Such protection may not be necessary in the CNS, as the composition of the interstitial fluid is governed by astroglia cells and by the blood–brain barrier.

Peripheral Nerves Are Built for Protection of the Axons

Fresh nervous tissue is soft, almost jellylike, with virtually no mechanical strength in itself. Protection of the CNS against external mechanical forces is afforded by its location within the skull and the vertebral canal and by its “wrapping” in membranes of connective tissue (see Fig. 6.1). For peripheral parts of the nervous system, the situation is different. Often located superficially, the peripheral bundles of axons and groups of nerve cells are exposed to various mechanical stresses. They are also subject to considerable stretching forces by movements of the body. Axons can be stretched only slightly before their impulse conduction suffers, and they may even break. To prevent this, peripheral nerves contain large amounts of dense connective tissue with numerous collagen fibers arranged largely longitudinally (Fig. 2.7). The collagen fibers, specialized to resist stretching, protect the axons effectively. The presence of connective tissue in peripheral nerves is the reason that the nerves become much thicker where they leave the skull or the vertebral canal.

The connective tissue components of peripheral nerves form distinctive layers. The **epineurium** is an external thick layer of mostly longitudinally running collagen fibers. Internal to this layer, the axons are arranged into smaller bundles, or **fascicles**, which are wrapped in the perineural sheath or **perineurium** (Fig. 2.8). The collagen fibers and fibroblasts within the fascicles constitute the **endoneurium**. The perineurium is special in that it contains several layers of flattened cells. The cells, which in some respects resemble epithelial cells, interconnect by various kinds of junctions. In addition, the capillaries within the endoneurium are unusually tight and prevent passage of many substances from reaching the axons, consistent with experimental data showing that the perineurium constitutes a **blood–nerve barrier** preventing certain substances from reaching the interior of the fascicles with the axons. It is not surprising that PNS tissue also needs extra protective mechanisms to ensure that its environment is kept optimal for conducting impulses. The protection is not as efficient as in the CNS, however, and may perhaps explain why peripheral nerves are often subject to diseases that affect their conductive properties.

MICROGLIA AND REACTIONS OF THE CNS TO INJURY

Microglial Cells Are Phagocytes

The third kind of glial cell, **microglia**, is so named because of its small size. Studies with immunocytochemical identification of specific membrane proteins show unequivocally that microglial cells constitute a distinct kind. Estimates indicate that microglia may constitute 5% to 20% of all glial cells, being fairly evenly distributed through all parts of the CNS. Microglial cells are of **mesodermal** origin. Thus, animal experiments indicate that **monocytes** invade the nervous system from the bone marrow during embryonic development and perhaps shortly after birth. This may correspond with periods of high rate of cell death (a surplus of neurons is formed in early embryonic life, with subsequent elimination of a large number). After invading nervous tissue, the monocytes undergo changes—such as development of processes—that transform them to microglial cells, as identified in the adult. Nevertheless, microglial cells retain the typical phagocytic capacity of monocytes. Further, several surface markers (antigens) are common to blood monocytes and microglia, and cells that express such antigens first occur in the CNS (of rodents) in late embryonic development.

The number of microglial cells is relatively stable after the prenatal invasion. Under normal conditions, the stock of microglial cells does not appear to be supplemented from the bloodstream. After **injury**, the number of cells with phagocytic activity (macrophages) increases in the CNS. The increase appears to be due both to invasion of monocytes from the bloodstream and to activation of local microglial cells. The invasion of monocytes after injury probably depends on damage to the blood–brain barrier (i.e., brain capillaries allow passage of elements of the blood they normally restrict).

In the **normal brain**, microglial cells are probably not solely in a “resting” state in anticipation of challenges (e.g., intruding microorganisms, trauma, ischemia, and so forth). Thus, their processes are steadily moving and renewed, and are therefore believed to constantly “scan” their immediate environment for foreign material and sick or dead cellular elements. In addition, microglial cells are equipped with receptors for several neurotransmitters, suggesting that they also may sense the state of neuronal activity in their vicinity. If they detect something unusual, more microglial cells move quickly to the site. They release inflammatory mediators and phagocytose foreign or dead material. These responses of microglial cells generally serve to minimize damage and protect neurons; that is, microglial cells serve to conserve **homeostasis**. For example, animal

experiments show that the presence of microglial cells reduces ischemic brain damage (after loss of blood supply). Removal of dead material by microglia seems to be necessary for regeneration of neuronal processes to occur. Nevertheless, in certain diseases with strong activation of microglial cells (and astrocytes) they promote tissue injury rather than repair. Activation of microglial cells in the spinal cord also seems to contribute to persistence of **pain** after nerve damage.

Reaction of Nervous Tissue to Injury and Inflammation

Tissue damage leads to an inflammatory reaction in which the invasion and activation of immunocompetent cells have a central role. The purpose of the invasion is to kill microorganisms, remove debris, and aid reparative processes. However, the inflammatory reaction is different in the CNS than in other tissues. Thus, there is often no invasion of neutrophil granulocytes, and the activation of microglia and invading monocytes to macrophages may take several days. Overall, immune reactions are weaker and slower in the CNS than elsewhere. This may be explained—at least in part—by the lack of lymphatic drainage from the CNS. The immune system, therefore, does not possess much information about nervous tissue conditions, in contrast to tissues of most other organs. Normally, only a small number of T lymphocytes, entering from the bloodstream, patrol the CNS.³ Perhaps these special conditions are necessary to prevent neuronal damage from the potent substances that are liberated from granulocytes and activated macrophages. For example, edema—a central component of inflammation—may become harmful and even life threatening when it occurs in the brain (because of the limited possibilities of expansion within the skull). Nevertheless, immune reactions *do* occur in the brain, sometimes with serious consequences, as in multiple sclerosis.

The main task of **astrocytes** after injury is probably to strengthen their normal function of keeping the ECF composition constant. Tissue damage—regardless of whether it is caused by bleeding, contusion, or circulatory arrest—increases the flow of ions and transmitters from the neurons to the ECF. Astrocytes increase their uptake to counteract such disturbances of the neuronal environment. Because the substances taken up are

osmotically active, the astrocytes may swell quickly: seconds or minutes after the damage (if the normal uptake capacity is surpassed). This may contribute to brain edema, a dangerous complication of head injuries. In the long term, astrocytes produce a kind of scar tissue at sites where neurons are lost.

In Chapters 9 and 11, we discuss the plastic processes of the nervous system that permit functional recovery after injuries (such as stroke).

Diseases of Peripheral Nerves

Diseases involving degeneration of peripheral nerves are called **neuropathies** and in humans can have various causes. In any case, the symptoms are due to transitory or permanent loss of impulse conduction. Neuropathy is a well known complication of metabolic diseases such as diabetes but can also be caused by toxic substances (e.g., lead). Some neuropathies are due to attacks of the immune system on axons or myelin. This sometimes occurs after an infectious disease or in the course of cancer, probably because the immune system produces antibodies that cross-react with normal antigens expressed by axonal or Schwann cell membranes. Axons express some antigens that are specific to whether the axons are motor or sensory, thick or thin, and so forth. Thus, it may be understandable why neuropathies often affect certain nerves only or certain kinds of axons only. Thus, when motor axons are affected, the patient presents with pareses in certain muscles, while affection of sensory axons might produce loss of cutaneous sensation and joint sense. Neuropathies may also affect subgroups of sensory axons, for example, affecting only the very thin axons mediating sensations of pain and temperature but sparing axons related to touch. In other cases only axons mediating joint sense are affected, whereas cutaneous sensation is spared (examples are described in Chapter 13, under “Clinical Examples of Loss of Sensory Information”).

A large group of neuropathies is **inherited**, among them, **Charcot–Marie–Tooth disease** (peroneal muscle atrophy). In most cases, the disease is inherited dominantly. The disease usually starts before the age of 20 years and leads to gradually increasing pareses and sensory loss, starting distally in the legs. Loss of myelin and degeneration of axons cause the symptoms. Most patients with Charcot–Marie–Tooth disease have a doubling of the gene coding for the peripheral myelin protein (PMP-22). Animal models with overexpression of PMP-22 suggest that this defect alone can cause deficient myelination and symptoms corresponding to Charcot–Marie–Tooth disease in humans.

³ HIV (human immune deficiency virus) can enter the CNS via infected T lymphocytes. Microglial cells then become infected because they express surface receptors that bind the virus. After being infected, microglial cells secrete toxic substances that kill neurons, thus producing the neurological symptoms occurring in AIDS (acquired immune deficiency syndrome).

3 | Neuronal Excitability

OVERVIEW

In Chapter 1 we considered some of the characteristic properties of neurons, such as their excitability and their ability to conduct impulses. The term **excitability** means that when a cell is sufficiently stimulated, it can react with a brief electrical discharge, called an action potential. The **action potential** (the nerve impulse) travels along the axon and is a major component in the communication among nerve cells and between nerve cells and other cells of the body. The action potential results from movement of charged particles—ions—through the cell membrane. A prerequisite for such a current across the membrane is an electric potential—the **membrane potential**—between the interior and the exterior of the cell, and the presence of **ion channels** that are more or less selective for the passage of particular ions. The opening of ion channels is controlled by neurotransmitters binding to the channel (**transmitter** or **ligand-gated channels**) or by the magnitude of the membrane potential (**voltage-gated channels**). The membrane potential results from an unequal distribution of positively and negatively charged particles on either side of the membrane.¹ Energy-requiring **ion pumps** are responsible for maintaining the membrane potential. The **resting potential**, that is, the membrane potential when the neuron is not receiving any stimulation, is due mainly to unequal distribution of K^+ ions and the fact that the membrane is virtually impermeable to all ions other than K^+ in the resting state. The resting potential, with the interior of the cell negative compared with the exterior, is typically approximately -60 mV. The **action potential** is a brief change of the membrane potential, caused by opening of channels that allow cations (especially Na^+) to enter the neuron, followed by an outward flow of K^+ ions. A net influx of cations reduces the membrane potential by making the interior less negative. This is called **depolarization**, and if it is sufficiently strong, an action potential is elicited due to opening of voltage-gated Na^+ channels. After the brief depolarization caused by influx

of Na^+ ions, the membrane potential is restored by the outward flow of K^+ ions. Restoration of the membrane potential is called **repolarization**. An increase of the membrane potential—hyperpolarization—makes the neuron less excitable (more depolarization is necessary to elicit an action potential). In a short period after an action potential, the membrane is in a **refractory state**, which means that another action potential cannot be elicited. This ensures that neurons can maintain the correct ion concentration balance. Once an action potential is elicited, it is conducted along the axon. This is not merely a passive movement of charged particles in the fluid inside the axon. Because axons are poor conductors (compared with a metal thread), the action potential has to be renewed along the axonal membrane by cycles of depolarization and repolarization. In **unmyelinated** axons, these cycles move along the axon as a continuous wave, while in **myelinated** axons renewal of the action potential occurs only at the **nodes of Ranvier**. Because the process of depolarization–repolarization takes some time, the speed of conduction is very much slower in unmyelinated axons than in myelinated ones. The action potential, when first elicited, is of the same magnitude. Neurons are nevertheless able to vary their messages because of the varying frequency and pattern of action potentials. Generally, the more synaptic inputs depolarize a neuron, the higher will be the frequency of axonal action potentials.

BASIS OF EXCITABILITY

Cell Membrane Permeability Is Determined by Ion Channels

Ions cross the cell membrane almost exclusively through specific, water-filled channels because their electrical charges prevent them from passing through the lipid bilayer (Figs. 3.1 and 3.2). The channels are more or less **selective** for particular ions, that is, some ions pass more easily through a channel than others. Some channels are very selective, allowing passage of only one kind of ion (e.g., Na^+ ions), whereas other channels are less selective (e.g., letting through several cations such as Na^+ , K^+ , and Ca^{2+}). It follows that the ease with which an ion can pass through the membrane—that is,

¹ Neither membrane potentials nor action potentials are properties unique to nerve cells. All cells have a membrane potential, although usually of less magnitude than that of neurons. Muscle cells and endocrine gland cells also produce action potentials in relation to contraction and secretion, respectively.

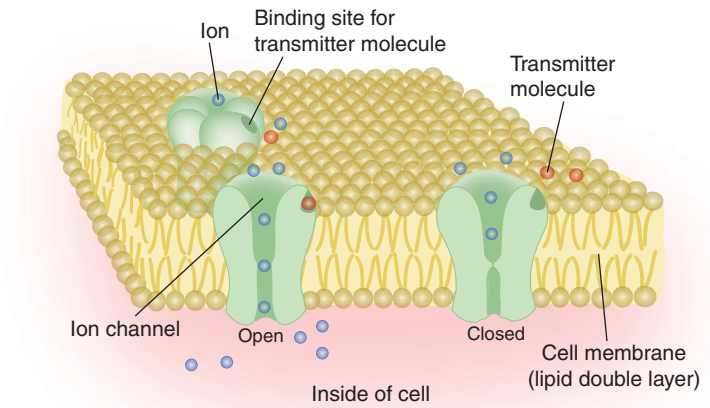


FIGURE 3.1 *Ion channels.* Schematic of a small part of the lipid bilayer of the cell membrane with interspersed ion channels. Binding of a transmitter molecule alters the opening state of the ion channel.

the **membrane permeability**² to that particular kind of ion—depends on (1) the presence of channels that let the ion through, (2) how densely these channels are distributed in the membrane, and (3) their opening state.

The current of ions through the membrane, however, does not depend solely on the density and opening of channels; an additional important factor is the **concentration gradient** across the membrane for the ion. That is, the steeper the gradient, the greater the flow of ions will be from high to low concentration (provided that the membrane is not totally impermeable to the ion). Further, because ions are electrically charged particles,

the **voltage gradient** across the membrane (i.e., the membrane potential) will also be important (Fig. 3.3). This means that if the interior of the cell is negative in relation to the exterior, the **cations** (positively charged ions) on the exterior will be exposed to a force that attracts them into the cell, while the interior cations will be subjected to forces that tend to drive them out. The strength of these attractive and expulsive forces depends on the magnitude of the membrane potential. Therefore, the concentration gradient and the membrane potential together determine the flow of a particular ion through the membrane (Fig. 3.3).

² The term **conductance** expresses the membrane permeability of a particular kind of ion more precisely. The conductance is the inverse of the membrane resistance. In an electrical circuit the current is $I = V/R$, where V is the voltage and R is the resistance (Ohm's law). This may be rewritten by using conductance (g) instead of R , as $I = g \cdot V$. In this way, one may obtain quantitative measures of membrane permeability under various conditions. For our purpose, however, it is sufficient to use the less precise term permeability.

The Membrane Potential

In a typical nerve cell, the potential across the cell membrane is stable at approximately 60 mV (millivolts) in the resting state, that is, as long as the cell is not exposed to any stimuli. We therefore use the term **resting potential**

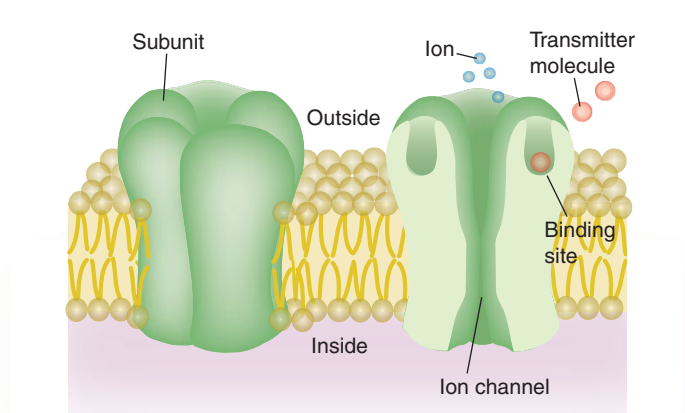


FIGURE 3.2 *Ion channels.* Five protein subunits are arranged around a central opening that can admit ions. At the outer side, the channel proteins are equipped with receptor sites for neurotransmitter molecules that regulate the opening of the channel. The figure shows the probable appearance of an acetylcholine receptor. (Based on Changeux 1993.)

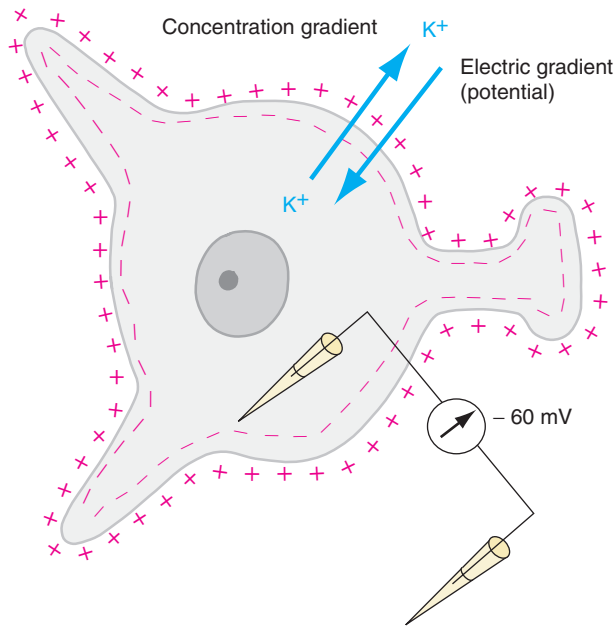


FIGURE 3.3 Forces acting on the K^+ ions. At the resting potential there is equilibrium between the inward and outward forces (large arrows) acting on the K^+ ions. One intracellular and one extracellular electrode (cones) measure the membrane potential.

in this situation (in different kinds of nerve cells, the resting potential may vary from about 45 mV to approximately 75 mV). The resting potential is due to a small surplus of negatively charged ions, **anions**, inside the cell versus the outside, and it has arbitrarily been decided to define the resting potential as negative, for example, -60 mV (Fig. 3.3).

The resting potential is caused primarily by two factors:

1. The **concentration of K^+ ions** is about 30 times higher inside than outside the cell (Figs. 3.4 and 3.5).
2. The cell membrane is **selectively permeable to K^+ ions** in the resting state (Fig. 3.5), that is, no other ions pass the membrane with comparable ease (the membrane, e.g., is about 50 times more permeable to K^+ than to Na^+).

Although the concentration differs greatly inside and outside the cell for ions other than K^+ (Fig. 3.4), the membrane is, as mentioned, almost impermeable to them (there are, e.g., very few open Na^+ channels in the resting state). Other ions therefore influence the resting membrane potential only slightly. Therefore, to explain the membrane potential we can, for the time being, ignore ions other than K^+ . The concentration gradient will tend to drive K^+ out of the cell, and further, K^+ ions can pass the membrane with relative ease through a particular kind of potassium channel that is open in the resting state. This means that positive charges are lost

from the interior of the cell, making the interior negative compared to the exterior, thereby creating a membrane potential. The membrane potential reaches only a certain value, however, because it will oppose the movement of K^+ ions out of the cell. Two opposite forces are at work: the concentration gradient tending to drive K^+ out of the cell and the electrical gradient (the membrane potential) tending to drive K^+ into the cell (Fig. 3.3). When the membrane potential is about -75 mV, these two forces are equally strong: that is, the flow of K^+ into the cell equals the flow out. This is therefore called the **equilibrium potential for K^+** , and its magnitude is determined by the concentration gradient for K^+ ions (the concentration gradient varies somewhat among neurons). The resting potential in most neurons, however, is lower than the equilibrium potential for K^+ because the cell membrane is slightly permeable to Na^+ (about 1/50th of the permeability to K^+). Therefore, some positive charges (Na^+) pass into the cell, driven by both the concentration gradient and the membrane potential, making the interior of the cell less negative than the equilibrium potential for K^+ . The membrane potential is consequently changed somewhat in the direction of the **equilibrium potential for Na^+** : that is, $+55$ mV. In the resting state, the inflow of positive charges is equal to their outflow, and the membrane potential is therefore stable. Even though the two opposite currents of K^+ and Na^+ are small, over time they would eliminate the concentration gradients across the membrane. This is prevented, however, by energy-requiring “pumps” in the cell membrane that actively transport ions through the membrane against a concentration gradient. This **sodium–potassium pump** expels Na^+ ions from the

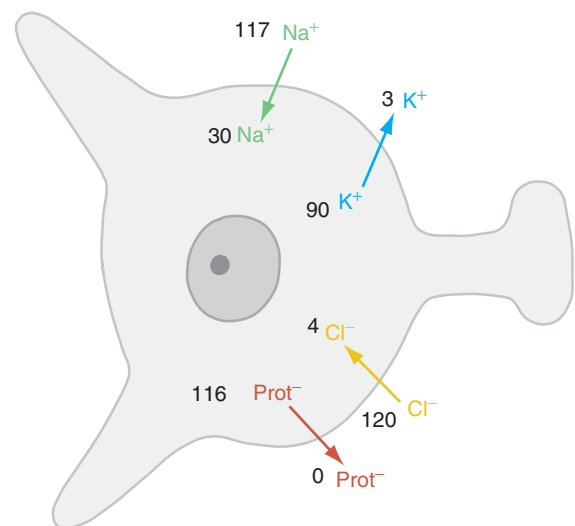


FIGURE 3.4 Distribution of ions of particular importance for the membrane potential. The exact concentrations depend on the resting potential (in this case -85 mV). Concentrations in mM.

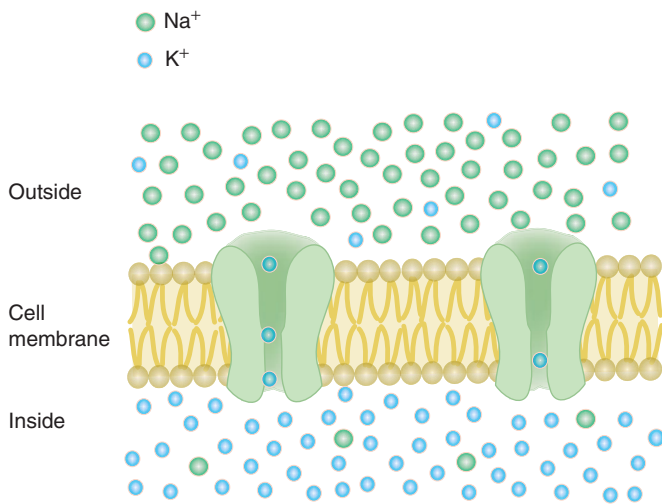


FIGURE 3.5 The unequal distribution of K^+ and Na^+ ions, together with open K^+ channels, largely explain the resting membrane potential.

interior, in exchange for K^+ , at the same rate that the ions leak through the membrane. In this way, the concentration gradients across the cell membrane are maintained.

Normally, the extracellular K^+ concentration is under tight control, as discussed in Chapter 2 (“Astroglia and the Control of Neuronal Homeostasis”). Such control is necessary because even small alterations in K^+ concentration influence the excitability of neurons significantly. For example, increased extracellular concentration moves more K^+ ions into the cell, thus depolarizing the neuron (making the membrane potential less negative) and lowering the threshold for eliciting action potentials.

Recording of Single-Cell Activity

Microelectrodes, with tips less than $1\ \mu\text{m}$ thick, can be used to record the activity of single neurons and their processes (single units) intracellularly. Among other things, this has made it possible to study in detail the electrical events at the synapses and how they are influenced by various experimental manipulations. The effects of different concentrations of intra- and extracellular ions have been studied, as have the synaptic effects of various transmitter candidates and drugs. The **voltage clamp** technique, which permits manipulation of the membrane potential, has been instrumental to our understanding of the properties of synapses and the basic mechanisms underlying their operations. Likewise, great progress has been made with the **patch clamp technique**, making possible measurements of ion currents limited to even a single ion channel. The study of the properties of ion channels and membrane receptors is today highly interdisciplinary. **Implanted extracellular**

electrodes can be used to record the activity of single neurons in relation to specific stimuli or behavioral tasks. This method has, for example, provided new insight into functional specializations within various areas of the cerebral cortex. By combining anatomic and physiological techniques, it has been possible to determine the functional properties of structurally defined cell types. After an intracellular recording has been made from a neuronal cell body or its axon, it can be filled with a tracer substance through the same pipette. Afterward, the neuron with all its processes can be visualized in sections.

Anions Are Also Unevenly Distributed

For simplicity, we have so far dealt with only two cations, K^+ and Na^+ , because they are the most important ones for the membrane potential and also for the action potential (discussed later in this chapter). Nevertheless, there are as many anions as cations. Chloride ions (Cl^-) and negatively charged protein molecules (Prot^-) are the major anions (Fig. 3.4). These ions are also unevenly distributed across the cell membrane: the concentration of Cl^- is 20 to 30 times higher outside than inside the cell, whereas the opposite situation exists for Prot^- . Therefore, **chloride** is the major extracellular anion, whereas **proteins** are the major intracellular ones. The proteins are so large that they cannot pass through the membrane; the membrane is impermeable to protein molecules. The membrane is somewhat permeable to Cl^- , however. The concentration gradient tends to drive chloride into the cell, whereas the membrane potential tends to drive it out, making the net flow of Cl^- small. In fact, the **equilibrium potential** for Cl^- , $-65\ \text{mV}$, is close to the resting potential of most nerve cells. Therefore, no active mechanism for pumping of chloride is needed.

The Sodium–Potassium Pump and Osmotic Equilibrium

All cells depend on the sodium–potassium pump to maintain the membrane potential and osmotic equilibrium between the intracellular and extracellular fluid compartments. Particular to neurons is their need for increased pumping in association with the firing of action potentials, which arise because of a current of Na^+ into the cell and of K^+ out of it. The speed of pumping increases with increasing intracellular Na^+ concentration. A significant part of our energy in the form of ATP is spent on driving the sodium–potassium pump. In the resting state of nerve cells, this may constitute approximately one-third of the total energy requirement, whereas after high-frequency trains of action potentials it may increase to two-thirds.

The unequal distribution of ions is of fundamental importance also for the ability of neurons to maintain

osmotic equilibrium. The distribution of ions must be such that the total concentrations of water-dissolved particles are equal inside and outside the cell. In other words, osmotic equilibrium means that the **water concentration** is equal inside and outside the cell (osmosis is the movement of water molecules from sites of high to sites of low water concentration). In case of osmotic imbalance, the cell will either swell or shrink (depending on whether the water concentration is lower inside or outside, respectively). An essential condition for osmotic balance is the low resting membrane permeability to Na^+ , as both the concentration gradient and the membrane potential tend to drive Na^+ into the cell. This situation changes dramatically when the cells fire action potentials, because the membrane then becomes highly permeable to Na^+ . Long trains of high-frequency action potentials may threaten the osmotic balance because more Na^+ ions enter the cell than can be pumped out. Fortunately, neurons have properties that limit their maximal firing rate and the duration of active periods. Under pathological conditions, however, these safeguards may fail. In severe **epileptic seizures**, for example, neurons fire with abnormal frequency for long periods, and this may probably contribute to cell damage by causing osmotic imbalance. Further, in situations with insufficient blood supply (ischemia), for example, after a **stroke**, ATP production suffers, resulting in slowing of the sodium–potassium pump. This, in turn, leads to osmotic imbalance and swelling of neurons. Such swelling is dangerous because neurons may be injured directly but also because swelling of the brain inside the skull (brain edema) reduces the blood supply.

Transmitter-Gated Ion Channels

Neurotransmitters control neuronal excitability by changing the opening state of ion channels (Figs. 3.1 and 3.2). A channel that is controlled by neurotransmitters (or other chemical substances) is called **transmitter-gated** or **ligand-gated** (the term “transmitter-activated” is also used). A large number of ion channels are now characterized that differ with regard to ion selectivity and transmitter specificity, that is, the ions that can pass a channel and the transmitter that controls it. The transmitter can either bind **directly** to the channel polypeptides (proteins) or act **indirectly** via chemical intermediates. In most known cases, the transmitter opens the channel to increase the permeability of the relevant ions. We consider here only the effects of directly acting neurotransmitters (indirect effects are discussed later in this chapter). Binding of a transmitter molecule to a specific **receptor site** at the external face of a channel polypeptide may change the form of the polypeptides, thereby changing the diameter of the channel (Figs. 3.1 and 3.2). Usually, the channel is open only briefly after binding of a transmitter molecule,

allowing a brief current of ions to pass through the membrane. In this way, a chemical signal from a presynaptic neuron—the neurotransmitter—elicits an electric current through the postsynaptic membrane.

As mentioned, ion channels are more or less **selectively permeable**, that is, they let certain kinds of ions pass through more easily than others. Some channels are highly selective, allowing the passage of one kind only (such as Ca^{2+} ions), whereas others are less selective and will allow passage of, for example, most cations. Channels that are permeable for anions in general are usually termed chloride (Cl^-) channels because Cl^- is the only abundant anion that can pass through the membrane. Size and charge of the ion influence its permeability. For example, the Na^+ ions are more hydrated (bind more water molecules) than the K^+ ion and therefore are larger (Fig. 3.5). This may explain some of their differences in permeability. By regulating the channel opening, the transmitter controls the flow of ions through the postsynaptic membrane. However, the transmitter only alters the **probability** of the channel being in an open state; it does not induce a permanent open or closed state.

Voltage-Gated Ion Channels

Many channels are not controlled primarily by chemical substances but by the magnitude of the membrane potential and are therefore called **voltage-gated**. Voltage-gated Na^+ and K^+ channels, for example, are responsible for the action potential and therefore also for the propagation of impulses in the axons. There are also several kinds of voltage-gated Ca^{2+} channels, which control many important neuronal processes, for example, the release of neurotransmitters. Voltage-gated channels are responsible for the activation of nerves and muscles by external electrical stimulation. Electrical stimulation of a peripheral nerve may produce muscle twitches by activating motor nerve fibers, as well as sensations due to activation of sensory nerve fibers.

The Structure of Ion Channels

The structure of several **ligand-gated** ion channels has now been determined. They consist of five polypeptide subunits arranged around a central pore. Three **families** of ligand-gated channels have been identified: the nicotinic receptor superfamily (GABA_A , glycine, serotonin, and nicotinic acetylcholine receptors), the glutamate receptor family, and the ionotropic ATP receptors. The subunits span the membrane and extend to the external and internal faces of the membrane (Fig. 3.2). Therefore, signal molecules inside the cell may also influence the opening of ion channels. As an example, members of the nicotinic receptor family consist of five equal subunits (Fig. 3.2), all contributing to the wall of the channel.

The subunits are large polypeptides with molecular masses of approximately 300,000. The transmitter binds extracellularly at the transition between two subunits but it is still unknown how the rapid binding (in less than 1 msec) produces conformational change in parts of the channel located, relatively speaking, far away.³

Voltage-gated channels resemble ligand-gated ones; they consist of four subunits arranged around a central pore. The amino-acid sequence has been determined for several of the subunits, although lack of three-dimensional data has prevented clarification of the mechanisms that control their opening and ion selectivity. Presumably, subtle differences between the subunits forming the channel explain their high selectivity to particular ions.

Inherited Channelopathies

Many different genes code for channel proteins. Because ion channels determine the excitability of neurons, it is not surprising that **mutations** of such genes are associated with dysfunctions of neurons and muscle cells. Common to many such **channelopathies** is that the symptoms occur in bouts. Of particular clinical interest is that many of the channelopathies affecting neurons are associated with **epilepsy**. Although channelopathies may not be the primary cause in the majority of patients with epilepsy, they may increase the susceptibility to other factors. For example, mutations associated with epilepsy affect ligand-gated channels that are receptors for the neurotransmitters γ -aminobutyric acid (**GABA**) and **acetylcholine**. Mutations affecting channels gated by **glycine** (an inhibitory transmitter) are associated with abnormal startle reactions. This may probably be related to the fact that glycine is preferentially involved in inhibition of motor neurons. Patients with a certain kind of headache—**familial hemiplegic migraine**—have a mutation of the gene coding for a specific Ca^{2+} -channel protein. Other mutations of the same gene are associated with other rare nervous diseases, for example, some that affect the cerebellum and lead to ataxic movements. Mutations of a kind of **voltage-gated potassium channel** ($\text{K}_v\alpha 1.1$)—expressed in highest density around the initial segment of axons—produce abnormal repolarization of motor axons and lead to repetitive discharges. This may explain the muscle cramps of such patients. Their episodes of ataxic movements are presumably caused by abnormal excitability of cerebellar neurons. Mutations of voltage-gated sodium channels (among other factors) cause bursts of intense pain (see also Chapter 15, under “Nociceptors, Voltage-Gated

Sodium Channels, and Channelopathies”). A number of mutations affect channels in **striated muscle membranes**, many of them associated with **myotonia** (inability to relax after a voluntary muscle contraction).

Different mutations of one gene can give different **phenotypes**, such as reduced density of channels or reduced opening probability. It is noteworthy, however, that the same mutation can produce different symptoms in different individuals, even within the same family. This strongly suggests that the genes coding for the proteins of a channel do not alone determine its final properties. Additional factors, such as the products of other genes and environmental factors, must also contribute. Many features of channelopathies are still unexplained—that they tend to occur episodically, that the symptoms often start at a certain age (in spite of the defect being present from birth), and that some forms remit spontaneously.

Alteration of the Membrane Potential: Depolarization and Hyperpolarization

As previously mentioned, in the resting state the membrane permeability for Na^+ is low. If for some reason Na^+ channels are opened so that the permeability is increased, Na^+ ions will flow into the cell and thereby reduce the magnitude of the membrane potential. Such a reduction of the membrane potential is called **depolarization**. The membrane potential is made less negative by depolarization. Correspondingly, one may predict that when the membrane permeability for K^+ is increased, more positive charges will leave the cell and the membrane potential will become more negative than the resting potential. This is called **hyperpolarization**. The same would be achieved by opening channels for chloride ions, enabling negative charges (Cl^-) to flow into the cell, provided that the membrane potential is more negative than the resting potential of Cl^- .

In conclusion, the membrane potential is determined by the **relative permeability** of the various ions that can pass through the membrane. At rest, the membrane is permeable primarily to K^+ , and the resting potential is therefore close to the equilibrium potential of K^+ . Synaptic influences can change this situation by opening Na^+ channels, thereby making the permeability to Na^+ dominant. This changes the membrane potential toward the equilibrium potential of Na^+ (at 55 mV). As shown in the following discussion, the action potential is caused by a further, sudden increase in the Na^+ permeability.

Markers of Neuronal Activity

Several methods can be used to visualize the activity of neurons. One method involves intracellular injection of a **voltage-sensitive fluorescent dye**. The intensity of fluorescence (as recorded with fluorescence microscopy

3 The binding of the transmitter most likely elicits a wave of conformational change in specific parts of the channel polypeptides. The actual opening of the channel may be caused by conformational change of just one, specific amino acid.

and advanced computer technology) gives an impression of neuronal activity at a given time. Thus, this (indirect) measure of activity can be correlated with experimental manipulation of a specific transmitter, the execution of specific tasks, and so forth. Another method takes advantage of the fact that **optic properties** of nervous tissue change with the degree of neuronal activity. This enables the recording of slow as well as rapid changes in neuronal activity in relation to experimental influences (it has been applied, e.g., in conscious persons during neurosurgery that necessitates exposure of the cerebral cortex). Other methods enable mapping of variations in neuronal activity at the time of death in experimental animals. Intravenously injected radiolabeled **deoxyglucose** is taken up by cells in the same way as glucose. It is not broken down, however, and therefore accumulates in the cells. Because glucose is the substrate for oxidative metabolism in the neurons, its uptake correlates with degree of neuronal activity. After exposing an animal to certain kinds of stimulation or eliciting certain behaviors, one can afterwards determine with autoradiography which neuronal groups were particularly active during stimulation or at the time of certain actions. Another method utilizes the fact that a few minutes with excitatory synaptic input induces expression of so-called **immediate early-genes** in many neurons. Most studied among such genes is *c-fos*. Without extra stimulation, *C-fos* mRNA and its protein product are present in only minute amounts in most neurons. Detection of increased levels of *c-fos* mRNA in tissue sections is therefore used as a marker of neurons that were particularly active in a certain experimental situation. This method is also used to determine where in the brain a drug exerts its effect. The method has its limitations, however. Thus, *c-fos* expression may be caused by nonspecific influences, and not all neurons express *c-fos* even when properly activated.

THE ACTION POTENTIAL

Voltage-Gated Sodium Channels Are Instrumental in Evoking an Action Potential

The basis of the action potential is found in the presence of **voltage-gated Na^+ channels**, which are opened by depolarization of the membrane (Fig. 3.6). Depolarization may be induced in several ways; for example, under artificial conditions by direct electrical stimulation. Normally, however, it is caused by neurotransmitters acting on transmitter-gated channels. The opening of transmitter-gated Na^+ channels often starts depolarization. Opening of the voltage-gated channels requires that the membrane be depolarized to a certain **threshold** value, that is, the threshold for

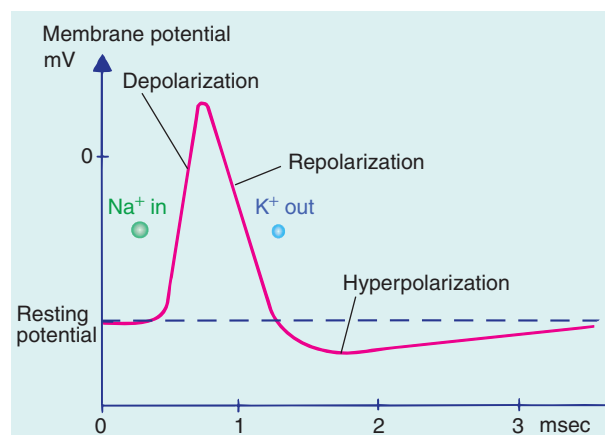


FIGURE 3.6 The action potential.

producing an action potential (Fig. 3.6). When voltage-gated channels are opened, the permeability to Na^+ is increased beyond what was achieved by the opening of transmitter-gated channels, and Na^+ flows into the cell driven by both the concentration gradient and the membrane potential. The membrane becomes more depolarized; in turn, this opens more voltage-gated channels, and so on. In this way, as soon as the membrane is depolarized to the threshold value, the permeability to Na^+ increases in an explosive manner. Even with all sodium channels fully open, however, the inward current of Na^+ ions stops when the membrane is depolarized to +55 mV; at that value the inward concentration force is equal to the outward electrical force (the membrane potential). As mentioned, +55 mV is the equilibrium potential of Na^+ . Figure 3.6 shows how, during an action potential, the membrane potential quickly changes to positive values and then returns almost as rapidly to approximately the resting value. This occurs because the membrane again becomes impermeable to Na^+ ; the Na^+ channels are closed or **inactivated**.⁴ Therefore, at the peak of the action potential and for a short time afterward, no Na^+ can pass through the membrane. In this situation with a positive membrane potential, K^+ is driven out by both the concentration gradient and the membrane potential (electrical force). Because no Na^+ can enter the cell, there is a net outward flow of positive charges, again making the interior of the cell negative. We say that the membrane is **repolarized**. The speed of repolarization is increased by the presence of **voltage-gated K^+ channels**, which open when the membrane is sufficiently depolarized. The opening of the voltage-gated K^+ channels is somewhat

⁴ **Inactivation** and closure involve different parts of the voltage-gated Na^+ channel. This is indicated by, among other findings, that whereas closure of the channel lasts as long as the membrane potential remains below threshold, inactivation is transitory and lasts only some milliseconds.

delayed compared with the Na^+ channels, but whereas the Na^+ channels inactivate after about 1 msec, the K^+ channels stay open for several milliseconds.

In summary, the action potential is caused by a brief inward current of Na^+ ions, followed by an outward current of K^+ ions. The whole sequence of depolarization–repolarization is generally completed in 1 to 2 msec. If the threshold is reached, an action potential of a certain magnitude arises, regardless of the strength of the stimulus that produced the depolarization.

Where Does the Action Potential Arise?

The action potential usually arises in the first part of the axon, the **initial segment** (Fig. 3.7; see also Fig. 2.6), where the density of voltage-gated Na^+ channels is higher than in the membrane of the dendrites and the cell soma. The current spreads electrotonically (passively) from dendritic and somatic synapses toward the initial segment. If the depolarization is sufficiently strong (reaches threshold), voltage-gated Na^+ and K^+ channels open and produce an action potential that is propagated along the axon. Although action potentials can be elicited in dendrites, their threshold is usually much higher than in the initial segment owing to lower density of voltage-gated channels.

The Action Potential and Changes of Ion Concentrations

One might think that an action potential would cause significant changes in the concentrations of Na^+ and K^+ on the two sides of the membrane, but this is not the case. The number of ions actually passing through the membrane during an action potential is extremely small compared with the total number inside the cell and in its immediate surroundings. Even in an axon with a diameter of about 1 μm , with a very small intracellular

volume compared to the membrane surface area, only 1 of 3000 K^+ ions moves out during the action potential. In addition, active pumping (the sodium–potassium pump) ensures that Na^+ is moved out and K^+ is moved in between each action potential and during periods of rest. Even when the sodium–potassium pump is blocked experimentally, a nerve cell can produce several thousand action potentials before concentration gradients are reduced so much that the cell loses its excitability.

The Refractory Period

After an action potential, some time must elapse before the neuron can again produce an action potential in response to a stimulus. The cell is said to be in a **refractory state**. This ensures at least a minimal rest for the cell between each action potential and thereby puts an upper limit on the frequency with which the cell can fire. The length of the refractory period, and therefore also the maximal frequency of firing, varies considerably among different kinds of nerve cells.

Two conditions are responsible for the refractory period. One is the aforementioned inactivation of the voltage-gated Na^+ channels, and the other is the fact that the membrane is hyperpolarized immediately after the action potential (Fig. 3.6). The inactivation of Na^+ channels means that they cannot be opened, regardless of the strength of the stimulus and the ensuing depolarization. Hyperpolarization occurs because the K^+ channels remain open longer than required just to bring the membrane potential back to resting value. These two different mechanisms can account for why the refractory period consists of two phases. During the first phase, the **absolute refractory period**, the cell cannot be made to discharge, however strong the stimulus may be; during the **relative refractory period**, stronger depolarization than normal is needed to produce an action potential.

Calcium and Neuronal Excitability

A cation other than Na^+ —namely, Ca^{2+} —may also contribute to the rising phase of the action potential. For Ca^{2+} , as for Na^+ , the extracellular concentration is much higher than the intracellular one, and there are voltage-gated calcium channels in the membrane.

Cellular influx of calcium can be visualized after intracellular injection of a substance that fluoresces when Ca^{2+} binds to it. During the action potential, calcium enters the cell—partly through Na^+ channels and partly through voltage-gated calcium channels, which have a more prolonged opening–closing phase than the sodium channels. There are also transmitter-gated calcium channels. In most neurons, the contribution of Ca^{2+} to the action potential is nevertheless small compared with that of Na^+ . In certain other cells such as

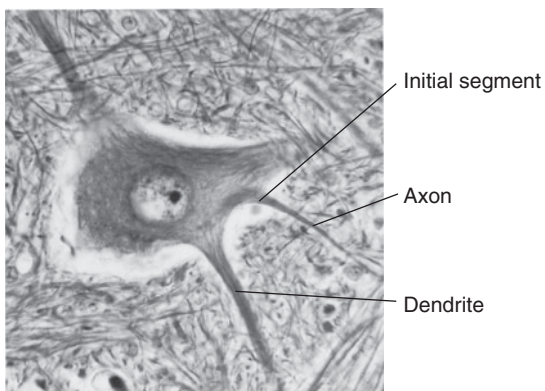


FIGURE 3.7 The initial segment of the axon is where the action potential usually arises. Photomicrograph of a motoneuron from the spinal cord stained with a silver-impregnation method.

heart muscle, however, calcium is the ion largely responsible for the action potential. Because the calcium channels open and close more slowly than the Na^+ channels, an action potential produced by calcium currents lasts longer than one produced by flow of Na^+ .

Another aspect of the functional role of calcium is that the extracellular calcium concentration influences the membrane excitability, which is most likely mediated through effects on the Na^+ and K^+ channels. Reducing the calcium concentration in the blood and interstitial fluid—**hypocalcemia**—lowers the threshold for evoking action potentials in neurons and muscle cells, whereas increasing the concentration—**hypercalcemia**—has the opposite effect. A typical symptom of hypocalcemia is muscle spasms—**tetany**—due to hyperexcitability of nerves and muscles. Severe hypercalcemia can cause drowsiness, nausea, and anorexia.

IMPULSE PROPAGATION

Electrical Properties of Axons

We now consider how the action potential moves along the axon. The ability of the axon to conduct electrical current depends on several conditions, some of which are given by the physical properties of axons, which are very different from, for example, those of copper wire. In addition, some conditions vary among axons of different kinds. An axon is a poor conductor compared with electrical conductors made of metal because the axoplasm through which the current has to pass consists of a weak solution of electrolytes (i.e., low concentrations of charged particles in water). In addition, the diameter of an axon is small (from <1 to $20\ \mu\text{m}$) with a correspondingly enormous **internal resistance** to the current in the axoplasm. Further, the axonal membrane is not a perfect insulator, so that charged particles are lost from the interior of the axon as the current passes along its length. The amount of current being lost is determined by the degree of **membrane resistance** (i.e., the resistance of the membrane to charged particles trying to pass). Finally, the axonal membrane (like all cell membranes) has an **electrical capacity**; that is, it can store a certain number of charged particles in the same way a battery does. This further contributes to the rapid attenuation of a current that is conducted along an axon: the membrane has to be charged before the current can move on.

The Action Potential Is Regenerated as It Moves Along the Axon

From the foregoing it can be concluded that how well the current is conducted in an axon depends on its internal

resistance (its diameter), the membrane resistance (how well insulated it is), and the capacity of the axonal membrane. If the propagation of the action potential along the axon occurred only by passive, electrotonic movement of charged particles, the internal resistance and loss of charges to the exterior would cause the action potential to move only a short distance before it “died out.” The solution to this problem is that the action potential is **regenerated** as it moves along the axon. Therefore, it is propagated with undiminished strength all the way from the cell body to the nerve terminals. As discussed, the strength of the action potential—that is, the magnitude of the changes of the membrane potential taking place—is the same regardless of the strength of the stimulus that produced it (as long as the stimulus depolarizes the membrane to threshold). Thus, increasing the strength of the stimulus increases the frequency of action potentials, whereas the magnitude of each action potential remains constant.

When the cell membrane at the initial segment (Fig. 3.7) is depolarized to threshold, an action potential is produced and is conducted passively a short distance along the axon. From then on, what occurs differs somewhat in myelinated and unmyelinated axons (Figs. 3.8 and 3.9).

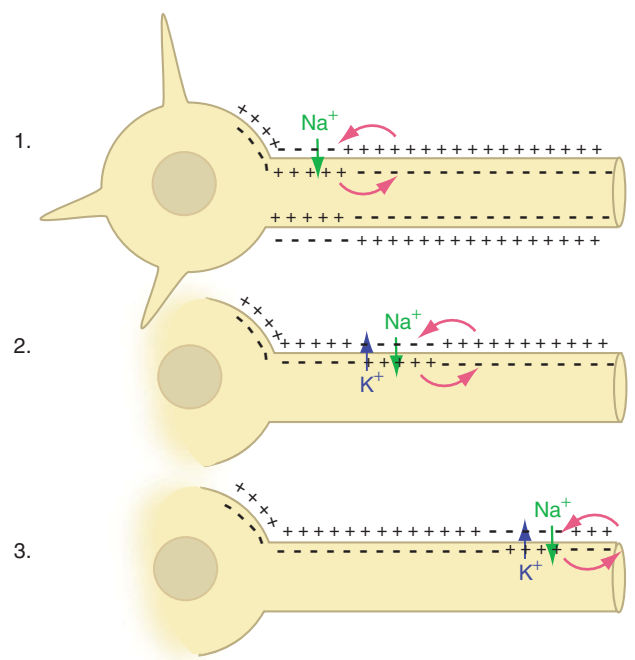


FIGURE 3.8 Impulse conduction in unmyelinated axons. Arrows show direction of movement of charged particles. The action potential is renewed continuously along the axonal membrane by a wave of depolarization–repolarization.

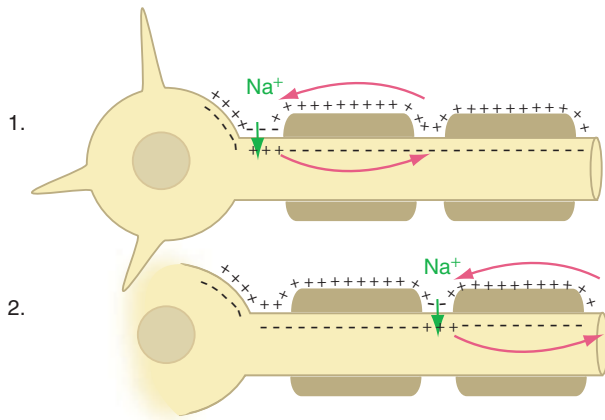


FIGURE 3.9 Impulse conduction in myelinated axons. Arrows show direction of movement of charged particles. The current moves electrotonically in the myelinated part of the axons, and the action potential is renewed only at the node of Ranvier, causing a small delay in impulse propagation.

Impulse Conduction in Unmyelinated Axons

The action potential is produced by positive charges penetrating to the interior of the axon, which at that point becomes positive relative to more distal parts along its length (Fig. 3.8). Positive charges then start moving in the distal direction (along the electrical gradient that has been set up). Outside the axon, a corresponding current of positive charges moves in the opposite direction, so that an **electrical circuit** is established. Movement of positive charges in the distal direction inside the axon means that the membrane is depolarized as the charges move along. This depolarization leads to the opening of enough voltage-gated Na⁺ and K⁺ channels to produce a “new” action potential. In this manner, the action potential moves along the axon at a speed that depends on the speed with which the charged particles (i.e., ions) move inside the axon and on the time needed for full opening of the ion channels. The membrane capacity represents a further factor slowing the propagation because the membrane has to be charged before there can be a net flow of charges through it.

In essence, the action potential is propagated as a wave of depolarization, followed closely by a corresponding wave of repolarization. When the membrane has just completed this cycle, it is in the refractory state for some milliseconds. This delay prevents the action potential from spreading “backward” toward the cell body (**antidromic** impulse conduction), and ensures that under normal conditions the impulse conduction is unidirectional. If, however, the axon is artificially stimulated (e.g., electrically) at some distance from the cell

body, the action potential spreads toward both the cell body and the end ramifications (**orthodromic** impulse conduction). Antidromic impulse conduction may occur in branches of peripheral sensory axons on natural stimulation and may play a part in certain disease symptoms (see Chapter 29, “Antidromic Impulses and the Axonal Reflex”).

Impulse Conduction in Myelinated Axons

In myelinated axons, the action potential is also regenerated along the axon (Fig. 3.9). However, in contrast to that in unmyelinated axons, the action potential is regenerated only at each **node of Ranvier**—that is, where the axon membrane lacks a myelin covering and is in direct contact with the extracellular fluid (see Fig. 2.6). As in unmyelinated axons, the action potential arises in the initial segment of the axon. The current then spreads passively (electrotonically) to the first node of Ranvier. Here, the depolarization of the membrane leads to opening of voltage-gated channels and a “new” action potential. The density of voltage-gated sodium channels is particularly high in the axonal membrane at the node of Ranvier. The current can flow electrotonically as far as the first node of Ranvier (and probably sometimes farther) because the axon is so well insulated by myelin, preventing loss of charges from the interior of the axon. (Myelin dramatically increases the resistance across the membrane and also reduces the membrane capacity.) In addition, the axonal diameter is larger in myelinated than in unmyelinated axons, thus reducing the internal resistance.

In conclusion, in myelinated axons the action potential does not move smoothly and slowly along, as in unmyelinated axons, but instead “jumps” from one node of Ranvier to the next. Although the impulse propagation is very rapid between nodes, at each node there is a delay due to the time required for opening of channels and establishment of sufficient flow of current.

Conduction Velocities in Myelinated and Unmyelinated Axons

The main reason myelinated axons conduct so much more rapidly than unmyelinated ones is that the action potential needs to be regenerated only at certain sites. A figure for conduction velocity (expressed in meters per second) in myelinated axons is obtained by multiplying the axonal diameter (in micrometers) by 6. An axon of 20 μm (the maximal diameter) therefore conducts at approximately 120 m/sec, whereas the thinnest myelinated axons of about 3 μm conduct at 18 m/sec. In comparison, a typical unmyelinated axon of about 1 μm conducts at less than 1 m/sec.

HOW NERVE CELLS VARY THEIR MESSAGES

Frequency Coding and Firing Patterns

So far we have treated the action potential as a unitary phenomenon. As mentioned, the strength of each action potential of a neuron does not vary: whenever depolarized to the threshold, the cell fires action potentials of constant magnitude. Therefore, the action potential is an **all-or-none** phenomenon, and one might think that each neuron would only be able to tell whether or not a stimulus occurs. We know, however, that the individual nerve cell can communicate to others about the strength of the stimulation it receives—such as the intensity of light or a sound, of something touching the skin, and so forth. It does so by varying the frequency and pattern of action potentials. To understand this, we need to know that a neuron is more or less continuously influenced by impulses from many other neurons. A sustained synaptic input that is strong enough to depolarize the cell to threshold does not merely produce one action potential but rather several in succession. The stronger the depolarization, the shorter the time required reaching the threshold after each action potential. Consequently, the firing frequency depends on the strength of depolarization (Fig. 3.10). We say that the neuron uses a **frequency code** to tell how strongly it has been stimulated. The **maximal frequency** of action potentials in some neurons is more than 100 per second (100 Hz), whereas in others it is much lower.

The average firing frequency is not the only way by which the neuron can alter its message. The **firing pattern** also carries information, and each neuronal type has its characteristic firing pattern that is caused by differences in membrane properties and synaptic

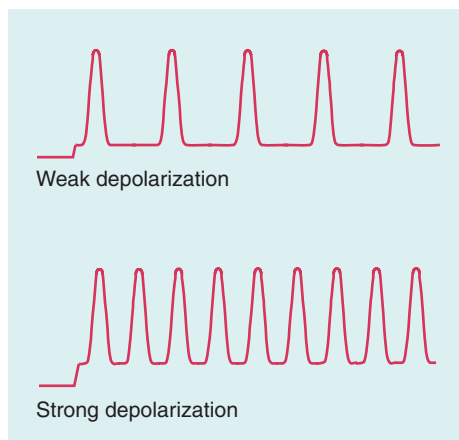


FIGURE 3.10 The frequency of action potentials depends on the magnitude of depolarization. Therefore, the frequency of action potentials reflects the total synaptic input to a neuron.

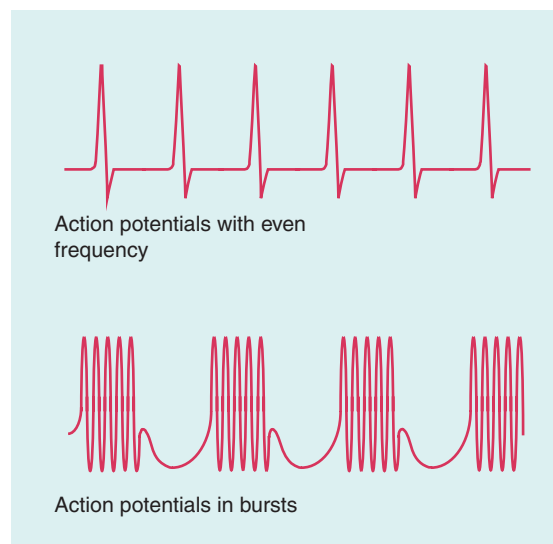


FIGURE 3.11 Different patterns of nerve impulses provide neurons with an additional means to vary the information they send to other neurons and muscle cells.

inputs (see earlier, “The Refractory Period”). Two neurons may both fire with an average frequency of, for example, 10 per second but nevertheless influence a postsynaptic cell differently. So-called **burst neurons** produce trains of action potentials with a high frequency and then pause for a while before a new train (burst) of impulses arises. Other neurons—so-called **single-spike neurons**—produce action potentials with regular intervals (Fig. 3.11).

Some neurons can switch between these two modes of firing. In such cases, the relationship is not linear between the strength of synaptic input and the firing frequency. The transition between the different firing patterns is evoked by a specific neurotransmitter, which does not in itself produce action potentials in the postsynaptic cell but changes its reactions to other inputs. For example, the neuron may change from burst to single-spike patterns or from a high firing frequency to no firing at all.

In conclusion, because the strength of action potential is constant, the code for the information carried by an axon is the frequency and pattern of action potentials.

Plateau Potentials

In some neurons, the occurrence of so-called plateau potentials causes the switch from low-frequency firing to high-frequency or bursting firing pattern. This has been shown for many neurons that control rhythmic muscle contractions. Plateau potentials are produced by a slow, depolarizing current, for example, by certain

voltage-gated Ca^{2+} channels that are open in a limited range of membrane potentials. Such a neuron can therefore change abruptly between two entirely different behaviors. The neurotransmitter **serotonin** can evoke plateau potentials in groups of spinal motor neurons

(see Chapter 21, under “Muscle Cramps and Plateau Potentials,” and Chapter 22, under “Monoaminergic Pathways from the Brain Stem to the Spinal Cord”). Release of this transmitter relates to motivation and attention rather than to specific information.

4 | Synaptic Function

OVERVIEW

In Chapter 3, we discussed the basis of nerve impulses and their conduction in axons. This chapter deals with the properties of synapses. We discuss mainly **chemical synapses**: synapses in which the signal is mediated by a neurotransmitter. Synapses with direct **electric coupling** (gap junctions) are common among glial cells but occur infrequently among neurons. The key events underlying signal transfer at chemical synapses are as follows: First, an action potential reaches the nerve terminal (bouton) and **depolarizes** it. This depolarization opens **Ca²⁺ channels**, enabling Ca²⁺ to enter the nerve terminal. Increase in intracellular Ca²⁺ concentration is a signal for release of **neurotransmitter** from vesicles by exocytosis. This produces a high concentration of neurotransmitter in the synaptic cleft. The released transmitter binds briefly to **receptors** in the **postsynaptic membrane**. After activation of the receptor, the transmitter must be **inactivated** quickly to reestablish a low background activity of the receptors, that is, to ensure a high **signal-to-noise ratio** at the synapse. Inactivation occurs partly by diffusion of the transmitter, partly by **enzymatic degradation**, and partly by specific uptake mechanisms (**transporter proteins**).

There are two main kinds of transmitter receptors. **Ionotropic receptors** are parts of ion channels and therefore influence the functional state of the channel directly. Therefore, transmitter actions elicited by ionotropic receptors are fast and precise. **Metabotropic receptors** are coupled indirectly (via intracellular second messengers) to ion channels. Their effects are therefore slower to start and longer lasting than effects mediated by ionotropic receptors. We also use the term **modulatory** of the synaptic effects of metabotropic receptors, because they adjust the excitability of the postsynaptic neuron so that it responds more or less vigorously to the precise effects of ionotropic receptors (in addition, metabotropic receptors may have effects on the growth and survival of the postsynaptic neuron).

The change of the membrane potential arising as a result of synaptic influence is called a **synaptic potential**. If the synaptic influence depolarizes the postsynaptic cell, the probability that the cell will fire action potentials is increased. This synaptic effect is called an **excitatory postsynaptic potential (EPSP)**. If the synaptic potential hyperpolarizes the cell, it is called an inhibitory postsynaptic potential (IPSP) because the probability of

the cell's firing is diminished. If the transmitter produces an EPSP, we use the terms **excitatory synapse** and **excitatory transmitter**. Likewise, an **inhibitory transmitter** produces an IPSP at an **inhibitory synapse**.

Because the depolarization caused by one EPSP is small, **summation** of many EPSPs is usually needed to reach a threshold for eliciting an action potential. This enables the neuron to integrate information from often many thousand synapses.

Synapses are **plastic**; that is, they can change their properties by use. This implies that certain kinds of activity can enhance or reduce the subsequent effect of a synapse on the postsynaptic neuron for a variable period (from milliseconds to years). Most likely, such **use-dependent** synaptic plasticity is the neuronal basis for **learning** and **memory**.

Unusual Synapses: Electrotonic and Dendrodendritic Transmission

Although it is rare, the pre- and postsynaptic elements are electrically rather than chemically coupled at some synapses. Electron microscopically, such electronic synapses differ from chemical synapses in that the synaptic cleft is only 2 nm compared to about 20 nm. This kind of cell contact is called a **nexus** or **gap junction**; it consists of channels that span the synaptic cleft. Through these channels ion currents can pass directly and quickly from one cell to another with no synaptic delay. In invertebrates and lower vertebrates, electrotonic synapses are formed between neurons mediating short-latency responses to stimuli (e.g., escape reactions). Electrotonic synapses may also provide electrical coupling between many neurons in a group, so that their activity may be **synchronized**. Chemical synapses may occur close to electrical ones and serve to uncouple the electrical synapse so that these apparently can be switched on and off. Even a small number of gap junctions between nerve cells—too small to produce efficient electric coupling—may be important by enabling transfer of small signal molecules, such as Ca²⁺, cyclic AMP, and inositol triphosphate (IP₃). In this way, one neuron may alter the properties of another without ordinary synaptic contact. Electrical coupling by gap junctions is much more common among glial cells than among neurons, and it occurs regularly among cardiac, smooth-muscle, and epithelial cells. There are also other unusual types of synapses. Contacts between dendrites

with all the morphological characteristics of synapses have been observed in several places in the central nervous system (CNS). Such dendrodendritic synapses are often part of more complex synaptic arrangements. Through dendrodendritic synapses, adjacent neurons can influence each other without being connected with axons. The function of such synapses, however, is not fully understood.

NEUROTRANSMITTER HANDLING AT THE SYNAPSE

Release of Neurotransmitters

We have previously described transmitter-containing synaptic vesicles, aggregated near the presynaptic membrane of boutons (Figs. 4.1 and 5.9). Depolarization of the presynaptic membrane by an action potential is the normal event preceding transmitter release. The depolarization opens voltage-gated calcium channels and allows a flow of Ca^{2+} ions into the bouton. The rise in Ca^{2+} concentration triggers the release of transmitter by exocytosis of vesicles (Fig. 4.1). The more calcium that enters, the more transmitter is released. By exocytosis, the membrane of synaptic vesicles fuses with the presynaptic membrane. The fusion opens the vesicle so that its content flows quickly into the cleft (Figs. 4.1 and 4.2). It takes only 0.1 to 0.2 msec from calcium inflow to the occurrence of release, which means that only vesicles already attached to the presynaptic membrane empty their contents. Further, although voltage-gated Ca^{2+} channels are present in all parts of the nerve terminal membrane, only those situated in the presynaptic membrane can influence the fusion of the vesicle with the presynaptic membrane.¹ The part of the synapse where the vesicles attach to the presynaptic membrane is called the active zone, and is characterized by cytoskeletal components that probably bind the vesicles to the calcium channels. That fusion really occurs during release is supported by, among other data, electron microscopic observations showing that the number of vesicles drops with long-term stimulation (trains of action potentials), while the number increases after a period of rest.

Exocytosis of vesicles is controlled by a large number of regulatory proteins that appear to be the same in all kinds of cells. Two features are nevertheless specific to exocytosis in neurons as compared with that in other cells: one is the speed of the process (<1 msec from arrival of the action potential to release); the other is that the release is restricted to a specific site (the synapse).

¹ This is because the fusion requires a very high concentration of Ca^{2+} , which occurs only close to the intracellular opening of the channel. In fact, there is evidence that the calcium channel constitutes a part of the protein complex that binds the vesicle to the presynaptic membrane. This ensures maximal Ca^{2+} concentration around the vesicle.

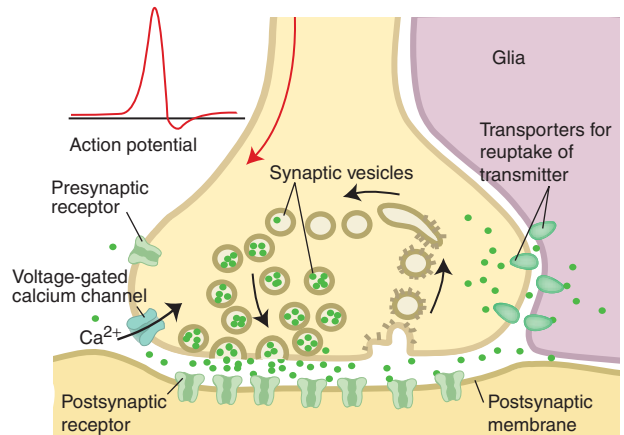


FIGURE 4.1 *Signal transmission at the synapse.* Schematic of some important features: calcium-dependent transmitter release, reuptake of transmitter by glia and neurons, and recycling of synaptic vesicles.

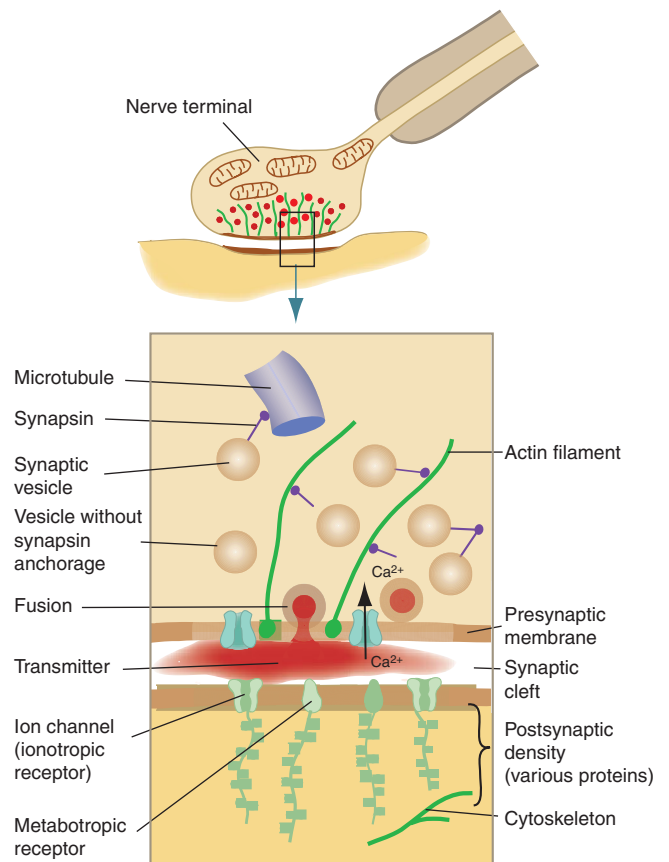


FIGURE 4.2 *Transmitter release and some of its machinery.* Calcium channels are located close to where the vesicles fuse with the presynaptic membrane. (Based on Walmsey et al. 1998.)

This indicates that some proteins are specific to the control of exocytosis in neurons. The fusion requires specific binding of vesicle-surface receptors to receptors in the presynaptic membrane. In addition, during fusion, various proteins dissolved in the cytoplasm participate by binding to the membrane-bound receptors, thus forming large complexes.

New, empty vesicles are formed by the opposite process of exocytosis, **endocytosis**. The endocytotic vesicles are **coated** with proteins (among them **clathrin** and **dynamin**) that are thought to help in budding of the vesicles from the membrane and in selecting their content. The recycled vesicles undergo a series of regulated steps until they are again filled with neurotransmitter (Fig. 4.1).

Several of the proteins involved in vesicle transport and fusion alter their activity in a use-dependent manner; that is, they may be involved in **synaptic plasticity** during development, recovery after brain damage, and learning in general. Some are also targets of drugs and toxins.

Mechanisms for Vesicle Transport and Fusion

Specific **transporter proteins** in the vesicle membrane fill the vesicles with neurotransmitter. After filling, the vesicles are moved toward the presynaptic membrane by a regulated process (Fig. 4.2). While some vesicles empty their contents, others move toward the presynaptic membrane and prepare for fusion. The synaptic vesicles can therefore be divided into two main groups: those situated close to the membrane that are ready for release when the Ca^{2+} concentration rises around them, and those that must move to the membrane before they can release their contents. The movement of vesicles requires the presence of **actin** filaments, and **microtubules** may also play a role. A group of proteins, **synapsins**, bind the vesicles to the actin filaments (Fig. 4.2), which probably serves to assemble the vesicles in positions for further movement and is triggered by the rise in the calcium concentration. Certain **protein kinases** (phosphorylating proteins) regulate the activity of the synapsins. Phosphorylation of synapsins increases mobility of the vesicles and is most likely another way of controlling the amount of transmitter released by an action potential, for example, in response to altered use of the synapse. Several proteins take part in the **docking** of the vesicle at the presynaptic membrane, and they probably also prepare the vesicles for fusion. Vesicle-bound receptors, such as **synaptobrevin/VAMP** (vesicle-associated membrane protein), mediate attachment to receptors in the presynaptic membrane (**syntaxin** is one such receptor). These receptors interact with several others—among them, **SNAP-25** that is free in the cytoplasm—thus forming large protein complexes that anchor the vesicles to the presynaptic membrane. The fusion appears

to require that the complex include **synaptotagmin**, which binds Ca^{2+} with low affinity (i.e., the concentration of Ca^{2+} must be high for bonding to occur). According to one hypothesis, synaptotagmin acts as a brake on fusion, and the binding of Ca^{2+} releases the brake. Mice lacking the gene for synaptotagmin have only reduced transmitter release, however, suggesting that other factors also play a role.

Neurotransmitters Are Released in Quanta

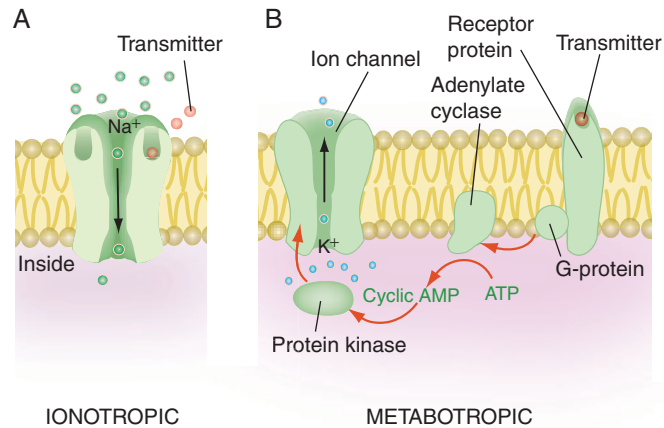
There is convincing evidence that transmitters are released in packets, or **quanta**, corresponding to the transmitter content of one vesicle. For synapses between motor nerve terminals and striated muscle cells, one vesicle contains on average 10,000 transmitter molecules. Only a few thousand molecules of each quantum are likely to bind to a receptor before they diffuse away or are removed by other means. Release of one quantum elicits a tiny excitatory postsynaptic potential (EPSP)—a **miniature EPSP**. If stimulation is increased, so that more transmitter is released, the depolarization of the muscle cell membrane increases in steps corresponding to one miniature EPSP. In the CNS, each bouton probably releases from none to a few quanta for each presynaptic action potential. This means that an action potential does not necessarily elicit transmitter release; it merely increases the **probability of release**. As discussed later, many presynaptic action potentials must coincide to fire a postsynaptic neuron.

Transmitters Act on Ionotropic and Metabotropic Receptors

The effects of a neurotransmitter depend primarily on the properties and localization of the receptors it can activate. There are two main kinds of transmitter receptors: ionotropic and metabotropic. **Ionotropic receptors** are parts of ion channels (Fig. 4.3A). Ionotropic receptors that are parts of Na^+ or Ca^{2+} channels evoke fast and brief **depolarizations** of the postsynaptic membrane, thus exerting **excitatory** actions. Ionotropic receptors coupled to Cl^- channels as a rule **hyperpolarize** the postsynaptic membrane and **inhibit** the postsynaptic neuron. Synapses equipped with ionotropic receptors mediate **fast** and **precise information**—for example, about “when,” “what,” and “where” concerning a sensory stimulus.

The other main kind—**metabotropic receptor**—is not coupled directly to ion channels but acts indirectly by way of **G proteins** and intracellular second messengers (Fig. 4.3B). G proteins may be regarded as universal translators, translating various kinds of extracellular signals to a cellular response (e.g., the “translation” of light and of gaseous and watery chemical substances to nerve impulses).

FIGURE 4.3 Two kinds of transmitter receptors. **A:** Ionotropic receptor with direct action on the ion channel. Note that the receptor is part of the channel proteins. **B:** Metabotropic receptor with indirect action on ion channels. Schematic. All indirectly coupled receptors act via G proteins, whereas other elements of the intracellular signal pathway may vary among different receptors. In this example cyclic AMP serves as the second messenger.



Most neurotransmitters act on both ionotropic and metabotropic receptors. That is, a neurotransmitter can exert both fast, direct synaptic effects and slow, indirect ones (at the same or different synapses). **Glutamate** and **GABA** (γ -aminobutyric acid) are by far the most abundant and ubiquitous transmitters acting on ionotropic receptors, although they also act on several kinds of metabotropic receptors. Several important neurotransmitters, such as **norepinephrine**, **dopamine**, and **serotonin**, exert their main actions on metabotropic receptors. We can conclude that to predict the actions of a neurotransmitter on a neuronal group, we must know the repertoire of receptors expressed by those neurons. Further, because the distribution of receptors differs, one transmitter may exert different actions in different parts of the brain.

Toxins Can Prevent Transmitter Release

Some of the proteins necessary for fusion are degraded by **tetanus toxin** and **botulinum toxin** (produced in certain foods if not treated properly). Both toxins are produced by anaerobic bacteria (i.e., they only grow in the absence of oxygen) and produce violent muscle spasms and paralysis, respectively. The toxins are proteases acting on the proteins that are involved in docking and fusion of synaptic vesicles. While tetanus toxin and some botulinum toxins degrade synaptobrevin, other botulinum toxins destroy SNAP-25, or syntaxin. Even extremely small amounts of the toxins produce muscle spasms (tetanus toxin) or paralysis (botulinum toxins) by preventing transmitter release. They evoke opposite effects because they act on different kinds of synapses: the tetanus toxin affects primarily a type of inhibitory synapse, whereas botulinum toxin acts at the neuromuscular junction, preventing release of the excitatory transmitter acetylcholine.

Inactivation of Neurotransmitters

Synaptic signal transfer is characterized by a precisely timed start and stop. We have looked into the mechanisms responsible for precise timing of transmitter release. It is also necessary, however, that the transmitter, once released, be quickly removed from the synaptic cleft after receptor activation. Simple **diffusion** of the transmitter seems to play an important part, especially during the first few milliseconds after release. Some transmitters (acetylcholine and neuropeptides) are degraded extracellularly by specific enzymes. The majority of transmitters, however, are removed from the extracellular fluid by **uptake** into glial cells or neurons (see also Chapter 2, under “Astroglia and the Control of Neuronal Homeostasis”). Specific transporter proteins in the cell membrane (see Figs. 2.5, 4.1, and 5.5) carry out the transmitter uptake. The transmitter transporters are driven by ion-concentration gradients across the cell membrane. There are two **families** of such transporter proteins. One is driven by the concentration gradients of Na^+ and Cl^- and transports the transmitters **GABA**, **glycine**, **dopamine**, **norepinephrine**, and **serotonin**. The other comprises five different transporters for **glutamate** and is driven by the concentration gradients of Na^+ and K^+ .

The task of the transporters is not to remove all traces of neurotransmitters from the extracellular fluid. Because both number and activity of the transporters are regulated, they rather serve to modulate up or down a certain baseline extracellular transmitter concentration. Even a small alteration of transporter activity can cause changes of transmitter–receptor activation. In areas with a high density of transporters, they also influence the ease by which neurotransmitters may activate receptors outside the synaptic cleft (on nerve terminals, dendrites, and cell bodies). In this way, the

transmitter transporters participate in the control of synaptic transmission and neuronal excitability.

Because the transporter proteins have important physiological roles, they are also interesting from a pharmacological aspect. Drugs that alter their function can be used therapeutically (such as **antidepressants** that are selective serotonin reuptake inhibitors), but some also have potential for abuse (such as **cocaine**, which inhibits the dopamine-reuptake transporter).

SYNAPTIC POTENTIALS AND TYPES OF SYNAPSES

Mechanisms of Postsynaptic Potentials (EPSPs and IPSPs)

Synaptic potentials arise when neurotransmitters activate ion channels. An **excitatory postsynaptic potential (EPSP)** arises at synapses where the transmitter **depolarizes** the postsynaptic membrane. An **inhibitory postsynaptic potential (IPSP)** arises at synapses where the transmitter **hyperpolarizes** the membrane.

Opening of cation channels allowing Na^+ to enter and K^+ to leave the cell produces an EPSP. Because the cations outside the cell are driven inward, by both the concentration gradient and the membrane potential, whereas K^+ inside the cell is driven out only by its concentration gradient, at first the inward current is largest (see Figs. 3.3, 3.4, and 3.5). As the membrane becomes more and more depolarized, however, the outward flow of K^+ increases and counteracts further depolarization (Fig. 4.4). Transmitter-gated channel opening is not subject to self-reinforcement, unlike the voltage-gated channels that produce the action potential. This means that the synaptic potentials rise and fall gradually (Fig. 4.4 and 4.5) and last longer than the action potential. We use the term **graded potential**, as opposed to the all-or-none behavior of the action potential. The current spreads passively (electrotonically) from the synapse outward in all directions along the cell membrane. In this way, the potential becomes gradually weaker, unlike the action potential that is constantly regenerated. Because typical EPSPs in neurons are small (<1 mV), and the membrane has to be depolarized at about 10 mV near the initial segment to reach **threshold** for an action potential, it follows that many EPSPs must be summated to fire the neuron. We return to summation of EPSPs later.

The mechanism behind an **IPSP** is usually the opening of transmitter-gated K^+ or Cl^- channels. This results in an outward flow of K^+ or an inward flow of Cl^- . In both cases, the inside of the cell becomes more negative, that is, the membrane is hyperpolarized. This is only true if the membrane potential is less negative than the equilibrium potentials of the ions in question, however. Although this is the normal situation for K^+ (equilibrium potential -90 mV), the equilibrium potential of

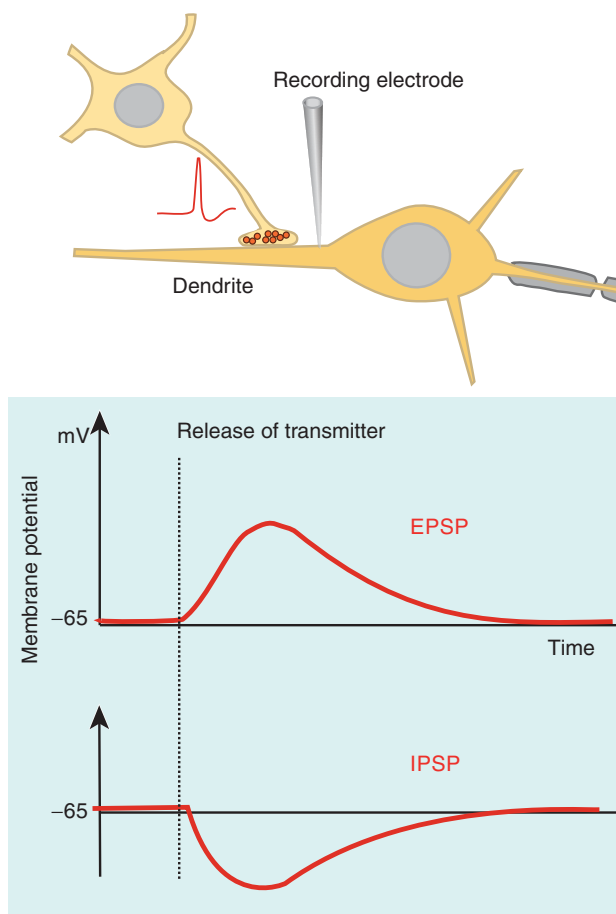


FIGURE 4.4 *Synaptic potentials.* Alterations of the membrane potential evoked by a single presynaptic action potential that releases a transmitter into the synaptic cleft. An EPSP (excitatory postsynaptic potential) is evoked by an excitatory transmitter (typically glutamate), while an inhibitory transmitter (typically GABA) produces an IPSP (inhibitory postsynaptic potential).

Cl^- is close to the resting potential in many neurons. If the resting potential is equal to the equilibrium potential of Cl^- , there is no net flow of Cl^- ions and, consequently, no IPSP is evoked.² Even in this case, however, opening of chloride channels can counteract the effects of excitatory transmitters. Thus, as long as the chloride channels remain open, even the slightest depolarization will cause Cl^- ions to flow into the cell and thereby minimize the change of the membrane potential. In this case, opening of chloride channels by an inhibitory transmitter **short-circuits** the depolarizing currents at nearby excitatory synapses.

2 If the resting potential is more negative than the equilibrium potential of Cl^- , opening of chloride channels causes a net outward flow of chloride ions and the cell is depolarized. This is the case in early embryonic development; the transmitter GABA, which is inhibitory in the adult nervous systems, has excitatory actions in the immature brain.

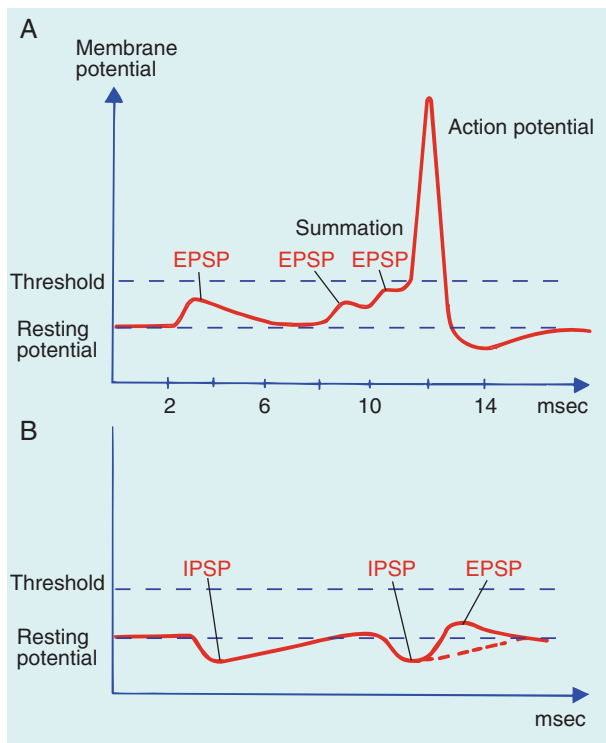


FIGURE 4.5 *Synaptic potentials.* **A:** The time course and polarity of an excitatory postsynaptic synaptic potential (EPSP). In this example, one EPSP alone does not depolarize the membrane to threshold for eliciting an action potential (AP), but if one EPSP (or more) follows shortly after the first one, the threshold is reached (summation). **B:** The time course and polarity of an inhibitory postsynaptic synaptic potential (IPSP) and how the hyperpolarization is reduced when an EPSP is added to an IPSP.

Summation of Stimuli Is Necessary to Evoke an Action Potential

One or a few presynaptic action potentials leading to transmitter release do not evoke an action potential in the postsynaptic cell. As previously mentioned, the membrane has to be depolarized to a **threshold** value (Fig. 4.5A) for an action potential to be evoked. Usually, the threshold is approximately 10 mV more positive than the resting potential, and the size of an EPSP is probably in most cases less than 1 mV. As previously mentioned, to produce an action potential the current produced at synaptic sites must be strong enough to reach the initial segment and depolarize the membrane to threshold (by opening voltage-gated Na^+ channels).

A subthreshold depolarization may nevertheless be of functional significance. If the synaptic potential is followed by another depolarization before the membrane potential has returned to resting value, the second depolarization is added to the first one so that threshold is reached. This phenomenon is called **summation** (Fig. 4.5A). The summation may be in time, as in the example above, and is then called **temporal summation**,

or it may be in space, and is then called **spatial summation**. In temporal summation, impulses may follow one another in rapid succession in one terminal, whereas in spatial summation, nerve terminals at different places on the cell surface release transmitter and depolarize the cell almost simultaneously. In addition, IPSPs are subject to spatial and temporal summation.

An EPSP increases the **probability** that the postsynaptic neuron produce an action potential: for a moment the neuron is more responsive to other inputs. Likewise, an IPSP decreases this probability.

Slow Synaptic Effects Modulate the Effect of Fast Ones

Because neurons are equipped with both ionotropic and metabotropic transmitter receptors, we may safely assume that every neuron receives both fast (direct) and slow (indirect) synaptic inputs. The slow effects modulate the effects of the fast ones, and we therefore use the term **modulatory transmitter actions**. A modulatory transmitter (when binding to an indirectly acting receptor) does not by itself evoke action potentials but alters the response of a neuron to fast, ionotropic transmitter actions. Usually, modulatory synaptic effects are mediated by altering opening states of K^+ or Ca^{2+} channels, thereby modulating both the membrane potential and the refractory period. The effects are nevertheless much more varied because there are several kinds of potassium and calcium channels, and several transmitters may influence each channel.

A brief train of impulses in axons releasing a transmitter that binds to indirectly acting receptors may keep the membrane depolarized or hyperpolarized for seconds after the train of impulses ends (slow EPSP or IPSP; Fig. 4.6). More intense stimulation may produce depolarization that lasts minutes in some neurons.

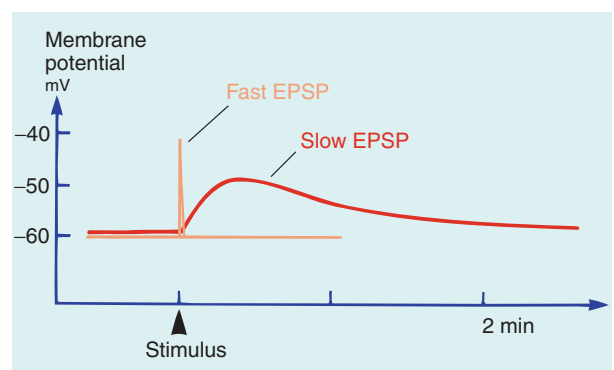


FIGURE 4.6 *Fast and slow synaptic actions.* Schematic. A fast EPSP lasts milliseconds and is caused by binding of transmitter molecules directly to channel proteins. A slow EPSP may last seconds or minutes and is due to activation of receptors indirectly coupled to ion channels.

An example may make this clearer: motor neurons in the cord receive fast, excitatory synaptic input from the cerebral cortex. These signals mediate the precise, voluntary control of muscle contraction. In addition, the motor neurons receive slow, modulatory synaptic inputs from cell groups in the brain stem whose activity is related to the degree of motivation for a particular movement. The modulatory input influences the strength of the response (frequency of action potentials) to signals from the cerebral cortex, that is, how fast the movement will be. However, the modulatory input does not initiate movements on its own.

Mechanisms of Modulatory Synaptic Effects

Slow EPSPs may be mediated by transmitters closing a kind of voltage-gated K^+ channel that is open at the resting membrane potential. This leads to lowered K^+ permeability and reduced flow of K^+ out of the cell, which results in depolarization. Because the membrane potential is shifted toward the threshold, fast depolarization is more likely to elicit an action potential. In addition, the effect on this kind of channel makes a fast EPSP larger and longer-lasting because the fast transmitter opens the K^+ channel during the repolarization phase of the EPSP. When the modulatory transmitter counteracts the opening of the channel in this phase, the depolarization becomes stronger, and the repolarization phase is prolonged. In this way the fast transmitter, rather than eliciting one, may produce a train of action potentials.

Modulatory synaptic effects may not change the resting membrane potential, if they are confined to channels that are not open at the resting potential. A kind of K^+ channel—closed at the resting potential—is opened by Ca^{2+} entering the cell during the action potential. This produces a relatively long-lasting hyperpolarization (the refractory period). A modulatory transmitter that reduces the opening of the K^+ channel would shorten the refractory period. As in the preceding example, a fast excitatory input might produce a train of impulses rather than only one, or the frequency of impulses during a train might be higher than without the modulatory influence.

Slow, long-lasting hyperpolarizing synaptic effects (slow IPSPs) are usually mediated by the indirect opening of K^+ channels. As we discussed later, the ubiquitous inhibitory transmitter GABA can act on receptors with such effect.

A Neuron Integrates Information from Many Others

We have seen that as a rule many impulses must reach a neuron almost simultaneously to make it fire, that is, to send an action potential through its axon. In other words, summation of excitatory synaptic effects is necessary.

The stronger the sum of excitatory effects, the shorter the time necessary to depolarize the cell to the threshold for eliciting another action potential. This means that the frequency of action potentials, or **firing frequency**, is an expression of the **total synaptic input** to a neuron. Total synaptic input here means the sum of both excitatory and inhibitory synaptic influences. Most neurons receive thousands of synapses; for example, large neurons in the motor cortex of the monkey may receive as many as 60,000 synapses. Often a neuron is strongly influenced (many synaptic contacts) by one neuronal group and weakly influenced by many others. This means that while such a neuron primarily transmits signals from one nucleus to another, many other cell groups facilitate or inhibit the efficiency of signal transmission.

Examples of Synaptic Integration

We will provide two examples of the integration of different synaptic inputs. The first concerns **motor neurons** of the spinal cord. Such a neuron—sending its axon to innervate hundreds of striated muscle cells in a particular muscle—is synaptically contacted by neurons in many parts of the nervous system. It may receive around 30,000 synapses, distributed over its dendrites and cell body. Some synapses inform the cell about sensory stimuli that are important for the movement produced by the muscle, others about the posture of the body, others about how fast an intended movement should be, and so forth. The sum of all these synaptic inputs—some of them excitatory, others inhibitory—determines the frequency of action potentials sent to the muscle and by that means the force of muscle contraction (each muscle, however, is governed by many such neurons, so that their collective activity determines the behavior of the whole muscle).

The other example concerns neurons in the spinal cord that mediate information about **painful stimuli**. Although such a neuron receives its strongest synaptic input (most synapses) from sensory organs reacting to painful stimuli, it is also contacted by thousands of synapses from other sources, such as cell groups that are active when the person is anxious. This means that the final firing frequency of this “pain-transmitting” neuron depends not only on the actual stimuli reaching the receptors but also on the activity within the CNS itself. This correlates well with the everyday experience that the pain we feel depends not only on the strength of the peripheral painful stimulus (such as dental drilling) but also on our state of mind. Although the main task of the neuron is to convey sensory information to the brain, this information is integrated in the spinal cord with signals from other sources conveying information the salience of the sensory information.

The Placement of Synapses Has Functional Significance

Where a synapse is located on the neuronal surface is obviously not a matter of chance (Fig. 1.8). There are several examples of axons arising from different cell groups that end on different parts: for example, some end only on proximal dendrites, others on distal dendrites or a particular segment of the dendrite. Further, **inhibitory** synapses are often located on or near the soma of the nerve cell, whereas **excitatory** ones are most abundant on dendrites. The placement can be of functional importance, because synapses close to the **initial segment** of the axon would be expected to have a greater chance of eliciting (or preventing) an action potential than synapses far out on the dendrites. (This is due to the loss of current during electrotonic spread of the synaptic potential over long distances.) In some neurons, powerful inhibitory synapses are even located on the initial segment itself, thereby forming a very efficient “brake” on neuronal firing.

In general, a synapse far out on a dendrite would be expected to exert a weaker effect on the excitability of the neuron than one placed close to the soma, and, consequently, that more summation would be needed for distal synapses than for proximal ones to fire the neuron. New findings suggest that this may not always hold true, however. Studies of pyramidal neurons (in the hippocampus) indicate that an EPSP recorded in the soma is of about equal magnitude regardless of whether it is evoked by a synapse that is proximal or distal on a dendrite. This means that a stronger depolarizing action at distal synapses compensates for their greater loss of current by electrotonic spread.

Another important point regarding the placement of synapses is that most excitatory synapses are located on dendritic **spines** (Figs. 1.1, 1.7, and 1.9). Most of the neurons in the cerebral cortex conform to this arrangement. Because cortical neurons constitute a large proportion of all neurons in the human brain, suggestions that about 90% of all excitatory synapses are located on spines may be realistic.

Spines: Crucial for Learning?

The functional significance of dendritic spines is still under debate. It is not simply a matter of increasing the receiving surface of the neuron because the dendritic membrane between spines may be virtually free of synapses. A spine typically consists of a narrow neck and an expanded part called the spine head (see Fig. 1.9). Since their microscopic identification more than 100 years ago, spines have been implicated in **learning** and **memory**, and recent animal experiments support this hypothesis. For example, the density of spines in the cerebral cortex is higher in rats that live in a challenging environment than in those that are confined to standard

laboratory cages. Further, the density of spines in the cerebral cortex is markedly reduced in individuals with severe mental retardation. Animal experiments with electric stimulation indicate that long-term increases in synaptic efficacy (long-term potentiation, LTP) are associated with changes of spine morphology and number. This effect is observed only among synapses affected by the increased stimulation and may be directly related to learning and memory.

An important function of spines may be to facilitate **local synaptic changes**. The narrow neck of the spines may ensure that the concentration of signal molecules responsible for LTP induction, such as Ca^{2+} , reach much higher levels in the spine head than in the dendrite. This would facilitate changes in those synapses contacting a particular spine, leaving other synapses unaffected.³ Spatial restriction of increased Ca^{2+} concentration may also serve a **protective role** since high intracellular Ca^{2+} concentration may damage the neuron.

Axoaxonic Synapses Enable Presynaptic Control of Transmitter Release

In axoaxonic synapses, the presynaptic bouton makes synaptic contact with a postsynaptic bouton, which, in turn, contacts a cell body or a dendrite (Fig. 4.7). Release of transmitter from the presynaptic bouton serves to regulate the amount of transmitter released by the postsynaptic bouton. This enables inhibition or facilitation of a subset of synaptic inputs to a neuron. The excitability of the postsynaptic neuron is unaltered, in contrast to the situation described above with postsynaptic inhibition by IPSPs.

In the best-studied kind of axoaxonic contacts, action potentials in the presynaptic bouton lead to reduced transmitter release from the postsynaptic bouton; that is, the effect is inhibitory with regard to the neuron contacted by the postsynaptic bouton (the postsynaptic bouton usually has an excitatory action). A prerequisite for this inhibitory effect to occur, however, is that the presynaptic bouton be depolarized (by an action potential) at the same time as or immediately before an action potential reaches the postsynaptic bouton. This phenomenon is termed **presynaptic inhibition** to distinguish it from postsynaptic inhibition. Presynaptic inhibition has been found most frequently among fiber systems that transmit sensory information; for example, sensory fibers entering the spinal cord are subject to

³ Indeed, experiments performed in slices of tissue from the hippocampus (a region involved in learning and memory)—enabling stimulation of single axospinous synapses—suggest that enduring changes (LTP) may be limited to the stimulated synapse. However, it seems that nearby synapses (10–20 synapses within a distance of 10 μm along the dendrite) have lowered thresholds for induction of LTP for some minutes after the stimulation. Such “crosstalk” among nearby synapses is presumably caused by dendritic spread of a diffusible substance produced in the stimulated spine.

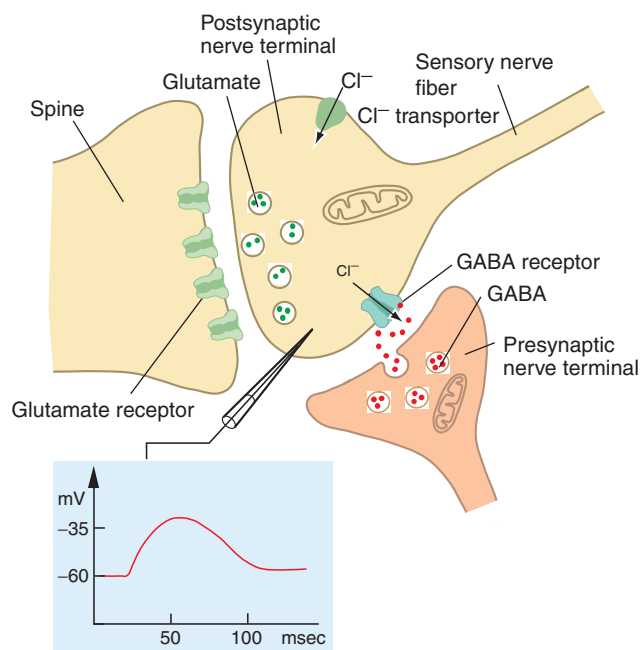


FIGURE 4.7 Presynaptic inhibition is mediated by axoaxonic synapses. The example is from the spinal cord where an inhibitory interneuron contacts the terminal of a sensory nerve fiber (from a spinal ganglion cell). The interneuron releases GABA that opens Cl⁻ channels and thereby depolarizes the postsynaptic nerve terminal (frame). This leads to release of less transmitter. See text for further explanations. (Based on Alvarez 1998.)

powerful presynaptic inhibition. In this way, signals to a sensory neuron from pain receptors can be selectively inhibited, while signals from other receptors are passed on unaltered.

Mechanisms of Presynaptic Inhibition and Facilitation

Several mechanisms may be involved in **presynaptic inhibition**. The phenomenon has been most studied in the spinal cord dorsal horn, where axoaxonic synapses are formed by inhibitory interneurons as they contact terminals of primary sensory afferents (Fig. 4.7). The transmitter released from the interneuron (usually GABA) opens chloride channels in the postsynaptic terminal (bouton). In most neurons, opening of chloride channels either hyperpolarizes or short-circuits the membrane, as described. In the sensory terminals in the cord, however, opening of chloride channels **depolarizes** the membrane, due to an unusually high intracellular chloride concentration. (To uphold this concentration gradient, these sensory neurons are equipped with a special transport mechanism for chloride coupled to the sodium–potassium pump). In this way, the equilibrium potential of Cl⁻ is more positive (–30 mV) than the resting potential (–65 mV) and, consequently, chloride ions move *out* of the nerve terminal

when chloride channels open. But how can depolarization of the presynaptic terminal reduce transmitter release? The answer seems to be that depolarization reduces the amplitude of action potentials as they invade the postsynaptic terminal; in turn, this leads to opening of fewer voltage-gated calcium channels. Because the amount of transmitter released is proportional to the influx of Ca²⁺, less transmitter will be released. It remains to be explained why depolarization of the postsynaptic terminal reduces the amplitude of the action potential. The most likely explanation is that depolarization inactivates some voltage-gated sodium channels in the postsynaptic terminal. In addition, direct influence on Ca²⁺ channels by the transmitter released from the presynaptic terminal may also contribute to presynaptic inhibition.

Presynaptic facilitation can be elicited by axoaxonic synapses by increasing Ca²⁺ influx in the postsynaptic bouton. Closure of K⁺ channels by the presynaptic transmitter prolongs the action potential by slowing the repolarization phase, thereby allowing more Ca²⁺ to enter the postsynaptic bouton.

Why Do We Need Inhibitory Synapses?

Inhibitory synapses are present everywhere in the CNS and are of vital importance for its proper functioning. For example, inhibition is necessary to suppress irrelevant **sensory information**, thereby enabling us to concentrate on certain events and leave others out. Inhibitory synapses also serve to increase the precision of sensory information by, for example, enhancing contrast between regions with different light intensity in visual images.

Inhibition is also necessary for another reason—namely, to interrupt or **dampen excitation**, which might otherwise lead to cell damage. As mentioned earlier, ions have to be pumped through the cell membrane in order to maintain osmotic equilibrium. If many neurons fire continuously with high frequency, even maximal pumping may be insufficient in this respect. Increased extracellular potassium concentration depolarizes neurons, thereby increasing further the neuronal excitation; this leads to more potassium extracellularly, and so on. **Epileptic seizures** are due to uncontrolled firing of groups of neurons, and drugs reducing the tendency for seizures generally increase the effect of inhibitory transmitters. Figure 4.8 shows how an excitatory neuron can limit its firing by way of an **inhibitory interneuron**. Although not shown in the figure, the interneuron is influenced by many other neurons that serve to adjust the brake, as it were. Such arrangements are common, for example, among the motor neurons that control muscle contractions (see Fig. 21.14). In general, inhibitory interneurons increase the **flexibility** of the nervous system.

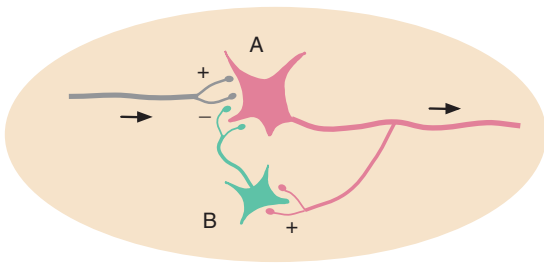


FIGURE 4.8 Inhibitory interneuron (B) mediating negative feedback to the projection neuron (A). Arrows show the direction of impulse conduction.

Signaling by Disinhibition

In some instances, inhibitory synaptic couplings may lead to increased rather than decreased excitation. This occurs when inhibitory neurons inhibit other inhibitory neurons that in their turn act on excitatory ones (Fig. 4.9). With such an arrangement, firing of the first inhibitory interneuron (green) inhibits the next inhibitory interneuron, which thereby reduces its activity. Thereby, the excitatory neuron (red) receives less inhibition and increases its firing. This is called **disinhibition**, and it plays an important role in diverse structures such as the retina and the basal ganglia (Fig. 23.14). If an inhibitory interneuron contacts both excitatory and inhibitory neurons, it might produce both inhibition and disinhibition at the same time in different neurons. By controlling the firing of such interneurons, central command centers—such as the motor cortex—can direct the signals in the desired direction. This occurs, for example, in the spinal cord where inhibitory interneurons serve to select the muscles that are best suited for a particular task.

SYNAPTIC PLASTICITY

Basis of Learning and Memory?

There is much evidence that synapses can alter their structure and function in an **activity-dependent** or **use-dependent** manner: that is, they are **plastic**. This means

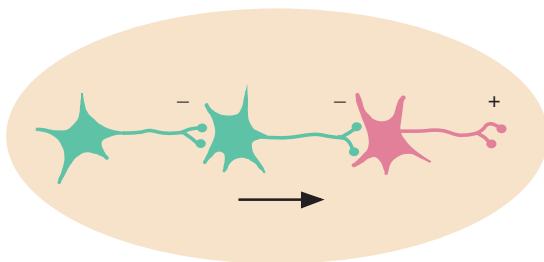


FIGURE 4.9 Disinhibition. Two inhibitory neurons (green) coupled in series increases the activity of an excitatory neuron (red).

that for a shorter or longer period, the postsynaptic action may be enhanced or reduced, as evidenced by altered amplitude of the postsynaptic potentials. Further, much experimental evidence supports the hypothesis that synaptic plasticity is the cellular basis of learning and memory. Martin, Grimwood, and Morris (2000, p. 650) express the hypothesis as follows: “Activity-dependent synaptic plasticity is induced at appropriate synapses during memory formation, and is both necessary and sufficient for the information storage underlying the type of memory mediated by the brain area in which that plasticity is observed.” When we learn, most likely numerous synapses change their efficacy within distributed neural networks; when we later can recall what we learned, it must mean that synaptic changes have been retained. When we forget, one reason may be the decay of learning-related synaptic changes. Obviously, the most difficult part of the hypothesis to prove is a causal relationship between specific synaptic changes and memory in behavioral terms.

Under Which Conditions do Synaptic Changes Occur?

Only a minute fraction of all the information reaching the brain is retained in memory, and correspondingly, use of a synapse does not always change its subsequent behavior. To induce a change, the presynaptic influences must conform to certain patterns. In general, plastic changes are likely to occur when the presynaptic activity is particularly strong and coincide in time with the postsynaptic neuron firing an action potential.⁴ This makes sense, as the immediate firing of an action potential after a synaptic input might be regarded as a sign of success. This situation is likely to arise only when several excitatory synaptic inputs reach the neuron at the same time. Looking for the functional meaning of simultaneous inputs, a crucial point may be the ability of neurons to detect **coincidences**. For example, a sensory input is significant only in a certain **context**, and only then should it be remembered. Accordingly, synaptic changes would occur only when information about the sensory event and its context coincide. However, contextual information should lead to synaptic change only if it signals that the sensory stimulus is important or unexpected. Indeed, in many situations, it appears that synaptic change depends on the coincidence of a specific input mediated by activation of ionotropic receptors and a modulatory input mediated by metabotropic receptors (Fig. 4.10). The first input provides fast and precise information—for example,

⁴ This was postulated by the Canadian psychologist Donald Hebb in 1949 (p. 62) in an influential attempt to explain the cellular basis of learning: “when an axon of cell A is near enough to excite a cell B and repeatedly or persistently takes part in firing it, some growth process or metabolic change takes place in one or both cells such that A’s efficiency, as one of the cells firing B, is increased.”

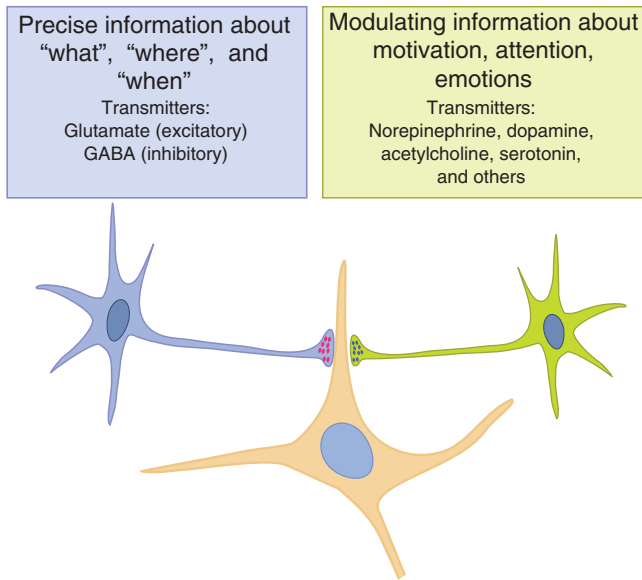


FIGURE 4.10 *Synaptic plasticity.* Learning—that is, synaptic change—depends in this example on simultaneous action of a specific synaptic input and a modulatory one. The latter provides a signal about the salience of the specific stimulus.

about the nature of a sensory stimulus—the other about the **salience** of the stimulus.

Mechanisms for Synaptic Plasticity

Broadly speaking, a change of synaptic efficacy may arise because:

- The **presynaptic** terminal releases more or less neurotransmitter in response to an action potential.
- The **postsynaptic** neuron has increased or reduced its response to the transmitter.

We have discussed the complex cellular processes that link an action potential to transmitter release (see, e.g., Figs. 4.1 and 4.2), and many of the factors involved have been shown experimentally to change their activity in a use-dependent manner. For example, there may be changes in the amount and activity of intracellular second messengers and protein kinases (among other things, regulating ion channels and receptor proteins). The properties, number, and distribution of transmitter receptors are also subject to activity-dependent modifications. Nevertheless, in spite of the large number of molecular mechanisms involved in induction and maintenance of synaptic plasticity, increased intracellular **calcium concentration** appears as a rule to **initiate** the process. The further pathways from a calcium signal to altered synaptic efficacy are multifarious and only partly known. With regard to structural correlates of synaptic plasticity, we mentioned above that **spines** undergo changes of size and form that are

correlated with altered synaptic efficacy. Further, the **formation** of new synapses (synaptogenesis) and **elimination** of old ones is very prominent during pre- and postnatal development but occur throughout adult life.

Enduring changes in neurons—at either a molecular or a structural level—require altered **protein synthesis**.⁵ Proteins have a restricted lifetime, however, and long-term change would therefore require long-term (in many instances life-long) alteration of **gene expression**. We now have much evidence that even synaptic activity lasting for minutes or less may produce altered expression of certain genes that encode for transcription factors (see Chapter 3, under “Markers of Neuronal Activity”). The experimental evidence so far mainly concerns transitory changes of gene expression; nevertheless, a number of genes have been shown to alter their expression in long-term synaptic changes. Although the complex signaling pathways that link neuronal activity to gene expression are thus starting to be revealed, many questions remain. It is not clear, for example, how the altered gene expression can be directed to certain synapses (those subjected to a “memorable” input) and not to others.

Kinds of Synaptic Plasticity

Several kinds of synaptic plasticity have been described on the basis of animal experiments, and more are probably yet to be discovered. It is customary to distinguish between short-term and long-term synaptic plasticity, without a sharp transition. **Short-term plasticity** lasts from less than a second to some minutes, whereas **long-term plasticity** can last for at least several weeks. For practical reasons it is not feasible to study the phenomenon for longer periods in experimental animals. Nevertheless, if synaptic plasticity underlies memories, we know from our own experience that some synaptic changes must last for a lifetime.

The ability of synapses to express plastic changes is subject to regulation by various signal substances. This phenomenon is called **metaplasticity**. Generally, metaplasticity may serve **homeostatic** purposes, by keeping plastic changes within certain limits. However, plasticity may probably also be up- or down-regulated by environmental challenges (“enriched” environment, stress), and in neurological diseases (e.g., Alzheimer’s disease, stroke, and Parkinson’s disease).

⁵ Brain-derived neurotrophic factor (BDNF) is an example of a growth factor that appears to be instrumental for induction of certain forms of long-term synaptic plasticity. Interestingly, BDNF plays a role in many brain processes in which synaptic plasticity plays a prominent role (from brain development to mental diseases).

Short-Term Plasticity

When action potentials reach the nerve terminal at relatively brief intervals, the amplitude of the ensuing postsynaptic potentials often increases gradually. This phenomenon is called **facilitation** and is due to increased transmitter release by each presynaptic action potential. The postsynaptic effect increases for each action potential until reaching a steady state after about 1 sec and then decays rapidly when stimulation stops. Further, at some synapses a series of presynaptic action potentials produce increased synaptic efficacy for minutes after the stimulation ends. This is called (synaptic) **potentiation**, and like facilitation, it is due to increased transmitter release from the nerve terminal. Potentiation can be particularly strong and long lasting after tetanic stimulation (action potentials with maximal frequency). This is called **posttetanic potentiation**. The presynaptic terminal “remembers” that it recently received unusually intense stimulation and alters its behavior accordingly. Facilitation and posttetanic potentiation are examples of short-term plasticity that are important for the nervous system’s capacity for storage of information. **Short-term depression**—that is, a weaker postsynaptic response with repeated presynaptic action potentials—is probably due to insufficient renewal of the releasable synaptic pool (Fig. 4.2).

Long-Term Plasticity: LTP and LTD

Long-term plasticity means changes of synaptic efficacy lasting for hours to weeks (years). **Long-term potentiation (LTP)** and **long-term depression (LTD)** can somewhat arbitrarily be defined as activity-dependent increases or decreases of synaptic efficacy that lasts for more than 1 hour. Presumably, LTP and LTD represent storage of information that is in some way meaningful to the individual (or interpreted as such). With these forms of long-term plasticity, cellular changes occur presynaptically and postsynaptically (not only presynaptically, as do facilitation and posttetanic potentiation). Different forms of both LTP and LTD have been described; they differ with regard to duration, the kind of activity that induces them, and molecular mechanisms. One important mechanism seems to be the **insertion of new receptors** in the postsynaptic membranes, which increases the receptor density and the effect of each quanta of transmitter released from the presynaptic terminal (see “Silent Synapses” later). Regardless of molecular mechanisms involved in the expression of LTP and LTD, the end results are input-specific alterations of synaptic efficacy (see also Chapter 32, under “LTP and Memories”).

LTP and LTD are evoked by different patterns of synaptic inputs. Whereas high-frequency firing or two simultaneous inputs induces LTP, low-frequency firing

or two inputs that are out of phase induces LTD. It seems reasonable that intense activity and synchronization of specific inputs strengthens connections within a network, whereas low activity or desynchronized inputs reduce strength of connectivity—the latter being interpreted as “noise” rather than meaningful information. Further, it appears that LTP is induced if a presynaptic action potential repeatedly precedes firing of the postsynaptic cell, whereas LTD occurs if the postsynaptic firing precedes the presynaptic action potential. In this way, inputs that cause postsynaptic firing (that is, “successful” ones) are strengthened, whereas inputs that do not contribute to postsynaptic firing are weakened.

It may seem paradoxical that the opposite phenomena—LTP and LTD—are both induced by an increase in intracellular Ca^{2+} . There is evidence, however, that intracellular responses to calcium transients can vary depending on their magnitude, time course, and site of origin.

Silent Synapses

It might seem unlikely that the brain contains a large number of synapses that are not in use. Nevertheless, there is now strong evidence that some synapses do not transmit a signal, even though the nerve terminal is invaded by action potentials (this is mainly studied in

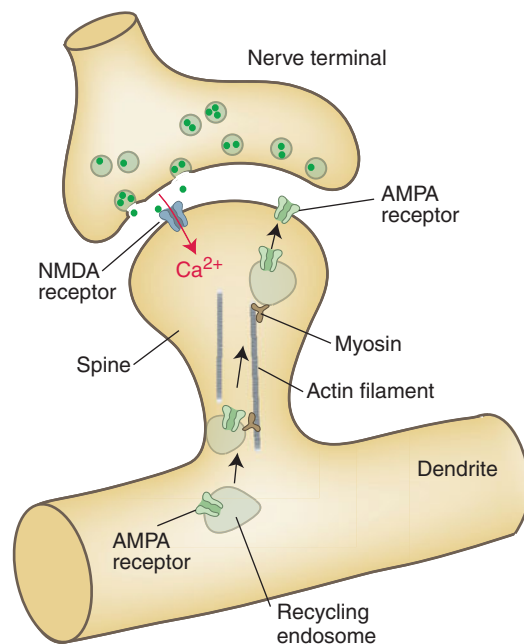


FIGURE 4.11 Getting silent synapses to “speak up.” Activation of NMDA receptors leads to rapid influx of Ca^{2+} ions into the spine. This induces transport of so-called recycling endosomes—containing AMPA receptors—from the dendrite to the spine head. The transport depends on interaction between actin filaments and a special form of myosin molecules in the spine shaft.

the cord and the hippocampus). This is not just a case of low release probability, because even repeated presynaptic firing is without effect (the probability that a presynaptic action potential releases the content of a synaptic vesicle varies enormously in the CNS). Further, in some areas stimulation of an axon evokes a weaker response than expected from the number of terminals. The reason for synapses being silent can be lack of either transmitter release or a postsynaptic response to the transmitter (due to lack of receptors or that the receptors are blocked). There is evidence of both mechanisms. For example, some glutamatergic synapses in the hippocampus lack the ionotropic amino-methylisoxazole-propionic acid (AMPA) receptors, while expressing voltage-dependent *N*-methyl-D-aspartate (NMDA) receptors (these are further described in Chapter 5, under “Glutamate Receptors”). Such synapses

are silent because “normal” presynaptic glutamate release does not activate them. This is because the opening of NMDA receptors requires a certain magnitude of depolarization in addition to binding of glutamate. If NMDA receptors are opened by particularly strong depolarization of the postsynaptic membrane, however, this may, in turn, lead to the insertion of AMPA receptors in the postsynaptic membrane (Fig. 4.11). The receptors are transported rapidly from endosomes to the postsynaptic density, as shown with fluorescence labeling methods. Afterward, the synapse is no longer silent but “speaks up” when glutamate is released. In many instances, LTP may be caused by silent synapses being activated by insertion of AMPA receptors. The finding that silent synapses appear to be most numerous shortly after birth (in rats) supports their possible role in learning and memory.

5 | Neurotransmitters and Their Receptors

OVERVIEW

Certain general properties of neurotransmitters were outlined in the preceding chapters. We recall that a signal is conveyed from one neuron to the next by release of a **neurotransmitter** (transmitter). “Conventional” or “classical” neurotransmitters are small molecules, such as amino acids and amines. Another important group of signal substances, released at synapses, are peptide molecules, called **neuropeptides**. Although the “typical” transmitter is released and acts at receptors in a synapse, many transmitter receptors are found **extrasynaptically**, that is, without connection to a synapse. Indeed, many transmitters act both at synapses and extrasynaptically. The latter action is called **volume transmission**, and is obviously less precise than synaptic transmission. Many receptors are located presynaptically on nerve terminals. Some of them are **autoreceptors** (binding the transmitter released from the terminal), and others are **heteroreceptors** (binding other transmitters released from neurons in the vicinity). Many nerve terminals contain more than one transmitter; often a classical transmitter is **colocalized** with one or more neuropeptides.

Synthesis of a transmitter usually depends on the activity of a **key enzyme**, which controls the amount of transmitter available at a synapse. Transmitter receptors far outnumber the transmitters; thus each transmitter usually acts on several **ionotropic** and **metabotropic** receptors. The effects of a transmitter on a certain neuron therefore depend on which receptors the neuron expresses. Both the amount of transmitter available for release and the postsynaptic receptors are subject to use-dependent plasticity.

The most important **amino acid transmitters** are **glutamate** and γ -aminobutyric acid (GABA). Both are present in virtually all parts of the central nervous system (CNS), and are responsible for most of the fast and precise synaptic transmission, by acting on ionotropic receptors. Glutamate is the dominant excitatory transmitter, whereas GABA is inhibitory. These transmitters mediate most of the spatially and temporally precise excitation and inhibition needed for perception, movements, and cognition. Glutamate binds to three families of receptors (amino-methylisoxazole propionic acid [AMPA], *N*-methyl-D-aspartate [NMDA], and metabotropic glutamate receptors). The **AMPA** receptors are typical ionotropic receptors with fast excitatory action,

whereas the **NMDA** receptors have properties that make them especially suited to detect coincidences and induce long-term potentiation (LTP). GABA acts on ionotropic **GABA_A** receptors and metabotropic **GABA_B** receptors.

Acetylcholine is used as transmitter by a limited number of neurons in the brain stem and basal forebrain, while axonal ramifications of the neurons occur in most parts of the brain. Acetylcholine acts on ionotropic **nicotinic** receptors and metabotropic **muscarinic** receptors. The latter type dominates in the CNS, and the actions of acetylcholine are especially related to motivation, sleep–wakefulness, and memory. The group of **biogenic amines** includes the monoamines **dopamine**, **norepinephrine** (epinephrine), and **serotonin** (and in addition histamine). Like acetylcholine, these transmitters are produced in a small number of neurons but nevertheless act in most parts of the brain. They bind mostly to metabotropic receptors, with actions related to arousal, mood (emotions), and synaptic plasticity. In general, acetylcholine and the biogenic amines have **modulatory** actions that serve to regulate the excitability of neurons, thus determining the magnitude of response to fast-acting transmitters such as glutamate and GABA. **Adenosine triphosphate** (ATP) and **nitric oxide** (NO) function as signal substances with transmitter-like actions in the CNS.

A large number of **neuropeptides** have modulatory and metabolic actions in the brain, and influence a variety of processes, from basic homeostasis to complex behaviors.

GENERAL ASPECTS

How to Prove that a Substance Functions as a Neurotransmitter

Many signal substances present in the brain are small molecules, including the **amino acids** glutamate, glycine, and GABA, and the **amines** norepinephrine, dopamine, serotonin, and histamine. Acetylcholine and ATP also belong to this group. Other signal substances are peptides and therefore fairly large. Such **neuropeptides** are chains of 5 to 30 amino acids. In general, the functions of the neuropeptides are far less clarified than those of the small-molecule transmitters. To prove that a substance present in a neuron actually functions as a

neurotransmitter is not easy. It is not sufficient that neurons express specific **binding sites** for a substance; hormones, growth factors, and other molecules also bind to neuronal membrane receptors. Neither are there clear-cut chemical differences between neurotransmitters and other intercellular signal molecules. Indeed, the same molecule may function in several roles; for example, norepinephrine is both a neurotransmitter and a hormone. Further, some molecules—such as glutamate and glycine—are intercellular signal molecules and have a role in cellular metabolism (e.g., as building blocks for proteins).

To **prove** that a substance functions as a signal molecule, one must show the presence of corresponding receptors and that the substance is released in sufficient amounts (under physiological conditions) to activate the receptors. Additional criteria must be met to classify such a signal substance as a neurotransmitter, however. The substance must be **produced** by neurons, it must be **stored** in nerve terminals and **released** by depolarization, and the release must be **calcium dependent**. In addition, the released substance must be directly responsible for the postsynaptic changes. Finally, there must exist mechanisms for **inactivation** of the transmitter after release. Only a few signal substances have met all of these criteria when tested experimentally. For several others, the probability that they function as transmitters is nevertheless high, and they are often described as transmitters without reservation. Strictly speaking, however, they should be termed “transmitter candidates” or “putative transmitters.” Acetylcholine was the first substance to be classified with certainty as a transmitter. The excitatory action of glutamate was discovered in the 1950s, but only toward 1990 was the neurotransmitter status of this ubiquitous amino acid verified.

Determination of Neuronal Content of Neurotransmitters and Distribution of Receptors

With biochemical methods, the content of transmitters in parts of the brain and in subcellular fractions can be determined. To obtain further knowledge, however, it is also necessary to link the transmitters to specific neurons with known connections and physiological properties. The first possibilities of studying the anatomy of neurons with known transmitters arose in the 1960s, when it was discovered that monoamine-containing neurons could be made fluorescent by a special treatment with formaldehyde. This marked the beginning of intense investigations, with other methods as well, to characterize neurons with regard to connections and at the same time with regard to their transmitters. The introduction of immunological methods, such as **immunocytochemistry**, to localize substances in nervous tissue has been of particular importance. By purifying a

potentially interesting substance present in nervous tissue, **antibodies** may be raised against it. The antibodies bind antigens where they are exposed in the tissue sections, and the antibodies can be visualized subsequently with the use of secondary antibodies. The secondary antibodies may be labeled with a fluorescent molecule, or they may be identified in other ways. Such methods have been widely used to demonstrate the localization of enzymes that are critical for synthesis or degradation of certain transmitters, such as tyrosine hydroxylase, which is necessary for the synthesis of dopamine and norepinephrine (Fig. 5.1A), choline acetyltransferase (ChAT) for synthesis of acetylcholine (Fig. 5.1B), and glutamic acid decarboxylase (GAD) for GABA. Even transmitter molecules that are themselves too small to serve as antigens can be specifically identified in tissue sections via immunocytochemical methods when conjugated to tissue proteins (with glutaraldehyde). This is the case for GABA (Fig. 5.1C), glutamate, and glycine. Immunocytochemical methods can also be applied to **ultrastructural** analysis, in order to determine the transmitter accumulated at specific synapses and also whether the transmitter is localized to certain organelles, such as presynaptic vesicles (Fig. 5.2). Combination of axonal transport methods and immunocytochemical procedures makes it possible to determine the connections, as well as the transmitter candidates and other neuroactive substances of specific neuronal groups.

Even though the determination of the transmitter candidates present in a neuron is of great importance, it is not always possible to know whether the substance has been synthesized in the cell or whether it has merely been taken up. Further, the concentration of a transmitter in parts of a neuron may be so low that it cannot be reliably detected with immunocytochemical methods. The use of ***in situ* hybridization** techniques helps to overcome this kind of problem. By these methods, it is not the neuropeptides or enzymes related to transmitter metabolism that are demonstrated but the presence of the corresponding **mRNA**.

Volume Transmission: Extrasynaptic Transmitter Release and Actions

A neurotransmitter (transmitter) is usually defined as a chemical substance that is released at a synapse and transmits a signal from one neuron to another (or to muscle cells or glandular cells). However, not all substances, otherwise behaving as neurotransmitters, are released at synapses (Fig. 5.3). In some places, neurotransmitters are released from **axonal varicosities** that do not form synapses. Further, many transmitter receptors are present **extrasynaptically**—that is, in the neuronal membrane outside synapses (Fig. 5.3). Therefore, some neurotransmitters act both at typical

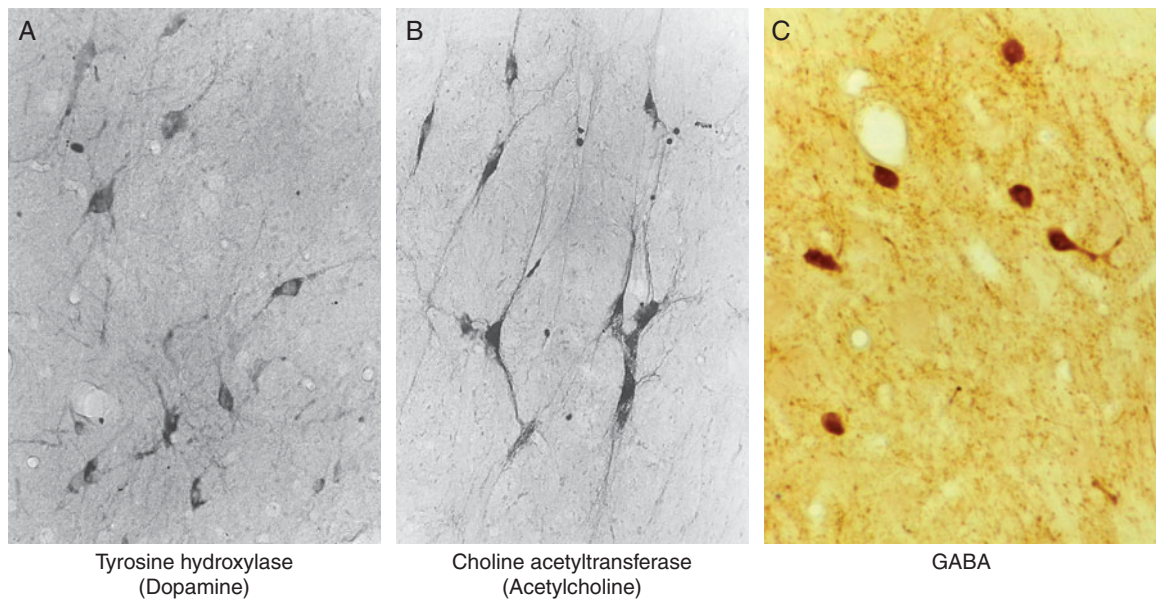


FIGURE 5.1 *Immunocytochemical demonstrations of neurotransmitters.* **A:** Dopaminergic neurons (substantia nigra) are visualized with an antibody against the enzyme tyrosine hydroxylase, which is involved in the synthesis of dopamine (Fig. 5.7). Neurons in between the black, labeled ones—not containing the enzyme—are not visible. Note that the method does not demonstrate the transmitter itself. **B:** Cholinergic

neurons (basal nucleus) are visualized with an antibody against the enzyme choline acetyltransferase **C:** GABAergic neurons are visualized with an antibody that binds to a GABA–protein complex in the section. The brown cell bodies, showing GABA-like immunoreactivity, are interneurons in a brain stem nucleus (monkey). The small brown dots are partly dendrites, partly axons and nerve terminals.

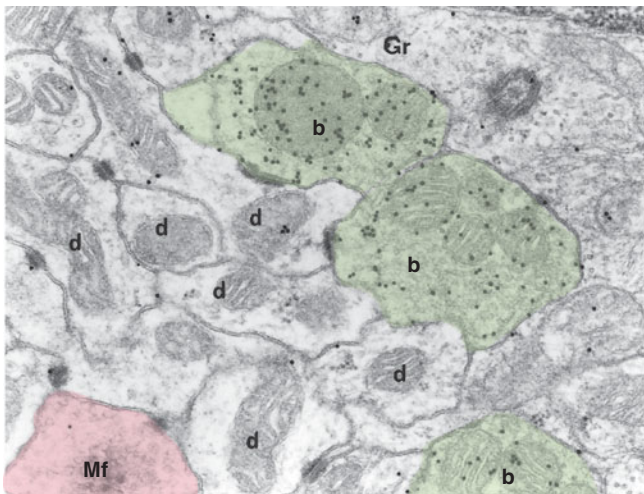


FIGURE 5.2 *Electron microscopic, immunocytochemical demonstration of neurotransmitters.* The small black dots represent gold particles bound in the tissue where GABA is present. The gold particles are conjugated to an antibody directed against a GABA–protein complex. The gold particles are concentrated over a particular kind of bouton (b), whereas dendrites (d) and part of a cell body (Gr) are not labeled. Another kind of nerve terminal (Mf) is also unlabeled and most likely contains a neurotransmitter other than GABA. Rat cerebellum. (Courtesy of Prof. O. P. Ottersen, Department of Anatomy, University of Oslo.)

synapses and more diffusely like local hormones—that is, on all receptors with the proper specificity within a certain distance from its release site. The distance depends on how quickly the transmitter is inactivated and how easily it diffuses. It appears that the effective distance is in the range of a few micrometers from the release site. We use the term **volume transmission** to distinguish this kind of transmitter action from the spatially precise action at synapses. The narrow definition of transmitters, focusing on their actions at synapses, fits well for transmitters such as glutamate and GABA with spatially and temporally precise actions. Other transmitters with mainly modulatory actions, such as norepinephrine, dopamine, and serotonin, act also by volume transmission. These different forms of transmitter actions complicate the analysis of how neurotransmitters control neurons and human behavior.

Colocalization of Transmitters

It was formerly assumed that each neuron releases only one transmitter. Recent studies have shown, however, that nerve terminals often contain more than one transmitter. Such **colocalization** of transmitters appears to be the rule rather than an exception. In most cases described so far, a small-molecule, “classical” transmitter is colocalized with one or several **neuropeptides**. In such cases, the terminal can evoke both fast synaptic

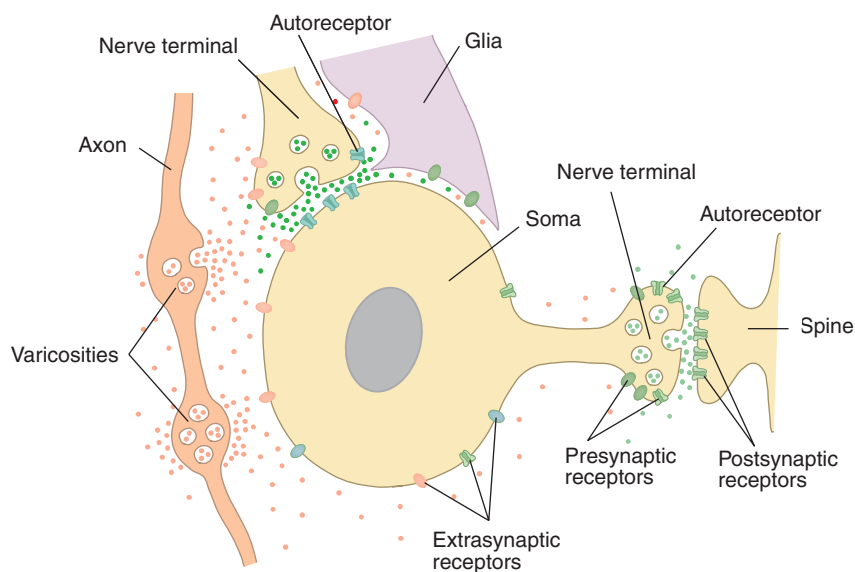


FIGURE 5.3 *Extrasynaptic receptors and transmitter release outside synapses.* Extrasynaptic receptors are localized both at the nerve terminals and on the somatic and dendritic surfaces of the neuron. Autoreceptors bind the transmitter released by the neuron itself. Note the release of transmitter from varicosities that do not form typical synaptic contacts.

actions mediated by ionotropic receptors and slow effects mediated by metabotropic receptors. Colocalization of two “classical” transmitters such as GABA and glycine—both acting on ionotropic receptors—has also been reported. GABA and glycine are both inhibitory, and it seems reasonable that colocalized transmitters exert similar postsynaptic actions. Even this may not be universally true, however. For example, certain spinal interneurons and neurons in the hypothalamus release ATP (excitatory) together with GABA (inhibitory); this means that both act on ionotropic receptors but apparently with opposite effects.

The widespread occurrence of colocalization complicates the interpretation of one particular transmitter’s contribution to synaptic effects, and accordingly, its contribution to certain behaviors. Hypotheses about neural functions and disease symptoms have often been based on the erroneous belief that a particular neuronal group or fiber system uses only one transmitter (the one that was first discovered).

Transmitter Receptors in General

The many transmitters and transmitter candidates (more than 50, including the neuropeptides) have an even larger variety of receptor types to act on. More than 200 different metabotropic (G protein-coupled) receptors have been identified in the CNS. (Not all bind neurotransmitters, however; many bind hormones and a variety of growth factors.)

Several requirements have to be met to conclude that a binding site for a transmitter functions as a receptor. The “final proof” requires that the amino-acid sequence

of the receptor site has been determined. After cloning of the protein, one can then determine whether it binds the transmitter (and agonists and antagonists) and produces the expected physiological effects.

As discussed in Chapter 3 (under “The Structure of Ion Channels”), all the **directly** acting, **ionotropic**, receptors—being part of ion channel proteins—are similarly built, with several subunits arranged around a central pore (see Fig. 3.2). In addition, the **indirectly** acting, **metabotropic** receptors share several structural features, although they are quite different from the ionotropic receptors. The metabotropic receptors usually consist of one large protein that makes several turns through the membrane with hydrophilic (water-soluble) groups on the interior and exterior of the membrane. The receptors mediate their effects via **G proteins** (see Fig. 4.2). Several receptor types with different postsynaptic actions have been identified for each of the best-known transmitters.

The most abundant transmitters, such as glutamate and GABA, act on both ionotropic and metabotropic receptors. The link between a neurotransmitter and its actions is made even more complex by the existence of subtypes of each main kind of receptor. Subtypes of the ionotropic (directly acting) GABA receptor (GABA_A) exemplify this. Each of the protein subunits forming the receptor (and the ion channel) comes in several varieties, and different combinations of them produce numerous subtypes of the GABA_A receptor. These subtypes are differently distributed in the brain. This may explain why drugs acting on different subtypes of the GABA_A receptor have different physiological and behavioral effects: they act on different neuronal networks.

Regulation of Receptor Density

The transmitter receptors are not static, immutable elements of the nervous system. We have discussed how changes in receptor density and activity may mediate **synaptic plasticity**. This means that learning would be expected to be associated with receptor changes. In animal experiments, for example, stressful psychological experiences leading to altered behavior also alter the activity of specific transmitter receptors. Drugs that interfere with transmitter actions often induce changes in the receptors. A drug that blocks the effect of a transmitter on a particular receptor type (antagonist) may indirectly produce increased postsynaptic receptor density. The reverse may occur with drugs that mimic the transmitter (agonist). Probably such **up- or down-regulation** of receptors represents an adaptive response: an abnormally high concentration of the transmitter (or an agonist) is counteracted by reduced receptor activity to maintain normal synaptic transmission. Down-regulation of receptors may explain many of the dramatic **withdrawal symptoms** that occur when an addicted individual abruptly discontinues a narcotic drug.

Presynaptic Transmitter Receptors

Transmitter receptors are also localized to the presynaptic membrane and can thereby modulate transmitter release (Fig. 5.3). We have discussed the axoaxonic synapses that mediate presynaptic inhibition by acting on receptors in the presynaptic membrane (see Fig. 4.7). In addition, the presynaptic membrane can express receptors for the transmitter released by the nerve terminal itself (see Fig. 4.1). Here we use the term **autoreceptors**. Often, autoreceptors inhibit transmitter release as a kind of **negative feedback**. Some neurons, for example, dopaminergic ones, are equipped with autoreceptors also on the cell body and dendrites. In addition to autoreceptors, nerve terminals may express **heteroreceptors**, that is, receptors for transmitters other than those they release themselves (often released by nearby terminals). Nerve terminals releasing **norepinephrine** can exemplify the complexity of presynaptic modulation. Such terminals can express α_2 adrenergic autoreceptors and muscarinic (acetylcholine), opiate, and dopamine receptors that inhibit release of norepinephrine from the terminal. In addition, the terminals also express β_2 adrenergic autoreceptors and nicotinic (acetylcholine) receptors that facilitate transmitter release. Thus, the amount of transmitter released by such a nerve terminal depends not only on the presynaptic activity of the neuron but also on the **local milieu** of the terminal (the concentration of various transmitters and other signal substances, as well as the presence of drugs or toxic substances). It should not come as a surprise

that the functional roles of presynaptic receptors are not fully understood.

Synthesis of Neurotransmitters

The small-molecule, “**classical**” neurotransmitters (Table 5.1) are synthesized in the nerve terminals, the synthesis being catalyzed by enzymes transported axonally from the cell body. As a rule, the rate of transmitter synthesis is determined by the activity of one **key enzyme**. Up- or down-regulation of the enzymatic activity represents one way of changing the properties of nerve cells—with regard to learning, for example. Activation of enzymes often requires that they be phosphorylated, and this may be a result of external stimuli that, via membrane receptors, induce increased intracellular concentration of second messengers (such as Ca^{2+} or cyclic AMP).

As the organelles necessary for protein synthesis are present almost exclusively in the cell body, the **neuropeptides** must be synthesized in the cell body and subsequently transported to the terminals. Accordingly, substances that block axonal transport, such as **colchicine**, lead to accumulation of neuropeptides in the cell body. Usually, the neuropeptides are produced as larger polypeptides (prepropeptides) that most likely are split into smaller units on their way to the terminals.

SPECIFIC NEUROTRANSMITTERS

Excitatory Amino Acid Transmitters: Glutamate and Aspartate

The amino acid group contains the ubiquitous excitatory transmitter, **glutamate** (Fig. 5.4). Neurons that release glutamate at their synapses are called **glutamatergic**. The dominant effect of glutamate in the CNS is fast excitation by direct action on ion channels, although it also acts on metabotropic receptors. Glutamate is responsible for fast and precise signal transmission in the majority (all?) of the large sensory and motor tracts, as well as in the numerous connections between various parts of the cerebral cortex that form the networks responsible for higher mental functions. The total concentration of glutamate in the brain is very high, although the distribution is uneven. Notably, effective uptake mechanisms keep the extracellular concentration very low (about 1/1000 of intracellular concentration). This is a prerequisite for glutamate’s function as a neurotransmitter—acting only on specific receptors after controlled release from nerve terminals. Low extracellular concentration is also mandatory because even a small increase is toxic to the neurons. Transporter proteins in astroglial membranes maintain the concentration gradient (probably with a

TABLE 5.1 *The Best-Known Small-Molecule (Classical) Neurotransmitters*

<i>Transmitter</i>	<i>Chemical</i>	<i>Receptor Name Mechanism, Ion Permeability*</i>	<i>Synaptic Action</i>	<i>Distribution of Receptors in the CNS[†]</i>	<i>Localization of Neurons Synthesizing the Transmitter in the CNS[†]</i>	<i>Localization of Neurons in the PNS</i>
Glutamate	Amino acid	AMPA, ionotropic, Na ⁺ NMDA, ionotropic, Ca ²⁺ mGluR1-5, metabotropic	Fast, excitation Fast, excitation Slow, excitation or inhibition, metabolic effects	"Everywhere"	"Everywhere"	Spinal ganglion cells
GABA	Amino acid	GABA ionotropic, Cl ⁻ GABA _B , metabotropic, K ⁺ , Ca ²⁺	Fast, inhibition Slow, inhibition	"Everywhere"	"Everywhere"	Gut, ganglia
Glycine	Amino acid	Ionotropic, Cl ⁻	Fast, inhibition		Brain stem, spinal cord, cerebellum	
Acetylcholine	Quaternary amine	Nicotinic, ionotropic, Na ⁺ Muscarinic, metabotropic K ⁺ , Ca ²⁺	Fast, excitation Slow, excitation or inhibition	Cerebral cortex, spinal cord (and other places) Cerebral cortex, hippocampus, thalamus (and other places)	Motoneurons, preganglionic autonomic neurons, basal nucleus, septal nuclei, nuclei in the reticular formation of the brain stem (and other places)	Parasympathetic ganglia
Norepinephrine	Amine	α (α_1 - α_2), metabotropic β (β_1 - β_2), metabotropic	Slow Slow	"Everywhere"	Locus coeruleus and diffuse cell groups in the reticular formation	Sympathetic ganglia
Dopamine	Amine	D ₁ (D ₁ , D ₅), metabotropic, increase cyclic AMP D ₂ (D ₂ , D ₃ , D ₄), metabotropic, decrease cyclic AMP	Slow Slow	"Everywhere" (especially basal ganglia and pre-frontal cortex)	Mesencephalon (Substantia nigra and ventral tegmental area)	
Serotonin (5-HT)	Amine	5-HT _{1A} , metabotropic, K ⁺ 5-HT ₂ , metabotropic 5-HT ₃ , ionotropic, Na ⁺	Slow, inhibition Slow, excitation Fast, excitation	"Everywhere"	Raphe nuclei (brain stem)	

*There are more receptor subtypes than shown here.

[†]The table is not complete regarding distribution of neurons and receptors.

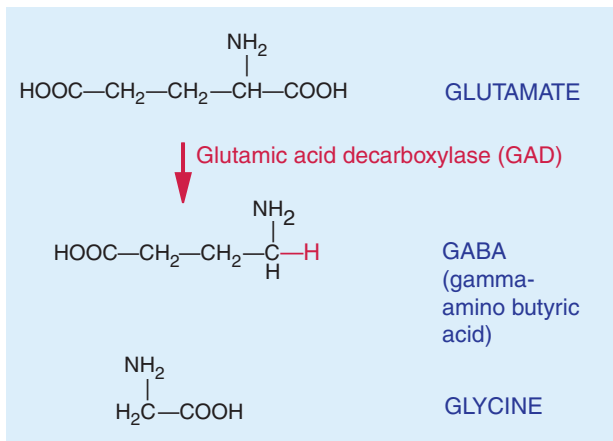


FIGURE 5.4 *Amino acid transmitters.* The enzyme glutamic acid decarboxylase (GAD) is specific for neurons that synthesize GABA.

smaller contribution from transporters in neuronal membranes). In situations with poor energy supply, such as reduced blood flow or low blood sugar, glutamate leaks from the cells because the uptake mechanisms fail. The ensuing rise in extracellular glutamate concentration contributes significantly to the rapid occurrence of neuronal damage—for example, in cases of cardiac arrest (see also Chapter 11, under “Ischemic Cell Damage”).

The amino acid **aspartate** is also highly concentrated in the CNS. It exerts an excitatory action by binding to glutamate receptors. Although decisive evidence is still lacking, recent studies speak in favor of aspartate being a neurotransmitter. For example, aspartate was found to be colocalized with glutamate in synaptic vesicles and released by exocytosis. Nevertheless, the distribution of possible aspartatergic synapses seems to be very limited in the brain. Therefore, with regard to excitatory synaptic transmission, aspartate must play a minor role compared with glutamate.

Synthesis of Glutamate

Glutamate is synthesized from two sources in the nerve terminals: **glucose** (via the Krebs cycle) and **glutamine** that is synthesized in glial cells and thereafter transported into the neurons (Fig. 5.5). Glutamine is converted in mitochondria to glutamate by the enzyme **glutaminase**. Specific transport proteins in the vesicle membrane fill the synaptic vesicles with glutamate. Glutamate is released as other transmitters by calcium-dependent exocytosis. Released glutamate is taken up by glia and converted to glutamine by the enzyme **glutamine synthetase**. Glutamine is then transported to nerve terminals, converted to glutamate, and so forth. Figure 5.5 shows the “glutamate–glutamine circuit,” which ensures reuse of the transmitter. Another advantage with the

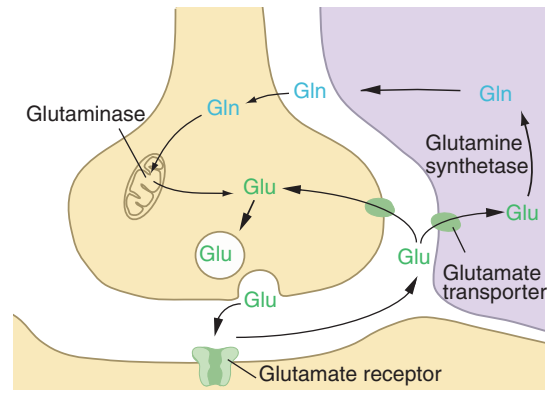


FIGURE 5.5 *The glutamate–glutamine “circuit.”* The enzyme glutamine synthetase converts glutamate (Glu) to glutamine (Gln) after uptake by glial cells. In contrast to glutamate, glutamine is neutral to neurons. Glutamine is transferred to nerve terminals where it is used to synthesize new glutamate that is concentrated in vesicles for release, and so forth.

circuit is that glutamine, unlike glutamate, does not influence neuronal excitability, and is not toxic in high concentrations. Therefore, its concentration need not be strictly controlled.

Glutamate Receptors

As mentioned, the dominant action of glutamate in the CNS is fast excitation by direct binding to ionotropic receptors. In the early 1980s, however, additional kinds of glutamate receptors (**GluRs**) were found. We now recognize three groups or families of receptors that glutamate can bind to: **AMPA/kainate receptors**,¹ **NMDA receptors**, and **metabotropic glutamate receptors (mGluRs)**. The two first groups are glutamate-gated ion channels (ionotropic receptors). The metabotropic glutamate receptors are G protein–coupled, like other metabotropic receptors. The various glutamate receptors were discovered via use of glutamate analogs that turned out to act only on certain kinds of receptors. The receptors were named after the analog that activated them selectively (amino-methylisoxazole propionic acid [AMPA]; *N*-methyl-*D*-aspartate [NMDA]). Cloning of the receptor proteins in the early 1990s showed that AMPA, NMDA, and metabotropic receptors belonged to different protein families. To date, 10

¹ **Kainate receptors** belong to the same family of ionotropic receptors as AMPA receptors. They have been identified in many parts of the CNS and can be localized both pre- and postsynaptically. The concentration of kainate receptors is low in most areas, and they have not been as extensively studied as AMPA and NMDA receptors. Their total contribution to fast excitation—as compared to that of AMPA receptors—is therefore still not clarified. Kainate receptors are expressed in spinal ganglion cells (sensory neurons), and animal experiments suggest that blockage of these receptors may alleviate chronic pain.

varieties of AMPA/kainate receptors, 5 NMDA receptors, and 8 metabotropic receptors have been cloned.²

AMPA receptors are ion channels admitting Na^+ (and K^+), which is typical of fast, excitatory synapses. The current view is that AMPA receptors are responsible for the majority of the fast excitatory signals in the CNS (mediating, e.g., precise sensory information and motor commands).

NMDA receptors have properties that distinguish them from other ionotropic receptors. They have attracted much interest due to their role in long-term potentiation (LTP), and, therefore, most likely in learning and memory. They have a much slower synaptic action than the AMPA receptors and are engaged in other tasks. One important feature of NMDA-gated ion channels is that they are much more permeable to Ca^{2+} than to Na^+ (in contrast to AMPA-gated channels). This makes possible many postsynaptic effects of glutamate binding in addition to depolarization, because Ca^{2+} can trigger a number of intracellular processes (e.g., related to synaptic plasticity). Another special feature of NMDA receptors is that they are **voltage-dependent** and remain closed at resting membrane potential. Binding of glutamate (or NMDA) to the receptor opens it only if the membrane is already depolarized—for example, by the opening of AMPA receptors in the vicinity. Depolarization removes Mg^{2+} ions that otherwise block the channel. A final characteristic feature is that when the NMDA channel is opened, the flow of Ca^{2+} through it lasts much longer than an ordinary EPSP (which is produced by opening the AMPA channels).

Metabotropic glutamate receptors (mGluRs) are located both postsynaptically and presynaptically.³ They fall into three groups, differing with regard to which intracellular signal pathway they activate. As to postsynaptic effects, it appears that **group I mGluRs** produce a long-lasting depolarization (slow EPSP), whereas **group II** has the opposite effect (slow IPSP). Obviously, the existence of glutamate receptors with inhibitory actions further complicates the analysis of glutamate as a transmitter. In addition, mGluRs have

metabolic effects that influence various neuronal processes—among them, the induction of LTP and LTD. As mGluRs are thought to be involved in several brain diseases (e.g., epilepsy, schizophrenia, and stroke), drugs modulating the function mGluRs are being developed and tested in animal models of human diseases.

NMDA Receptors: Mediators of Both Learning and Neuronal Damage

Long-term potentiation (LTP) was described in Chapter 4. The transmission at an excitatory synapse can be changed for a long time when the synapse is active simultaneously with other excitatory synapses in the vicinity (associative LTP). In many areas in the CNS, the induction of LTP depends on activation of NMDA receptors, and the NMDA receptors appear to have just the right properties for this task because they require both postsynaptic depolarization and glutamate binding. (Not all LTP depends on NMDA receptors, however.) NMDA receptors have binding sites for substances other than glutamate, also. The amino acid **glycine** (otherwise acting as an inhibitory transmitter) binds to the NMDA receptor, and such binding is necessary for glutamate to open the NMDA channel. Other substances that occur naturally in the brain also influence the activity of the NMDA receptors, and thereby presumably determine how plastic many synapses are (**metaplasticity**). Changes in the concentrations of such substances might be relevant for learning and memory in general and for recovery after brain damage.

The NMDA receptor is also one among several candidates for mediating **cell damage** after abnormal excitatory activation (see also Chapter 11, under “Ischemic Cell Damage”). This occurs when nervous tissue receives insufficient oxygen (hypoxia), as in severely reduced blood pressure, stroke, cerebral bleeding, and so forth. Abnormally intense excitation may also occur during **epileptic seizures**. In such circumstances, extracellular glutamate concentration rises steeply and presumably, all kinds of glutamate receptors are activated (see later, “Glutamate Transporters”). Activation of NMDA receptors may nevertheless be especially important because it can lead to a rapid increase of the intracellular Ca^{2+} concentration. There is evidence that increased calcium concentration is crucial for cell death, among other things by increasing depolarization and initiating a vicious cycle that activates proteolytic enzymes and induces large concentration changes of ions. In turn, this causes osmotic imbalance with cell swelling and potential destruction. If activation of glutamate receptors has a crucial role for cell death after a stroke, blockage of glutamate receptors might be effective in reducing the damage (if started within 1–2 hours after onset of the symptoms).

2 Glutamate receptors are expressed also in **peripheral tissues** such as bone (osteoblasts and osteoclasts), in taste cells, in some ganglion cells, and in insulin-producing cells in the pancreas where they modulate insulin secretion. NMDA receptors (and other kinds of glutamate receptors) are expressed in the membrane of unmyelinated axons that lead from nociceptors (pain receptors) in the skin. The functions of glutamate receptors in peripheral tissues are less understood than in the brain.

3 Studies with immunogold labeling indicate that **metabotropic glutamate receptors** (mGluR1) are located at the periphery of synapses, whereas AMPA receptors occupy the central region. In addition, mGluRs are found without relation to synapses (enabling extrasynaptic transmission, see Fig. 5.1). Such segregation might allow the receptors to respond differentially to glutamate: the AMPA receptors would be activated by normal presynaptic stimulation (low frequency of action potentials), whereas mGluRs would be activated only by high-frequency stimulation that releases large amounts of glutamate at the synapse, or by spillover of glutamate from nearby synapses.

Animal experiments have yielded positive results, but so far they have not been confirmed in humans. Excitatory amino acid transmitters and the NMDA receptor may also be involved in the cell damage that occurs in various **neurodegenerative disorders** of the nervous system, such as amyotrophic lateral sclerosis (ALS) and Huntington's disease.

While excessive NMDA-receptor activation can harm neurons, **blockage** can also produce dramatic symptoms. This is exemplified by the drugs **ketamine** (Ketalar, used as a short-acting anesthetic) and **phencyclidine** (PCP, or “angel dust”) that block NMDA receptors. Both drugs influence consciousness and disturb thought processes. Side effects of ketamine used for anesthesia are nightmares and hallucinations during awakening. The thought disturbances resemble those occurring in schizophrenia, and this led to the “**glutamate hypothesis**” for this disease. Ketamine in low doses may be effective for treating chronic pain, probably by blocking NMDA receptors on sensory neurons in the cord. **Alcohol** (ethanol) also influences NMDA receptors (besides many other actions in the nervous system; see later, “GABA Receptors Are Influenced by Drugs, Alcohol, and Anesthetics”).

Glutamate Transporters

Five structurally different glutamate transporter proteins are identified, which also differ with regard to their distribution in the brain. The quantitatively dominant transporters are concentrated in glial membranes apposing neurons, particularly around synapses (see Fig. 4.1; see also Fig. 2.5). It is not unexpected that the concentration of transporters is highest in parts of the brain with a high density of glutamatergic nerve terminals.

The transport of glutamate into glial cells is driven by concentration gradients of Na^+ and K^+ . That is, the electrochemical gradient is crucial for the activity of the transporters. Other factors also influence their activity, however. Thus, the expression of transporter proteins increases with activation of glutamate receptors, while the expression decreases after removal of glutamatergic innervation.

Glutamate Transporters and Brain Damage

Under pathologic conditions with insufficient energy supply (e.g., low blood flow) the electrochemical gradient cannot be maintained. Because of the high intracellular concentration of glutamate, the transporters reverse their direction of transport so that glutamate is released into the extracellular space instead of being removed from it. In this way, the extracellular glutamate concentration can reach levels several hundred times that of the normal resting level. This leads to massive receptor activation and high flow of Na^+ and Ca^{2+}

into the neurons, probably initiating a vicious cycle leading to cell death (see also earlier, “NMDA Receptors: Mediators of Both Learning and Neuronal Damage,” and Chapter 11, under “Ischemic Cell Damage”).

Inhibitory Amino Acid Transmitters: GABA and Glycine

GABA is the dominant inhibitory transmitter, being present in nearly all parts of the CNS (Figs. 5.1 and 5.2). As many as 20% of all synapses in the CNS may be GABAergic. GABA is used as a transmitter mainly by **interneurons** (most projection neurons are glutamatergic). It is synthesized from glutamic acid in a single step by the enzyme **glutamic acid decarboxylase** (GAD, Fig. 5.4). GABA acts on ionotropic **GABA_A receptors**, and metabotropic **GABA_B receptors**. GABA is removed from the extracellular space by high-affinity transporters (GAT), which are mainly localized to neuronal membranes (differing in this respect from glutamate transporters).

GABA appears to play an important role during **development** of the nervous system, and occurs very early—even before synapses are established. When synapses start to occur, GABA acts as an excitatory transmitter because it depolarizes rather than hyperpolarizes the postsynaptic neuron (by acting on GABA_A receptors).⁴ GABA is possibly the first excitatory transmitter to shape neuronal networks (before glutamate has taken over as the dominant excitatory transmitter). In addition, GABA influences proliferation, migration, and maturation of neurons.

Glycine (Fig. 5.4) is also an inhibitory transmitter, although with a much more limited distribution than GABA. Glycine is used as transmitter by a population of spinal interneurons and by some brain stem and cerebellar neurons. **Glycine receptors** are parts of chloride channels and have fast excitatory actions. **Strychnine** blocks glycine receptors (thereby blocking inhibition of spinal and brain stem motor neurons), and this explains why strychnine poisoning produces muscle cramps. Likewise, the **tetanus toxin** provokes violent muscle spasms because it inhibits synaptic release of glycine.

GABA Receptors

In most areas, GABA acts by opening **chloride channels**, thereby producing a brief hyperpolarizing current (IPSP) or short-circuiting excitatory currents (see Chapter 3, “Mechanisms of Postsynaptic Potentials (EPSPs and IPSPs)”). The receptor that forms the

⁴ This is probably due to high intracellular chloride concentration in embryonic neurons. The equilibrium potential for Cl^- is therefore more negative than in mature neurons, so that opening of chloride channels leads to net flow of Cl^- out of the neuron. The same situation appears to arise in the adult spinal cord in certain conditions with chronic pain, that is, GABAergic interneurons may lose their normal inhibitory action on pain transmission.

Cl⁻ channel is termed **GABA_A**; it consists of five membrane-spanning subunits (similar to the acetylcholine receptor shown in Fig. 4.2). There are several subtypes of the GABA_A receptor, as mentioned (see earlier, “Transmitter Receptors in General”). Besides the binding site for GABA, the GABA_A receptor has several others, which are targets of alcohol and several common drugs. GABA is normally present in low concentrations extracellularly, and may bind to **extrasynaptic** GABA_A receptors (Fig. 5.3). This would mediate a tonic, fairly diffuse inhibition (in addition to the phasic one produced at GABAergic synapses). Although the function of this tonic inhibition is unknown, it is modulated by anesthetics, certain drugs, and alcohol.

GABA can also produce **slow IPSP** (long-lasting hyperpolarization) by indirectly opening K⁺ channels or blocking Ca²⁺ channels. The receptors producing these effects are termed **GABA_B**, and are G protein-coupled, metabotropic receptors. GABA_B receptors are found both presynaptically and postsynaptically.⁵ GABA_B receptors inhibit several **reflexes**, such as the spinal stretch reflex (contraction in response to rapid stretch of a muscle) and the cough reflex. Although GABA_B receptors are present in many parts of the CNS, the concentration in most places is much lower than that of GABA_A. Accordingly, blockage of GABA_B receptors produces fewer behavioral effects than does blockage of GABA_A. It is possible that GABA_B receptors are activated only under special circumstances, whereas GABA_A receptors are more continuously active.

GABA Receptors Are Influenced by Drugs, Alcohol, and Anesthetics

Benzodiazepines, barbiturates, and some anesthetics bind to different sites at the GABA_A receptor, but all potentiate the synaptic effect of GABA. The **benzodiazepines** (e.g., diazepam) act by increasing the opening frequency of the Cl⁻ channels, whereas the barbiturates prolong their opening time. Benzodiazepines are used to reduce anxiety and provide muscle relaxation, and some derivatives are used as hypnotics. The **barbiturates** have similar effects and were formerly widely used as sedatives and hypnotics (they are now mainly used to induce general anesthesia and to treat epilepsy). Another drug, **baclofen** (Lioresal), binds selectively to the GABA_B receptor and potentiates the effect of GABA. It is used to reduce abnormal muscle tension occurring after damage to descending motor pathways (see Chapter 22,

under “Spasticity”).⁶ Certain steroid hormones, among them the female sex hormone **progesterone**, also bind to GABA_A receptors and produce actions similar to barbiturates. An anesthetic drug (alphaxalone) was developed on this basis.

The central nervous effects of both **alcohol** and **inhalation anesthetics** (i.e., gases used for general anesthesia, such as halothane) were formerly ascribed to unspecific membrane influences. It now seems, however, that they act mainly by way of receptor binding. Both alcohol and gaseous anesthetics bind to (among others) GABA_A receptors and increase their activity by increasing inhibition. Chronic alcohol consumption down-regulates GABA_A receptors, and this may contribute to the development of tolerance (i.e., the dose must be increased to achieve the same effect) and the heightened excitability by abstinence. As mentioned, alcohol also binds to **NMDA receptors**. There is evidence that chronic alcohol intake leads to up-regulation of NMDA receptors in the frontal lobes, probably because alcohol inhibits NMDA-receptor activation. Alcohol also increases the amount of the transmitter **dopamine** in parts of the brain (especially in the nucleus accumbens), perhaps as a consequence of reduced NMDA-receptor activation (glutamate is believed to reduce dopamine release). Other transmitters, such as serotonin and neuropeptides, as well as several intracellular signal pathways, are influenced by alcohol. Genetic variations in transmitter receptors and enzyme systems may at least partly explain why people react so differently to alcohol.

Acetylcholine

The actions of acetylcholine in the CNS are especially important with regard to **attention**, **learning**, and **memory** (see Fig. 4.10; see also Chapter 26 under “Multiple Pathways and Transmitters Are Responsible for Cortical Activation”).⁷ In **Alzheimer’s disease**—with memory impairment as a key feature—acetylcholine and acetylcholine receptors in the cerebral cortex are severely reduced (see also Chapter 10, under “Alzheimer’s Disease and Frontotemporal Dementia [Pick’s Disease],” and Chapter 31, under “Cholinergic Neurons Projecting to the Cerebral Cortex”).

5 Some **sensory neurons** express GABA_B receptors in their peripheral ramifications, where GABA inhibits release of other transmitters. Peripheral release of neuropeptides can cause increased local blood flow, edema, and stimulation of pain receptors (see Chapter 13, under “Release of Neuropeptides from Peripheral Branches of Sensory Neurons”).

6 It was initially assumed that **baclofen** reduces muscle tension by increasing presynaptic inhibition of sensory nerve terminals in the spinal cord, thereby reducing the depolarization of motor neurons. Recent data indicate that axoaxonic synapses on sensory terminals act mainly on GABA_A receptors, however, and that baclofen acts directly on the motor neurons and interneurons rather than presynaptically.

7 Acetylcholine also influences the **microcirculation** of the brain, by producing vasodilation via muscarinic receptors and release of nitric oxide (see later, “Nitric Oxide and Blood Vessels”). The cholinergic fibers innervating brain vessels arise in the basal forebrain and not in the peripheral parts of the autonomic nervous system.

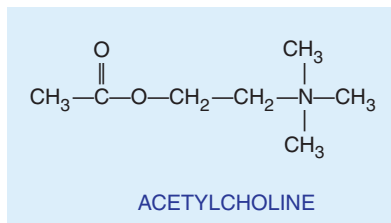


FIGURE 5.6 Structure of the neurotransmitter acetylcholine.

Acetylcholine is a quaternary amine (Fig. 5.6) **synthesized** through the binding of choline to acetyl-coenzyme A by the enzyme **choline acetyltransferase** (ChAT). The presence of this enzyme is characteristic of **cholinergic neurons** (neurons containing acetylcholine, Fig. 5.1). Acetylcholine is present in somatic and autonomic motor neurons in the spinal cord and brain stem and in autonomic (parasympathetic) ganglia. In addition, cholinergic neurons make up several diffusely organized cell groups in the brain stem (parabrachial nucleus) and in the basal forebrain (the basal nucleus and the septal nuclei; see Fig. 10.1).

After release, acetylcholine is broken down to choline and acetate by the enzyme **acetylcholine esterase** (AChE). The enzyme is very efficient: one molecule can hydrolyze 5000 molecules of acetylcholine per second. Choline (but not acetylcholine) is taken up into nerve terminals by high-affinity transporter proteins. Uptake of choline appears to be the most important factor in regulating the synthesis of acetylcholine.

An important feature of cholinergic neurons with axon ramifications within the CNS is that they to only a limited extent release transmitter at typical synapses. In the cerebral cortex, for example, axons from cholinergic neurons form widespread ramifications with **varicosities** (Fig. 5.3). Only about 10% of the varicosities were estimated to form typical synapses when examined via serial sections and electron microscopy. Therefore, acetylcholine must be expected to act rather diffusely on all neurons in the vicinity equipped with the appropriate receptors--that is, it acts mainly via **volume transmission**. This fits with the fact that both nicotinic and muscarinic receptors in the cerebral cortex are localized extrasynaptically on dendrites, neuronal somata, and nerve terminals (presynaptically). The same arrangement holds for other cell groups with widespread axon ramifications that release modulatory transmitters (such as monoaminergic ones, discussed later).

Acetylcholine Receptors (AChRs)

Acetylcholine can bind to two kinds of receptors: ionotropic **nicotinic receptors** (nAChR; see Fig. 4.2), and metabotropic **muscarinic receptors** (mAChR). The names

derive from the early observations that nicotine and muscarine mimicked the effects of acetylcholine (nicotine and muscarine are plant alkaloids; muscarine is present in certain kinds of poisonous mushrooms). The nicotinic receptors produce fast, excitatory synaptic actions, whereas the muscarinic receptors mediate indirect, modulatory effects on neuronal excitability. Depending on the subtype of muscarinic receptor present, the effect may be inhibitory or excitatory. Nicotinic and muscarinic receptors are distributed differently in the nervous system, and there are several subtypes of both (which can be distinguished pharmacologically by use of different agonists and antagonists). For example, different subtypes of nicotinic receptors are expressed in striated muscle cells, in autonomic ganglia, and in the CNS.

Seven **subtypes** of the **nicotinic receptor** have been identified. There are three main groups: receptors in skeletal muscle, in autonomic ganglia, and in the CNS. The functional role of nicotinic receptor in the CNS is not well understood, but they are present in many places and are localized both presynaptically and postsynaptically. They therefore may influence neuronal excitability both directly and indirectly by modulating release of other transmitters (e.g., glutamate). Much better known are the actions of acetylcholine at the cholinergic synapses between motor neurons and **skeletal muscle** cells (see Figs. 21.4 and 21.5). Binding of acetylcholine opens the channel for cations, but the permeability is largest for Na^+ . Opening of such channels elicits an action potential that spreads out in all directions to reach all parts of the muscle cell membrane. This is the signal that leads to muscle contraction. (Whereas skeletal muscle cells contain only nicotinic receptors, **smooth muscle cells** are equipped only with muscarinic receptors.)

Muscarinic receptors are quantitatively the dominant acetylcholine receptors in the CNS. So far, five subtypes (**m1–m5**) have been cloned (all are blocked by atropine) but m1 and m2 are the quantitatively most important ones. Whereas m1 receptors are located postsynaptically, m2 receptors are found predominantly presynaptically. In the **cerebral cortex**, which receives many cholinergic nerve terminals, a major effect of acetylcholine is to reduce the permeability of a K^+ channel by acting on m1 receptors. This makes the neurons more susceptible to other excitatory inputs so that, for example, a neuron will react more easily to a specific sensory stimulus. Another kind of muscarinic receptor opens a K^+ channel, thereby producing long-lasting hyperpolarization, while a third type closes a Ca^{2+} channel.

Blockers of Acetylcholine Receptors

Curare is an antagonist (blocker) of the nicotinic receptors and was used as an arrow poison by South American

natives to paralyze victims. Derivatives of curare are used to achieve muscle relaxation during surgery. **Atropine** and **scopolamine** are relatively unselective antagonists of muscarinic receptors (i.e., they block all subtypes). The snake venom α -**bungarotoxin** binds specifically to muscle nicotinic receptors and blocks the effect of acetylcholine (and kills the victim by paralyzing all skeletal muscles). The French neuroscientist Jean-Pierre Changeux and coworkers achieved the first isolation and characterization of a transmitter receptor by use of α bungarotoxin. To obtain sufficient amounts of the receptor, **electric eels** (*Torpedo*) were chosen for study because they are equipped with electric organs that produce strong electric shocks by activating nicotinic receptors.

Nicotine Addiction

Genetic variability among AchR subunits appears to be related to nicotine dependence; that is, persons with genes for a certain subunit might have an increased risk of becoming nicotine dependent. The development of addictive behavior depends, at least partly, on nicotine-receptor-mediated stimulation of **dopaminergic** neurons (that release dopamine in the nucleus accumbens; see Chapter 23, under “The Ventral Striatum, Psychosis, and Drug Addiction”). However, the relation between nicotine and addictive behavior is complex and several transmitters other than dopamine are involved (e.g., glutamate and GABA). Further, chronic nicotine consumption induces plastic changes in several neuronal groups. At the cerebral cortical level, a region called **insula** (see Chapter 34 under “The Insula”) may be particularly important. For example, activation of neuronal groups in the insula was found in a brain imaging study to increase in relation to the person feeling an urge for a drug. Further, smokers suffering a stroke that damaged the insula were much more likely to quit smoking than were smokers suffering lesions in other parts of the brain.

Biogenic Amines

The **biogenic amines** constitute a subgroup of the small-molecule transmitters. The group includes the **monoamines** norepinephrine, epinephrine, dopamine, and serotonin (one amine group), and in addition **histamine** (two amine groups). Neurons that use monoamines as transmitters are called **monoaminergic**. The monoamines are **synthesized** by enzymatic removal of the carboxylic group from an aromatic amino acid (Figs. 5.7 and 5.8). Norepinephrine, epinephrine, and dopamine are **catecholamines**.⁸ Neurons that contain the monoamines

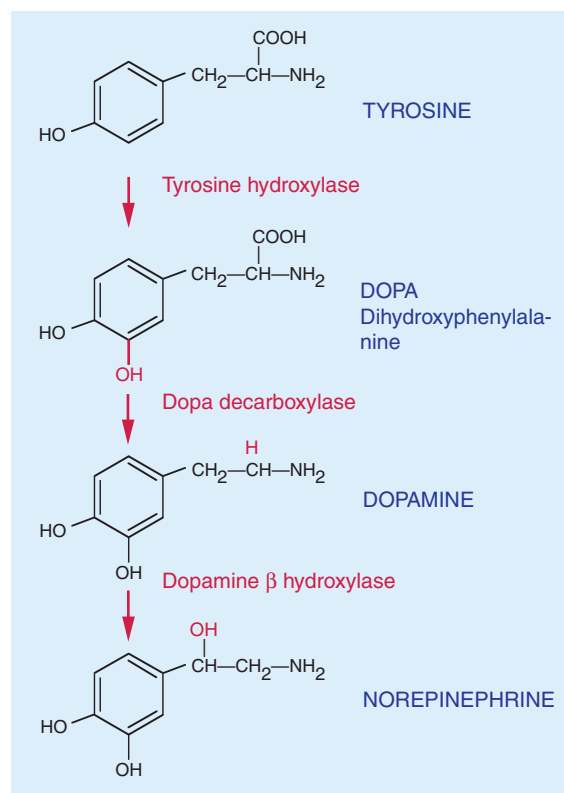


FIGURE 5.7 The catecholamines dopamine and norepinephrine and the key enzymes in their synthesis.

norepinephrine, dopamine, serotonin, and histamine are said to be noradrenergic, dopaminergic, serotonergic, and histaminergic, respectively (the same terminology is used for the receptors corresponding to these transmitters).

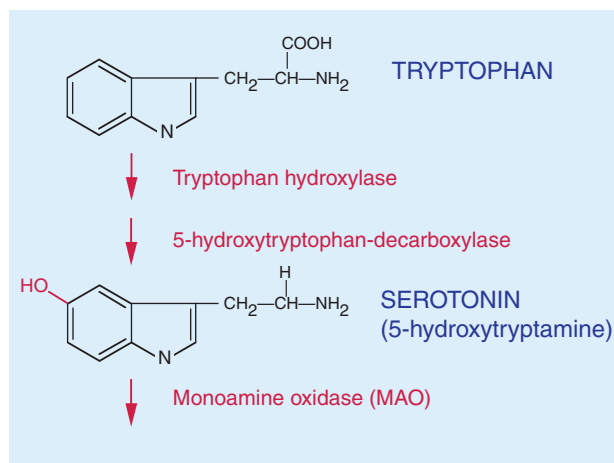


FIGURE 5.8 Serotonin. This neurotransmitter synthesized from tryptophan in two enzymatic steps. It is broken down by monoamine oxidase in the nerve terminals.

⁸ Catecholamines: Compounds consisting of a catechol group (benzene ring with two hydroxyl groups) with an attached amine group.

A common feature of the biogenic amines is that they are synthesized only in a small number of neurons, which, however, have widely branching axons. In this way, these few neurons ensure that the transmitters can act in most parts of the CNS. Like acetylcholine, the biogenic amines are to a large extent released from varicosities without typical synaptic contacts (Fig. 5.3); therefore, their effects are presumably mainly mediated by **volume transmission** and binding to extrasynaptic receptors. Localization and functions of monoaminergic cell groups are discussed in Chapters 23 (dopamine) and 26 (norepinephrine and serotonin).

Actions of the Biogenic Amines

The biogenic amines act (with one exception) on **metabotropic receptors**; that is, they exert mainly slow, **modulatory** actions. Monoaminergic (like cholinergic) neurons are therefore suited to modulate simultaneously the specific information processing mediated by glutamate and GABA in many anatomically separate neuronal groups. This is related to regulation of **attention, motivation, and mood**. In addition, the monoamines play important roles with regard to **plasticity** and **learning**.

Drugs used to treat diseases such as schizophrenia, Parkinson's disease, and depression alter the functioning of monoamines.

Synthesis of Biogenic Amines

Norepinephrine (noradrenaline), epinephrine (adrenaline), and dopamine are all synthesized from the amino acid tyrosine (Fig. 5.7). Tyrosine is taken up from the bloodstream by active transport mechanisms and concentrated in the nervous tissue. The synthesis of catecholamines goes through several enzymatic steps. The first is the conversion of tyrosine to dihydroxyphenylalanine (DOPA) by the enzyme tyrosine hydroxylase (Fig. 5.7), which appears to be rate limiting for the synthesis of catecholamines under most conditions. The activity of tyrosine hydroxylase is regulated by negative feedback from released catecholamines by way of presynaptic autoreceptors (other factors also influence the activity, however). DOPA is converted to dopamine by the enzyme aromatic amino acid decarboxylase (DOPA decarboxylase). The reaction is so rapid that very little DOPA can be detected in the brain normally. Therefore, the synthesis of dopamine can be increased by artificial supply of DOPA (in the form of levodopa, Sinemet, Parcopa), as done in Parkinson's disease in which parts of the brain have very low levels of dopamine. (Dopamine itself does not pass the blood–brain barrier and therefore cannot be used therapeutically.) Norepinephrine is synthesized from dopamine by the enzyme dopamine β -hydroxylase (Fig. 5.7). The detection

of this enzyme in tissue sections is used to identify noradrenergic neurons.

Serotonin (5-hydroxytryptamine [5-HT]) is synthesized from the amino acid **tryptophan** (Fig. 5.8). The reaction is catalyzed by the enzyme **tryptophan hydroxylase** (and the same aromatic amino acid decarboxylase that catalyzes DOPA to dopamine). Tryptophan hydroxylase is rate limiting for the synthesis of serotonin, although other factors also contribute.⁹ The serotonin concentration is much higher in certain cell types peripherally than in the brain (blood platelets, mast cells, chromaffin cells in the gut, and others). It is therefore mandatory that serotonin—like other transmitters—cannot pass the blood–brain barrier. Peripherally, serotonin produces contraction of **smooth muscle cells**, among other effects.

Histamine hardly passes the blood–brain barrier, and is synthesized in the brain from the amino acid **histidine**.

Monoamine Receptors

Like the other neurotransmitters discussed so far, each of the monoamines bind to several different receptors (see Table 5.1). Monoamine receptors are found both presynaptically and postsynaptically (as mentioned, the majority are most likely localized extrasynaptically). Their effects are correspondingly diverse: some monoaminergic receptors inhibit transmitter release, others increase it; some produce slow EPSPs, others produce slow IPSPs. Consequently, each monoamine can have diverse effects, depending on the receptor repertoire of the target neurons.

Norepinephrine (and epinephrine, which mainly functions as a hormone) binds to two main receptor types, α - and β -**adrenergic receptors**. Their effects are often opposite, and they are differentially distributed (these receptors are treated in more detail in Chapter 28). In the cerebral cortex, norepinephrine produces a slow depolarization by binding to β receptors (by indirect, G protein–coupled closure of a K^+ channel).

Dopamine acts on two main types of receptors— D_1 and D_2 —with different properties and different distribution in the CNS. Common to the dopamine receptors in the D_1 group is that they act by increasing the synthesis of cyclic AMP, whereas the D_2 receptors decrease it. Usually, one uses the terms D_1 -like (D_1 and D_5) and D_2 -like receptors (D_2 , D_3 , and D_4). Dopamine receptors are discussed further in Chapter 23 (under “Actions of Dopamine in the Striatum”).

Serotonin receptors fall into seven groups— $5-HT_1$ to $5-HT_7$ —with further subtypes of each (see Table 5.1).

⁹ In contrast to catecholamines, the synthesis of serotonin appears not be regulated by negative feedback by way of autoreceptor stimulation; that is, increased extracellular transmitter concentration does not inhibit the synthesis.

All of the serotonin receptors are metabotropic except the 5-HT₃ receptor, which is ionotropic (belonging to the same protein family as the other ion channel receptors). The metabotropic serotonin receptors act through different intracellular signal pathways and can have either excitatory or inhibitory effects (evoking slow EPSPs and IPSPs). For example, activation of 5-HT₁ receptors causes inhibition by opening certain K⁺ channels. The same neurons may also be equipped with another kind serotonin receptor that closes K⁺ channels, making it difficult to sort out the total effect of serotonin. Serotonin is further discussed in Chapter 13 (under “Nociceptors”), Chapter 22 (“Monoaminergic Pathways to the Brain Stem and Spinal Cord”), and Chapter 26 (under “The Raphe Nuclei” and “Pathways and Transmitters Responsible for Cortical Activation”).

Histamine Receptors—Homeostasis and Wakefulness?

Histamine receptors occur extrasynaptically and on varicosities in many parts of the CNS. In the **thalamus** and the **cerebral cortex**, histamine actions appear to be involved in **arousal** and **wakefulness**. Histamine receptors in the **hypothalamus** are most likely involved in **homeostatic processes** (e.g., food and water intake, temperature regulation, and hormone secretion). The axon ramifications releasing histamine contact not only neurons but also **glial cells** and small **blood vessels**. Presumably, this also relates to homeostatic control.

Three varieties of the histamine receptor—H₁-H₃—have been identified in the CNS; all acting indirectly via G proteins. H₁ receptors mediate the effect of histamine on wakefulness, partly directly on neurons in the cerebral cortex, partly on subcortical cholinergic and monoaminergic neurons, which in their turn influence the cerebral cortex. The **antihistamines** used against travel sickness and allergy are H₁ antagonists, and this probably explains why drowsiness is a frequent side effect of such drugs. Activation of H₁ receptors in the hypothalamus reduces food intake in experimental animals. Many psychoactive drugs have antihistaminergic side effects, which may be one reason why weight gain is common among patients with long-term treatment with such drugs.

Monoamine Removal

Transporter proteins in the membrane of nerve terminals end the transmitter action and control the extracellular concentration of monoamines. Glial cells do not appear to play an important part in uptake of monoamines (in contrast to glutamate). There are specific transporters for norepinephrine, dopamine, and serotonin—all belonging to the same protein family (called NAT, DAT, and SERT/5-HTT, respectively). The two catecholamine transporters have low selectivity,

however, so that, for example, the dopamine transporter also can take up norepinephrine, if it is present in the vicinity. After uptake into nerve terminals, the monoamines are partly transported into vesicles, partly broken down by the enzyme **monoamine oxidase (MAO)**.

Monoamine Oxidase, Serotonin Transporters, and Behavior

There are two varieties of monoamine oxidase (MAO). MAO-A has the highest affinity to the monoamine transmitters and is particularly concentrated in catecholaminergic neurons. MAO-B is concentrated in serotonergic and histaminergic neurons and in glia. Mutation of the gene coding for MAO-A (located on the X chromosome) was found in a Dutch family with lack of MAO-A, and abnormal aggressiveness among the male members. Correspondingly, knockout mice that lack the MAO-A gene behave aggressively and have increased brain levels of monoamines. These and other findings have led some to suggest that individual variations among the genes for monoamine oxidase may dispose for excessive aggressiveness and violent behaviors.

Individual differences in personality and behavior have also been associated with genetic **polymorphism** in monoamine metabolism. For example, certain varieties of the gene coding for the **serotonin transporter** are associated with high levels of anxiety and depression. These traits appear to be associated with a heritable reduced functioning of the serotonin transporter (not increased as one might have expected, because antidepressants inhibit reuptake of monoamines).¹⁰ Further, animal experiments indicate that symptoms in adult animals depend on reduced transporter activity during a short period after birth. Only animals with genetic vulnerability *plus* experience of psychological trauma in this period (such as separation from the mother) developed behavioral disturbances as adults. Therefore, it seems that normal serotonin transmission in early postnatal development is necessary for the development of neuronal networks handling emotions and stress. This assumption is further supported by other animal experiments showing that the presence of the 5-HT_{1A} receptor in early development is necessary and sufficient for normal anxiety-related behavior as an adult, regardless of whether or not the receptor is expressed in the adult animals.

When interpreting findings such as those described here, one should bear in mind that they say more about

10 This seeming paradox can probably be explained by complex compensatory processes initiated by inhibition of the transporter, such as up- and down-regulation of receptors, enzymes, and feedback loops. Therefore, reduced transporter activity does not necessarily lead to increased transmitter activity.

the behavior of a brain with disturbed monoamine functions than about the normal functions of monoamines. Serotonin is, for example, certainly not the substrate of aggression but may—together with many other transmitters—be necessary for normal signal processing in the complex networks that generate and control certain emotions and their behavioral expressions. Further, genetic vulnerability of the kind described above increases the probability of developing certain mental disorders or personality traits, but does not determine their development. How easily a person becomes mentally disturbed as an adult depends on how certain brain networks developed in early childhood. This, in turn, depends on a complex interaction between inherited traits (e.g., variety of the serotonin transporter) and the environment.

Localization of Monoaminergic Neurons

Neurons that synthesize **catecholamines** are largely restricted to some small cell groups in the brain stem, hypothalamus, and peripheral nervous system. Most **noradrenergic neurons** in the CNS occur in somewhat diffuse cell groups in the brain stem reticular formation, the **locus coeruleus** being the largest and most distinct one (see Fig. 26.7). The majority of **dopaminergic neurons** are localized to the mesencephalon, in one large nucleus called the **substantia nigra** (see Fig. 23.5) and more diffuse cells groups in the vicinity (**ventral tegmental area**, VTA). In addition, smaller numbers of dopaminergic neurons are found in the retina, in the olfactory bulb, and the hypothalamus.

Serotonergic neurons lie almost exclusively in a group of nuclei near the midline of the brain stem reticular formation, called the **raphe nuclei** (see Fig. 26.1).

Histamine is synthesized only in the small **tubero-mammillary nucleus** in the hypothalamus.

Modulatory Transmitter “Systems”

We described some features shared by monoaminergic and cholinergic neuronal groups (with the exception of cholinergic motor neurons). First, a small number of neurons send axons to large parts of the CNS—that is, the cell bodies producing the enzymes necessary for transmitter synthesis are very restricted in distribution, whereas the transmitters and their receptors are present almost everywhere. Second, each axon ramifies extensively, and the terminal branches are equipped with numerous **varicosities** (Fig. 5.3). As mentioned, these varicosities only infrequently form typical synapses but release transmitters more diffusely (**volume transmission**), presumably by acting largely on extrasynaptic receptors. Finally, the transmitters act predominantly via **metabotropic** receptors—exerting slow, modulatory effects on neuronal excitability.

These anatomic and physiological features imply that the monoaminergic and cholinergic cell groups do not mediate precise temporal or spatial information. They are well suited, however, to modulate the functions of specific glutamatergic and GABAergic systems, for example, by improving the **signal-to-noise** ratio. In this way, they may increase the precision of information handling in many parts of the brain—for example, in the cerebral cortex during processing of complex cognitive tasks. Their widespread connections furthermore ensure that many neuronal groups receive a similar modulatory input, as would be important for control of consciousness, awareness, different phases of sleep, emotions, motivation, and so forth. Monoamines furthermore set the level of excitability of spinal neurons to control voluntary movements and of neurons that are transmitting specific sensory information. For example, brain stem serotonergic neurons with axonal ramifications in the cord appear to both facilitate movement and inhibit sensory transmission.

Trying to bind together seemingly disparate actions, one might speculate that the modulatory transmitter “systems” ensure that sensory, motor, and cognitive processes are coordinated toward a common goal. The modulatory transmitters would do this by mediating a signal about the **value** of specific events. These speculations are supported by the well-established roles of monoamines and acetylcholine in **synaptic plasticity** (Fig. 4.10).

In spite of their homogeneities, it is an oversimplification to regard each transmitter-specific group as a functional entity and to use terms such as “the serotonin system,” “the dopamine system,” and so on. First, the large number of receptors for each transmitter, with different distributions and effects in the brain, indicate that each transmitter-specific cell group has complex relations to behavior. For example, dopaminergic neurons influence neuronal networks engaged in quite different behavioral tasks. Second, even if localizations are not sharp, each transmitter-specific group contains subgroups that differ in where they send their axons, and from where they receive afferents. For example, the various serotonergic raphe nuclei (see Fig. 26.6) send axons to different parts of the CNS. Third, most or all monoaminergic neurons also contain one or more neuropeptides. The effects obtained by stimulation of one of these cell groups therefore cannot be ascribed to one transmitter alone.

Adenosine Triphosphate and Adenosine

The purines adenosine triphosphate (ATP) and adenosine can exert marked effects on neuronal excitability, both in the CNS and peripherally. Only ATP, however, appears to act as a transmitter (it is, e.g., concentrated in vesicles and its release is calcium dependent).

The effects are mediated by purinoceptors, localized both pre- and postsynaptically. (Many other cell types than neurons express purinoceptors—e.g., smooth muscle cells.) P_1 receptors bind adenosine, while P_2 receptors bind ATP. After release, ATP is enzymatically degraded. The transmitter role of ATP is best known in the autonomic nervous system, where it is usually colocalized with acetylcholine or norepinephrine and. ATP and the “classic” transmitter are found in the same vesicles, and they are released together (see Chapter 28, under “Noncholinergic and Nonadrenergic Transmission in the Autonomic Nervous System”). As a rule, ATP excites neurons and smooth muscle cells by acting on ionotropic receptors. There are, however, examples of inhibitory effects of ATP (probably by way of metabotropic receptors) on smooth muscle cells—for example, in the longitudinal muscle layer of the large intestine. In the inner ear, efferent nerve fibers to the sensory cells (hair cells) release ATP together with acetylcholine, and modulate the sensitivity of the sensory cells.

So far, little is known with certainty about the transmitter role of ATP in the CNS, in spite of the wide distribution of purinoceptors. Some brain stem neuronal groups appear to use ATP as transmitter (notably **locus coeruleus**, where it colocalizes with norepinephrine). It is probably also used as transmitter in a subgroup of **spinal ganglion cells**—that is, sensory neurons conducting impulses from peripheral receptors to the spinal cord (see Chapter 13, under “Primary Sensory Fibers and Neurotransmitters”). A population of **spinal interneurons** seems to release ATP in parts of the cord that receives signals from pain receptors (laminae I and II; see Fig. 6.10). Ionotropic P_2 receptors increase the release of glutamate and substance P from terminals of spinal ganglion cells in the dorsal horn. This may contribute to the heightened excitability of spinal neurons, which is typical of chronic pain conditions.

ATP and purinoceptors mediate signals between neurons and **glial cells**, and may participate in the interaction between the immune system and neurons.

As mentioned, **adenosine** has marked effects on neurons, even though it is unlikely to act as a transmitter. Injection of minute amounts of adenosine inhibits spinal cord neurons that mediate signals from **pain** receptors to the brain. Adenosine also appears to be involved—in some yet unknown way—with the analgesic effects of morphine and the morphine-like substances produced in the brain (opioids).

Purinoceptors

There are two main groups of P_2 receptors. One group (P_{2X}) consists of six **ionotropic** receptors with fast, excitatory action. The channels are most permeable to Ca^{2+} and have a longer opening time than, for example AMPA receptors (but much shorter than NMDA receptors).

The other group (P_{2Y}) consists of seven **metabotropic** receptors. Both groups are widely distributed in the CNS, although the transmitter role of ATP is reasonably certain only in a few areas. The study of ATP as a neurotransmitter has so far been hampered by the lack of specific receptor antagonists. P_1 receptors are blocked by **xanthenes**, such as **caffeine** and **theophylline**. Theophylline inhibits bronchial smooth muscle cells and is used therapeutically in **asthma**.

Nitric Oxide

Nitric oxide (NO) is a gas that diffuses freely through biologic membranes. It functions as a signal molecule in many organs of the body, among them the nervous system. Its functional role in the nervous system is only partly clear, however. Even though NO often is called an “unconventional neurotransmitter,” it does not meet the criteria used to define a transmitter: it is not stored in vesicles, it is not released by calcium-dependent exocytosis, and it does not bind to membrane-bound receptors.

In the **peripheral nervous system**, autonomic nerve fibers release NO that acts on smooth muscle cells. In the CNS, NO has a number of cellular effects and appears to take part in the control of various systems; it also influences behavior. It has quite different effects in the brain depending on its concentration, however: in low concentrations, it functions as a signal molecule that modulates neuronal behavior, whereas it is toxic in higher concentrations (which is perhaps not surprising because NO is a free radical and as such very reactive).

NO is **synthesized** from the amino acid **arginine** by specific enzymes, **NO synthases**, of which there are several different types. Synthesis of NO in nerve terminals is probably induced by the increase of intracellular Ca^{2+} concentration that is triggered by a presynaptic action potential. After synthesis, NO diffuses freely in all directions. It is not only delivered into the synaptic cleft but also enters all cells that are near the nerve terminal. Calculations from the cerebral cortex indicate that NO can diffuse more than 100 μm from its release site and reach about 2 million synapses. The most abundant **NO receptor** is a water-soluble, cytoplasmic enzyme, **guanylyl cyclase**, which controls synthesis of cyclic GMP. This is an intracellular messenger with various effects, such as activating protein kinases and acting directly on ion channels. In this way NO can modulate neuronal excitability and firing frequency.

NO synthases are present in neurons in several parts of the central and peripheral nervous system. In the cerebral cortex, they are found in GABAergic neurons, which, in addition, contain neuropeptides (substance P and somatostatin). In other parts of the brain, NO synthases occur in cholinergic and monoaminergic neurons. Although we do not know the functions of

NO liberated from such neurons, there is some evidence that NO is one of numerous factors that are involved in the induction of **LTP**, and therefore presumably also in neuronal **plasticity**. NO also appears to play a special role in neurons with **rhythmic firing**—such as the brain stem and thalamic neurons that regulate sleep–wakefulness, and the hypothalamic neurons that control bodily functions with daily variations (circadian rhythms). Presumably, NO, due to its fast and wide diffusion, is particularly suited to **synchronize** the activity of many neurons.

In situations with insufficient energy supply (most often due to reduced blood flow), increased intracellular Ca^{2+} concentration leads to increased NO synthesis. This seems likely to contribute to the neuronal damage in such situations (see Fig. 10.2). On the other hand, release of NO might improve the blood supply by producing local vasodilatation.

NO and Blood Vessels

NO was first discovered in endothelial cells and named **endothelium-derived relaxing factor**. In fact, its main effect in most organs is relaxation of smooth muscle cells, causing vasodilatation and increased blood flow. NO released from autonomic (parasympathetic) nerve fibers, for example, is responsible for penile erection. **Nitroglycerin** and similar drugs give vasodilatation by inducing synthesis of NO.

Vessels in the CNS are also affected by NO. There is evidence that NO, together with other signal molecules, mediates the increased **local blood flow** associated with increased neuronal activity (e.g., the blood flow in the cortical motor area increases when the neurons increase their firing during execution of voluntary movements).

Neuropeptides

A large number of neuroactive peptides have so far been identified in the CNS, but many of them were first found in the peripheral nervous system and in the gut. Although there is firm evidence of a transmitter function for only a few neuropeptides, many of them have marked effects on physiological processes and behavior when administered locally in the CNS. Several of the neuropeptides elicit slow inhibitory or excitatory synaptic potentials when administered in minute amounts close to neurons, suggesting a **modulatory** transmitter role. This assumption is supported by the identification of several G protein–coupled neuropeptide receptors. Neuropeptides can also elicit intracellular responses related to **growth** and **development**. Normally, the concentrations of neuropeptides are low in neurons. The synthesis increases, however, when the homeostasis of the nervous system is challenged (e.g., in infections, stroke, trauma). The way neuropeptides are released

may perhaps fit an “emergency” role: it seems that neuropeptide release requires a high firing frequency or burst firing, whereas low-frequency firing suffices to release small molecule (classical) transmitters.

The duration of action appears to be longer for neuropeptides than for other neuromodulators (in spite of the presence of extracellular peptidases). Thus, the half-life after release can be very long (about 20 minutes for vasopressin, for example)—giving ample time for extracellular diffusion. In addition, at least some neuropeptides are released from dendrites. Thus, it seems that the actions of the neuropeptides are, as a rule, rather diffuse. Some of the neuropeptides may thus function more like local hormones than like neurotransmitters, as also suggested by a mismatch between the distribution of receptors for a neuropeptide and nerve terminals containing the neuropeptide.

A characteristic feature of neuropeptides is that they are **colocalized** in nerve terminals with small-molecule (classical) transmitters. Whereas small-molecule transmitters are stored in small, electron-lucent vesicles, neuropeptides are found in larger vesicles with an electron-dense center (**dense-core vesicles**; Fig. 5.9). Two or more peptides may also coexist, and there may be more than one small-molecule transmitter together with the peptides.

Several neuropeptides will be mentioned in subsequent chapters in relation to various cell groups and neuronal systems.

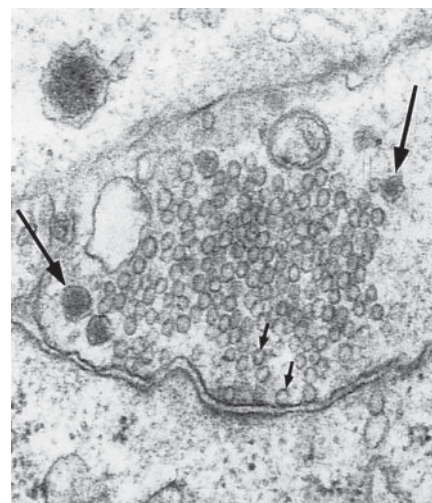


FIGURE 5.9 Nerve terminal containing both a “classical” transmitter and neuropeptides. Large arrows show the dense-core vesicles that contain neuropeptides with transmitter actions. Small arrows show the small, clear vesicles that contain small molecule “classical” transmitters.

Examples of Colocalization of Neuropeptides and Small-Molecule Transmitters

Terminals of nerve fibers innervating **salivary glands** contain both **acetylcholine** and the neuropeptide **vasoactive intestinal polypeptide (VIP)**. Both substances are released when the nerves are stimulated, but they act on different target cells: acetylcholine elicits secretion from glandular cells, whereas VIP produces vasodilatation by relaxing smooth muscle cells, thereby increasing blood flow to the organ at the same time as its secretion is increased. Another example concerns **primary sensory neurons** (spinal ganglion cells; see Figs. 6.9 and 13.16) mediating information from peripheral sense organs to the spinal cord. Some of these—particularly those related to painful stimuli—contain both **glutamate** and the peptides **substance P** and **calcitonin gene-related peptide**. When released with glutamate, these peptides appear to have several actions: they can bind to specific postsynaptic receptors, they can act presynaptically to increase their own release and the release of glutamate, and they can sensitize the postsynaptic glutamate receptors to enhance the effect of glutamate.

Spinal interneurons that receive sensory information exhibit different transmitter combinations. One group of interneurons contains **GABA** and the peptide **enkephalin**. Another group contains both **enkephalin** and **substance P** in combination with (most likely) **glutamate**. Therefore, one neuropeptide (enkephalin) may be colocalized with either an inhibitory or an excitatory transmitter with fast synaptic actions. Apparently, spinal interneurons may express a variety of transmitter combinations with correspondingly complex synaptic actions. When we also think of the variety of receptors each neuron expresses, it is easy to understand why unraveling the functional role of a single transmitter can be difficult.

ACTIONS OF DRUGS ON THE NERVOUS SYSTEM

Most Drugs Influence Synaptic Function

Several drugs have been mentioned in connection with discussions of synaptic function and neurotransmitters. Most drugs acting on the nervous system do so by influencing synaptic transmission directly or indirectly, regardless of whether their aim is to alleviate disorders of mood, cognition, movements, memory, or behavior. The drugs may interfere presynaptically in the synthesis, release, degradation, or reuptake of transmitters or postsynaptically in the activity, numbers, or localization of receptors. Certain drugs have one or more of these actions regarding one or several transmitters. Some drugs also seem to influence synaptic functions indirectly by changing the expression of neurotrophic factors that (among other tasks) govern synaptic plasticity.

Examples: Drugs Altering Monoamine Activity in the Brain

The most common **antipsychotic drugs** (neuroleptics) all block **dopamine receptors**. Their therapeutic effects are primarily related to altering the transmitter functions of dopamine, although they also influence other transmitters (they exhibit, for example, more or less anticholinergic and antihistaminergic effects). In addition, dopamine receptors are present in neuronal groups related to movement control and muscle tone, and this explains why motor dysfunctions are common side effects of antipsychotics. Conversely, patients with Parkinson's disease who are treated with levodopa to increase dopamine levels in the brain may develop psychotic symptoms during treatment. Nevertheless, the "dopamine hypothesis of schizophrenia," which states that the symptoms can be explained as a result of dopaminergic overactivity, is too simplistic. Not all symptoms can be explained in this way, and dopaminergic overactivity does not appear to be present in all patients. Further, psychotic symptoms can also be treated effectively with drugs that block serotonin receptors (so-called atypical antipsychotic drugs, such as **clozapine**, have higher affinity for serotonin receptors than for dopamine receptors). **MAO inhibitors** were the first drugs with significant effects on major **depressions**. Later, **tricyclic antidepressants (TCAs)** were introduced, which inhibit relatively selectively the norepinephrine transporter. The new antidepressants, such as **fluoxetine** (Prozac), inhibit rather selectively the serotonin transporter (**selective serotonin-uptake inhibitors [SSRIs]**). Although the clinical effects of antidepressants may appear to be caused simply by increased levels of monoamines available at the synapses, this cannot be the whole story. For example, while inhibition of the transporters occurs immediately, the clinical effect comes after days or weeks with treatment. Therefore, clinical effects were sought in compensatory processes, such as up- and down-regulation of receptors. Recent research has revealed a number of effects of antidepressants other than altering monoamine levels—for example, on expression of **neurotrophins** and neuronal **plasticity**. The fact that antidepressants, in spite of acting on different monoamine transporters, have similar clinical effects prompted the hypothesis that they act by regulating **transcription** of the same set of genes. This has received some experimental support. For example, one study identified gene transcripts that were equally regulated by treatment with different classes of antidepressants.

Cocaine inhibits the dopamine transporter very selectively; that is, its affinity to the dopamine transporter is much higher than to other monoamine transporters. The dopamine transporter has even been called a "cocaine receptor." **Amphetamine** inhibits the dopamine

transporter but also other monoamine transporters, although relatively weakly compared with cocaine. In addition, amphetamine increases the release of catecholamines and has other synaptic effects as well.

Drug Effects Are Often Multifarious

Most transmitters are present in several parts of the nervous system, which differ anatomically and physiologically. Therefore, it should not be surprising that even drugs that apparently influence the actions of only one transmitter nevertheless have multifarious effects. Some of these are desired therapeutic effects, whereas others are disturbing or even dangerous side effects. Development of more specific drugs—for example, drugs acting on only one receptor subtype—may reduce the side effects but will hardly eliminate them. This is because each receptor type (even one among several subtypes for one transmitter) occurs in functionally different parts of the nervous system.

Drugs Interact with Dynamic Processes in the Brain

Another important point is that storage, release, and uptake of transmitters, as well as the expression of receptors, are **dynamic processes**. The acute perturbations caused by a drug are usually counteracted quickly

by feedback loops that up- or down-regulate the transmitter release, the transporters, the receptors, and so forth. For example, when the activity of **tryptophan hydroxylase** (the rate-limiting enzyme in serotonin synthesis) is reduced, the neurons respond by increasing synthesis of the enzyme and its axonal transport to the nerve terminals.¹¹ Finally, we should keep in mind that alteration of one transmitter's activity as a rule leads to alterations of other transmitters as well. One mechanism behind this is the action on presynaptic receptors: a cholinergic terminal in the cerebral cortex may be equipped with serotonin receptors, so that a drug with specific action on serotonergic transmission may nevertheless also alter cholinergic transmission. Another mechanism is postsynaptic: many dopaminergic neurons receive glutamatergic synapses, so that drugs altering the action of glutamate will change dopamine release as well.

¹¹ **Lithium**, used prophylactically for major bipolar disorder (manic depression), increases tryptophan uptake and initially increases serotonin synthesis (this is only one of several cellular effects of lithium). After 2 to 3 weeks with treatment, however, the tryptophan uptake is still increased, but the tryptophan hydroxylase activity is reduced so that the rate of serotonin synthesis is normal. It is possible that the therapeutic effect of lithium is due to stabilization of serotonin metabolism, which makes it less vulnerable to psychological stress or spontaneous chemical changes in the brain.

6 | Parts of the Nervous System

OVERVIEW

In this chapter, we describe the main features of the anatomy of the central nervous system (CNS), with brief mention of the functional significance of the various parts. We will treat structure and function of many of these parts in more depth in later chapters dealing with functional systems. It will then be assumed, however, that the reader is familiar with the names and the locations of the major cell groups and tracts of the CNS.

The CNS can be subdivided anatomically into different parts. The **spinal cord** lies in the vertebral canal, whereas the **brain** is located in the cranial cavity. The brain is further subdivided into the **brain stem**, which constitutes the upward continuation of the spinal cord, the **cerebellum** (“little brain”), and the **cerebrum** or **cerebral hemispheres**. The cerebellum and cerebrum largely cover the brain stem and constitute the major part of the brain in higher mammals and particularly in humans.

The **spinal cord** consists of a central region with gray matter surrounded by white matter. The **gray matter** contains neurons that may be subdivided based on where they send their axons. **Motor neurons** send their axons out of the cord to reach muscles and glands. **Sensory neurons** receive signals from sense organs in the body and transmit this information to the brain. The spinal **interneurons** or **propriospinal** neurons ensure communication among neurons in the spinal cord. The **white matter** contains ascending tracts carrying signals to higher levels, and descending tracts enabling the brain to control spinal cord neurons (e.g., motor neurons). In addition, many propriospinal axons interconnect neurons in different parts of the cord. The cord consists of **segments**, each giving rise to a pair of **spinal nerves**. The spinal nerves and their peripheral branches transmit sensory signals from the tissues of the body and motor signals to muscles and glands. Each spinal nerve connects with the cord through a dorsal and a ventral **nerve root**. The **dorsal roots** carry only sensory nerve fibers, and the cell bodies of the sensory neurons of each root form a **spinal ganglion**, which appears as an ovoid enlargement of the dorsal root. Functionally, the cord enables fast, automatic responses to signals from the body (e.g., withdrawal of the hand from a hot object). Nevertheless, most tasks carried out by the cord are controlled or modulated by higher levels of the CNS.

The **brain stem** is a rostral continuation of the spinal cord but has a more complex internal organization. On a purely anatomical basis, the brain stem is divided (starting caudally) into the **medulla oblongata**, the **pons**, the **mesencephalon**, and the **diencephalon** (often, however, we do not include the diencephalon). Twelve **cranial nerves** emanate from the brain stem being numbered from rostral to caudal. With some exceptions, the cranial nerves innervate structures in the head (“spinal nerves of the head”) and contain sensory and motor nerve fibers. The first cranial nerve—the **olfactory nerve**—serves the sense of smell. The second—the **optic nerve**—transmit signals from the retina, while the third (**oculomotor**), fourth (**trochlear**), and sixth (**abducens**) nerves control the movements of the eye ball. The fifth nerve—the **trigeminal**—emanates from the pons and brings sensory signals from the face as well as motor signals to the masticatory muscles. The seventh nerve—the **facial**—innervates the mimic muscles of the face. The eighth nerve—the **vestibulocochlear**—carries signals from the vestibular apparatus (recording movements and positions of the head in space) and the cochlea (recording sound waves). The ninth nerve—the **glossopharyngeal**—is concerned mainly with sensations and movements of the pharynx, including taste impulses from the back of the tongue. The tenth nerve—the **vagus** nerve—participates in the innervation of the pharynx but in addition innervates the larynx, and sends (autonomic) motor signals to the heart, the lungs, and the most of the gastrointestinal tract. The eleventh nerve—the **accessory**—innervates two muscles in the neck (the trapezius and the sternocleidomastoid). The twelfth nerve—the **hypoglossal**—is the motor nerve of the tongue. The cranial nerves arise in **cranial nerve nuclei**; some of which are sensory, others are motor.

Diffuse collections of neurons in between the cranial nerve nuclei are collectively termed the **reticular formation**. The reticular formation consists of neuronal groups with different tasks. Some groups are concerned with control of **circulation** and **respiration**; some regulate **sleep** and **wakefulness**, while others control **eye movements**.

The thalamus and the hypothalamus make up the bulk of the **diencephalon**. The **thalamus** is a large, egg-shaped collection of nuclei in the centre of the cerebral hemispheres. The thalamic nuclei transmit sensory information (of all kinds except smell) to the cerebral cortex. In addition, thalamic nuclei transmit information

to the cerebral cortex from subcortical motor regions, notably the basal ganglia and the cerebellum. The **hypothalamus** is concerned mainly with the control of autonomic and endocrine functions that serve to maintain bodily homeostasis (e.g., circulation, digestion, and temperature control).

The **cerebral cortex** is a folded sheet of gray matter covering the cerebral hemispheres. It consists of six layers of neurons, each layer characterized by the morphology and connectivity of its neurons. In addition, the cortical mantle is divided into many **areas**, differing with regard to, among other things, their thalamic connections. Even though each area is to some extent specialized, most tasks of the cerebral cortex—whether they are motor, sensory, or cognitive—are carried out by **distributed networks** interconnecting neurons in many areas of the cerebral cortex. Among the many descending tracts from the cerebral cortex, the **pyramidal tract** targets motor neurons in the spinal cord and brain stem. The **corpus callosum**, consisting of commissural fibers, enables cooperation between the two cerebral hemispheres.

The **basal ganglia** consist of several large nuclei in the interior of the cerebral hemispheres. The **striatum** consists of **putamen** and the **caudate nucleus**, and receive its main afferents from the cerebral cortex. The striatum send efferents to the **globus pallidus** and the **substantia nigra**. From these, signals are directed to the brain stem, and back to the cerebral cortex via the thalamus. By influencing motor neuronal groups in the frontal lobe of cerebral cortex (and in the brain stem), the basal ganglia contribute to the **control of movements**. Connections with other frontal areas in cerebral cortex (and certain subcortical nuclei) enable the basal ganglia to influence **cognitive functions**.

The **cerebellum**, situated dorsal to the brain stem in the posterior cranial fossa, consists of a thin sheet of highly folded gray matter, and a group of centrally located, **deep cerebellar nuclei**. The cerebellum is divided anatomically into a narrow middle part called the **vermis**, and more bulky lateral parts called the **cerebellar hemispheres**. In addition, a deep fissure divides the cerebellum into an **anterior lobe** and a **posterior lobe**. These anatomic subdivisions correspond largely to differences with regard to connections. Thus, the vermis has reciprocal connections with the spinal cord and motor nuclei in the brain stem, whereas the cerebellar hemispheres are reciprocally connected with the cerebral cortex. These connections enable the cerebellum to play a decisive role in **coordination** of voluntary movements by acting on motor neurons in the cerebral cortex, the brain stem, and the spinal cord.

Some Anatomic Terms Used in this Book

The terms **medial**, toward the midline, and **lateral**, away from the midline, are used to describe the relative

position of structures in relation to a midsagittal plane of the body. The terms **cranial** (or **rostral**), toward the head or nose, and **caudal**, toward the tail, are used to describe the relative position of structures along a longitudinal axis of the body. Thus, for example, nucleus A in the brain stem lies medial to and rostral to nucleus B, which in turn lies lateral to and caudal to A. The terms **ventral** and **dorsal** are used to describe the relative position of structures in relation to the front (*venter* means belly) and the back (*dorsum*) of the body, respectively. **Anterior** (front) and **posterior** (rear) are used interchangeably with ventral and dorsal, except for the human forebrain, where anterior means toward the nose and ventral means toward the base of the skull.

The Living Human Brain Can Be Studied with Computer-Based Imaging Techniques: CT, MRI, and DWI

Techniques for computer-based image analysis of the living human brain have revolutionized the possibilities for localizing disease processes in the brain and for studying normal structure and function.

The first of the imaging techniques that enabled us to identify smaller parts of the living brain is **computer tomography (CT)**. This method makes it possible to see X-ray pictures of thin slices through the brain. The examiner may choose the plane of sectioning. CT affords much more precise visualization of brain structures than conventional X-ray examination, which includes all tissue between the X-ray tube and the film. It also provides good visualization of the ventricular system, which previously could be visualized only by replacing the cerebrospinal fluid with air and then making an X-ray examination. CT can also visualize the distribution of a radioactive substance in the living brain, enabling the study of the distribution of neuroactive substances and also the comparison of blood flow in different parts of the brain.

Magnetic resonance imaging (MRI) represents a further technical development. This technique is based not on X-rays but on signals emitted by protons when they are placed in a magnetic field. Depending on the proton concentration in different tissue components, the pictures may show clearly, for example, the contrast between gray and white matter (Figs. 6.1 and 6.28). The bone of the skull gives very little or no signal and is seen as black in the pictures. With this technique, the brain can be visualized in slices with a resolution not far from that of a corresponding section through a fixed brain. Areas with changes in the tissue—for example, infarction, bleeding, or tumor—can be identified. In addition, blood vessels can be visualized to advantage (see Fig. 8.3 and 8.8). Apart from the diagnostic advantages, the MRI technique also improves the correlation of the functional disturbances with the actual damage

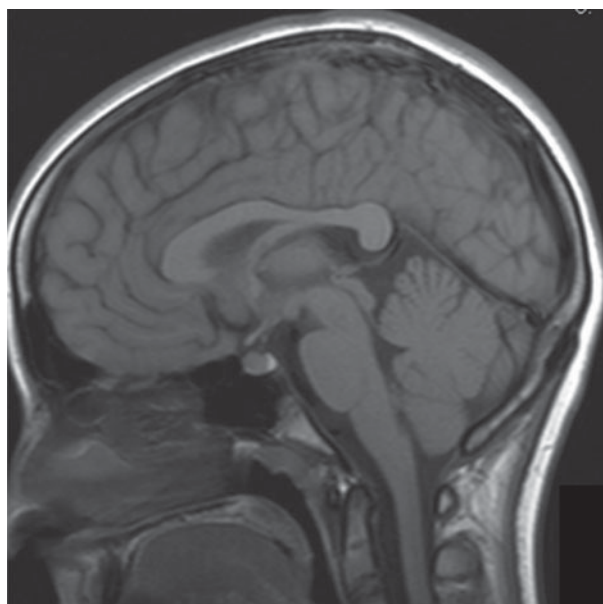


FIGURE 6.1 *Magnetic resonance image (MRI).* This is taken at a level corresponding to the drawing in Fig. 3.1. Most of the structures seen in Fig. 3.1 can be identified in this picture. (Courtesy of Dr. S. J. Bakke, Rikshospitalet University Hospital, Norway.)

of the brain. MRI can also be used to study dynamic processes in the brain (see Chapter 14, under “Methods to Study Neuronal Activity and Metabolism in the Living Human Brain”).

Further developments of the MRI technology have added new applications. For example, with **diffusion-weighted imaging (DWI)** a region devoid of blood supply can be detected only minutes after occlusion of an artery (much earlier than with conventional functional [fMRI]). Further, DWI also enables study of myelination during normal brain development, and even the visualization of major fiber tracts.

THE SPINAL CORD

In humans, the spinal cord is a 40 to 45 cm-long cylinder of nervous tissue of approximately the same thickness as a little finger. It extends from the lower end of the brain stem (at the level of the upper end of the first cervical vertebra) down the vertebral canal (Figs. 6.2 and 6.3) to the upper margin of the second lumbar vertebra (L_2). Here the cord has a wedge-shaped end called the **medullary conus** (or simply conus). In **children**, the spinal cord extends more caudally, however, and reaches to the third lumbar vertebra in the newborn. This difference between the position of the lower end of the spinal cord in adults and infants is caused by the vertebral column growing more rapidly than the spinal cord. In early embryonic life, the neural tube and the

primordium of the vertebral column are equally long (see Fig. 9.12).

The spinal cord is somewhat flattened in the antero-posterior direction and is not equally thick along its length. In general, the thickness decreases caudally, but there are two marked **intumescences** (Fig. 6.3): the **cervical and lumbar enlargements** (intumescencia cervicalis and lumbalis). The intumescences supply the extremities with sensory and motor nerves, hence the increased thickness.

In the midline along the anterior aspect of the cord, there is a longitudinal furrow or fissure, the **anterior (ventral) median fissure** (Fig. 6.4). Some of the vessels of the cord enter through this fissure and penetrate deeply into the substance of the cord. At the posterior aspect of the cord, there is a corresponding, but shallower, furrow in the midline—the **posterior (dorsal) median fissure**. In addition, on each side there are shallow, longitudinal sulci anteriorly and posteriorly—the **anterior and posterior lateral sulci**. These laterally

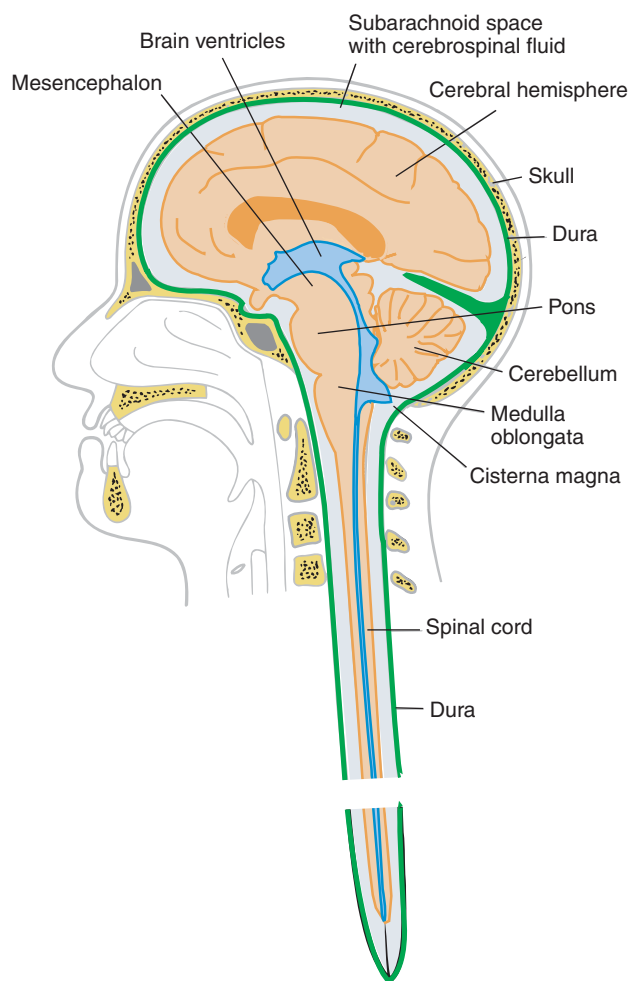


FIGURE 6.2 *The central nervous system, as viewed in a midsagittal section.*

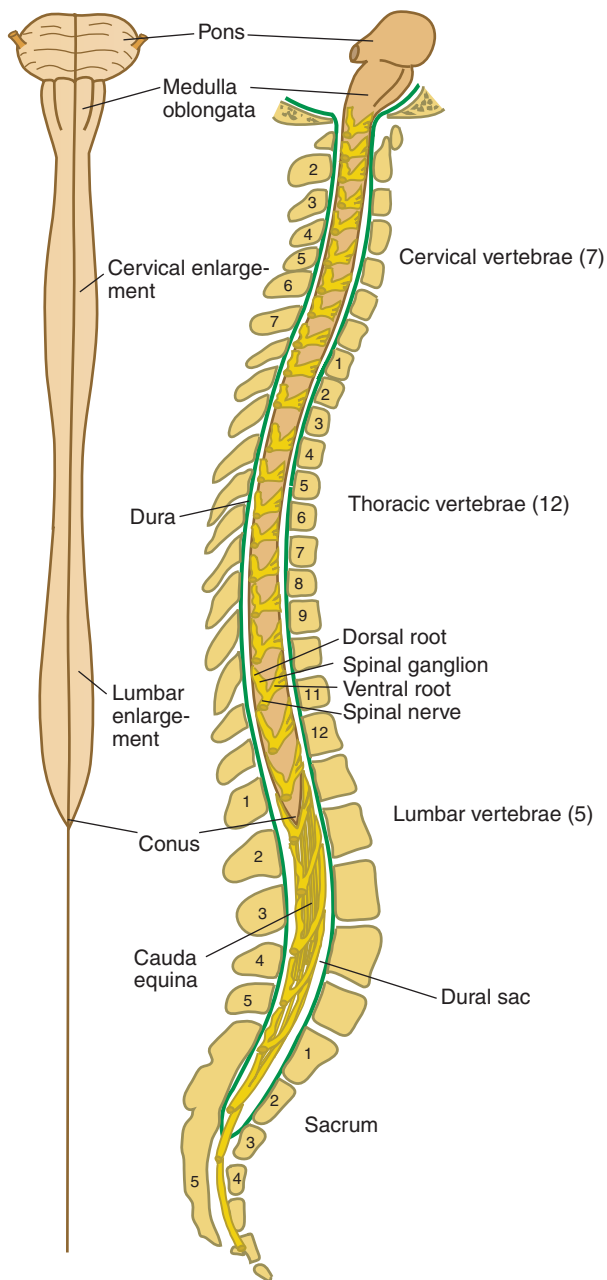


FIGURE 6.3 *The spinal cord.* **Left:** The cord from the ventral side, with the cervical and lumbar enlargements. **Right:** The cord and the vertebral column from the side, but the right halves of the vertebrae have been removed to expose the vertebral canal with its contents. Below the first and second lumbar vertebrae, the vertebral canal contains only nerve roots (the cauda equina), which then unite to form the spinal nerves.

placed sulci mark where the spinal nerves connect with the cord.

The **color** of the spinal cord is whitish because the outer part consists of axons, many of which are myelinated. The **consistency** of the cord, as of the rest of the CNS, is soft and jellylike.

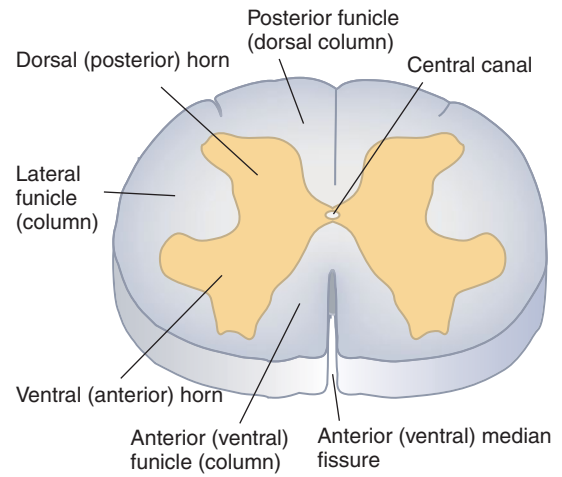


FIGURE 6.4 *Cross section of the spinal cord at a lumbar level.* There is a central H-shaped region of gray matter, and the surrounding white matter is subdivided into funiculi.

Spinal Nerves Connect the Spinal Cord with the Body

Axons mediating communication between the CNS and other parts of the body make up the **peripheral nerves**. The axons (nerve fibers) leave and enter the cord in small bundles called **rootlets** (Fig. 6.5). Several adjacent rootlets unite to a thicker strand, called a root or **nerve root**. In this manner, rows of roots are formed along the dorsal and ventral aspects of the cord: the **ventral (anterior) roots** and the **dorsal (posterior) roots**, respectively. Each dorsal root has a swelling, the **spinal ganglion**, which contains the cell bodies of the sensory axons that enter the cord through the dorsal root (Figs. 6.3 and 6.5). A dorsal and a ventral root unite to form a **spinal nerve**. The spinal ganglion lies in the

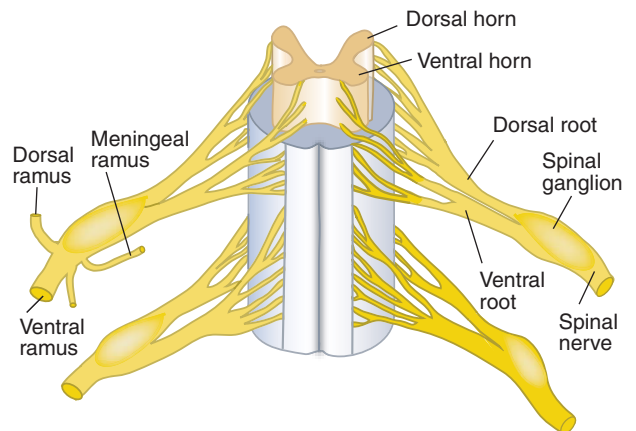


FIGURE 6.5 *Two segments of the spinal cord, as viewed from the ventral aspect.* In the upper part, the white matter has been removed. The dorsal and ventral roots emerge from the posterior and anterior lateral sulci, respectively, and unite to form spinal nerves. The spinal ganglion is located at the site where the roots unite.

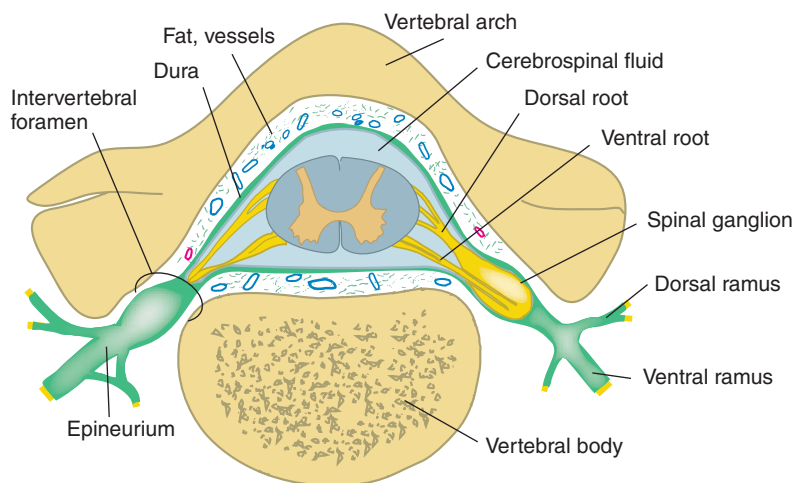


FIGURE 6.6 Cross section of the vertebral column showing the positions of the spinal cord and the spinal nerves. The spinal ganglion is located in the intervertebral foramen. The spinal cord is surrounded by the cerebrospinal fluid contained within the dura. Outside the dura there is fat and a venous plexus, which also serve as soft padding for the cord and the spinal nerves.

intervertebral foramen just where the dorsal and ventral roots unite (Fig. 6.6). There is an important functional difference between the ventral and dorsal roots: the ventral roots consist of efferent (motor) fibers, and the dorsal roots of afferent (sensory) fibers.

In total, 31 spinal nerves are present on each side, forming symmetrical pairs (Fig. 6.3). They all leave the vertebral canal through the intervertebral foramina on each side. As mentioned, the ventral and dorsal roots unite at the level of the intervertebral foramen to form the spinal nerves. The spinal nerves are numbered (as a general rule) in accordance with the number of vertebrae above the nerve. We therefore have 12 pairs of **thoracic nerves**, 5 pairs of **lumbar nerves**, and 5 pairs of **sacral nerves**. In humans, there is only 1 pair of coccygeal nerves. There are seven cervical vertebrae but 8 pairs of **cervical nerves** because the first cervical nerve leaves the vertebral canal above the first cervical vertebra (Fig. 6.3). Therefore, the numbering of the cervical nerves differs from the numbering of the other spinal nerves.

The spinal cord extends caudally only to the level of between the first and the second lumbar vertebrae. Whereas the upper spinal nerves pass approximately horizontally from the cord to the intervertebral foramen, the lower ones have to run obliquely downward in the vertebral canal to reach the corresponding intervertebral foramen, and the distance between the site of exit from the cord and the site of exit from the canal increases steadily (Fig. 6.3). Below the conus, the vertebral canal contains only spinal nerve roots running longitudinally. This collection of dorsal and ventral roots is called the **cauda equina** (the horsetail).

The Spinal Cord Is Divided into Segments

The part of the spinal cord that gives origin to a pair of spinal nerves is called a **spinal segment**. There are,

therefore, as many segments as there are pairs of spinal nerves. They are numbered accordingly, the first cervical segment giving origin to the first cervical nerves, and so on. There are no surface markings of the cord to indicate borders between the segments, but the rootlets nevertheless outline them rather precisely (Fig. 6.5).

The cervical **enlargement** (intumescence) corresponds to the fourth cervical (C_4) segment through the second thoracic (T_2) segment, the lumbar enlargement to the

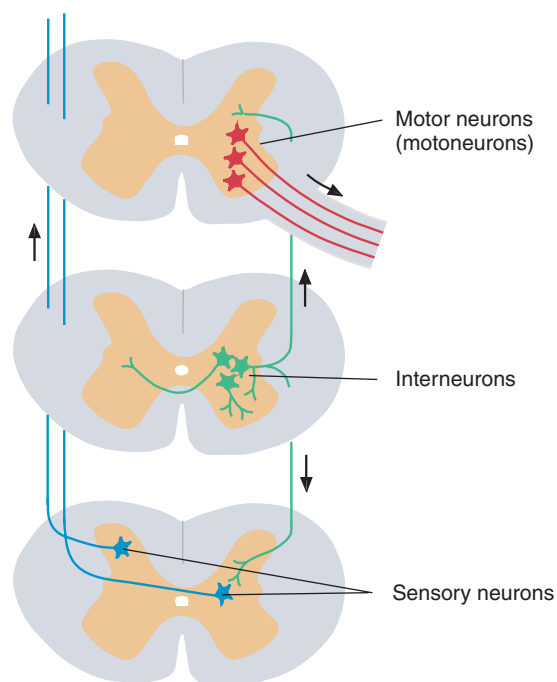


FIGURE 6.7 Three main types of neurons in the spinal cord. Schematic of neuronal types classified in accordance with where their axons terminate: *Motor neurons* supply skeletal muscles, smooth muscles, and glands. The *interneurons* ensure communication among neurons in the cord, and the *sensory neurons* send their axons to higher levels of the central nervous system. (See also Fig. 6.9.)

first lumbar (L_1) segment through the second sacral (S_2) segment.

The difference in rostrocaudal level between the spinal segments and the exit from the vertebral canal of the spinal nerves is of practical importance. Thus, identical symptoms may be provoked by a process close to the cord at one level and by one close to the intervertebral foramen at a considerably lower level (as should be clear from the preceding description; however, this does not concern the nerves in the cervical region).

The Spinal Cord Consists of Gray and White Matter

When cut transversely, the fresh spinal cord can be seen to consist of an outer zone of white matter and a central, H-shaped region of **gray matter**. The arms of the H, extending dorsally and ventrally, are called the **dorsal horn** and **ventral horn**, respectively (Figs. 6.4 and 6.8). The gray matter extends as a column through the length of the spinal cord (Fig. 6.5). The **central canal** is seen as a narrow opening in the center of the cord (Fig. 6.4). The central canal ends blindly in the caudal end of the cord, whereas it continues rostrally into the ventricular system of the brain (Fig. 6.1).

The **white matter** of the cord contains axons running longitudinally. Some of these axons convey signals from the cord to higher levels of the CNS; others, from higher levels to the cord. Finally, a large proportion of the fibers serve the signal traffic, and hence cooperation, between the segments of the cord. Because the first two groups of axons become successively more numerous in

the rostral direction, the proportion of white to gray matter increases from caudal to rostral.

The white matter is divided into **funicles**, or **columns**, by drawing lines in the transverse plane from the sulci on the surface of the cord to the center (Fig. 6.4). Thus, in each half of the cord, the white matter is divided into a **ventral** or **anterior funicle** (funiculus), a **lateral funicle**, and a **dorsal** or **posterior funicle**. For the latter, the term **dorsal column** is used most frequently.

The Spinal Gray Matter Contains Three Main Types of Neurons

Among neurons in the gray matter of the spinal cord, there are both morphological and functional differences. Three main types may be identified according to where they send their axons (Fig. 6.7):

1. Neurons sending their axons out of the CNS
2. Neurons sending their axons to higher levels of the CNS (such as the brain stem)
3. Neurons sending their axons to other parts of the spinal cord

Often neurons of the same kind lie together in the gray matter of the cord. We will now consider in some detail each of these three groups.

Efferent Fibers from the Cord Control Muscles and Glands

The cell bodies of the first kind of neuron listed above are located in the ventral horn and at the transition between the dorsal and ventral horns. The **somatic motor neurons** or **motoneurons** have large, multipolar cell bodies and are located in the ventral horn proper (Fig. 6.8; see also Figs. 1.2 and 21.3). The dendrites extend for a considerable distance in the gray matter (Fig. 6.12). The axons leave the cord through the ventral root, follow the spinal nerves, and end in skeletal muscles (muscles that are controlled voluntarily). The motoneurons are discussed further in Chapter 21.

There is also another group of neurons that sends its axons out of the cord through the ventral root—the **autonomic motor neurons**. These supply smooth muscles and glands with motor signals. They belong to the autonomic nervous system, which controls the vascular smooth muscles and visceral organs throughout the body. These neurons are termed **preganglionic** because they send their axons to a ganglion (see Figs. 28.1 and 28.2). The cell bodies lie in the **lateral horn** (Fig. 6.8). Most of them form a long, slender column, the **intermediolateral cell column**. This column is present only in the thoracic and upper two lumbar segments of the cord and belongs to the sympathetic part of the autonomic nervous system. A corresponding, smaller group of neurons is present in the sacral cord (S_3 – S_4) and

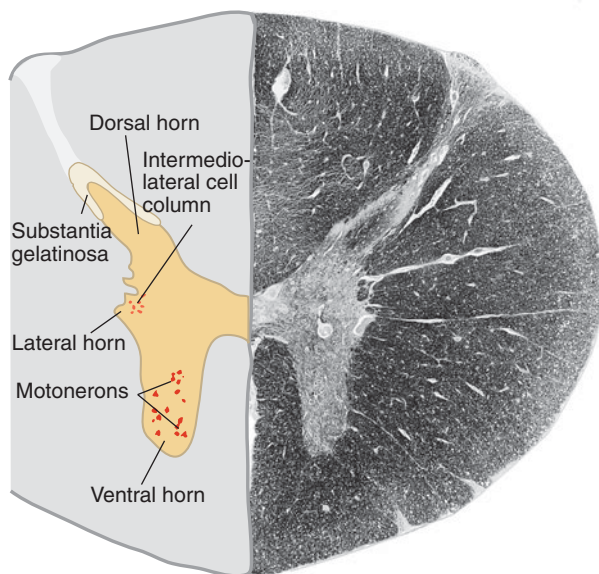


FIGURE 6.8 Cross section of the spinal cord at the thoracic level. Photomicrograph of a section stained so that myelinated axons appear dark. The large motoneurons in the ventral horn have also been stained (Nissl staining) and are just visible. Owing to shrinkage, the motoneurons are surrounded by a clear zone.

belongs to the parasympathetic part of the autonomic nervous system.

Both the somatic and the autonomic motor neurons are under synaptic influence from higher levels of the CNS.

Sensory Neurons in the Cord Are Influenced from the Dorsal Roots and Convey Signals to the Brain

The second main type of spinal neuron sends axons to higher levels of the CNS. Their cell bodies lie mainly in the dorsal horn and in the transition zone between the dorsal and ventral horn (Figs. 6.7 and 6.9). Their job is to inform the brain of the activities of the spinal cord, and especially about what is going on in the body. To fulfill the latter task, the neurons must receive signals from **sense organs**—receptors—in various parts of the body (in the skin, muscles, viscera, and so on). **Sensory**, or **afferent**, nerve fibers conducting impulses from the receptors enter the spinal cord through the dorsal roots and ramify, forming terminals in the gray matter of the cord (Fig. 6.9; see also Fig. 13.12). The sensory neurons have their cell bodies in the **spinal ganglia** (Fig. 6.5; see also) and are therefore called **spinal ganglion cells** (see Fig. 13.13). These are morphologically special, as they have only one process, which divides shortly after leaving the cell body: One peripheral process connects with the sense organs, and the other extends centrally and enters the spinal cord (pseudounipolar neuron; see Fig. 1.5). In accordance with the usual definition of

axons and dendrites, the peripheral process (conducting toward the cell body) should be called a dendrite, whereas the central process is an axon. Both processes are, however, morphologically and functionally axons (e.g., by conducting action potentials and by being myelinated).

The dorsal root fibers form synaptic contacts—in part directly, in part indirectly through the interneurons—with neurons in the spinal cord, sending their axons to various parts of the brain. Such axons, destined for a common target in the brain, are grouped together in the spinal white matter, forming **tracts** (Latin: *tractus*). Such tracts are named after the location of the cell bodies and after the target organ. A tract leading from the spinal cord to the cerebellum, for example, is named the “spinocerebellar tract” (tractus spinocerebellaris).

Interneurons Enable Cooperation between Different Cell Groups in the Spinal Cord

The axons of the third type of spinal neuron do not leave the spinal cord. Usually, the axons ramify extensively and establish synaptic contacts with many other neurons in the cord, within the segment in which the cell body is located, and in segments above and below (Fig. 6.7). Such neurons are called **spinal interneurons**, to emphasize that they are intercalated between other neurons.¹ Many spinal interneurons are found in an intermediate zone between the dorsal and ventral horns, where they receive major synaptic inputs from sensory fibers in the dorsal roots. Many of these interneurons establish synaptic contacts with motoneurons in the ventral horn, thus mediating motor responses to sensory stimuli. Most interneurons, however, receive additional strong inputs from other spinal interneurons and from the brain.

As mentioned, spinal interneurons also send collaterals to terminate in segments of the cord other than the one in which their cell body and local ramifications are found. Such collaterals enter the white matter, run there for some distance, and reenter the cord at another segmental level, to ramify and establish synaptic contacts in the gray matter (Fig. 6.7). Axons of this kind in the white matter are called **propriospinal fibers** (i.e., fibers “belonging” to the spinal cord itself), to distinguish them from the long ascending and descending fibers that connect the cord with the brain.

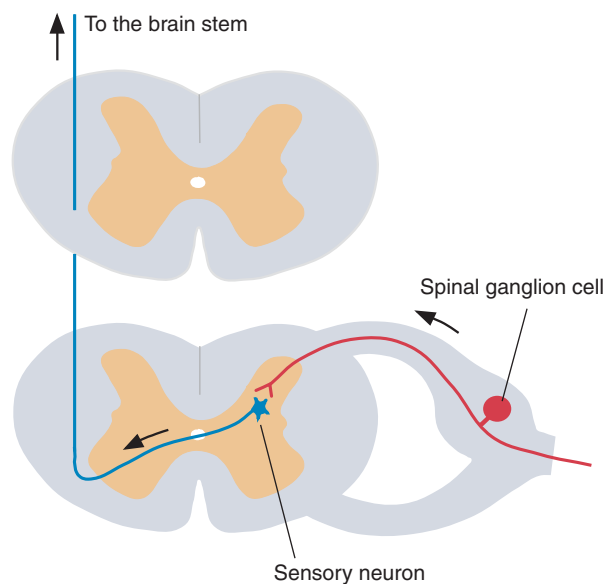


FIGURE 6.9 Sensory neuron in the gray substance of the spinal cord. The neuron, which sends its axon to the brain stem, is synaptically contacted by sensory afferents that enter the cord through the dorsal root (pseudounipolar ganglion cell). The presentation is very simplified; in reality, every sensory neuron is contacted by numerous dorsal root afferents.

¹ Strictly speaking, most neurons in the CNS are interneurons according to this definition—including, for example, spinal neurons with axons ascending to the brain. In practice, the term “interneurons” is nevertheless restricted to neurons with an axon ramifying near the cell body, thus synaptically coupling neurons within one nucleus.

Propriospinal neuron is, therefore, another term used for a spinal interneuron.²

Propriospinal fibers provide opportunities for cooperation among the spinal segments. Most of movements necessitate close coordination of the activity in many segments, each controlling different groups of muscles. Some propriospinal connections are very long and interconnect segments in the cervical and lumbar parts of the cord that control muscles in the forelimb and hindlimb, respectively. Cooperation between the forelimbs and hind limbs is necessary, for example, during walking.

Spinal interneurons are discussed further in Chapter 13, under “Sensory Fibers Are Links in Reflex Arcs: Spinal Interneurons,” and in Chapter 22, under “The Pyramidal Tract Can Open and Close Spinal Reflex Arcs.”

The Spinal Gray Matter Can Be Divided into Zones Called Rexed's Laminae

Systematic observations with the microscope of transverse sections of the spinal cord stained to visualize cell bodies show that neurons with different sizes and shapes are also differently distributed (Fig. 6.10; see also Fig. 13.16). Essentially, neurons with common morphological features are collected into transversely oriented bands or zones. What appear as bands in the transverse plane are, three-dimensionally, longitudinal slabs or sheets. This laminar pattern is most clear-cut in the dorsal horn, whereas in the ventral horn neurons are collected into regions that form longitudinal columns rather than plates (see Fig. 21.2). Nevertheless, the columns in the ventral horn, as well as the slabs in the dorsal horn, are termed **laminae**. This pattern was first described in 1952 by the Swedish neuroanatomist Bror Rexed and has since proved to be of great help for investigations of the spinal cord. Altogether, Rexed described 10 laminae; **laminae I–VI** constitute the dorsal horn, whereas **lamina IX** is made up of columns of motoneurons in the ventral horn. **Lamina II (substantia gelatinosa;** Figs. 6.8, 6.10, and 6.11) is of particular importance for the control of signals from pain receptors, and thus how much a painful stimulus hurts. **Lamina VII** constitutes the transition between the dorsal and ventral horns and contains mainly interneurons. **Lamina VIII**, located medially in the ventral horn, contains many neurons that send axons to the other side of the cord (commissural fibers).

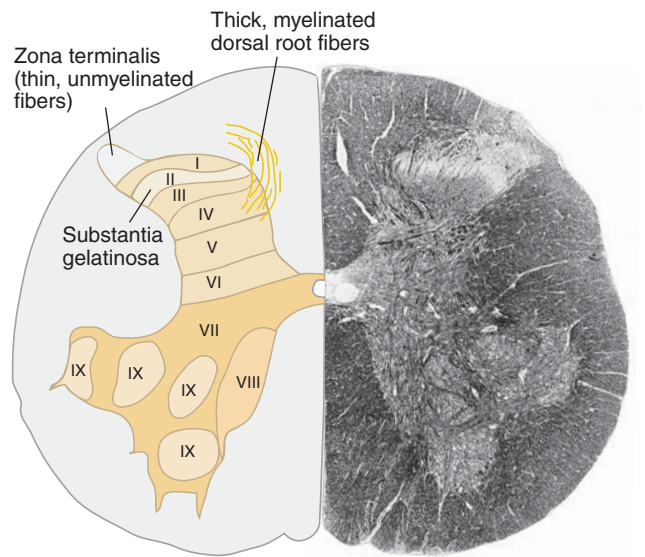


FIGURE 6.10 Cross section of the spinal cord at the lumbar level (lumbar intumescence). **Left:** Photomicrograph of section stained to show myelin and neuronal cell bodies. **Right:** The borders between Rexed's laminae. The groups of motoneurons in the ventral horn (α and γ motoneurons) constitute lamina IX; the zona terminalis (tract of Lissauer) consists primarily of thin dorsal root fibers.

Even though the cell bodies are arranged in laminae, the dendrites extend much wider, as shown in Fig. 6.12. Thus, a neuron belonging to a particular lamina receives synaptic inputs from nerve terminals in neighboring laminae. Nevertheless, experimental studies of the connections of the spinal cord and of the functional properties of single spinal neurons have shown that the various laminae differ in these respects. Thus, the laminae may be regarded, at least to some extent, as the nuclei of the spinal cord. We will return to Rexed's laminae when dealing with the functional organization of the spinal cord in later chapters.

The Spinal Nerves Divide into Branches

The dorsal (sensory) and ventral (motor) roots join to form spinal nerves, as described. Each spinal nerve then divides into several **branches (rami)** just outside the intervertebral foramen (Figs. 6.5 and 6.6). The thickest one, the **ventral ramus** (ramus ventralis), passes ventrally. A thinner branch, the **dorsal ramus** (ramus dorsalis), bends in the dorsal direction. In contrast to the spinal roots, the ventral and dorsal rami contain both sensory and motor fibers. This is caused by mixing of fibers from dorsal and ventral roots as they continue into the spinal nerves.

The dorsal rami innervate muscles and skin on the back, whereas the ventral rami innervate skin and muscles on the ventral aspect of the trunk and neck and, in addition, the extremities. Thus, the ventral rami

² It was formerly believed that propriospinal neurons—that is, spinal neurons with axons entering the white matter but not leaving the spinal cord—and interneurons represented two distinct cell groups. Recent studies have shown that spinal interneurons have local (intra-segmental) branches, as well as collaterals destined for other segments (intersegmental).



FIGURE 6.11 Cross section of the spinal cord at the cervical level (*cervical intumescence*). The broad ventral horns contain the motoneurons that supply muscles of the arm and shoulder girdle. Compared with the lumbar cord (Fig. 6.10), the cervical cord contains more white matter in proportion to gray matter; this is because all descending and ascending fibers to and from the lower levels pass through the cervical cord.

innervate much larger parts of the body than the dorsal ones, which explains why the ventral rami contain more nerve fibers and are considerably thicker than the dorsal ones. Some of the ventral rami join each other to form **plexuses** (see Fig. 21.1).

Each spinal nerve also sends off a small branch, the **meningeal ramus**, which passes back through the intervertebral foramen to reenter the vertebral canal (Fig. 6.5). These branches supply the meninges of the spinal cord with sensory and autonomic (sympathetic) fibers.

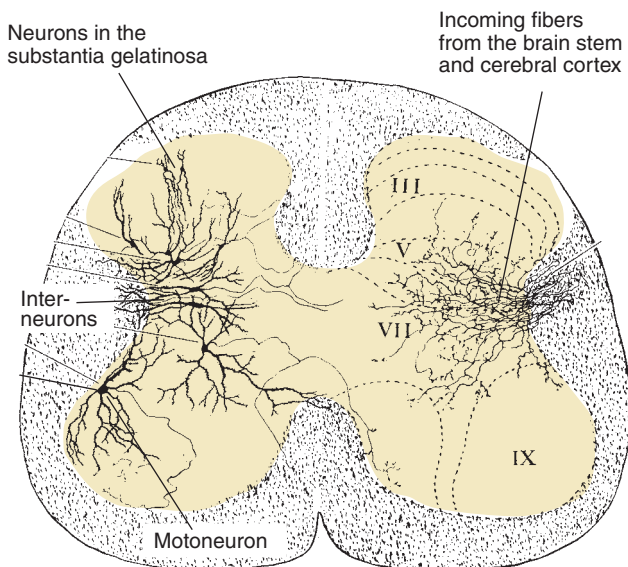


FIGURE 6.12 Dendritic arborizations of spinal neurons extend beyond the laminae of their cell bodies. Composite drawing based on observations in many Golgi-impregnated transverse sections from the spinal cord of the cat. The dendrites extend far, not only in the transverse plane as shown here but also longitudinally. To the right (at A) are the terminal ramifications of axons descending to the cord from higher levels of the CNS. (From Scheibel and Scheibel 1966.)

The Spinal Cord Consists of Subunits that Are Controlled from the Brain

Each segment of the spinal cord is to some extent a functional unit, since, as we will see in Chapter 13, a pair of spinal nerves relates to a particular “segment” of the body (see Fig. 13.14). A spinal segment may be regarded as the “local government” of a part of the body: it receives sensory information from its own district, processes this information, and issues orders through motor nerves to muscles and glands to ensure adequate responses. However, just as local governing bodies in our society must take orders from higher ones (e.g., county versus national governments), the spinal segments have only limited independence. Many of the functional tasks of the spinal cord are under strict control and supervision from higher levels of the CNS. This control is mediated by fibers from the brain stem and cerebral cortex, which descend in the white matter of the cord and terminate in the gray matter of the spinal segments that are to be influenced. The brain also ensures that the activity of the various spinal segments is coordinated, so that it serves the body as a whole and not only a small part. To be able to carry out this coordination, the brain must continuously receive information about conditions in all the “local districts” of the body and in the spinal segments related to them. This information is mediated by long, ascending fibers (forming various tracts) in the white matter of the cord that terminates in the brain stem. The local cooperation among spinal segments is taken care of by the numerous propriospinal fibers coming from spinal interneurons.

THE BRAIN STEM

The brain stem represents the upward (rostral, cranial) continuation of the spinal cord (Fig. 6.1). It consists of several portions with, in part, clear-cut surface borders

between them (Figs. 6.13 to 6.15). Whereas the lower-most (caudal) part of the brain stem is structurally similar to the spinal cord, the upper parts are more complicated. The subdivisions of the brain stem are as follows (from caudal to rostral): the **medulla oblongata** (often just called medulla), the **pons** (the bridge), the **mesencephalon** (the midbrain), and the **diencephalon**. Usually, however, often we use a more restricted definition including only the parts that extrude from the base of the brain: the medulla, pons, and mesencephalon.

The Brain Stem Contains the Third and Fourth Ventricles

A continuous, fluid-filled cavity varying in diameter stretches through the brain stem (Fig. 6.2). It is the upward continuation of the thin central canal of the spinal cord, and it continues rostrally into the cavities of the cerebrum (see Figs. 7.3 and 7.5). Together, these cavities constitute the ventricular system of the brain and spinal cord. The cavity in the brain stem has two dilated parts: one, the **fourth ventricle**, is at the level of the medulla and pons, whereas the **third ventricle** is

situated in the diencephalon. We return to the ventricular system in Chapter 7.

The Cranial Nerves

Examination of the internal structure of the brain stem shows that it is more complicated than that of the spinal cord (see, e.g., Fig. 6.16). Even though gray matter is generally located centrally, surrounded by a zone of white matter in both the brain stem and the cord, the gray matter of the brain stem is subdivided into several regions separated by strands of white matter. The white matter of the brain stem consists of myelinated fibers, as in other parts of the CNS. The regions with gray matter contain various nuclei, or groups of neurons with common tasks.

Many of the nuclei belong to the **cranial nerves**. In total, there are 12 pairs of cranial nerves, which, with the exception of the first, all emerge from the brain stem (Figs. 6.13 and 6.15). They correspond to the spinal nerves but show a more complex and less regular organization. Thus, there is no clear separation of

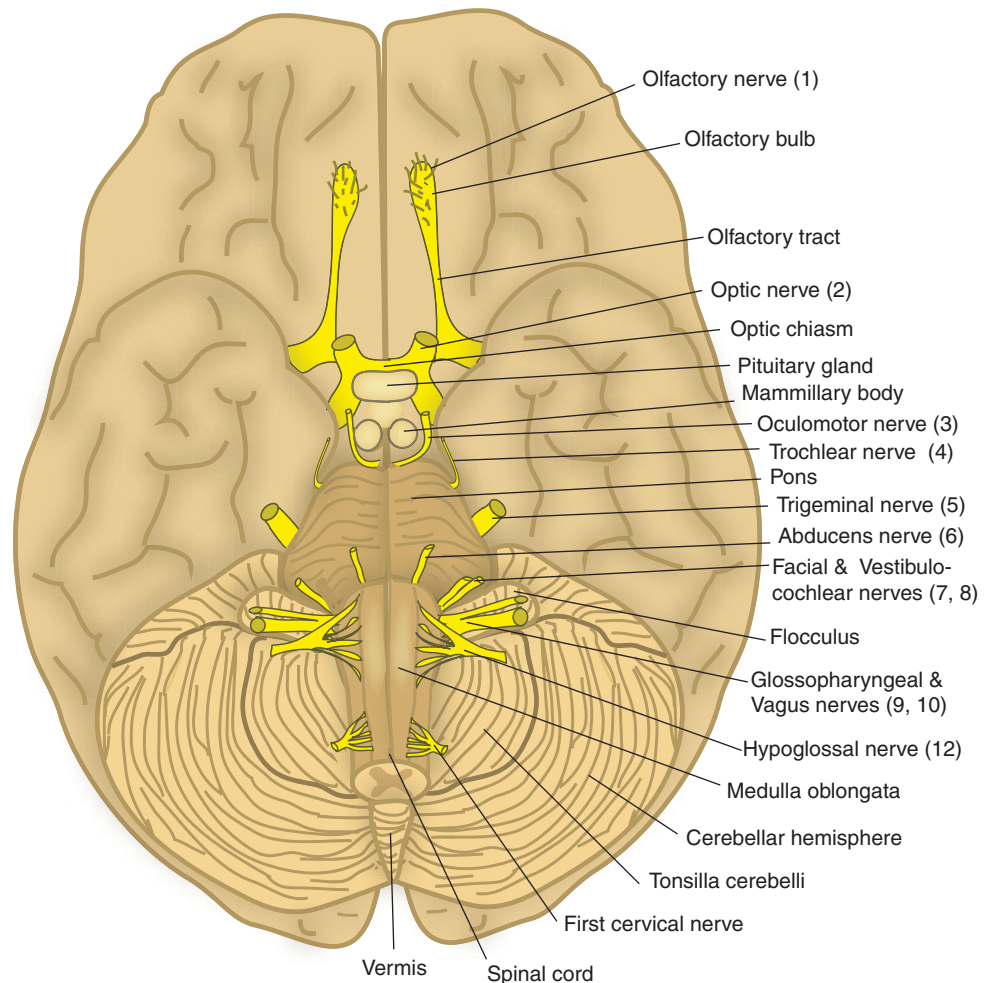


FIGURE 6.13 *The basal aspect of the brain.* Only some of the cranial nerves are shown.

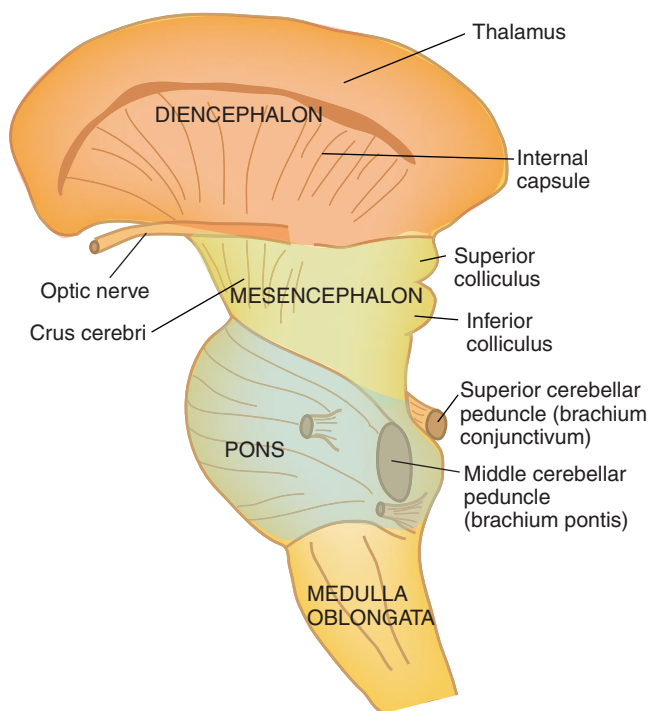


FIGURE 6.14 The brain stem, as viewed from the left side. The levels of sections shown in several of the following figures are indicated.

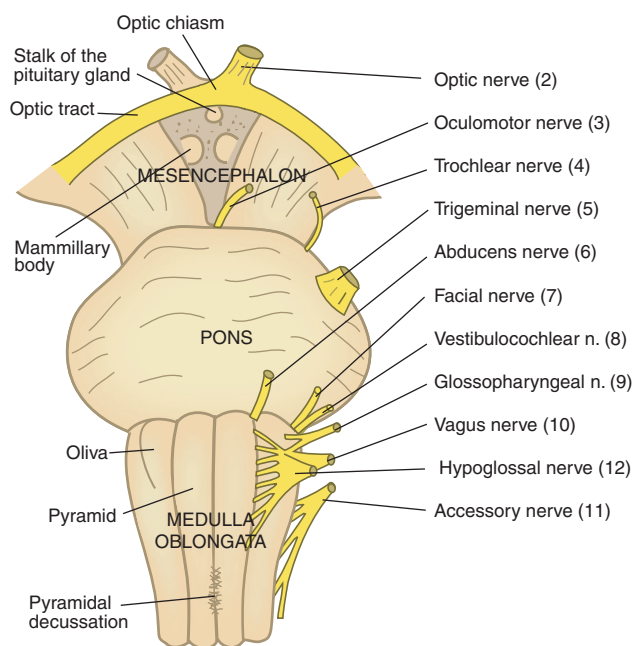


FIGURE 6.15 The cranial nerves. The brain stem is seen from the ventral side.

sensory (dorsal) and motor (ventral) roots. The cranial nerves are numbered from rostral to caudal, in accordance with where they emerge on the surface of the brain stem. Figure 8.9 shows the places on the base of the skull where the cranial nerves leave through small holes or fissures.

Many of the cranial nerves contain fibers that conduct impulses out of the brain stem; that is, the fibers are efferent, or motor. These fibers belong to neurons with their cell bodies in nuclei that are called **motor cranial nerve nuclei**. They correspond to the groups (columns) of neurons in the ventral and lateral horns of the spinal cord. The cranial nerves, like the spinal nerves, also contain sensory, afferent fibers that bring impulses from sense organs. The brain stem cell groups in which these afferent fibers terminate are, accordingly, called **sensory cranial nerve nuclei**; they correspond to the laminae of the spinal dorsal horn. The sensory fibers entering the brain stem have their cell bodies in ganglia just outside the brain stem, **cranial nerve ganglia**, corresponding to the spinal ganglia of the spinal nerves. Most cranial nerves are mixed—that is, they contain both motor and sensory fibers—but a few are either purely sensory or purely motor.

The only cranial nerve not emerging from the brain stem is the first cranial nerve, the **olfactory nerve** (nervus olfactorius). This consists of short axons coming from sensory cells in the roof of the nasal cavity, which, immediately after penetrating the base of the skull, terminate in the **olfactory bulb** (bulbus olfactorius) (Fig. 6.13). The other cranial nerves are briefly mentioned in the next section in connection with a description of the main structural features of the brain stem. The cranial nerves and their central connections are treated more thoroughly in later chapters, particularly Chapter 27.

The Reticular Formation Extends through Central Parts of the Brain Stem

Among the cranial nerve nuclei and other clearly delimited cell groups, there are more diffuse collections of neurons. In microscopic sections stained to visualize the neuronal processes, a network-like pattern is seen. The old anatomists therefore termed this part—present in the core of most of the brain stem—the **reticular formation** (formatio reticularis; Figs. 6.16–6.18; see also Fig. 26.1). In reality, however, the reticular formation is not one homogeneous structure but, rather, a conglomerate of cell groups with different connections and functional tasks. For example, some parts of the reticular formation are primarily concerned with control of **circulation** and **respiration**; other parts regulate **sleep** and **wakefulness**. However, the collective term, the reticular formation, is still in common use, and it may be practical to retain it to denote parts of the brain stem

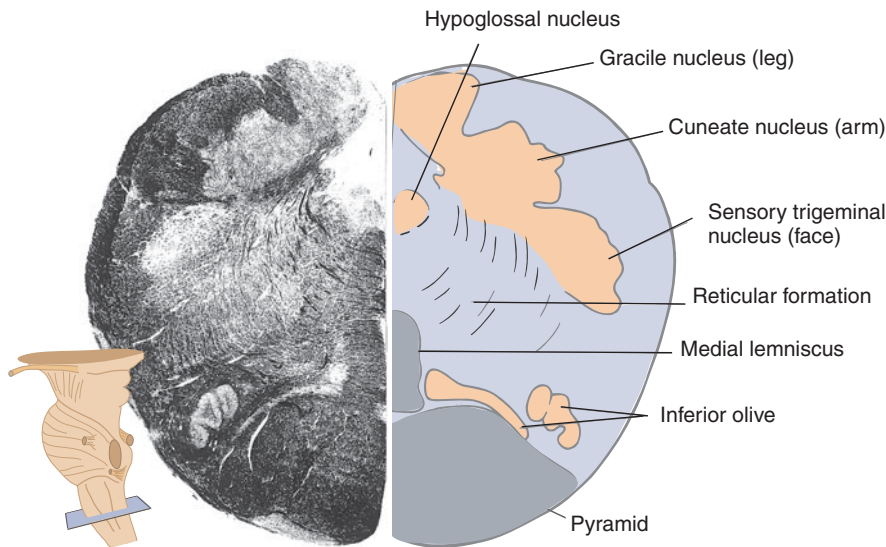


FIGURE 6.16 *Lower part of the medulla oblongata.* Cross section. Inset shows level and plane of the section. Left half shows photomicrograph of a section with darkly stained myelinated fibers (Woelke method); that is, white matter appears dark and gray matter appears light.

with certain common anatomic features, without implying that they have common functional tasks. The reticular formation is treated more comprehensively in Chapter 26.

The Medulla Oblongata

Ventrally in the midline, the medulla has a longitudinal sulcus, which is a continuation of the ventral median fissure of the cord (Fig. 6.15). The sulcus ends abruptly at the lower end of the pons. The so-called **pyramids**

protrude on each side of the longitudinal sulcus. Each pyramid is formed by a thick bundle of axons belonging to the **pyramidal tract**, which conveys signals from the cerebral cortex to the spinal cord and is essential for our control of voluntary movements (the pyramidal tract is discussed in Chapter 22). Close to the lower end of the medulla, on the transition to the cord, bundles of fibers can be seen to cross the midline, forming the **pyramidal decussation** (Fig. 6.15). Lateral to the pyramid is an oval protrusion (the olive), which is formed by a large nucleus, the **inferior olivary nucleus**

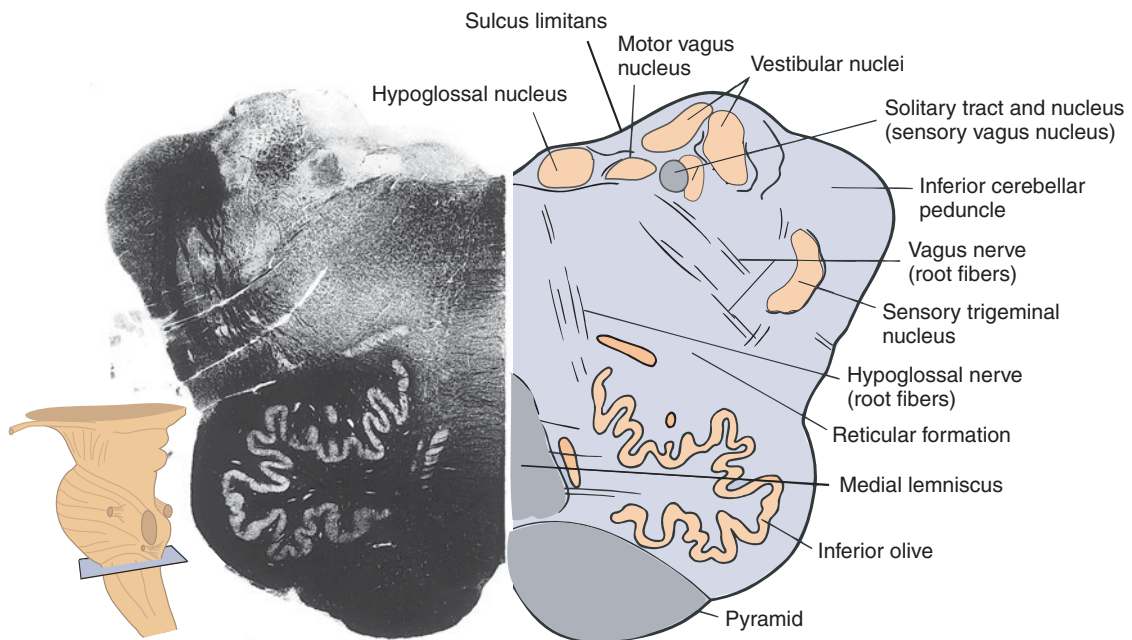


FIGURE 6.17 *Upper part of the medulla oblongata.* Cross section; myelin stained.

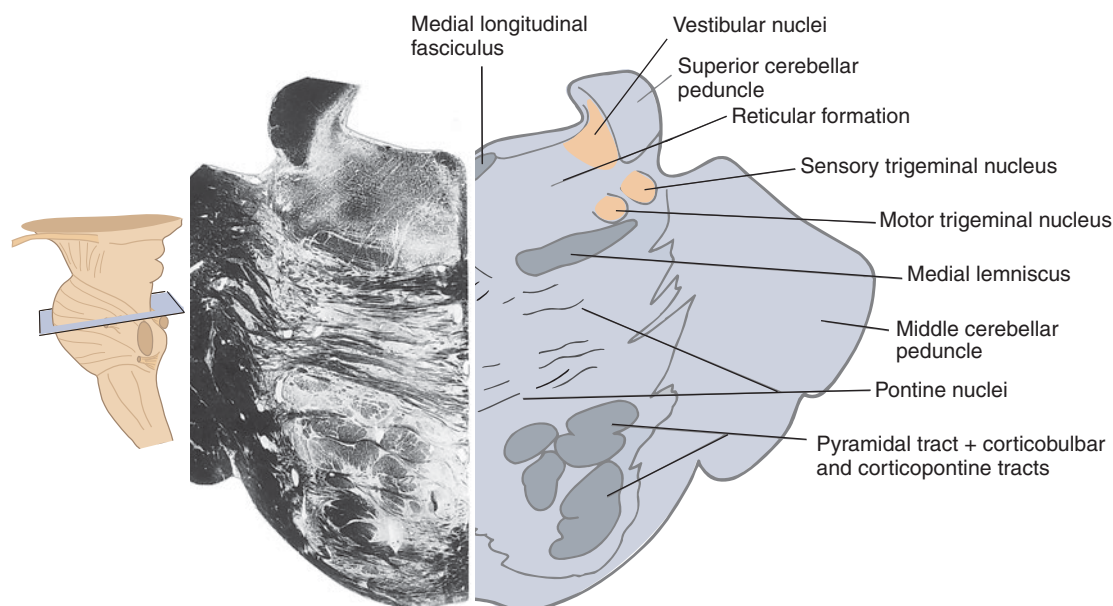


FIGURE 6.18 Upper part of the pons. Cross section; myelin-stained.

(inferior olive), which sends its efferents to the cerebellum. Between the olive and the pyramid is a row of small bundles of nerve fibers (Fig. 6.15; see also Fig. 27.1), which are the rootlets of the **hypoglossal nerve** (the twelfth cranial nerve, *nervus hypoglossus*). This nerve supplies the striated muscles of the tongue with motor fibers. Lateral to the olive, the rootlets of the **glossopharyngeal** and **vagus nerves** (ninth and tenth cranial nerves, *nervus glossopharyngeus* and *nervus vagus*) leave the brain stem. These two nerves supply the pharynx, the larynx, and most of the viscera with motor and sensory fibers. The **accessory nerve** (eleventh cranial nerve, *nervus accessorius*) runs cranially along the lateral aspect of the medulla. Most of its fibers come from the upper cervical spinal segments but enter the cranial cavity to leave the skull together with the glossopharyngeal and vagus nerves (see Fig. 27.8). The accessory nerve supplies two muscles in the neck with motor fibers.

Figure 3.16 shows a cross section through the **lower part of the medulla**, at a level below the caudal end of the fourth ventricle. The section is stained so that regions with white matter (the myelinated fiber tracts) are dark, whereas gray matter (the nuclei) appears light. The ventrally located bundle of cross-sectioned fibers is the **pyramidal tract**, forming the pyramid (Fig. 6.15), and containing about 1 million fibers. Dorsal to the pyramid lies a highly convoluted band of gray matter, the **inferior olivary nucleus**. The **dorsal column nuclei**, the **gracile** and **cuneate** (nucleus gracilis and nucleus cuneatus), are located dorsally in the medulla. The efferent fibers from these nuclei arch ventrally and take up a position close to the midline dorsal to the

pyramids, where they form a triangular area of cross-sectioned fibers. This is an important sensory tract, the **medial lemniscus** (*lemniscus medialis*), that leads from neurons in the dorsal column nuclei to nuclei in the diencephalon (see Fig. 13.17). The dorsal column nuclei receive afferent fibers that ascend in the dorsal columns (or dorsal funicles) and convey impulses from sense organs in the skin and muscles and around joints. Close to the midline, just ventral to the central canal, lies the **hypoglossal nucleus**, consisting of the cell bodies of the motor fibers that form the hypoglossal nerve. The efferent fibers of the hypoglossal nucleus pass ventrally and leave the medulla at the lateral edge of the pyramid (Fig. 6.15). Lateral to the motor cranial nerve nuclei are found several sensory cranial nerve nuclei, among them the big **sensory trigeminal nucleus** that receives sensory impulses from the face, carried in the **trigeminal nerve** (the fifth cranial nerve, *nervus trigeminus*). Note how the nuclei that are transmitting sensory impulses from the leg, arm, and face are distributed from medial to lateral in the dorsal part of the medulla (Fig. 6.16).

A cross section through the **upper part of the medulla** (Fig. 6.17) shows partly the same fiber tracts and nuclei as the section at a lower level (Fig. 6.16). In addition, we may notice the big **vestibular nuclei** situated dorsally and laterally under the floor of the fourth ventricle (these nuclei also extend cranially into the pons; see Fig. 19.7). They receive sensory impulses from the vestibular apparatus in the inner ear via the **vestibular nerve** (the eighth cranial nerve). One of the main efferent pathways from the vestibular nuclei forms a distinct tract, the **medial longitudinal fasciculus** (*fasciculus longitudinalis medialis*), close to the midline under the

floor of the fourth ventricle (Fig. 6.18). This tract terminates in the motor nuclei of the cranial nerves supplying the eye muscles, thus conveying influences on eye movements from the receptors for equilibrium. Further, Fig. 3.17 shows the motor and sensory **nuclei of the vagus**, and some of the fibers of the **vagus nerve**, which pass laterally and ventrally, to leave the medulla lateral to the olive (Fig. 6.15).

The Pons

The pons forms a bulbous protrusion at the ventral aspect of the brain stem, with clear-cut transversely running fiber bundles (Figs. 6.13 and 6.14). It is sharply delimited both caudally and cranially. The transverse fiber bundles are formed by fibers from large cell groups in the pons, the pontine nuclei, and terminate in the cerebellum. The fiber bundles join at the lateral aspect of the pons to form the **middle cerebellar peduncle** (brachium pontis) (Figs. 6.14 and 6.18). Several cranial nerves leave the brain stem at the ventral aspect of the pons. At the lower (caudal) edge, just lateral to the midline, a thin nerve emerges on each side. This is the **abducens nerve** (the sixth cranial nerve, *nervus abducens*) that carries motor fibers to one of the external eye muscles (rotates the eye laterally). Still at the lower edge of the pons, but more laterally, two other cranial nerves leave the brain stem. Most ventrally lies the **facial nerve** (seventh cranial nerve, *nervus facialis*), which brings motor impulses to the mimetic muscles of the face (it also contains some other kinds of fibers that are considered in Chapter 27). Closely behind the facial nerve lies the **vestibulocochlear nerve** (the eighth cranial nerve, *nervus vestibulocochlearis*), which carries sensory impulses from the sense organs for equilibrium and hearing in the inner ear. The **trigeminal nerve** (the fifth cranial nerve, *nervus trigeminus*) leaves the brain stem laterally at middle levels of the pons. The largest portion of the nerve consists of sensory fibers from the face, whereas a smaller (medial) portion contains motor fibers destined for the masticatory muscles.

In a **cross section** through the pons, the large **pontine nuclei** can be seen easily (Fig. 6.18). As mentioned, the neurons of the pontine nuclei send their axons to the cerebellum. Because their main afferent connections come from the cerebral cortex, the pontine nuclei mediate information from the cerebral cortex to the cerebellum. The **medial lemniscus** borders the pontine nuclei dorsally and has turned around and moved laterally compared with its location in the medulla (Fig. 6.16). In the lower part of the pons, the nucleus of the abducens nerve, the **abducens nucleus**, is located dorsally and medially, whereas the **facial nucleus** lies more ventrally and laterally (see Fig. 27.11; Fig. 17.11 also shows the course taken by the efferent fibers of the abducens and facial nuclei, forming the sixth and seventh cranial nerves, respectively).

Figure 6.18 also shows the **sensory trigeminal nucleus** laterally (this nucleus extends as a slender column through the medulla, pons, and mesencephalon; see also Figs. 6.16–6.18, and 27.2). Medial to the sensory nucleus lies the **motor trigeminal nucleus** (the masticatory muscles), but this nucleus is present only in the pons.

The Medulla and Pons Seen from the Dorsal Side

At the dorsal side of the medulla oblongata, at caudal levels, there are two longitudinal protrusions, the **gracile** and **cuneate tubercles** (Fig. 6.19). These are formed by the **dorsal column nuclei**, mentioned earlier (they are relay stations in pathways for sensory information from the body to the cerebral cortex). The most medial of these nuclei, the **gracile nucleus**, receives impulses from the leg and lower part of the trunk, whereas the laterally situated **cuneate nucleus** receives impulses from the arm and upper part of the trunk. Further laterally, another oblong protrusion (tuberculum cinereum) is formed by the **sensory trigeminal nucleus**, the relay station for sensory impulses from the face.

Rostral to the upper end of the dorsal column nuclei, there is a flattened, diamond-shaped area, the **rhomboid fossa**, extending rostrally onto the posterior face of the pons (Fig. 6.18). This constitutes the “floor” of the fourth ventricle (Fig. 6.19). Laterally and rostrally, the **cerebellar peduncles** delimit the rhomboid fossa (these have been cut in Fig. 6.19). Some of the cranial nerve nuclei and some fiber tracts form small protrusions medially at the floor of the fourth ventricle—notably the hypoglossal nucleus (hypoglossal trigone), the vagus nucleus (vagal trigone) and more rostrally the root fibers of the facial nerve—the latter forming the **facial colliculus**. (Figure 27.11 explains how the facial colliculus is formed.)

The Mesencephalon (Midbrain)

The part of the brain stem rostral to the pons, the **mesencephalon**, is relatively short (Figs. 6.14 and 6.15). Ventrally, an almost half-cylindrical protrusion is present on each side of the midline—the **crus cerebri**, or **cerebral peduncle** (Figs. 6.15 and 6.20).³ Crus cerebri consists of nerve fibers descending from the cerebral cortex to the brain stem and spinal cord; among these fibers are those of the pyramidal tract. The fibers continue into the pons, where they spread out into several smaller bundles among the pontine nuclei (Fig. 6.18).

³ Strictly speaking, the term cerebral peduncle denotes both the crus cerebri and parts of the mesencephalon dorsal to the crus except the colliculi (the latter collectively termed the “tectum”). The region between the crus cerebri and the tectum is called the **tegmentum** and includes the periaqueductal gray, the red nucleus, and the substantia nigra. Previously, however, the crus cerebri and the cerebral peduncle were both applied to the ventralmost, fiber-rich part.

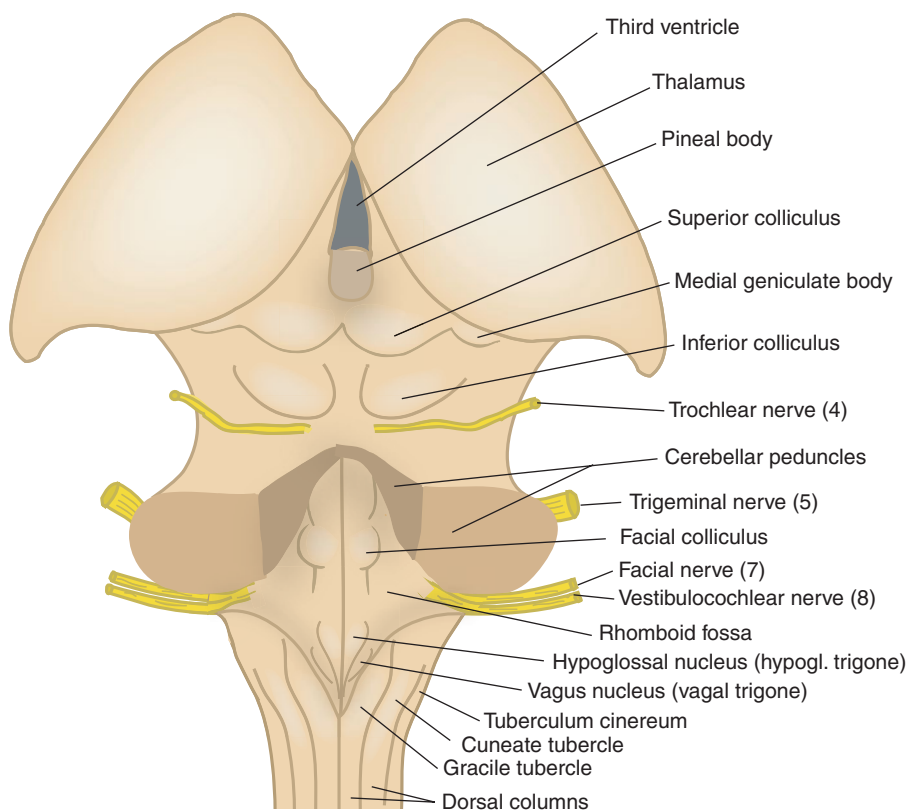


FIGURE 6.19 The brain stem. Viewed from the dorsal aspect.

In the furrow between the two crura, the **oculomotor nerve** emerges (third cranial nerve, *nervus oculomotorius*) (Figs. 6.13 and 6.15). As the name implies, the nerve carries motor impulses to muscles that move the eye. The oculomotor nerve innervates four of the six extraocular (striated) muscles and, in addition, two smooth internal muscles that regulate the diameter of the pupil and the curvature of the lens.

At the **dorsal side** of the mesencephalon, there is a characteristic formation of four small, rounded protrusions, two on each side of the midline (Fig. 6.18). These are called the **colliculi** (*corpora quadrigemina*) and consist of, on each side, the **superior colliculus** and the **inferior colliculus**. The superior colliculus consists of cell groups that control reflex movements of the eyes and the head, while the inferior colliculus is a relay station in the pathways that bring auditory impulses to awareness.

A thin fiber bundle, the **trochlear nerve** (fourth cranial nerve, *nervus trochlearis*), emerges on each side below the inferior colliculi (Fig. 6.19). This is the only cranial nerve that emerges on the dorsal side of the brain stem. It supplies one of the extraocular muscles with motor fibers.

In a **cross section** of the mesencephalon (Fig. 6.20), the crus cerebri can be recognized ventrally, and the superior colliculus dorsally. In the midline just ventral to the colliculi there is a small hole, which is a cross section of the **aqueduct** (*aquaeductus cerebri*), a narrow

canal that interconnects the third and fourth ventricles (Figs. 6.23 and 7.5). Surrounding the aqueduct is a region of gray matter called the **periaqueductal gray substance** (*substantia grisea centralis*), which coordinates behavioral responses to stressful events and influences pain perception. Ventral to the periaqueductal gray, close to the midline, we find the **oculomotor nucleus** (or nucleus of the oculomotor nerve). Further ventrally lies the large **red nucleus** (*nucleus ruber*), so named because of its slightly reddish color. Just dorsal to the crus and ventral to the red nucleus is the **substantia nigra** (the black substance). The neurons of the substantia nigra contain a dark pigment, making the nucleus clearly visible macroscopically. The red nucleus and the substantia nigra are both important for the control of movements.

The Diencephalon Contains the Thalamus and the Hypothalamus

Figure 6.21 shows how the diencephalon merges with the mesencephalon caudally without any clear transition. Neither rostrally is the diencephalon clearly delimited (Fig. 6.13) because in early embryonic life it fuses with the primordium of the cerebral hemispheres. The **optic nerve** (the second cranial nerve, *nervus opticus*) carrying afferent fibers from the retina ends in the diencephalon.

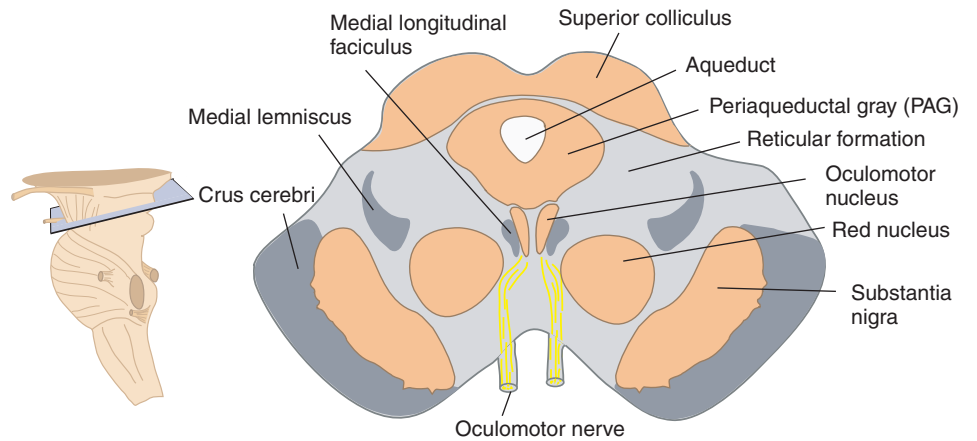


FIGURE 6.20 *The mesencephalon.* Cross section. The major nuclei are indicated in red and the major tracts are in gray.

The largest part of the diencephalon is occupied by the **thalamus**, situated on each side of the third ventricle (Figs. 6.22, 6.24, and 6.27). The thalamus consists of many smaller nuclei and is a relay station for almost all information transmitted from the lower parts of the CNS to the cerebral cortex (notably most kinds of sensory information). Each thalamus is approximately egg-shaped with a flattened side toward the third ventricle (Fig. 6.22). Lateral to the thalamus lies a thick sheet of white matter, the **internal capsule** (capsula interna). It consists mainly of fiber tracts connecting the cerebral cortex with the thalamus, the brain stem, and the spinal cord, among them the pyramidal tract (Figs. 6.14, 6.24, 6.27, and 6.30). The crus cerebri is a caudal continuation of fibers of the internal capsule.

In a frontal section of the brain (Fig. 6.24) the thalamus is subdivided by narrow bands of white matter forming a Y, called the **internal medullary lamina** (see also Fig. 6.22). This divides the thalamic gray matter in

three main parts: an **anterior** nuclear group (or complex), a **medial** nuclear group, and a lateral part or region made up of a **dorsal** and a **ventral** nuclear group.⁴ The **pulvinar**, continuous with the lateral part, makes up most of the posterior part of the thalamus (Figs. 6.21 and 6.22). In addition, the posterior part of the thalamus includes two nuclei partly covered by the pulvinar, the **lateral geniculate body** (corpus geniculatum laterale) and the **medial geniculate body** (corpus geniculatum mediale). Each of the thalamic nuclei connects to

⁴ The nomenclature of thalamic nuclear subdivisions may appear bewildering, and matters are not made easier by lack of agreement among leading investigators. For example, the nomenclature presented in the *Terminologia Anatomica* (1998) differs from that used in the scholarly book *The Human Nervous System*, edited by Paxinos and Mai (2004). Throughout the present book when dealing with the thalamus, I have tried to simplify matters to provide just enough anatomical detail to help the reader understand the functional organization of the thalamus.

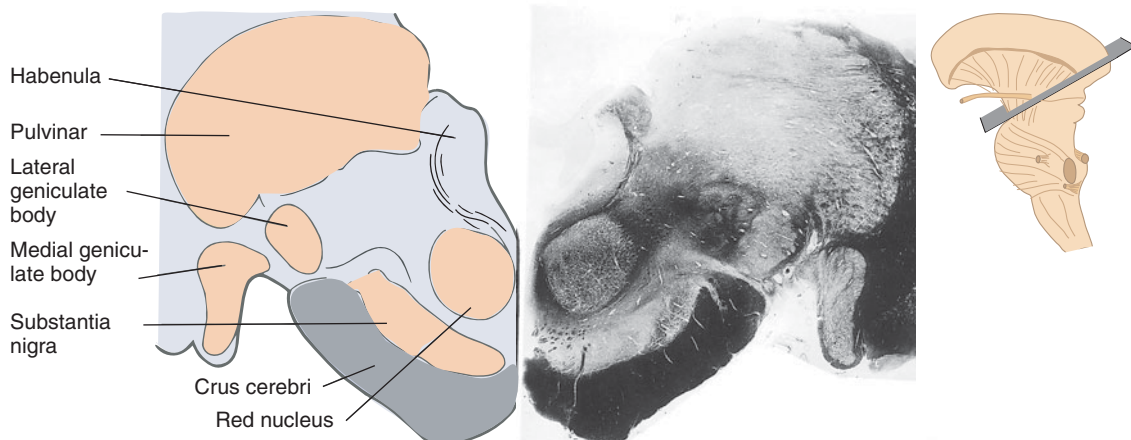


FIGURE 6.21 *Transition between the mesencephalon and the diencephalon.* Oblique frontal section; myelin stained. Inset shows plane and level of the section.

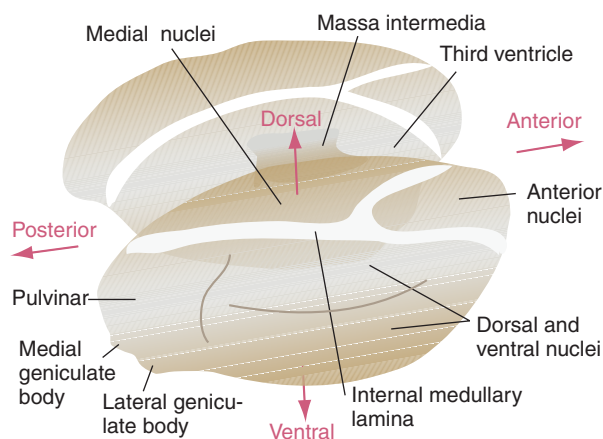


FIGURE 6.22 *The thalamus.* Drawing of the thalami of both sides, to indicate their three-dimensional form.

different parts of the cerebral cortex (see Fig. 34.7). For example, the lateral and medial geniculate bodies are relay stations for visual and auditory impulses, respectively. Thus, fibers of the optic nerve end in the lateral geniculate body while fibers from the inferior colliculus end in the medial geniculate body.

The **optic nerves** from the two eyes unite just underneath the diencephalon (Figs. 6.13 and 6.15) to form the **optic chiasm** (chiasma opticum, or just chiasma), in which there is a partial crossing of the optic nerve fibers (see Fig. 16.14). In their further course from the optic chiasm to the lateral geniculate body, the fibers form the **optic tract** (Figs. 6.15 and 6.24). The fibers from the lateral geniculate body to the visual cortex form the **optic radiation** (Fig. 16.15).

Anterior and inferior to the thalamus lies the **hypothalamus** (Fig. 6.23), which exerts central control of the autonomic nervous system—that is, with control of the visceral organs and the vessels. The hypothalamus forms the lateral wall of the anterior part of the third ventricle. The border between the thalamus and the hypothalamus is marked by the shallow **hypothalamic sulcus** (Figs. 6.23 and 6.24). The **mammillary body** (corpus mammillare), a special part of the hypothalamus, protrudes downward from its posterior part (Figs. 6.13, 6.23, and 6.29). The **fornix** is a thick, arching bundle of fibers originating in the cerebral cortex (in the so-called hippocampal region in the temporal lobe) and terminating in the mammillary bodies (see Fig. 32.2). The major efferent pathway of the mammillary body goes to the thalamus, forming a distinct fiber

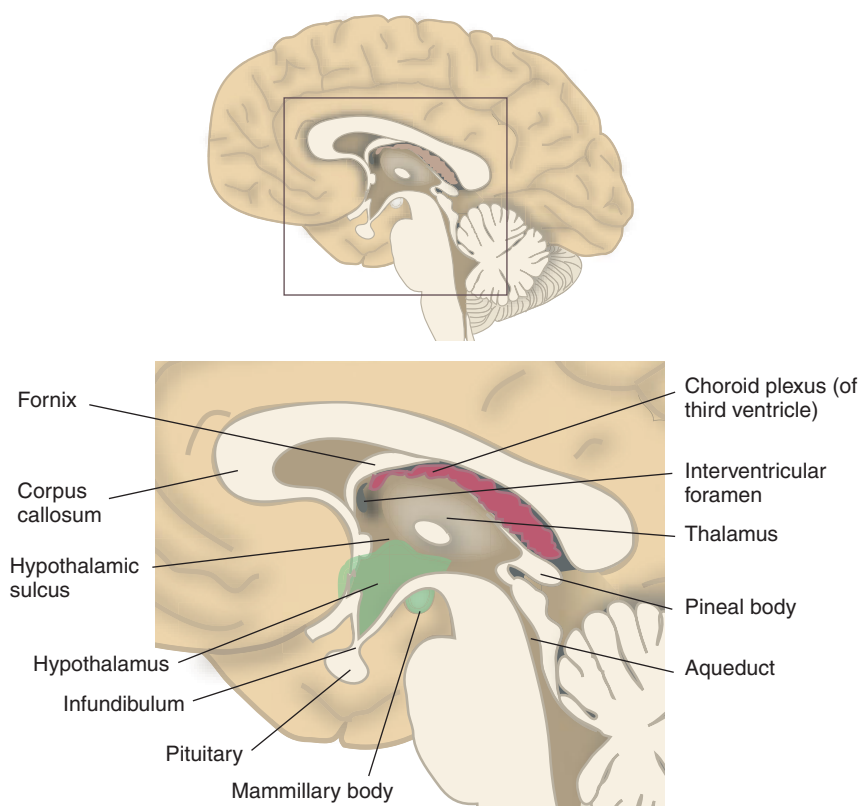


FIGURE 6.23 *The hypothalamus.* Drawing of midsagittal section showing the upper parts of the brain stem. The hypothalamus is indicated in red.



FIGURE 6.24 *The diencephalon.* Frontal section; myelin stained.

bundle, the **mammillothalamic tract** (fasciculus mammillothalamicus) (Figs. 6.24 and 30.6). In front of the mammillary bodies, the floor of the third ventricle bulges downward like a funnel and forms the stalk of the pituitary gland, the **infundibulum**. The region between the mammillary bodies and the infundibulum is called the **tuber cinereum** (see Fig 30.3). It contains neuronal groups that influence the activity of the pituitary gland.

The **pituitary** (Figs. 6.13 and 6.23) consists of a **posterior lobe**, developed from the CNS, and an **anterior lobe**, developed from the epithelium in the roof of the mouth. The anterior lobe, secreting several hormones that control important bodily functions, is itself under the control of the hypothalamus. This is discussed further in Chapter 30.

Epithalamus and the Pineal Body

The diencephalon also includes a small area called the **epithalamus**, located posteriorly in the roof of the third ventricle. In addition to a small nucleus, the **habenula** (Fig. 6.21), the epithalamus contains the **pineal body** or gland (*corpus pineale*) (Figs. 6.19, 6.23, and 6.31). This peculiar structure lies in the midline (unpaired) and is formed by an evagination of the roof of the third ventricle. It contains glandular cells, **pinealocytes**, which produce the hormone **melatonin** (and several neuropeptides). It also contains large amounts of **serotonin**, which is a precursor of melatonin. Melatonin influences several physiological parameters, especially those that

show a cyclic variation. This is discussed more thoroughly in Chapter 30, under “Hypothalamus and Circadian Rhythms” and under “Melatonin.”

The **habenula** (Fig. 6.21) lies just underneath the pineal body (one on each side). This small nucleus (composed of several subnuclei) receives afferents from the hypothalamus and the septal nuclei, among other sources. Its main efferents go to nuclei in the mesencephalon. The functional role of the habenula is not known, but the pathway from the hypothalamus via habenula to the mesencephalon may be engaged in the bodily expressions of strong **emotions**—for example, rage or fear. The habenula is one among several neuronal groups that are altered in severe **depression**.

THE CEREBRUM

The cerebrum has an ovoid shape and fills most of the cranial cavity. Whereas its convexity—that is, its upper and lateral surfaces—are evenly curved, the basal surface is uneven. In the center of the basal surface, the brain stem emerges (Fig. 6.13). The cerebrum is almost completely divided in two by a vertical slit, the **longitudinal cerebral fissure** (*fissura longitudinalis cerebralis*), so that it consists of two approximate half-spheres or **cerebral hemispheres** (Figs. 6.25 and 6.26). Each of the cerebral hemispheres contains a central cavity, the **lateral ventricle** (Figs. 6.29 and 7.3). The lateral ventricles are continuous, with the third ventricle through a small opening, the **interventricular foramen** (Fig. 6.37).

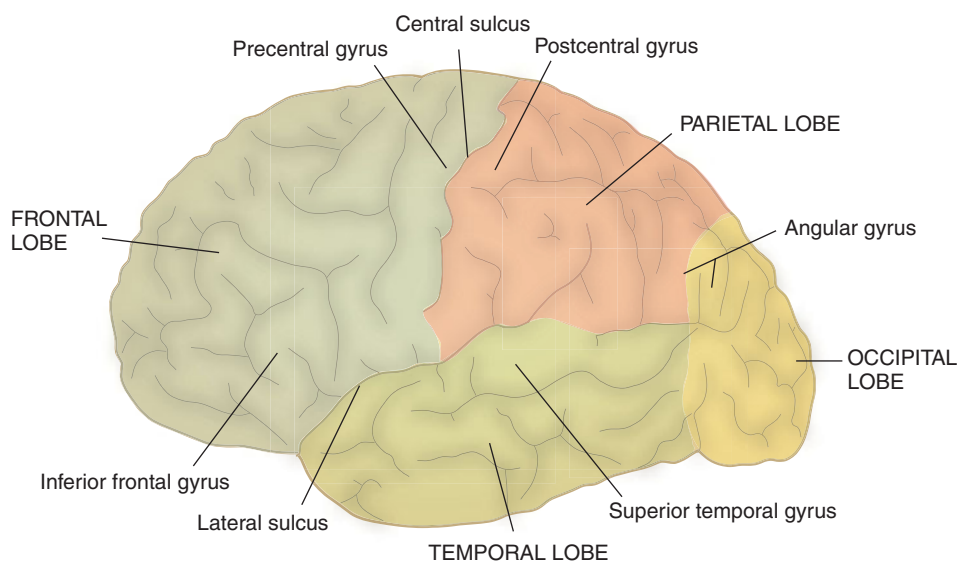


FIGURE 6.25 The left cerebral hemisphere. Viewed from the lateral aspect.

The lateral ventricles are surrounded by masses of white matter with some embedded nuclei. The surface of the hemisphere is covered everywhere by a 3 to 5 mm-thick layer of gray substance, the **cerebral cortex** (cortex cerebri). The structure, connections, and functions of the cerebral cortex are covered most completely in Chapters 33 and 34, but some main features are briefly described here because knowledge of them is necessary for the chapters dealing with sensory and motor systems.

The neurons of the cerebral cortex receive impulses from lower parts of the CNS, most of which are relayed through the **thalamus**. In addition, there are numerous **association fibers**—that is, fibers interconnecting

neurons in various parts of the cerebral cortex of one hemisphere. Finally, a vast number of **commissural fibers** interconnect neurons in the two hemispheres. Most of the commissural fibers are collected into a thick plate of white matter, the **corpus callosum**, which joins the two hemispheres across the midline (Figs. 6.26, 6.27, 6.28, and 6.29). The fibers in the corpus callosum enable impulses to travel from one hemisphere to the other and thus ensure that the right and left hemispheres can cooperate (see Chapter 34, under “The Function of the Commissural Connections”). A few commissural fibers pass in the **anterior commissure** (commissura anterior) (Fig. 6.26).

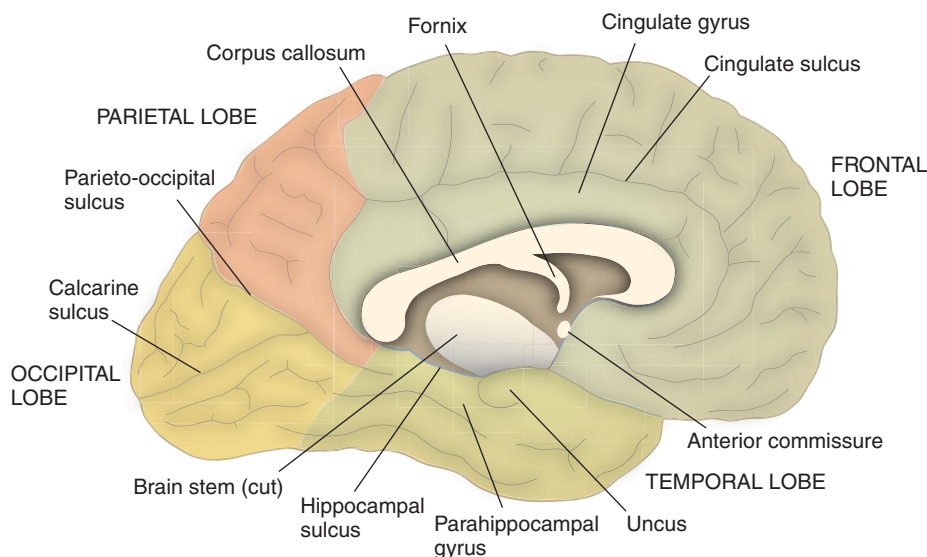


FIGURE 6.26 The left cerebral hemisphere. Viewed from the medial aspect. The brain stem has been cut off.

The Surface of the Hemisphere Is Highly Convoluted and Forms Gyri and Sulci

During embryonic development, the cerebral hemispheres fold as they grow in size (see Fig. 7.4). This leads to a great increase in their surface area and thus in the amount of cortex relative to their volume (only about one-third of the total cortical surface is exposed). The furrowed, walnut-like appearance of the cerebral hemispheres of humans and some higher mammals is highly characteristic (Figs. 6.25, 6.26, and 6.28). The folding of the hemisphere produces deep **fissures** and more shallow **sulci**. Between the sulci, the surface of the cortex forms rounded **gyri**. Apart from the fissure that divides the two hemispheres along the midline, the longitudinal cerebral fissure, the largest fissure in each hemisphere is the **lateral cerebral sulcus** (fissure) or the Sylvian fissure (Fig. 6.25). This fissure follows a course upward and backward and extends deep into the hemisphere (Fig. 6.27). The small gyri in the bottom of the lateral sulcus form the **insula** (the island) (Figs. 6.27 and 6.30). More sulci and gyri will be mentioned when dealing with the lobes of the cerebrum.

The pattern of the fissures and the larger sulci in the human brain is fairly constant. Nevertheless, variations are great enough to make it impossible to know exactly where a certain sulcus is located only from landmarks on the outside of the skull. The smaller sulci and gyri are subject to considerable individual variation.

The Hemisphere Can Be Divided into Four Lobes

With more or less sharply defined borders (formed by fissures and sulci), one can distinguish four lobes of the cerebral hemispheres (Figs. 6.25 and 6.26). They are named in accordance with the bone of the skull under which they are located. The **frontal lobe** (lobus frontalis) lies in the anterior cranial fossa above the orbit. The frontal lobe is separated from the **parietal lobe** (lobus parietalis) by the **central sulcus**, which extends from the medial edge of the hemisphere laterally to the lateral sulcus. Below the lateral sulcus lies the **temporal lobe** (lobus temporalis). Neither the parietal nor the temporal lobe has any clearly defined border posteriorly toward the occipital lobe. The **occipital lobe** lies above the cerebellum, which is located in the posterior cranial fossa (Fig. 6.1). On the medial aspect of the hemisphere, the border between the parietal and the occipital lobe is marked by the **parieto-occipital sulcus** (sulcus parieto-occipitalis) (Fig. 6.26).

Some Functional Subdivisions of the Cerebral Cortex

Anatomically and functionally, we divide the cerebral cortex into different regions, which do not, however, coincide with the different lobes. Here, only a few points are mentioned. The gyrus in front of the central sulcus, the **precentral gyrus** (gyrus precentralis) (Fig. 6.25), coincides with the **motor cortex** (MI), which

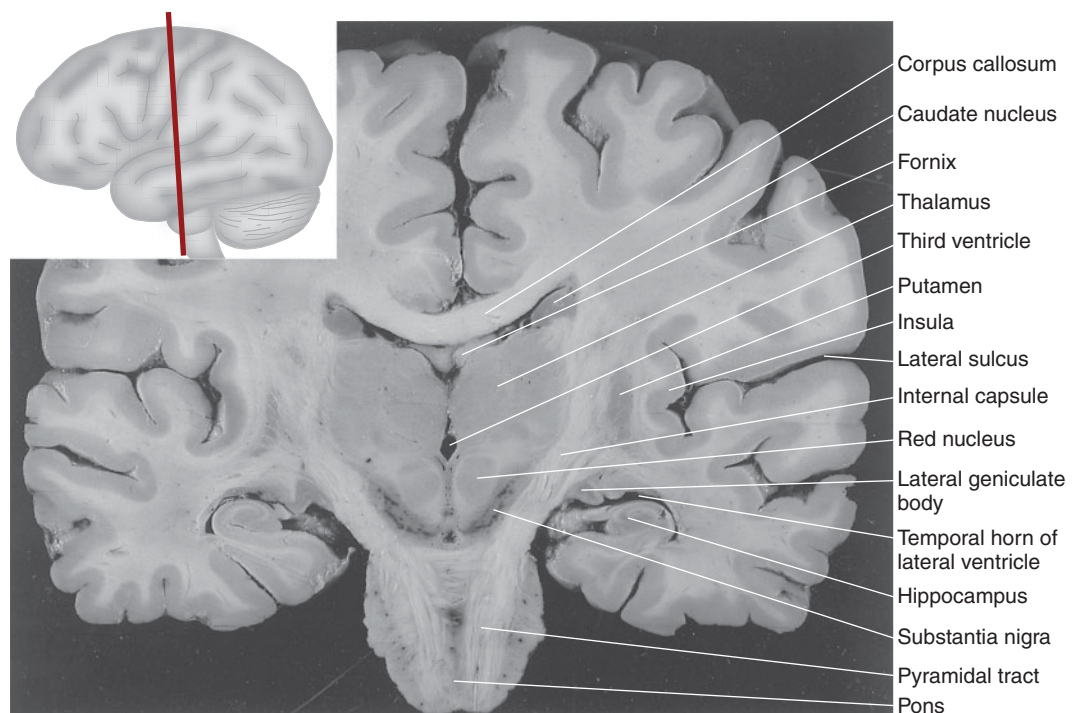


FIGURE 6.27 The cerebral hemispheres and upper part of the brain stem. Photograph of frontal section. Compare with Fig. 6.28.

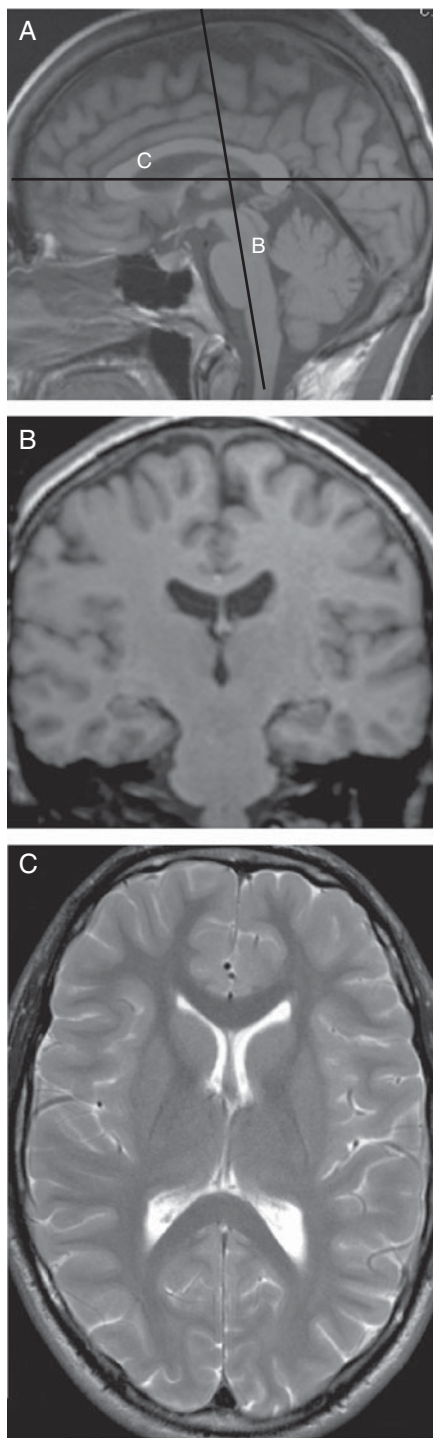


FIGURE 6.28 Magnetic resonance images (MRI) in the three conventional planes. **A:** Midsagittal plane. **B:** frontal (coronal) plane; corresponding to Fig. 3.27. **C:** Horizontal (transverse) plane. **A** and **B** are T1 weighted with water appearing dark. **C** is T2 weighted with water appearing white. (Courtesy of Dr. S. J. Bakke, Rikshospitalet University Hospital, Norway.)

is of special significance for the execution of voluntary movements. Destruction of this gyrus in one hemisphere, cause pareses in the opposite side of the body. Many of the fibers in the pyramidal tract come from the precentral gyrus, and, as mentioned, most of these fibers cross the midline on their way to the spinal cord. The **postcentral gyrus**, situated just posterior to the central sulcus, is the major receiving region for sensory impulses from the skin, the musculoskeletal system, and the viscera. This region is called the **somatosensory cortex (SI)**. The tracts that conduct impulses from the sense organs to the cortex are also crossed. Thus, destruction of the postcentral gyrus on one side leads to lowered sensibility (for example, of the skin) on the opposite side of the body. We have previously mentioned the **medial lemniscus**, which is part of the pathways from the sense organs to the postcentral gyrus. The fibers of the medial lemniscus terminate in a subdivision of the **lateral thalamic nucleus** (Fig. 6.24), and the neurons there send their axons to the postcentral gyrus.

The **visual cortex**—the main cortical region receiving information from the eyes—is located in the occipital lobe around a deep sulcus (fissure), the **calcarine sulcus** (sulcus calcarinus) (Fig. 6.26). The impulses start in the retina and are conducted in the optic nerve and the optic tract to the **lateral geniculate body** (Figs. 6.21 and 6.22), and from there to the visual cortex (see Fig. 16.14). The visual cortex can be distinguished from the surrounding parts of the cortex in sections perpendicular to the surface: it contains a thin whitish stripe running parallel to the surface (caused by a large number of myelinated fibers). Because of the stripe, this part of the cortex was named the **striate area** by the early anatomists (see Fig. 16.18).

The **auditory cortex**—the cortical region receiving impulses from the cochlea in the inner ear—is located in the superior temporal gyrus of the temporal lobe (Fig. 6.25; see also Fig. 17.11). The pathway for auditory impulses is synaptically interrupted in the **medial geniculate body** (Figs. 6.21 and 6.22).

The **olfactory cortex** is a small region on the medial aspect of the hemisphere near the tip of the temporal lobe. It is part of the so-called **uncus** (Fig. 6.26; see also Fig. 18.3) and extends somewhat onto the adjoining **parahippocampal gyrus**. The olfactory cortex receives fibers from the **olfactory bulb** (bulbus olfactorius) through the **olfactory tract** (tractus olfactorius) (Fig. 6.13). The cortex of the parahippocampal gyrus extends into the **hippocampal sulcus** (fissure), forming a longitudinal elevation, the **hippocampus** (Figs. 6.27 and 6.31; see also Fig. 32.2). The hippocampus belongs to the phylogenetically oldest parts of the cerebral cortex and has a simpler structure than the newer parts. The hippocampus and adjoining cortical regions in the medial part of the temporal lobe are of particular interest

with regard to **learning** and **memory** (this is treated further in Chapter 32).

The Cerebral Cortex Consists of Six Cell Layers

Examination of a microscopic section cut perpendicular to the surface of the cerebral cortex shows that the neuronal cell bodies are not randomly distributed (see Figs. 34.1 and 34.2). They are arranged into **layers** or **laminae** parallel to the surface. Each layer is characterized by a certain shape, size, and packing density of the cell bodies (compare with the Rexed's laminae of the spinal cord). We number the layers from the surface inward to the white matter. **Layer 1** is cell poor and consists largely of dendrites from neurons with cell bodies in deeper layers and of axons with terminals making synapses on the dendrites. **Layers 2 and 4** are made up of predominantly small, rounded cells and are therefore called the **external** and **internal granular layer**, respectively. These two layers have in common that they largely have a receiving function: many of the afferent fibers to the cerebral cortex terminate and form synapses in layers 2 and 4. Fibers conveying sensory information from lower levels of the CNS end predominantly in layer 4, and consequently this layer is particularly well developed in the sensory cortical regions mentioned above. **Layers 3 and 5** contain cells that are larger than those in layers 2 and 4 are, and the cell bodies tend to be of pyramidal shape, hence the name **pyramidal cells**. Many of the pyramidal cells in layer 5 send their axons to the brain stem and spinal cord, where they influence motor neurons. Layer 5 is therefore especially well developed in the motor cortex in the precentral gyrus. The pyramidal neurons in layer 3 send their axons primarily to other parts of the cerebral cortex, either in the same hemisphere (association fibers) or to cortex in the hemisphere of the other side (commissural fibers). The cell bodies of **layer 6** are smaller and more spindle-shaped than those in layer 5 are. Many of the neurons send their axons to the thalamus, enabling the cerebral cortex to influence the impulse traffic from the thalamus to the cortex (feedback connections).

There are also numerous **interneurons** in the cerebral cortex, providing opportunity for cooperation between the various layers (interneurons with “vertically” oriented axons) and between neurons in different parts of one layer (interneurons with “horizontally” oriented axons). The layers are obviously not independent units.

The Cerebral Cortex Can Be Divided into Many Areas on a Cytoarchitectonic Basis

We mentioned that some layers are particularly well developed in certain regions of the cortex—for example, layer 4 in the sensory receiving areas and layer 5 in the motor cortex. There are many more differences

when all layers all over the cortex are systematically compared. Such **cytoarchitectonic** differences (i.e., differences in size, shape, and packing density of the cell bodies) form the basis of a parcellation of the cerebral cortex of each hemisphere into approximately 50 **cortical areas** (areae). Maps of the cerebral cortex showing the positions of the various areas were published by several investigators around year 1900 and are still in use. The German anatomist Brodmann (see Fig. 20.3) published the most widely used map. Such cytoarchitectonically-defined areas have later been shown, in many cases, to differ also with regard to connections and functional properties. The numbering of the cortical areas may appear illogical. For example, the motor cortex in the precentral gyrus corresponds to area 4 of Brodmann. This borders posteriorly on area 3 but on area 6 anteriorly. Area 3 borders on area 1, which borders on area 2, and so on (see Fig. 34.3). It is not necessary to learn the position of more than a few of the cortical areas, however, and these will be dealt with in connection with the various functional systems in Parts II–VI of this volume.

The Basal Ganglia

The interior of the hemispheres contains large masses of gray substance. Largest among these are the so-called **basal ganglia**, which perform important tasks related to the control of movements.⁵ Other nuclear groups (the amygdala, the septal nuclei, and the basal nucleus) are discussed in Chapter 31, and the basal ganglia are discussed in Chapter 23. Here we mention only a few main points with regard to the anatomy of the basal ganglia.

The basal ganglia receive massive afferent connections from the cerebral cortex and acts, by way of their efferent fibers, primarily back on motor regions of the cortex. Sections through the hemispheres show that the basal ganglia consist of two main parts (Figs. 6.29 and 6.30). In a horizontal section (Fig. 6.30) one large part lies lateral to the internal capsule, and a smaller part lies medial to the internal capsule and anterior to the thalamus. The largest part is called the **lentiform nucleus** (nucleus lentiformis) because of its shape. It consists of two closely apposed parts: the lateral or external part is the **putamen**, and the medial or internal part is the **globus pallidus**. The part of the basal ganglia situated medial to the internal capsule is the **caudate nucleus** (nucleus caudatus). The name describes its form: a large part of the nucleus forms a long, curved “tail” (Fig. 7.4, see also Fig. 23.1). The putamen and the caudate

⁵ The name basal ganglia has been retained from a time when all collections of neurons were called ganglia, regardless of whether they were located inside or outside the CNS. Today we use the term ganglion only for collections of neurons outside the CNS, as discussed in Chapter 1. However, the name basal ganglia is so well established that it is not practical to exchange it.

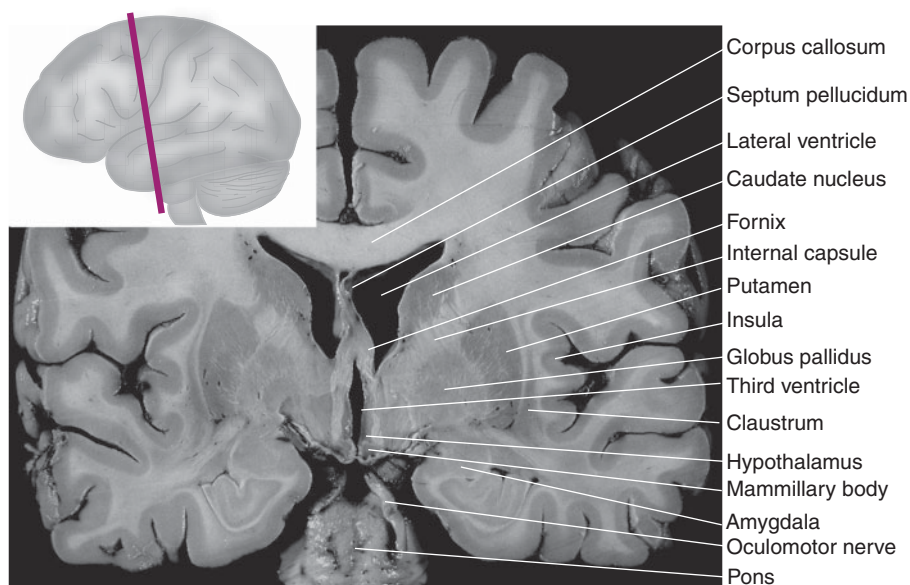


FIGURE 6.29 *The cerebral hemispheres.* Frontal section, placed more frontally than as shown in Fig. 3.27, showing the basal ganglia and amygdala.

nucleus together are called the **striatum** (or neostriatum). The caudate nucleus consists of an anterior bulky part, the **caput** (head), and a progressively thinner **cauda** (tail). The cauda extends first backward and then down and forward into the temporal lobe, located in the wall of the lateral ventricle. Figure 7.4 shows how

this peculiar form can be explained on the basis of the embryonic development of the cerebral hemispheres.

The **claustrum** forms a sheet of grey matter lateral to the putamen (Fig. 6.29). It has reciprocal connections most parts of the cerebral cortex. Little is known about its function or clinical significance, but based on its

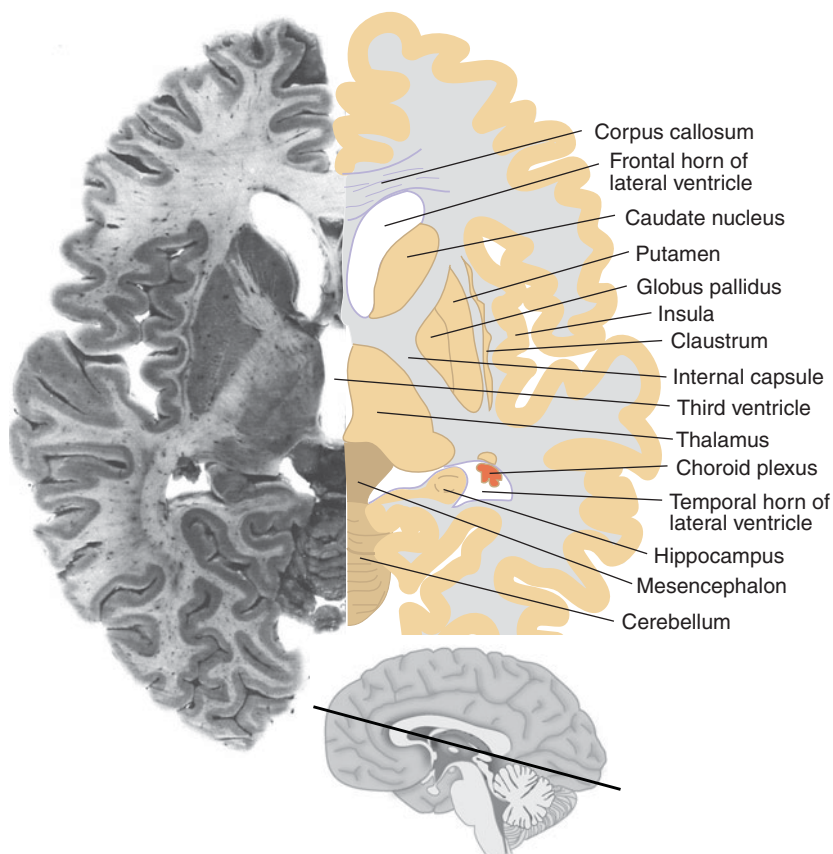


FIGURE 6.30 *The internal structure of the cerebral hemisphere.* The left half is a photograph of a horizontal section through the left hemisphere. Compare with Fig. 6.28.

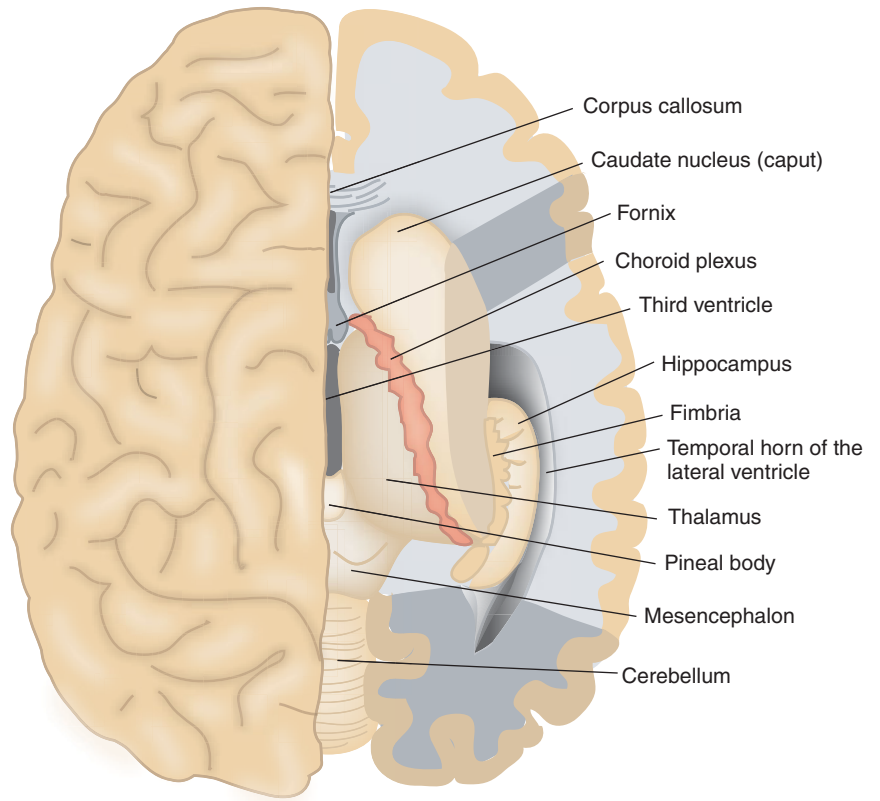


FIGURE 6.31 *The internal structure of the cerebral hemispheres.* The upper parts of the right hemisphere has been cut away to open the lateral ventricles. In the anterior part, the caudate nucleus forms the bottom and lateral wall of the ventricle. In the temporal horn of the ventricle, the hippocampus forms the medial wall.

connections it has been suggested to deal with sensory integration. One hypothesis proposes that by virtue of its integrative potential, claustrum may be linked with the formation of conscious percepts.

THE CEREBELLUM

The cerebellum (the “little brain”) is first and foremost of importance for the execution of movements; like the basal ganglia, it belongs to the motor system. The cerebellum is located in the posterior cranial fossa, dorsal to the brain stem (Figs. 6.2 and 6.32). It is connected with the brain stem anteriorly by way of three stalks of white matter on each side: the inferior, middle, and superior cerebellar peduncles (Figs. 6.14 and 3.19). In general, the **inferior cerebellar peduncle**, or **restiform body** (*corpus restiforme*), contains fibers that carry impulses from the spinal cord to the cerebellum, whereas the **middle cerebellar peduncle**, or the **brachium pontis**, conveys information from the cerebral cortex. The **superior cerebellar peduncle**, or the **brachium conjunctivum**, contains most of the fibers conveying impulses out of the cerebellum—that is, the cerebellar efferent fibers.

Like the cerebrum, the cerebellum is covered by a layer of gray substance, the **cerebellar cortex** (cortex cerebelli), with underlying white matter. Enclosed in the white matter are regions of gray matter, the **central (deep) cerebellar nuclei** (see Fig. 24.14). From the neurons in these nuclei come the majority of efferent fibers that convey information from the cerebellum to other parts of the CNS (the cerebral cortex, various brain stem nuclei, and the spinal cord).

The cerebellar surface is extensively folded, forming numerous narrow sheets, or **folia**, that are predominantly oriented transversely (Fig. 6.32). The fissures and sulci between the folia are partly very deep; the deepest among them divide the cerebellum into **lobes** (Fig. 6.32A; see also Fig. 24.2). In addition, the cerebellum can be subdivided macroscopically on another basis. In the posterior part of the cerebellum a narrow middle region is situated deeper than the much larger lateral parts (Fig. 6.32B). This middle part of the cerebellum is called the **vermis** (worm) and is present also in the anterior part of the cerebellum, although not as clearly distinguished from the lateral parts as posteriorly. The lateral parts are called the **cerebellar hemispheres**. A small bulbous part on each is connected medially with a thin stalk to the vermis. This part is

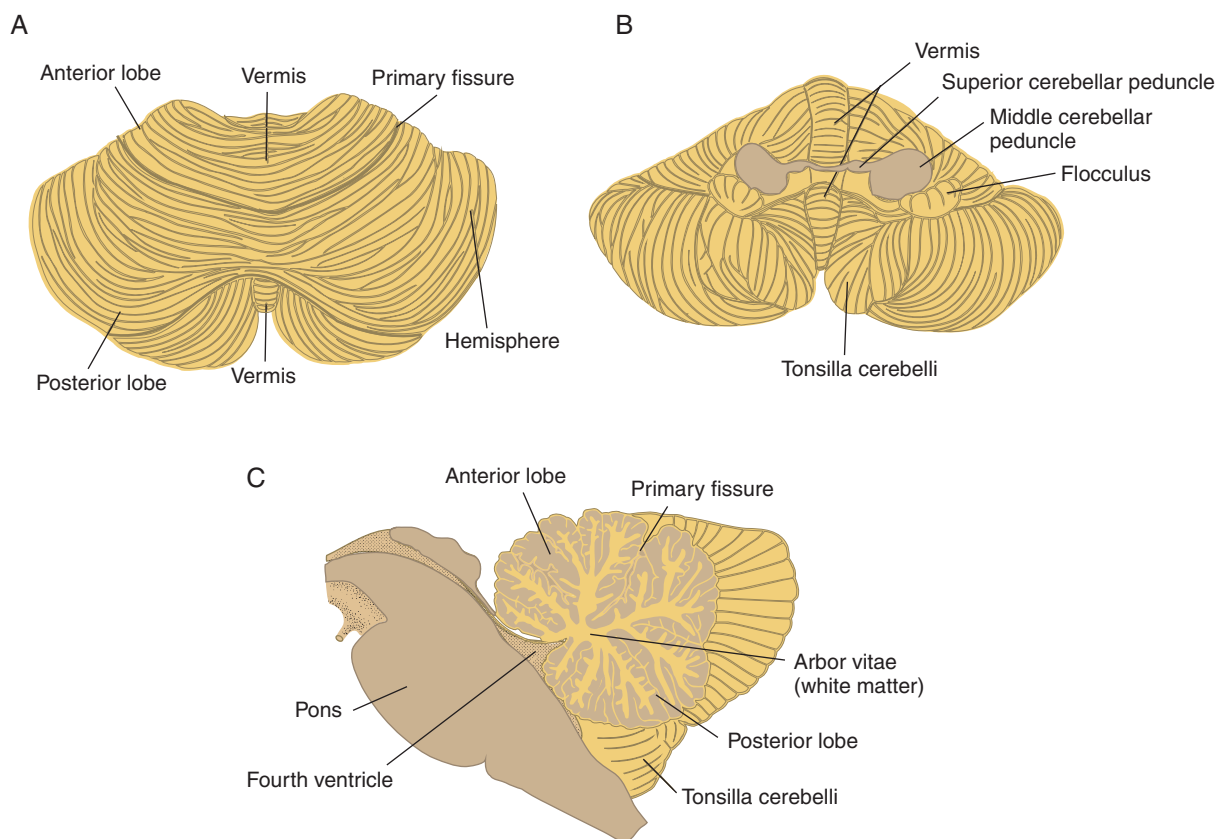


FIGURE 6.32 *The cerebellum.* A: Seen from the rostoposterior aspect. B: Seen from the ventral aspect (the side facing the brain stem). C: Midsagittal section showing the cerebellar cortex as a thin layer

with white matter underneath (white matter indicated in black in the drawing).

called the **flocculus** (Fig. 6.31B) and lies close to the middle cerebellar peduncle, just posterior to the seventh and eighth cranial nerves (Fig 6.13). A midsagittal section through the cerebellum (Fig. 6.32C) shows clearly the deep sulci and fissures. The white substance forms a

treelike structure called the **arbor vitae** (the tree of life, which is not very fitting since the cerebellum is not necessary for life). The fourth ventricle, extending into the cerebellum like the apex of a tent, is also evident in the midsagittal section.

7 | The Coverings of the Brain and the Ventricular System

OVERVIEW

The central nervous system (CNS) is well protected against external forces as it lies inside the skull and the vertebral canal. In addition to this bony protection, the CNS is wrapped in three membranes of connective tissue—the **meninges**—with fluid-filled spaces between the membranes. In fact, it is loosely suspended in a fluid-filled container. The innermost membrane—**pia**—is thin and adheres to the surface of the brain at all places. The outermost membrane—**dura**—is thick and fibrous and covers the inside of the skull and spinal canal. The **arachnoid** is a thin membrane attached to the inside of the dura. The **subarachnoid space** is with **cerebrospinal fluid (CSF)** and lies between the pia and the arachnoid.

The **ventricular system** consists of irregularly shaped, fluid-filled cavities inside the CNS. There are four dilations or ventricles. The two lateral ventricles are largest and are located in the cerebral hemispheres. They communicate with the third ventricle lying between the two thalami. The third ventricle, situated between the cerebellum and the brain stem, communicates with the fourth ventricle via the narrow **cerebral aqueduct**. Vascular **choroid plexuses** in the ventricles produce about 0.5 liters of CSF per day. The CSF leaves the ventricular system through openings in the fourth ventricle and enters the subarachnoid space. Drainage of CSF occurs through small evaginations—**arachnoid villi**—of the arachnoid emptying into venous sinuses and along cranial and spinal nerve roots and then into extracranial lymphatic vessels. The CSF has a protective function, it minimizes accumulation of harmful substances in the nervous tissue, and it probably serves as a signal pathway.

THE MENINGES

The Pia Mater, the Arachnoid, and the Subarachnoid Space

The innermost one is the vascular **pia mater** (usually just called pia). It follows the surface of the brain and spinal cord closely and extends into all sulci and depressions of the surface (Figs. 7.1 and 7.2). Thin vessels

pass from the pia into the substance of the brain and supply the external parts, such as the cerebral cortex, with blood (the deeper parts of the brain are supplied by vessels entering the brain at its basal surface). The next membrane, the **arachnoid**, does not follow the uneven surface of the brain but extends across depressions, fissures, and sulci. Between the pia and the arachnoid exists a narrow space, the **subarachnoid space**, which is filled with **CSF** (Figs. 6.1, 7.1, and 7.5). Numerous thin threads of connective tissue connect the pia with the arachnoid, thus spanning the subarachnoid space. The depth of the subarachnoid space varies from place to place because the arachnoid, as mentioned, does not follow the surface of the brain. Where it crosses larger depressions, the subarachnoid space is considerably widened, forming so-called **cisterns** filled with CSF. Several cisterns are found around the brain stem, but the largest one, the **cisterna magna**, or **cerebellomedullary cistern**, is located posterior to the medulla below the cerebellum (Figs. 6.1 and 7.5). The CSF enters the cisterna magna from the fourth ventricle (Fig. 7.5).

The subarachnoid space is continuous around the whole CNS. Substances released into the subarachnoid space at one place therefore quickly spread out. A **subarachnoid hemorrhage**, for example, most often caused by rupture of a vessel at the base of the brain, quickly leads to mixing of the CSF with blood. Thus, a sample of CSF taken from the dural sac at lumbar levels will be bloody.

The Dura Mater

The outermost membrane, the **dura mater** (usually just called the dura), is thick and strong because it consists of dense connective tissue (Figs. 7.1, 7.2, and 7.5). The dura covers closely the inside of the skull, and its outermost layers constitute the periosteum. The arachnoid follows the inside of the dura closely so that there is only a very narrow space between these two meninges, the **subdural space** (Figs. 7.1 and 7.2). The dura extends down into the vertebral canal to enclose the spinal cord. It extends further down than the cord, however, forming a sac around the roots of the lower spinal nerves (the cauda equina). This **dural sac** extends down to the

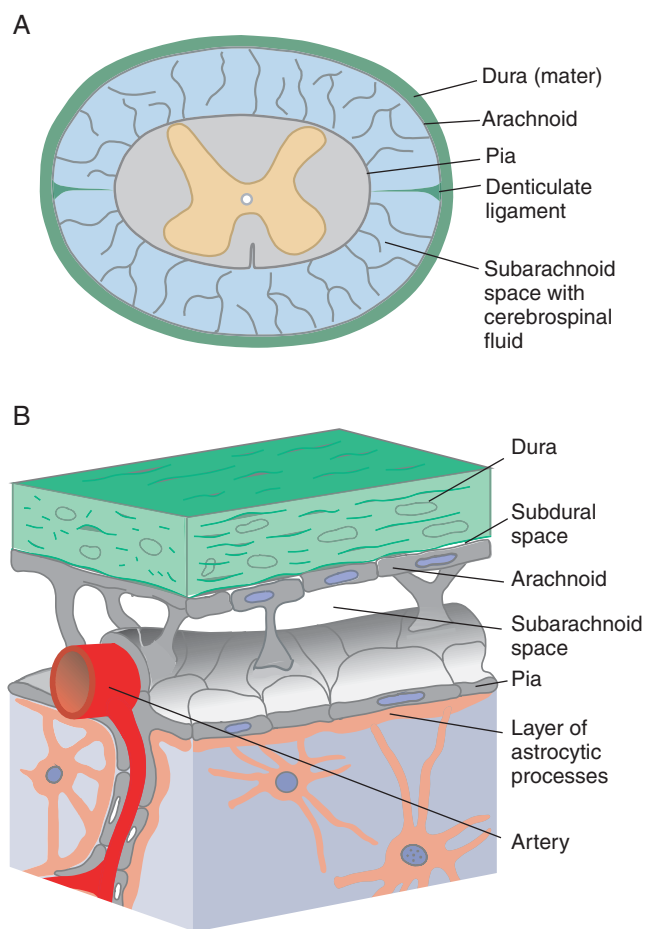


FIGURE 7.1 The meninges and the subarachnoidal space of the spinal cord.

level of the second-third sacral vertebra (Fig. 6.3). Thus below the level of the first and second lumbar vertebrae (the lower end of the cord), the dural sac contains only spinal nerve roots and CSF (Fig. 6.3). This is a safe place to perform a **lumbar puncture**—that is, enter the subarachnoid space with a needle to take samples of the CSF, as there is no danger of harming the cord.

Anchoring of the Brain and Spinal Cord

In a few places, the dura forms strong infoldings, serving to restrict the movements of the brain within the skull. Large movements can stretch and damage vessels and nerves connecting the brain with the skull (one of the possible effects of head injuries). From the midline, the **falx cerebri** extends down between the two hemispheres (Figs. 7.2 and 8.7). Posteriorly, the falx divides into two parts that extend laterally over the superior face of the cerebellum and attach to the temporal pyramid. These two folds meet in the midline and form the **cerebellar tentorium** (see Fig. 8.9). In the anterior part of the tentorium, there is an elongated opening for the brain stem. If the pressure in the skull above the tentorium increases (due to bleeding, a tumor, or brain edema), part of the temporal lobe may be pressed down or **herniate** between the tentorium and the brain stem, harming the brain stem temporarily or permanently.

The **spinal cord** is anchored to the meninges partly by the spinal nerves and partly by two thin bands, the **denticulate ligaments** (Fig. 7.1A), extending laterally from the cord to the arachnoid and dura (this is not a continuous ligament but one that forms 21 lateral extensions from the cord to the dura). The spinal cord nevertheless moves considerably up and down in the dural sac with movements of the vertebral column (the length of the vertebral canal varies by almost 10 cm from maximal flexion, when it is longest, to maximal extension).

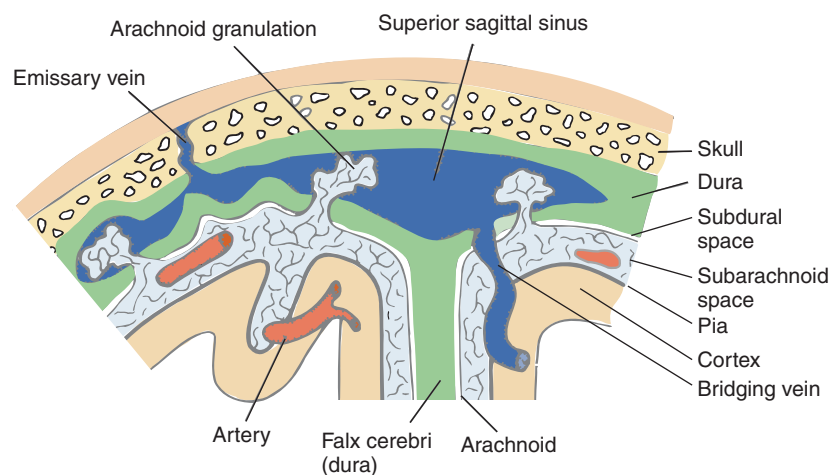


FIGURE 7.2 The meninges, the subarachnoidal space, and the superior sagittal sinus. Schematic of a frontal section through the head, with the skull and the brain. See also Figs. 1.27 and 3.37.

The Meninges and Stiffness of the Neck

Infection of the meninges, **meningitis**, typically produces stiffness of the neck. Every attempt to bend the neck or the back forward evokes an immediate reflexive muscular resistance. When the doctor tries to lift the patient's head off the pillow, the neck is kept straight. The infection causes inflammation of the meninges, which also extends onto the vessels and the nerve roots in the subarachnoid space. Forward bending (flexion) of the vertebral column elongates the spinal canal, as mentioned, and that stretches the meninges, the vessels, and the nerve roots. Most likely, this accounts for the intense pain associated with any effort to flex the back or neck (similar to the pain felt by stretching an inflamed area of the skin or a joint capsule). Straining or coughing also causes pain in patients with meningitis. However, other means than infections by bacteria and viruses can produce such **irritation of the meninges**. For example, one strong irritant is blood in the subarachnoid space. Thus, stiffness of the neck is by itself only a sign of meningeal irritation and does not indicate a specific cause.

Meningeal irritation usually causes a strong **headache**. This occurs most dramatically with **subarachnoid hemorrhage**, in which an intense headache starts abruptly the moment the bleeding starts and blood flows into the subarachnoid space. In such instances, the cause is usually spontaneous rupture of an **aneurysm** (a sac-like dilatation) on one of the arteries at the base of the skull.

THE CEREBRAL VENTRICLES AND THE CEREBROSPINAL FLUID

We have mentioned the ventricular system several times in connection with treatment of the various parts of the brain. Here we consider the ventricular system as a whole, along with the CSF.

The Location and Form of the Ventricles

The thin central canal of the cord continues upward into the brain stem. There the canal widens to form the **fourth ventricle** at the posterior aspect of the medulla and pons (Figs. 6.1, 7.3, and 7.5). The ventricle has a tentlike form with the apex projecting into the cerebellum and two **lateral recesses** (recessus lateralis). The diamond-shaped **rhomboid fossa** at the dorsal aspect of the brain stem (Fig. 6.19) forms the “floor” of the fourth ventricle, while the cerebellar peduncles form the lateral walls.

The **third ventricle** is a thin slitlike room between the two thalami (see Figs. 6.27, 6.30, 6.31, 7.3, and 7.5). During embryonic development, the primordia of the

hemispheres become closely apposed to the diencephalon (see Fig. 9.13). The loose masses of connective tissue between the hemispheres form an approximately horizontal plate that constitutes the roof of the third ventricle. The choroid plexus is attached to the inside of the roof (see Figs. 6.23 and 7.5).

The two **lateral ventricles** represent the first and second ventricles, but these terms are not used. From a central part in the parietal lobe, the lateral ventricles have processes called horns into the three other lobes: an **anterior (frontal) horn**, a **posterior (occipital) horn**, and an **inferior (temporal) horn** extending downward and anteriorly into the temporal lobe (Fig. 7.3; see also Figs. 6.29 and 6.31). The anterior horn is the largest and is bordered medially by the **septum pellucidum**,¹ whereas the head of the caudate nucleus bulges into it from the lateral side (Figs. 6.29 and 7.5). The central part of the ventricle lies just above the thalamus (Fig. 6.31). The inferior horn starts at the transition between the central part and the posterior horn and follows the temporal lobe almost to its tip (Fig. 7.3). Medially in the inferior horn there is an elongated elevation, the **hippocampus** (Figs. 6.27 and 6.31), formed by invagination of the ventricular wall from the medial side by the **hippocampal fissure**.

The **curved form** of the lateral ventricles can be understood on the basis of the embryonic development of the cerebrum (Fig. 7.4). Both the lateral ventricles and the nervous tissue in its walls (such as the caudate nucleus and the hippocampus) eventually obtain a curved shape.

The Cerebrospinal Fluid Is Produced by Vascular Plexuses in the Ventricles

All of the ventricles are filled with a clear, watery fluid, the **CSF**. Most of the fluid is produced by vascular tufts, the **choroid plexus**.² This is present in all four ventricles (Fig. 7.5), but the largest amount of choroid plexus is in the lateral ventricles (Fig. 6.31). The plexuses arise in early embryonic life by invaginations of the innermost membrane (the pia mater) at sites where the wall of the neural tube is very thin (Fig. 7.6). An elaborate structure of thin, branching protrusions, or **villi**, arises here (Fig. 7.7). The choroid plexuses attach to the wall of the ventricles with a thin stalk (**tela choroidea**). The surface of the villi is covered by **simple cuboid**

1 There is a thin slit between the septum pellucidum of the two sides that has nothing to do with the ventricles. In embryonic development, the slit is continuous with the room between the two hemispheres but is later closed when the corpus callosum grows across the two hemispheres. Downward, the fornices close the slit (Fig. 6.29).

2 Early observations in experimental animals suggested that the choroid plexus is not solely responsible for CSF production. Thus, removal of the choroid plexus did not prevent development of hydrocephalus after blocking the CSF drainage. It is now assumed that in humans 10% to 30% of the CSF is produced by brain interstitial fluid passing through the ependyma.

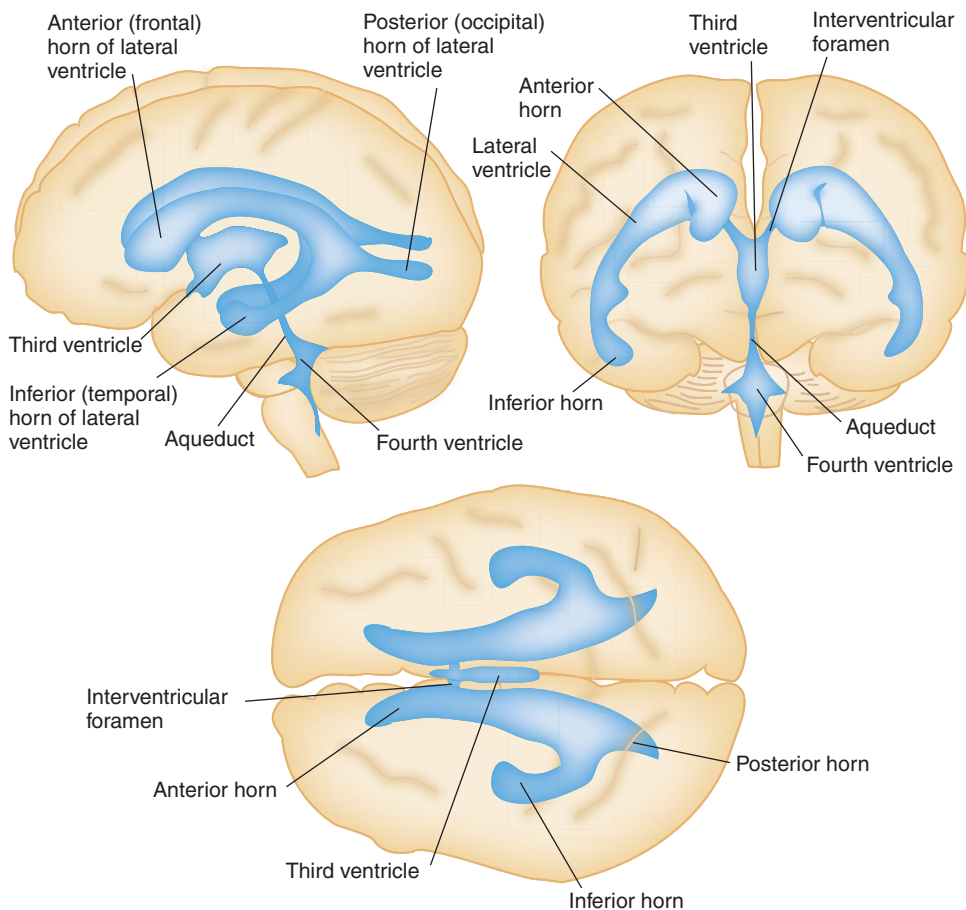


FIGURE 7.3 The ventricular system of the brain.

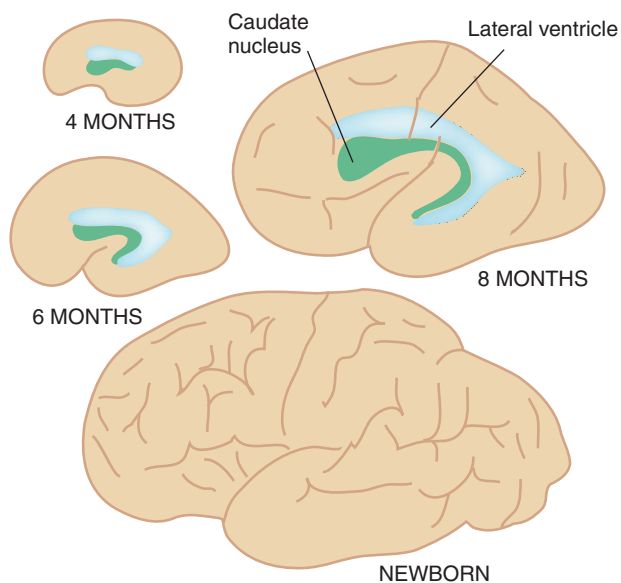


FIGURE 7.4 Development of the cerebral hemispheres and the lateral ventricles. The characteristic arched shape of the ventricles can be explained by the manner in which the hemispheres fold during their growth in embryonic life. The caudate nucleus, located in the wall of the lateral ventricle, is indicated in blue. All structures in the wall of the ventricles attain the curved shape. The figure also shows the development of gyri and sulci. Compare the pattern in the newborn and in the adult (Fig. 6.25). (Based on Hamilton, Boyd, and Mossman 1972.)

epithelium (which is continuous with the ependyma covering the inside of the ventricles). The epithelial cells have microvilli that increase their surface in contact with the CSF. The interior of the villi is composed of loose connective tissue with numerous capillaries of the **fenestrated** type, which are rather leaky. Therefore, the hydrostatic pressure inside the capillaries produces a net flow of water with solutes (and a fair amount of proteins) into the interstitial space of the villi. This protein-rich fluid cannot leave the villi directly, however, because the epithelial cells covering the villi attach to each other with **tight junctions** (Fig 7.7). The transport of water through the epithelium is caused by active transport of sodium. Thus, the pumping of sodium produces an osmotic gradient so that water diffuses from the interior of the villi into the ventricles. Other ions, such as chloride and bicarbonate, follow the water passively. The epithelial cells of the choroid plexus are equipped with water channels or **aquaporins** (AQP1) on the apical surface (that is, facing the CSF). Presumably, the aquaporins are important for the rapid transport of water. Indeed, the **rate of CSF secretion** by the choroid plexus (0.2 mL/min/g tissue) is much higher than in other secretory epithelia in the body.

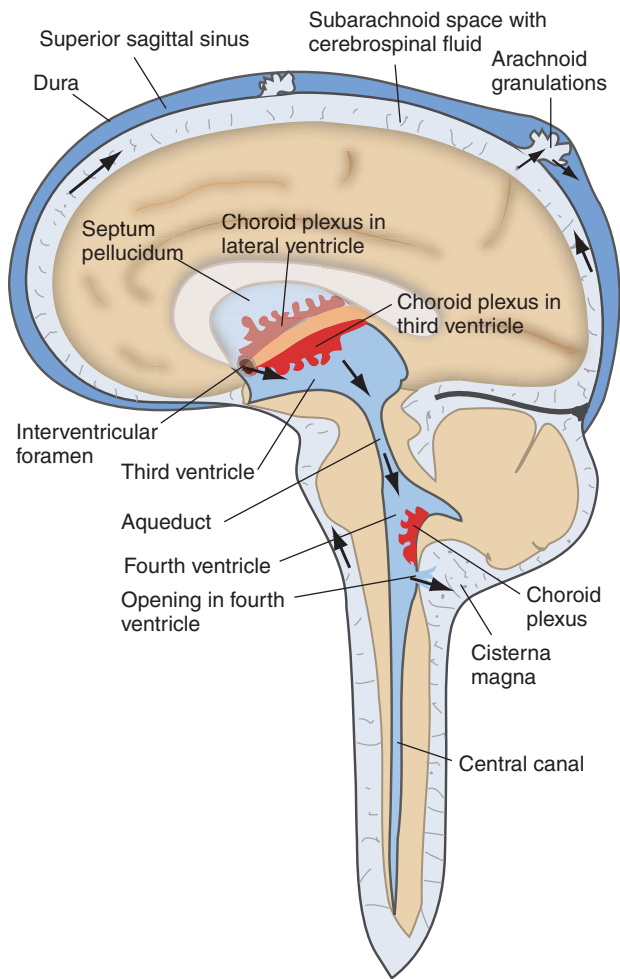


FIGURE 7.5 The ventricular system, the subarachnoid space, and the choroid plexus. All ventricles contain choroid plexus. Arrows indicate the flow of the cerebrospinal fluid from the ventricles into the subarachnoid space through openings in the walls of the fourth ventricle.

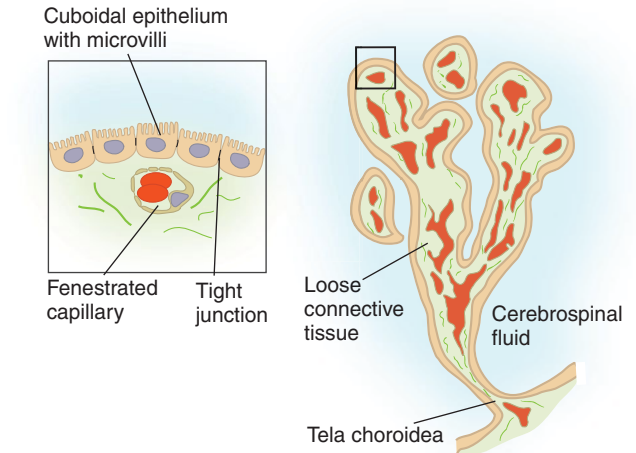


FIGURE 7.7 Structure of the choroid plexus, exemplified by one villus (inset). Arrows indicate the flow of fluid from the capillaries to the ventricles.

The epithelium of the choroid plexus represents a barrier between the blood and the CSF, the **blood–CSF barrier**. Thus, many substances that can leave the capillaries of the choroid plexus cannot enter the CSF. This is obviously important, because neurons are extremely sensitive to changes in the composition of their environment. We will later describe a similar but even more important barrier between the blood in the brain capillaries and the brain interstitial fluid.

Composition and Functions of the Cerebrospinal Fluid

The concentration of sodium, potassium, and several other **ions** is about the same in the CSF as in the blood (there are some minor differences, however). The concentration of **glucose** is about two-thirds that in the blood. A major difference concerns **proteins**: there is normally very little protein in the CSF (less than 0.5% of the plasma protein concentration).

Water and soluble substances are freely exchangeable between the CSF and the interstitial fluid of the nervous tissue because the ependyma is freely permeable to water and even small protein molecules. It is not surprising, therefore, that many neurotransmitters, peptides, and other neuroactive substances occur in the CSF, and their presence there does not by itself signify a functional role. Some substances, however—notably hormones synthesized in the anterior pituitary—are apparently actively secreted into the CSF, not simply accepted by passive diffusion. Some other substances appear to use the CSF as a means to reach specific receptors close to the ventricles.

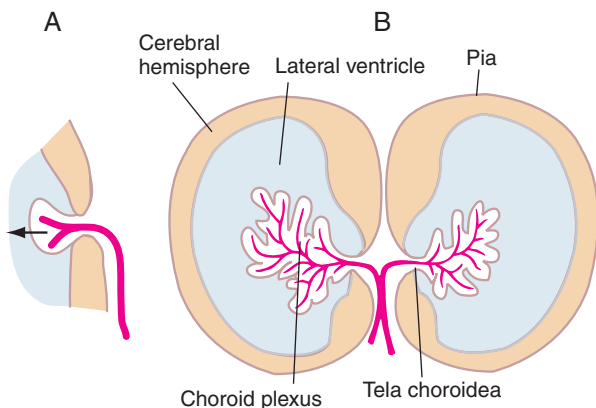


FIGURE 7.6 Embryonic development of the choroid plexus. Schematic of a frontal section through the cerebral hemispheres at an early stage, showing how the choroid plexus is formed by invaginations of the pia into the ventricles.

The CSF has an important **protective function** because the brain almost floats in it. Thus, theoretically **buoyancy** reduces the weight of the brain to about 50 g, which means less traction on vessels and nerves connected to the CNS. Further, the effect on the brain of blows to the head is dampened because water has to be pressed aside before the brain hits its hard surroundings (the skull).

Another possible functional role of the CSF can be deduced from the fact, mentioned above, that water and solutes pass freely between it and the extracellular fluid (interstitium) of the nervous tissue. This means that diffusion into the CSF may minimize **accumulation of harmful substances** in the nervous tissue (such as potassium ions during prolonged periods of intense neuronal activity). This would be of significance, however, only for neurons that are fairly close to the ventricles, as the diffusion of molecules in the labyrinth-like brain interstitium is slow (much slower than in free water). Thus, after injecting representative substances in the brain, the concentration is reduced by 90% some 1 to 3 mm away from the injection site.

Regardless of the normal functions of substances in the CSF, examination of the CSF composition gives valuable information of the extracellular fluid of the brain. This fluid compartment is difficult to examine directly, but because the ependyma is freely permeable, one can safely assume that the composition of the CSF matches fairly well the environment of the neurons.

Cerebrospinal Fluid as a Signal Pathway

The concentration of the hormone **melatonin** (which influences sexual functions and circadian rhythms by binding to receptors in the hypothalamus) is twice as high in the third ventricle as in the lateral ventricles and 100 times higher than in cerebral arteries. This suggests that melatonin is secreted into the CSF and uses this as its main transport medium, whereas the bloodstream is of minor importance.

Some studies indicate that substances in the CSF might be of importance for **sleep**. Thus, by transferring CSF from a sleep-deprived animal, the recipient becomes sleepy. The responsible substance has not been identified but **interleukin 1 β** (IL-1 β) is a likely candidate. Thus, its concentration in the CSF increases by sleep deprivation and has also been shown to induce sleep. The further signal pathway of IL-1 β from CSF to relevant neurons is not known, but it finally influences the neurons in the reticular formation that are responsible for regulation of sleep and wakefulness.

Several **growth factors** are synthesized and secreted by the choroid plexus. While the choroid plexus epithelium expresses receptors for such growth factors—suggesting an autocrine function—they may also be expected to act on nervous tissue close to the ventricles.

Circulation and Drainage of the Cerebrospinal Fluid

About **one-half liter** of CSF is produced each day, yet the total volume of fluid in the ventricles and the subarachnoid space is only 130 to 140 mL (approximately 20 mL is in the ventricles).³ In addition, approximately 75 mL surrounds the spinal cord. Thus, the total amount is renewed several times a day. This means that effective means of drainage must exist.

The fluid produced in the lateral ventricles flows into the third ventricle through the **interventricular foramen** (of **Monro**) (Fig. 7.5; see Fig. 6.23). From there the fluid flows through the narrow **cerebral aqueduct** to the fourth ventricle. The choroid plexuses in the third and fourth ventricles add more fluid. The fluid leaves the ventricular system and enters the subarachnoid space (more specifically, the cisterna magna) through three openings in the fourth ventricle: one in the midline posteriorly (the **foramen of Magendie**) and two laterally (the lateral recesses or **foramina of Luschka**) (Fig. 7.5). The fluid then spreads out over the entire surface of the brain and spinal cord. From the base of the brain, there is an upward stream along the lateral aspects of the hemispheres toward the midline (Fig. 7.5). The flow velocity is uneven, however, as shown by following the spread of injected substances. After injection into the lateral ventricles labeled substances are detectable after a few minutes in the subarachnoid space, but then moves very slowly over the cerebral hemispheres (several hours) and down to the lumbar cistern (one hour).

The routes of CSF **drainage** are not completely known, although emptying into venous sinuses by way of **arachnoid villi** is usually assumed to constitute the main route. The arachnoid villi are small evaginations of the arachnoid. Several villi together form macroscopically visible protrusions called **arachnoid granulations**, which are particularly prominent along the superior sagittal sinus (Figs. 7.2 and 7.5). Passage of fluid from the subarachnoid space to the venous sinuses is presumably caused by a difference in hydrostatic pressure, the pressure being higher in the subarachnoid space (about 15 cm of H₂O) than in the sinuses (7–8 cm of H₂O). Nevertheless, we do not actually know how much of the CSF that is drained in this way.⁴

3 The total volume of the ventricles has been determined mainly by use of plastic casts in fixed brains. The average total volume is probably some less than 20 cm³. The individual variations are surprisingly large, however (the normal range is probably from 7 to 30 cm³). Most studies have examined only the lateral ventricles, finding an average volume of about 7 cm³ for each. The volumes of the third and fourth ventricles appear to be some less than 1 cm³ for each. Even though the ventricular size increases somewhat with age, this can explain only a small fraction of the individual variations. In the same individual, however, the two lateral ventricles are quite similar in volume.

4 One reason for questioning the role of the arachnoid granulations in CSF drainage is that they are not present prenatally while the choroid plexuses are well developed by the end of the embryonic period (about 8 weeks after fertilization). Therefore, in the fetus and newborn other routes seem to be responsible for most or all of the CSF drainage.

Another, and perhaps quantitatively more important route of drainage, is along cranial and spinal **nerve roots** and then into **extracranial lymphatic vessels** (the nervous tissue contains no lymphatics). Animal experiments indicate that drainage through the **cribriform plate** (where the olfactory nerve fibers enter the skull, see Fig. 8.10) may be of particular significance.

Brain Edema, Herniation, and Hydrocephalus

Because the CNS and the CSF are located in a closed container with rigid walls, the pressure inside the container increases if, for some reason, the amount of substance inside it increases. The intracranial pressure is the same for the nervous tissue and the CSF, but it can be measured most conveniently in the latter. For each heartbeat, for example, the pressure inside the cerebral ventricles increases slightly because more blood is pumped into the brain. Correspondingly, the pressure increases on coughing and straining because the drainage of venous blood from the cranial cavity and the vertebral canal is reduced.

Regardless of cause, any **intracranial expansive process** leads to increased intracranial pressure. When the pressure increases, the blood pressure increases to maintain cerebral blood flow. Thus, abnormally elevated blood pressure, usually combined with slowing of the heart rate, is one of the signs of severely increased intracranial pressure. However, an increase above a certain level reduces blood flow and the functioning of the brain suffers (with signs of confusion and eventually loss of consciousness). This occurs, for example, in patients with **brain edema**. This is a dangerous complication of acute brain damage caused by, for example, head injuries. The edema is caused by extravasation of fluid from the brain capillaries. Similarly, an **intracranial hemorrhage**, which may arise from vessels inside or outside the brain substance, can also cause a dangerous increase of the intracranial pressure. If the expansion process is located in the posterior fossa (below the cerebellar tentorium; see Fig. 8.9), the **cerebellar tonsils** (see Figs. 6.13 and 6.32C) may be dislocated downward into the foramen magnum and thus compress the medulla (tonsillar herniation). If the expansive process is located above the tentorium (in the middle or anterior fossa), the **uncus** of the temporal lobe (see Fig. 6.26) can be dislocated downward beneath the edge of the tentorium and compress the brain stem at the level of the mesencephalon (see Figs. 6.29 and 8.9). Both forms of **herniation** may lead to serious brain damage or death.

A condition with increased amount of CSF and dilatation of the ventricles is called **hydrocephalus**. It is usually assumed that the condition arises because of

increased intraventricular pressure due to obstruction of the drainage of the CSF—for example, by closure of the aqueduct (inflammation, tumor, bleeding) or the outlets from the fourth ventricles. Even if this may be the cause in some, it cannot explain all cases of hydrocephalus. Thus, some persons develop hydrocephalus in spite of apparently normal intraventricular pressure and no sign of obstruction (communicating or normal pressure hydrocephalus).

If obstruction of CSF flow from the ventricles occurs in an adult, in whom the sutures of the skull have grown firmly together, the intracranial pressure increases markedly, with dramatic symptoms. If the condition arises in early childhood before closure of the sutures, however, the size of the head grows abnormally, with the skull yielding to the increased intracranial pressure. This may continue for some time with surprisingly few signs of cerebral dysfunction, but the development of the brain will eventually suffer, with results such as mental retardation. Hydrocephalus in children may be treated by shunting the CSF directly into a big vein outside the skull.

Examination of the Cerebrospinal Fluid and the Ventricles

Examination of the CSF can provide valuable information about neurological diseases. A sample of the fluid is usually withdrawn with a thin needle from the subarachnoid space in the dural sac—that is, below the level of the second lumbar vertebra (lumbar cistern). This way there is no risk of damaging the spinal cord (see Fig. 6.3). When the tip of the needle is in the subarachnoid space, the intracranial pressure can also be measured. Examination of the fluid, in part under the microscope, can give information about possible bleeding into the subarachnoid space and about infections and inflammations of the brain itself (**encephalitis**) or the meninges (**meningitis**). In the latter case, there are numerous leukocytes in the CSF. The concentration of proteins, normally very low, may increase in certain diseases (e.g., in **multiple sclerosis**, in which the proteins are antibodies against components of the nervous tissue).

The **shape** and **size** of the ventricles can be determined noninvasively in living subjects by use of computer tomography (CT) and magnetic resonance imaging (MRI) (see Figs. 6.2, 6.28, 23.4, and 32.11). Atrophy of the nervous tissue of the brain—for example, atrophy of the cerebral cortex, which occurs in dementia—leads to dilatation of the ventricles, whereas expansive, space-occupying processes like hemorrhages and tumors may distort and compress the ventricles.

8 | The Blood Supply of the CNS

OVERVIEW

Of all cell types in the body, neurons are the most sensitive to interruption of their supply of oxygen (anoxia). Only a few minutes' stop in the blood flow may cause neuronal death. The oxygen consumption of the brain is high even at rest. Therefore, the blood supply of the central nervous system (CNS) is ample, and the brain receives about 15% of the cardiac output at rest. Regulatory mechanisms ensure that the brain gets what it needs—if necessary, at the expense of all other organs. The arteries of the brain lie within the cranial cavity and are mostly devoid of anastomoses (connections) with arteries outside the skull. Therefore, other arteries cannot take over if the intracranial ones are narrowed or occluded.

The cerebral blood flow is largely controlled by the local conditions in the nervous tissue, that is, there is a high degree of **autoregulation**. Local changes in the concentrations of ions, oxygen, carbon dioxide, and various signal substances determine the resistance offered by the arterioles. Brain vessels receive sensory innervation from the trigeminal nerve.

In most organs, small-molecule substances pass the capillary wall, and their concentration is therefore similar in the blood plasma and in the interstitial fluid. In contrast, the CNS exerts strict control of what is let in. The **blood–brain barrier** (a similar barrier exists between the blood and the cerebrospinal fluid [CSF]) is due mainly to special, selective properties of the brain capillaries.

In a few small regions adjoining the ventricles, the capillaries are fenestrated and hence let substances from the blood pass through easily. At such places, neurons are exposed to substances of the blood that do not enter other parts of the brain. These regions are called the **circumventricular organs**.

Broadly speaking, the **internal carotid artery** supplies most of the cerebral hemispheres, whereas the **vertebral artery** supplies the brain stem and the cerebellum. **Communicating arteries** at the base of the skull establish anastomoses between the posterior (vertebral) and the anterior (internal carotid) cerebral circulations.

CEREBRAL MICROCIRCULATION AND THE BLOOD–BRAIN BARRIER

The brain has a very high density of capillaries, and neurons are seldom more than 10 μm from the

nearest capillary. The total length of all capillaries in the brain are said to be 400 miles, and their total surface area more than 20 m^2 . To understand the normal properties and the responses to disease of brain capillaries, however, they cannot be studied in isolation. The term **neurovascular unit** serves to emphasize the close structural and functional relationship between neurons, glial cells, associated capillary-endothelial cells, basal lamina, and pericytes (Fig. 8.1).

Regulation of Cerebral Circulation

Regulation of the blood flow is one among several factors governing the composition of the brain's extracellular milieu—that is, the concentration of ions, neuroactive substances, nutrients, and water (osmolarity). Control of the properties of astrocytes and the blood–brain barrier are other important factors (see Chapter 2, under “Astroglia and the Control of Neuronal Homeostasis”).

The cerebral circulation exhibits a high degree of **autoregulation**—that is, conditions in the brain itself determine the blood flow. If the blood pressure falls, the arteries dilate. This reduces vascular resistance so that the blood flow is upheld. If the blood pressure rises, the opposite happens: the arteries constrict. This is an important control mechanism, since increased capillary hydrostatic pressure may cause brain edema. The brain maintains almost constant blood flow as long as the systolic pressure is between 60 and 160 mm Hg. If the pressure falls below 60 mm, however, the flow falls steeply and the person becomes unconscious.

Among the many factors that control cerebral blood flow, local changes in the immediate surroundings of the neurons have an important role. These are changes in concentrations of ions (among them H^+ ions), CO_2 and O_2 . **Hypoxia** (abnormally low concentration of O_2) and **hypercapnia** (above-normal concentration of CO_2) both cause marked vasodilatation and increased cerebral blood flow. Increased local blood flow is closely coupled to increased neuronal activity, and this phenomenon is utilized in studies of correlations between changes in brain activity and behavior (see “Regional Cerebral Blood Flow and Neuronal Activity”).

Autonomic circulatory control seems to play a minor part in the brain (in contrast to in most other organs), even though **sympathetic fibers** innervate brain vessels. Such fibers release norepinephrine, neuropeptide Y (NPY) and possibly ATP, and stimulation causes vasoconstriction.

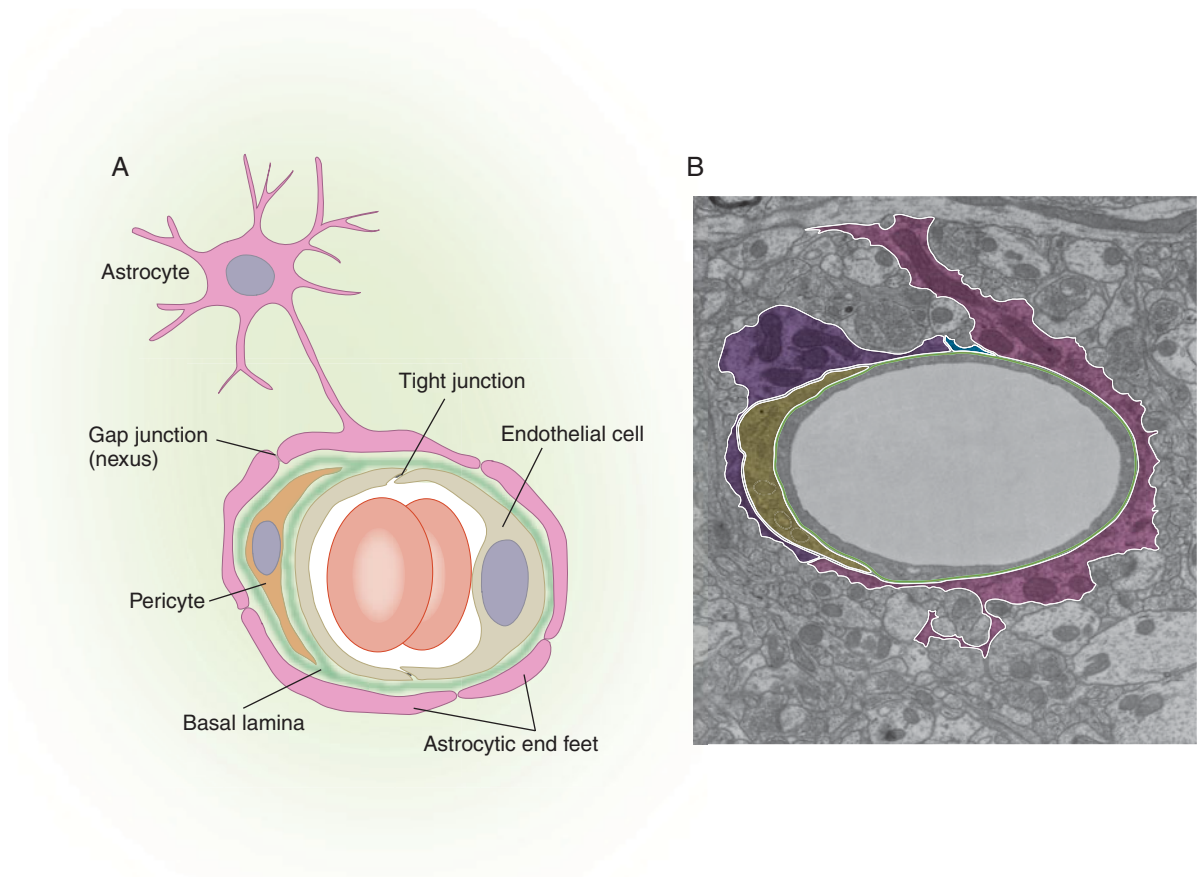


FIGURE 8.1 *The blood–brain barrier/neurovascular unit.* **A:** Schematic drawing showing the main features. Important elements are tight junctions between the endothelial cells and a continuous layer of astrocytic end feet. Gap junctions establish low-resistance connections between the astrocytes. The basal lamina also contributes to the barrier properties of the neurovascular unit. Not shown in the figure, although included in the term neurovascular unit, are neurons that are in contact with processes of the astrocyte (see Figs. 2.3 and 2.5). **B:** Electron micrograph showing a brain capillary (hippocampus) and

its relationship to processes of three astrocytes (the processes are marked with different colors). The electron micrograph is one among many in a true series of ultrathin sections, used for three-dimensional reconstruction of the astrocytic processes. This makes it possible to decide that the marked processes belong to different astrocytes. (Courtesy of Drs. Thomas Misje Mathiesen and Ole Petter Ottersen, Department of Anatomy, Institute of Basic Medical Sciences, University of Oslo, Norway.)

Nevertheless, under normal conditions, activity of sympathetic fibers appears to have only marginal effects on blood flow. Thus, although sympathetic activity can constrict large brain arteries, peripheral small vessels dilate (probably because of local control). Brain arteries are also supplied by **serotonergic fibers** that arise in the raphe nuclei of the brain stem and that, on stimulation, mainly cause vasoconstriction.

Finally, brain arteries receive **sensory innervation** from the **trigeminal nerve**. These fibers contain substance P, calcitonin gene-related peptide (CGRP), and neurokinin. Electric stimulation of such sensory fibers causes vasodilatation, presumably by peripheral release of neuropeptides. The trigeminal arterial innervation, along with the vascular serotonin receptors, is an important factor in the pain of **migraine** attacks. Thus, the most effective drugs to prevent the pain of migraine are serotonin-receptor agonists. The drugs are

thought to act, at least in part, by preventing release of neuropeptides from trigeminal vascular nerve endings, thereby reducing perivascular inflammation.

Regional Cerebral Blood Flow and Neuronal Activity

Neuroimaging techniques, such as positron emission tomography (PET) and functional magnetic resonance imaging (fMRI), permit the visualization of brain regions activated in conjunction with specific behaviors and mental operations (see Chapter 11, under “Methods to Study Neuronal Activity and Connectivity in the Living Brain: PET and fMRI”). This is especially valuable for the study of speech, abstract problem solving, and other functions that can be studied only in humans. PET and fMRI are based on the existence of a relationship between the **regional cerebral blood flow** (rCBF) and the neuronal activity in that region. Thus, increased

neuronal activity is regularly followed in a few hundred milliseconds by increased local blood flow (hyperemia).

The mechanisms responsible for the coupling of neuronal activity to blood flow are not clear, partly due to conflicting data. One important mechanism, however, appears from animal experiments to be activation of **astrocytes** by synaptic release of **glutamate** (by binding to metabotropic glutamate receptors in the astrocyte membrane). Thus, elevated Ca^{2+} in astrocytic end feet (Fig. 8.1) would produce vasodilatation by release of vasoactive substances. Although there are several possible mediators (e.g., nitric oxide [NO] and adenosine), in vivo experiments suggest that metabolites of **arachidonic acid**, such as **prostaglandins**, may be of particular importance. Nevertheless, signals from astrocytes do not seem to be responsible for all the hyperemia caused by synaptic activity. Possibly, NO released from neurons may contribute. NO would seem well suited: it is produced in many neurons on synaptic activation, it diffuses freely, and it causes vasodilatation. A further question concerns whether the control of regional blood flow is exerted *only* on the smooth muscle cells of arterioles. Thus, the **pericytes** (Fig. 8.1) surrounding capillaries are contractile, and may contribute to the change of blood flow induced by synaptic activity.

Even though a close relationship between neuronal activity and blood flow seems to be the rule, exceptions may exist. For example, it was recently shown that blood flow may increase in **anticipation** of an event that is expected to require increased neuronal activity in a certain neuronal population. In such cases, the arterial dilatation is local and apparently not related to a global regulation of cerebral blood flow. Although the mechanism for this anticipatory vascular response is not known, it most likely involves the innervation of small cerebral vessels. If anticipatory vascular control is a regularly occurring phenomenon, it represents a source of error to the interpretation of fMRI studies, since these assume that increased local blood flow is a sign of increased neuronal activity.

Why do we Need a Blood–Brain Barrier?

In most organs of our body, small-molecule substances pass the capillary wall with relative ease, and their concentration is therefore similar in the blood plasma and in the interstitial fluid. In contrast, the CNS exerts strict “immigration” control. The **blood–brain barrier** (we have previously mentioned a similar barrier between the blood and the CSF) is mainly due to special, selective properties of the brain capillaries. Consequently, the composition of the interstitial (extracellular) fluid of the brain differs from that in most other organs. The need for such a barrier may seem obvious: substances that would disturb the delicate balance of neuronal

excitability must be kept out. Many neuroactive substances (such as glutamate, monoamines, and many neuropeptides) are produced also in peripheral tissues and are present in the blood plasma in varying concentrations. Neuronal excitability cannot be subject to incidental blood plasma variations of neuroactive substances. It should be emphasized, however, that there is not only a barrier intercalated between the blood and the brain but also **specific transport mechanisms** for certain substances that the brain needs.

Physiological Properties of the Blood–Brain Barrier

Several factors together constitute the blood–brain barrier, which is more than just a barrier since some substances are actively transported into the brain:

1. Brain capillaries are much less permeable than capillaries in most other tissues so that even many small molecules cannot pass the capillary wall. This results mainly from very extensive **tight junctions** between the endothelial cells.¹ Substances therefore have to pass through the plasma membrane of the capillaries to enter the brain interstitium. (In the capillary walls of most other organs, there are slits—as a result of less extensive tight junctions between the endothelial cells—where water can flow.) Water-soluble substances are thus effectively prevented from passing (except via specific uptake mechanisms), whereas small, lipid-soluble molecules can pass the plasma membrane with ease. Indeed, all drugs used for treating diseases of the CNS are of this kind.

2. There are very few **endocytotic (pinocytotic) vesicles** in the cytoplasm of endothelial cells in the CNS (in contrast to, e.g., muscle capillaries). Such vesicles are believed to transport small and large molecules through the endothelial wall.

3. Brain endothelial cells are equipped with **transporter molecules (P glycoprotein)** that actively expel lipid-soluble molecules—that is, such that enters by passive diffusion through the plasma membrane. Knockout mice that lack the gene for P glycoprotein illustrate its importance: they show increased sensitivity to drugs and toxins in blood plasma. Indeed, P glycoprotein transports many drugs out of the brain.

4. Brain endothelial cells appear to be able to actively pump **ions** that are present in different concentration in

¹ Brain endothelial tight junctions contain a number of **adhesion molecules** (which are transmembrane proteins). Several of these show altered expression in various neurological diseases. **Occludin** is one such protein of importance for tight junction properties. Occludin shows decreased expression in diseases with breakdown of the blood–brain barrier (e.g., multiple sclerosis). Alterations of various **claudins** have been implicated in disruption of the blood–brain barrier caused by epileptic seizures, ischemia, dementia, and HIV-1 infections.

the brain extracellular fluid and in the blood plasma (Na^+ , K^+ , Ca^{2+} , and others).

5. **Organic acids** are actively pumped out of the brain by a specific transporter (primarily expressed in the choroid plexus).

6. Water-soluble substances, such as **glucose** and some **amino acids**, are taken up by active transport after binding to specific receptors in the endothelial membrane (carrier-mediated transport). Glucose is an example of a substance with high water solubility that nevertheless reaches high concentrations in the brain (this is necessary because the neurons depend almost solely on glucose as a source of energy). The glucose transporter **GLUT1** is specific to brain capillaries.²

7. **Macromolecules**, such as some growth factors and cytokines, are to a limited extent carried from blood plasma into the brain, probably by receptor-mediated transport.

The properties of the blood–brain barrier are of consequence for whether a drug may gain access to the brain; as a rule, only lipid-soluble drugs can reach therapeutic concentrations in the brain. Certain drugs such as barbiturates used for induction of anesthesia are highly lipid-soluble and act rapidly. Other drugs, such as penicillin, have low lipid solubility and pass the blood–brain barrier only with difficulty. In serious infections of the CNS, penicillin (or another drug with low lipid solubility) must be injected directly into the CSF, usually in the cisterna magna (see Fig. 7.5). The drug then easily enters the brain tissue because there is no barrier between the CSF and the brain interstitial fluid.

Induction and Maintenance of the Blood–Brain Barrier

The special structural and functional properties of brain capillaries depend primarily on influence from surrounding elements. If peripheral tissue is transplanted into the brain, the in-growing capillaries attain the properties characteristic of the peripheral tissue: that is, no blood–brain barrier forms. The opposite happens if brain tissue is transplanted into another organ. The astrocytic processes that surround all brain capillaries in the adult (Fig. 8.1; see also Figs. 2.3 and 2.5), are certainly important for maintaining several features of the blood–brain barrier. Their exact roles are not clear, however, partly due to conflicting data. For example, experiments show that astrocytes can induce tight junctions in brain endothelial cells. Nevertheless, other cell types (probably neuronal progenitor cells) appear to be responsible for the first formation of tight junctions in

embryonic development, as it occurs before the appearance of astrocytes.³

Modulation of the Blood–Brain Barrier in Health and Disease

In cell cultures, a number of physiological and pharmacological substances affect the properties of tight junctions. Normally, for example, a limited number of **lymphocytes** are allowed to enter the brain to “patrol” the tissue for foreign molecules. This requires specific receptor-mediated mechanisms that (most likely) transiently open tight junctions to let in lymphocytes. Unfortunately, also some **metastatic cancer cells** pass the blood–brain barrier. **Malnutrition** alters the blood–brain barrier, and substances like **histamine** and **bradykinin** make brain capillaries leaky (just as they do in other organs). According to animal experiments, even conditions with **inflammatory pain** (due to injection of irritants in the paw) can alter the permeability of the blood–brain barrier, presumably by circulating cytokines acting on tight junction proteins. **Stress** can increase the permeability of the blood–brain barrier to drugs, according to animal experiments. Presumably, this is so also in humans. For example, **pyridostigmine** (an acetylcholinesterase inhibitor used to treat myasthenia gravis) normally acts only in peripheral tissues. During the first Gulf War (1990–1991) soldiers that were given prophylactic pyridostigmine (in case of nerve gas exposure) exhibited much more central side effects than are observed when the drug is given to soldiers in peacetime.

In several different, unrelated neurological diseases, the blood–brain barrier becomes less effective, mainly due to disturbed interactions between glia and endothelial cells. At the ultrastructural level, the neurovascular unit undergoes changes, such as loss of endothelial-cell tight junctions, and changes of the astrocytes. In **multiple sclerosis**, for example, the blood brain–barrier is partly opened during exacerbations of the disease. Furthermore, the neurovascular unit is altered in **neurodegenerative diseases** (such as Alzheimer’s disease and Parkinson’s disease), in **infections** of the CNS (such as meningitis and septicemia), and in **ischemia** (due to stroke or traumatic brain injury). In the latter case, the disruption of the blood–brain barrier contributes to brain edema.

It is not clear, however, whether alterations of the neurovascular unit are causal to the diseases or are

2 Mutations of the human *GLUT1* gene cause a syndrome with infantile seizures, delayed development, and microcephaly.

3 Growth factors of the **Wnt** family (Wnt7 and Wnt8) are secreted by the neuroepithelial cells in the earliest phases of nervous system development. Binding of Wnts to specific receptors in endothelial cells induce vessels to grow into the nervous tissue and develop their characteristic properties, such as the expression of the glucose transporter GLUT1. Mice lacking the genes for Wnt7 and Wnt8 show abnormal vascular growth, impaired blood brain barrier properties, and lack of GLUT1 expression. (Wnt signaling also influences a number of other developmental processes in the brain.)

merely responses to the disease processes (which may nevertheless contribute to the disease manifestations).

Some Parts of the Brain Lack a Blood–Brain Barrier

In a few small regions adjoining the ventricles, the capillaries are fenestrated and hence let substances from the blood pass through easily. At such places, neurons are exposed to substances of the blood that do not enter other parts of the brain. These regions are called the **circumventricular organs** (Fig. 8.2). Among these, the **area postrema** is found in the lower end of the fourth ventricle, whereas the **subfornical organ** lies in the roof of the third ventricle just underneath the fornix close to the interventricular foramen. Both the area postrema and the subfornical organ contain many neurons that send their axons to other parts of the CNS and can thus mediate various specific influences on the nervous system. Neurons in the area postrema are involved in the **vomiting reflex** when this is elicited by toxic substances of the blood (see also Chapter 27, under “The Vomiting Reflex”). Neurons in the subfornical organ monitor the **salt concentration** of the blood. They send signals to the hypothalamus that can initiate responses necessary to maintain the fluid balance of the body. Further, neurons in the subfornical organ respond to circulating peptides involved in regulating **energy balance** (see Chapter 30, under “Regulation of Digestion and Feeding”). The subfornical organ and other parts of the circumventricular organs are probably also targets of substances in the blood that induce **fever** and other symptoms of infections (see Chapter 30, “Hypothalamic Neurons Are Influenced by Hormones and Fever”).

The **median eminence** (eminencia mediana) in the hypothalamus (see Fig. 30.6C) also lacks a blood–brain barrier. This region does not contain neuronal cell bodies but receives nerve fibers from other parts of the hypothalamus. Hormones released from nerve terminals in the median eminence are transported by the

bloodstream to the anterior pituitary (see Chapter 30, under “The Influence of the Hypothalamus on the Anterior Pituitary”).

The two endocrine glands that are from the brain, the **posterior pituitary** and the **pineal body**, also lack a blood–brain barrier. As with the median eminence, this is related to their release of hormones directly into the bloodstream.

ARTERIAL SYSTEM

The Brain Receives Arterial Blood from the Internal Carotid and the Vertebral Arteries

Broadly speaking, the internal carotid artery supplies most of the cerebral hemispheres, whereas the vertebral artery supplies the brain stem and the cerebellum.

The **internal carotid artery** (arteria carotis interna) enters the cranial cavity through a canal at the base of the skull (the carotid canal in the temporal bone) and then divides into three branches (Fig. 8.3):

1. The **ophthalmic artery** passes to the orbit through the optic canal and thus does not supply the brain itself (although, strictly speaking, the retina is part of the CNS). The **central retinal artery** (a. centralis retinae) enters the eye through the optic nerve and supplies the retina.
2. The **anterior cerebral artery** runs forward over the optic nerve and along the medial aspect of the hemisphere (Figs. 8.3 and 8.5). It supplies most of the cortex on the medial aspect of the hemisphere (except the most posterior parts and the inferior aspect of the temporal lobe). Its branches reach only a short distance onto the convexity of the hemispheres, supplying the leg representations of the motor and somatosensory areas. Shortly after its origin, the anterior cerebral artery gives off thin branches that penetrate the base of the hemisphere to supply anterior portions of the basal ganglia and hypothalamus.
3. The **middle cerebral artery** is the largest branch. It curves laterally into the lateral cerebral fissure and follows this backward and upward (Figs. 8.3, 8.4, and 8.5). On its way, numerous branches are given off that supply most of the cerebral cortex on the convexity of the hemispheres, notably the motor and somatosensory cortical areas, except for their most medial parts (with neuronal groups concerned with the motor and sensory functions of the legs; see Figs. 22.5 and 22.9), which are supplied by the anterior cerebral artery. The deep parts of the cerebrum, such as most of the basal ganglia and the internal capsule, receive their own branches from the middle cerebral artery, **lenticulostriate arteries** (Fig. 8.4), and by a separate branch of the internal carotid artery, the **anterior choroid artery** (Fig. 8.3).

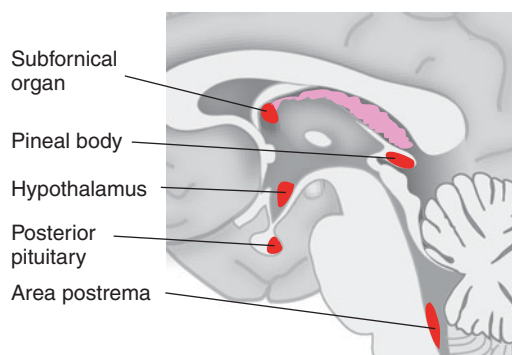


FIGURE 8.2 Regions of the brain devoid of blood–brain barrier.

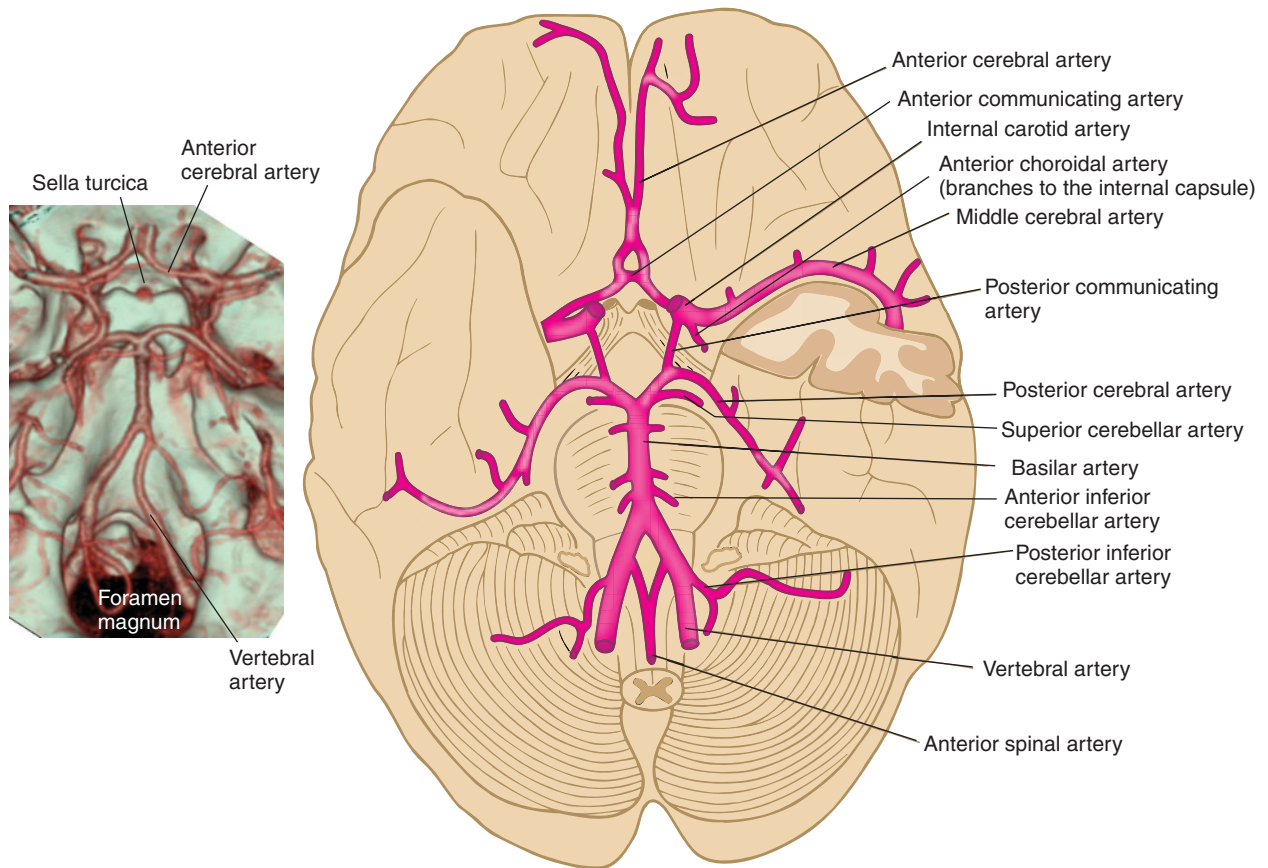


FIGURE 8.3 *The main arteries of the brain.* Anastomoses occur between the branches of the internal carotid artery and the vertebral artery and between the two anterior cerebral arteries, forming an

arterial circuit at the base of the brain. **Left:** Arteries at the base of the brain as reconstructed from MRIs. (Courtesy of Dr. S.J. Bakke, Rikshospitalet University Hospital, Oslo, Norway.)

The **vertebral artery** (Fig. 8.3) enters the posterior fossa after ascending through the transverse foramina of the cervical vertebrae (see Fig. 27.7). When passing from the atlas through the foramen magnum, the artery is highly convoluted, enabling it to follow the large movements in the upper cervical joints without being overstretched or compressed. Nevertheless, extreme movements—particularly in elderly people with sclerotic arteries—may temporarily occlude the vertebral artery. This may lead to loss of consciousness due to lack of blood supply to the brain stem. Backward bending of the head combined with rotation is particularly apt to compress the vertebral artery. Further, the vertebral arteries supply the following branches:

1. The **basilar artery** arises when the vertebral arteries of the two sides unite at the lower level of the pons to form (Fig. 8.3).
2. The vertebral arteries and the basilar artery send off numerous branches to supply the medulla

oblongata, the pons, the mesencephalon, and the cerebellum. The largest branches are the posterior inferior cerebellar, anterior inferior cerebellar, and the superior cerebellar arteries (Fig. 8.3):

- a. The **posterior inferior cerebellar artery** supplies the lateral part of the medulla and inferior parts of the cerebellar hemispheres. It usually originates from the extracranial portion of the vertebral artery; more seldom from the basilar artery.
- b. The **anterior inferior cerebellar artery** supplies lateral parts of the pons and parts of the cerebellum (see Chapter 27, under “Lateral Pontine Infarction”).
- c. The **superior cerebellar artery** supplies the dorsal aspect of the cerebellum and parts of the pons and the mesencephalon.
- d. The **labyrinthine artery** is a small branch from the anterior inferior cerebellar artery (or, less frequently, from the basilar artery). It supplies the labyrinth in the inner ear (its occlusion will therefore cause deafness on the side of occlusion and vertigo).

e. Several thin branches—called **perforator arteries**—penetrate the brain stem from the basilar artery along its course. Some arise from the dorsal aspect of the basilar artery and supply midline structures in the pons and mesencephalon. Others course laterally for a variable distance before they penetrate the brain stem.

3. The **posterior cerebral arteries** are the two end branches of the basilar artery and arise at the upper end of the pons. The posterior cerebral artery curves posteriorly around the mesencephalon and continue at the medial side of the hemisphere to the occipital lobe (Figs. 8.3 and 8.5). The posterior cerebral artery supplies large parts of the occipital lobe, notably the visual cortex, and the inferior aspect of the temporal lobe containing higher visual association areas (e.g., necessary for the recognition of objects). Perforating branches leave the first part of the posterior cerebral artery to supply the cerebral peduncle (the crus cerebri with descending tracts as well as nuclei dorsal to the crus, such as the oculomotor nucleus). Some branches supply the dorsal parts of the mesencephalon and posterior parts of the diencephalon.

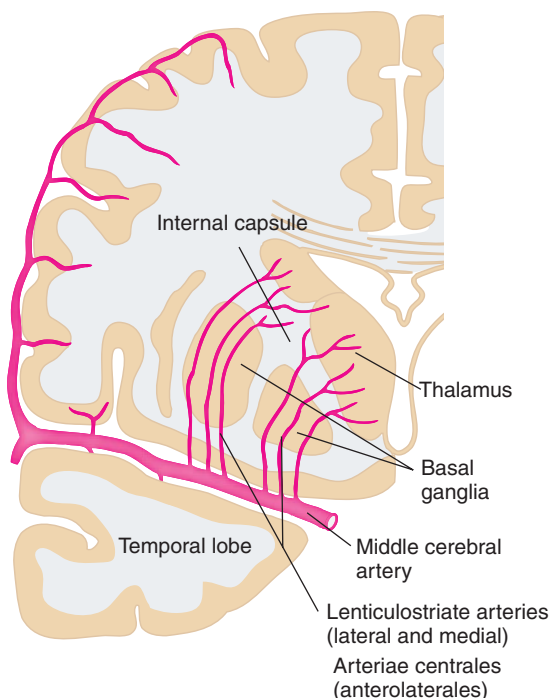


FIGURE 8.4 Course of the middle cerebral artery. In the depth of the lateral fissure, this artery gives off branches to the internal capsule and to the basal ganglia.

Communications between the Major Brain Arteries

At the base of the brain, there is a connection on each side between the middle and the posterior cerebral arteries, the **posterior communicating artery** (arteria communicans posterior) (Fig. 8.3). This means that, as long as the communicating artery is open, if one of the two main arterial trunks (the internal carotid or the vertebral artery) is narrowed or even occluded, the other may compensate for the loss. There is also a corresponding communicating artery between the two anterior cerebral arteries, called the **anterior communicating artery** (Fig. 8.3). In this manner a circle of anastomosing arteries is formed at the base of the skull, the **circle of Willis** (circulus arteriosus cerebri), which may be of great clinical significance. It explains, for example, how some people may have a totally occluded internal carotid artery on one side without any neurological signs. In addition, the communicating arteries can most likely become wider when the occlusion of the internal carotid artery develops slowly over the years.

The Spinal Cord Receives Arteries at Many Levels

In general, the arteries of the cord are arranged with one artery running in the midline anteriorly, the **anterior spinal artery**, and one on each side running along the rows of posterior roots, the **posterior spinal arteries** (Fig. 8.6). All three arteries begin cranially as branches of the vertebral arteries but receive contributions from the small arteries that enter the vertebral canal along with the spinal nerves.

Individual Variations in Size and Distribution of Cerebral Arteries Have Clinical Significance

The **symptoms** produced by occlusion of one arterial branch are variable because of individual differences in the size and exact distribution of the various arterial branches. Thus, if one artery is small, another supplying a neighboring territory is usually large. The anterior and posterior inferior cerebellar arteries are examples of this phenomenon: sometimes one of them supply the total territory normally supplied by both. Even the two vertebral arteries often differ markedly in size in one person, explaining why symptoms caused by occlusion of the artery may vary from minimal to life threatening.

Less than 50% of the population appears to have a “typical” **circle of Willis**, that is, the communicating arteries are symmetrical and with a certain cross-sectional diameter. In some persons the anterior communicating artery is very thin or missing; in others this may concern the posterior communicating artery on one or both sides. Therefore, the ability of the carotid artery of one side to compensate for the loss of the corresponding artery of the other side would be expected to vary

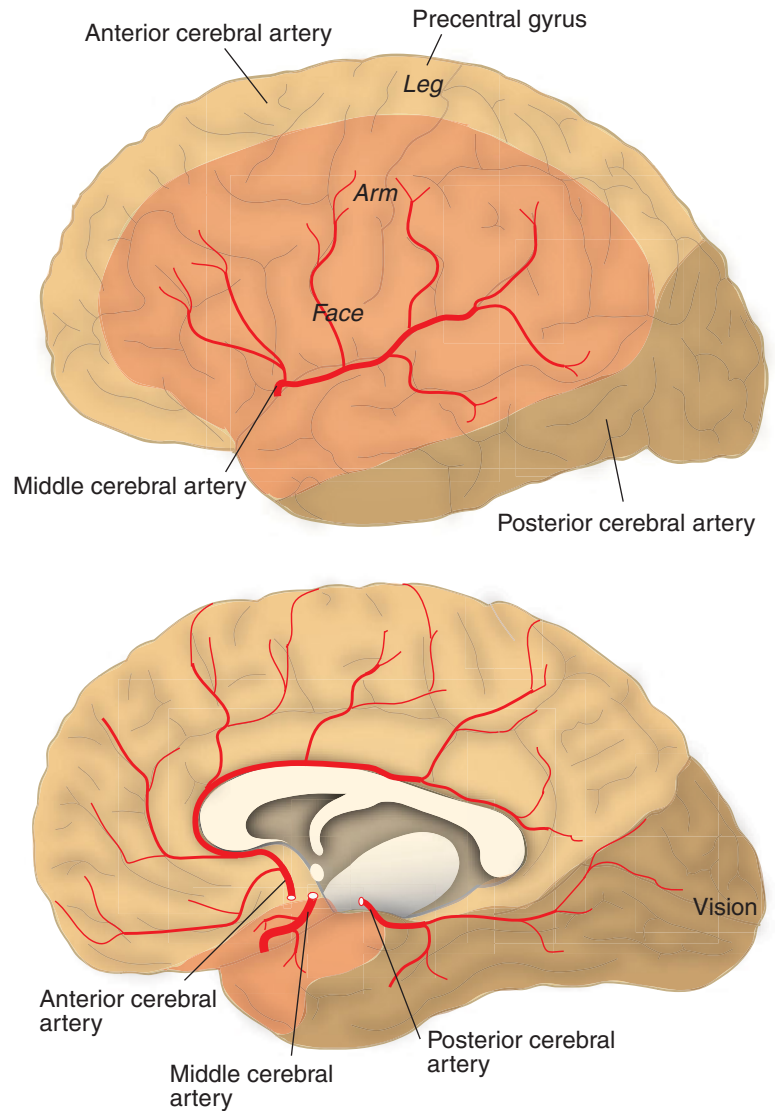


FIGURE 8.5 The parts of the brain supplied with blood from the main arterial branches.

considerably among individuals. Similar individual variations exist with regard to the ability of the anterior (carotid) and posterior (vertebral) circulations to compensate each other.

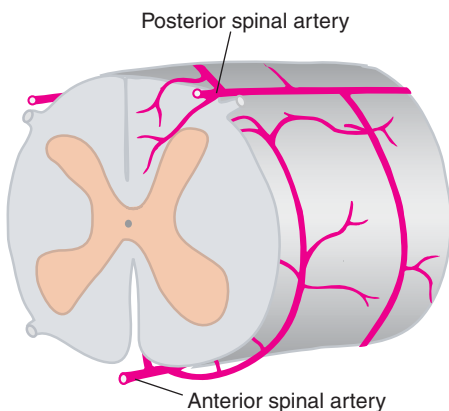


FIGURE 8.6 The arterial supply of the spinal cord.

VENOUS SYSTEM

The Venous Blood Is Collected in Sinuses

The cerebral veins can be divided into deep and superficial types. The latter partly accompany the arteries on the surface of the brain. All of the veins empty into large **venous sinuses** that are formed by folds of the dura (Figs. 8.7–8.9).

The **superficial veins** at the dorsal parts of the hemispheres run upward and medially and empty into the large **superior sagittal sinus** in the upper margin of the falx cerebri. Where the falx cerebri meets the tentorium

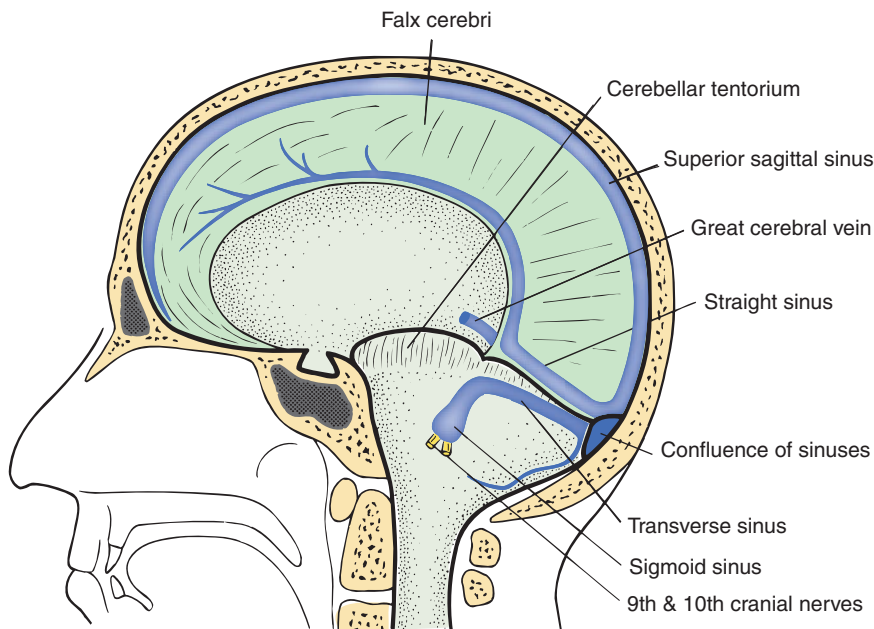


FIGURE 8.7 *Folds of the dura and the venous sinuses.* The folds minimize the movements of the brain and contain venous sinuses (in blue).

cerebelli, the superior sagittal sinus divides into two parts, the **transverse sinuses**, so named because they follow a transverse course laterally along where the tentorium is attached to the occipital bone. The **sigmoid sinus**—forming the direct continuation of the transverse sinus—empties into the **internal jugular vein** at the jugular foramen. The internal jugular vein leaves the skull and continues downward into the neck.

Most of the blood in the **deep cerebral veins** collects into the **great cerebral vein of Galen** (vena cerebri magna; Fig. 8.7). This comes out from the inferior side of the posterior end of the corpus callosum and empties into the **straight sinus** (sinus rectus) in the midline of the tentorium. The straight sinus drains into the superior sagittal sinus at the **confluence region**, from which the transverse sinus originates. Unlike the arteries, the cerebral veins have numerous **anastomoses**. At some locations, there are also **emissary veins** that form connections between intracranial and extracranial veins (see Fig. 6.34).

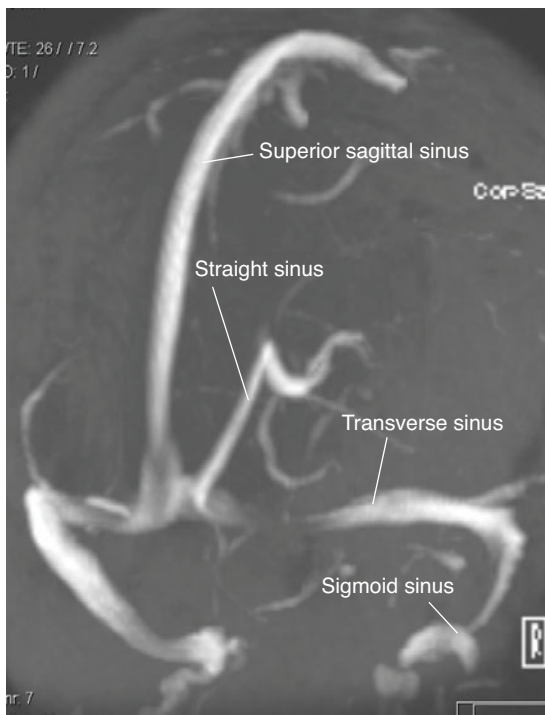


FIGURE 8.8 *Venous sinuses as visualized via MRI.* (Courtesy of Dr. S.J. Bakke, Rikshospitalet University Hospital, Oslo, Norway.)

Bridging Veins

Veins draining into the sinuses from the subarachnoid space—for example, along the superior sagittal sinus—are called **bridging veins** (see Fig. 6.34). They constitute some of the few attachments that exist between the convexity of the hemispheres and the skull. Head injuries that lead to sudden displacement of the brain inside the skull may cause bridging veins to tear, with venous bleeding as a result. This results in a chronic **subdural hematoma**, which expands very slowly (due to an osmotic effect of the decomposing blood of the hematoma). The name implies that the blood collects between the dura and the arachnoid, in the subdural space. The blood remains localized, in contrast to what happens when the bleeding occurs in the subarachnoid space. The symptoms arising from a subdural hematoma may be due to pressure on the underlying parts of the brain (e.g., paresis if the hematoma overlies the motor cortex), or they may be due to increased intracranial

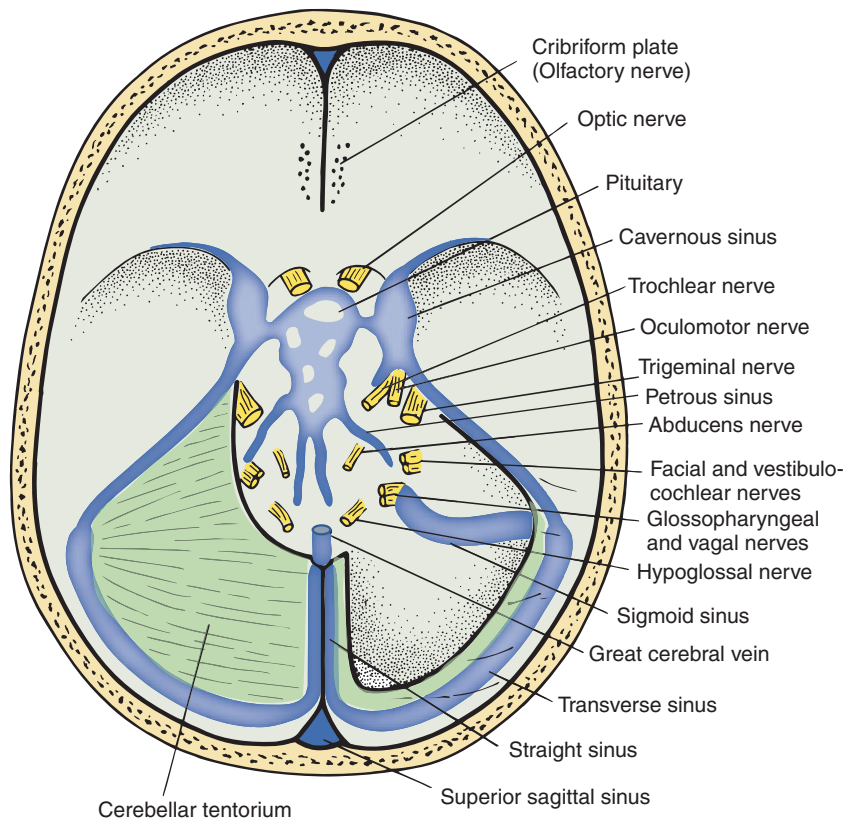


FIGURE 8.9 *The large venous sinuses at the base of the skull. On the right side, the tentorium cerebelli is partly removed. The cranial nerves and their sites of exit from the skull are also shown.*

pressure with more unspecific symptoms, such as headache and confusion.

The Cavernous Sinus Has Relations to the Pituitary and Cranial Nerves

One of the sinuses at the base of the skull is the **cavernous sinus** lateral to the pituitary gland (Fig. 8.9). Blood from the pituitary (among other structures) flows into the cavernous sinus and continues posteriorly in the **inferior petrosal sinus** to the internal jugular vein. Some cranial nerves, particularly those moving the eyeball, pass through the cavernous sinus on their way to the orbit. Infections can occasionally spread from the face or the nasal sinuses to the cavernous sinus (by way of

venous anastomoses). In such cases a dramatic and life-threatening condition ensues. Symptoms such as headache, confusion, and loss of consciousness are caused by elevated intracranial pressure, while double vision (diplopia), and pain and loss of sensibility in the face are due to affection of the third, fourth, fifth, and sixth cranial nerves.

Venous Drainage of the Spinal Cord

The **venous blood** is collected in a venous plexus at the surface of the cord. This plexus empties into another, larger plexus at the surface of the dura, the **epidural plexus** (see Fig. 6.6). From there the blood is emptied into veins outside the vertebral canal.

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II

DEVELOPMENT, AGING, AND PLASTICITY

THE fully developed nervous system consists of incredibly complex networks of connections. There are billions of neurons in the human brain, and each one is probably, on average, connected with several thousand others. The number of possible combinations of synaptic contacts is therefore astronomic. Indeed, a section of nervous tissue stained to reveal all neuronal processes might appear as a chaotic jungle. Nevertheless, we have ample evidence that this is very far from the case: order exists everywhere, and the mutual connections between neuronal groups are far from random. This leads to a number of questions, such as: How do the thousands of individual cell groups find their highly specific positions in the brain? How do the

complicated and precisely organized networks arise during development of the individual? What roles do genetic and environmental factors play in the final structure and performance of the brain? Such questions are dealt with in **Chapter 9**. In **Chapter 10**, we discuss the changes taking place in the aging brain and their consequences for function. A common theme in Chapters 9 and 10 is nervous system plasticity—that is, its ability to adapt structurally and functionally to altered demands. In **Chapter 11**, we discuss plastic changes in the nervous system as the basis for recovery of function after damage to the central nervous system. We will argue that a common theme in all rehabilitative efforts is to facilitate learning.

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9 | Prenatal and Postnatal Development

OVERVIEW

Development of the nervous system starts by growth of parts of the cellular layer covering the dorsal aspect of the embryo. Diffusible substances produced by the underlying tissue induce growth of the cells. This phase of **induction** is an example of how the cells and tissues interact during development. Then follows a more prolonged period during which the primitive neurons and glial precursors divide mitotically. This is called the phase of **cell proliferation**. The number of divisions is genetically determined. To get from the site of proliferation to their final position in the nervous system, the primitive neurons have to **migrate**. After reaching their final location, they start to aggregate into groups, representing the future nuclei.

For most neurons, differentiation starts with the outgrowth of processes after migration. By **differentiation**, we mean the development of structural and functional features that makes cells different from each other. Very early in differentiation the dendritic tree starts to develop. Fully grown, this pattern is often highly characteristic. While the neuronal processes are developing, the neuronal membranes undergo changes, and the neurotransmitters and other specific synaptic properties of the neurons begin to be expressed.

The next step in the development of the nervous system is the **outgrowth of axons** and establishment of **specific synaptic connections**. During this phase, the axons must find their way and recognize the neurons with which they are going to establish synaptic contact. Several factors have a role in this process. Neurons can be equipped with specific recognition molecules, the target region can produce a diffusible substance that “attracts” the axons, and the time of birth of a neuron may determine the synaptic contacts to be established.

Many more neurons are formed during the proliferation phase than the number present in the mature nervous system. In fact, many neurons are eliminated at the time their axons are establishing synaptic contacts. This is called the phase of **selective cell death** and has been observed in many parts of the nervous system. Usually, about half of the neurons are eliminated during this phase, although there are large regional differences.

In some systems, the **elimination of axon collaterals** occurs at a later stage than selective cell death. This phenomenon, which perhaps may be compared with the pruning of a tree so that only the most viable

branches remain, has been most intensively studied with regard to the innervation of skeletal muscles.

Neuronal activity is necessary for the proper development of axonal ramifications and for the formation of synapses. During this process, certain connections survive and possibly expand, whereas others are eliminated. Impulse activity alone is not enough for proper development in the central nervous system (CNS), however. The pattern of impulses has to be one that occurs in an intact, normal animal—which probably means that it must contain meaningful information—to be of consequence for synapse formation. We use the term **use-dependent synaptic plasticity** to characterize developmental processes that require proper use of the relevant neural networks.

PRENATAL DEVELOPMENT

Assembly of neurons into nuclei and establishment of their interconnections are largely **genetically** determined. The outer shape of the brain, the position of thousands of neuronal groups (nuclei), and the main neural connections arise prenatally: our genes contain the “recipe” for the building of the nervous system in considerable detail. During embryonic development, precursor cells are triggered to differentiate—first to neurons and glial cells, and then to numerous specialized subtypes of each. This occurs by modifications of the cell’s chromatin, for example, by methylation of specific sites of the DNA, thereby modifying the expression of certain genes (without altering the DNA itself). Largely, our genome possesses the instructions that determine final size of neurons, the shape of their dendritic trees, and the types of neurotransmitters expressed, as evidenced by growth of neurons in culture. Nevertheless, the full functional development of the brain depends critically on proper use of the neurons and their interconnections (discussed later in this chapter). Consequently, the normal development and performance of the nervous system depend on interactions of genetic and environmental factors.

The Central Nervous System Develops as a Long Tube

A few days after fertilization of the egg, differentiation of the cells of the embryo to the main kinds of tissue has started. From being a round lump of cells, after about a

week, the embryo resembles an elongated disc. In the second week the disc is covered on the upper or dorsal side and along the edges by primitive epithelium, the ectoderm, whereas the under or ventral side is covered by an epithelium—the endoderm—which later forms the intestinal tract and its glands. Between these two epithelial layers develops the mesoderm that later differentiates into the musculoskeletal system. In the third week (after 18 days) the development of the nervous system starts, with formation of a thickening, the neural plate, in the prospective cranial end of the disc (Figs. 9.1 and 9.2). The thickening is due to growth of the ectodermal cells so that they form a tall, simple, columnar neuroepithelium. Diffusible substances¹—called morphogens¹—from the underlying mesodermal cells induce formation of the neural plate (see also “Neuromeres and Hox Genes” later). The differentiation of ectodermal cells to neuroepithelium proceeds in a cranial-to-caudal direction. A longitudinal infolding of the ectoderm occurs by the end of the third week (Fig. 9.1). This neural groove is subsequently closed in the fourth week; the closing starts in the middle part (Fig. 9.2) and forms the neural tube, which is soon covered by the ectoderm dorsally (Figs. 9.1 and 9.2). The wall of the tube is formed by primitive neuroepithelial cells, which proliferate enormously and develop into neurons and glial cells. The lateral edge of the neural plate forms a distinct cell group, the neural crest, which later forms a longitudinal column on each side of the neural tube (Fig. 9.1). The neural crest produces the neurons of the peripheral nervous system, among them spinal ganglion cells and autonomic ganglion cells. The neural crest also produces Schwann cells and satellite cells (a kind of glia) in the ganglia.

The Neural Crest Produces More Than Neurons and Glia

Experiments with labeling of neural crest cells before they migrate (Fig. 9.1) show that some differentiate into non-nervous structures. These include the smooth muscles of the eye (intrinsic eye muscles) and, most likely, the **pia** and the **arachnoid** (the dura mater is believed to originate from the mesoderm). Migrated neural crest cells also form the dermis and subcutis of the **face** and the cartilaginous skeleton of the **visceral arches**. Finally, neural crest cells produce endocrine

1 **Morphogens** are substances that spread by diffusion from a localized source and govern the embryological development and patterning of organs and body parts. Their effects depend on their concentrations—often so that high and low concentrations exert opposite effects. Among several morphogens involved in patterning of the human nervous system, the protein **sonic hedgehog** plays an important role at very early stages. For example, sonic hedgehog is expressed by the notochord when the dorsal–ventral differentiation of the neural tube begins. It also acts at later stages to guide axonal growth, attracting outgrowing axons in low concentrations, and repelling them in high concentrations (as shown for retinal ganglion cell axons growing from the eye toward the brain).

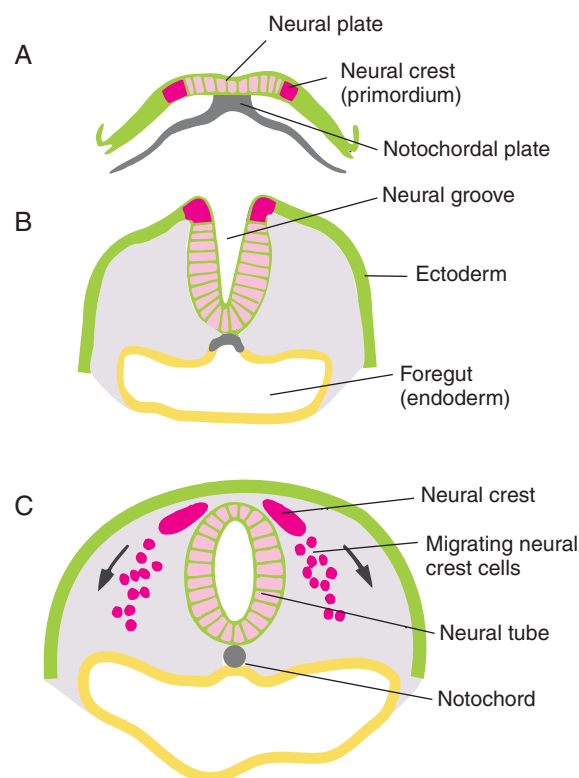


FIGURE 9.1 Formation of the neural tube and the neural crest. Schematic cross sections through embryos at different stages of development. **A:** The formation of the neural tube is induced by substances diffusing from the underlying mesoderm (notochordal plate). Approximately 17 days. **B:** The neural tube is formed by growth and folding of the neural plate. Approximately 21 days. **C:** The neural crest gives origin to spinal ganglion cells and autonomic ganglion cells (and some other cell types). The neural crest cells migrate into the body to form ganglia. Approximately 24 days.

cells of the **adrenal medulla**, **melanocytes** of the skin, and parts of the **septum** that divides the pulmonal artery and the aorta.

Early Development of the Cranial End of the Neural Tube

Whereas the caudal end of the neural tube—which develops into the spinal cord—retains its simple tubular form, the expanded cranial end undergoes marked changes. This occurs because different parts grow at different rates, and because the tube bends, forming **flexures** (Fig. 9.3). In the fourth week three swellings or **primary vesicles** take shape (Figs. 9.3 and 9.4). The cavities inside the primary vesicles are continuous and develop into the ventricular system of the brain (see Fig. 7.3). The most cranial vesicle is called the **prosencephalon** (forebrain), the middle one the **mesencephalon** (the midbrain), and the caudal-most one the **rhombencephalon** (hindbrain). A ventrally directed bend—the **cervical flexure**—arises at the junction between the rhombencephalon and the spinal cord (Fig. 9.3). Later, a **mesencephalic flexure**

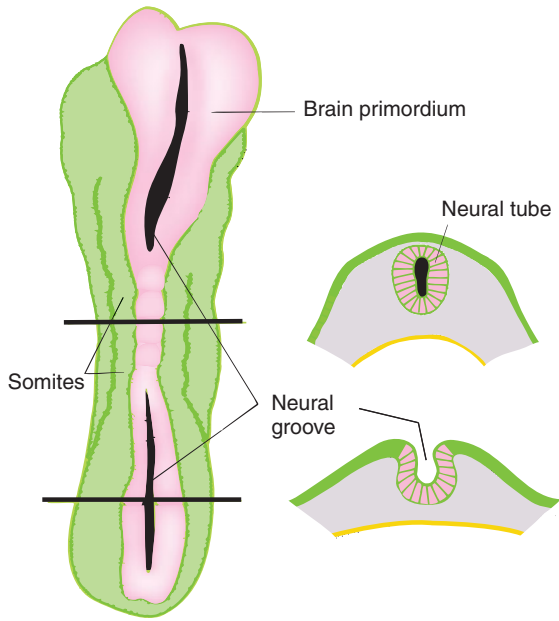


FIGURE 9.2 The neural plate and closure of the neural groove. Drawing of a 22-day-old human embryo, approximately 1 mm long. The central nervous system is shown in pink. (Based on Hamilton, Boyd, and Mossman 1972.)

arises between the rhombencephalon and the mesencephalon. A dorsally directed bend—the **pontine flexure**—later divides the rhombencephalon into two parts (Fig. 9.5).

Early in the fourth week, the ventral aspect of the prospective brain exhibits shallow, transverse grooves. These are external signs of segmentation of the cranial

end of the neural tube, and each segment is called a **neuromere**. The segmentation is most obvious in the rhombencephalon, and we use the term **rhombomere** in this region. Although their external signs disappear by the sixth week, the neuromeres are important because they represent the first segregation of neurons that later differentiate into the various nuclei of the brain stem. Thus, several cranial nerves and their nuclei are first laid down according to a segmental pattern, like the spinal nerves, although later development makes the cranial nerve pattern less regular.

The mesencephalon changes little during further development, in contrast to the two other primary vesicles. The **prosencephalon** develops into the diencephalon and the cerebral hemispheres, whereas the **rhombencephalon** differentiates into the medulla oblongata, the pons, and the cerebellum (Figs. 9.4 and 9.5). The rostral end of the prosencephalon produces two more vesicles (one on each side), called the **telencephalon** (Figs. 9.4 and 9.5), which later forms the cerebral cortex and basal ganglia. In addition, the **olfactory bulbs** (see Fig. 3.13) arise as evaginations from the ventral aspect of the telencephalon. The remaining caudal part of the prosencephalon forms the diencephalon, which includes the thalamus and hypothalamus. At an early stage (Fig. 9.3), cuplike evaginations—the **eye vesicles**—are formed from the prosencephalon (the part later to become the diencephalon). The eye vesicles develop into the retina and the optic nerve. The rhombencephalon develops two parts: the **myelencephalon** forming the medulla oblongata, and the **metencephalon** forming the pons and most of the cerebellum (the cerebellum also develops from the mesencephalon, as discussed next).

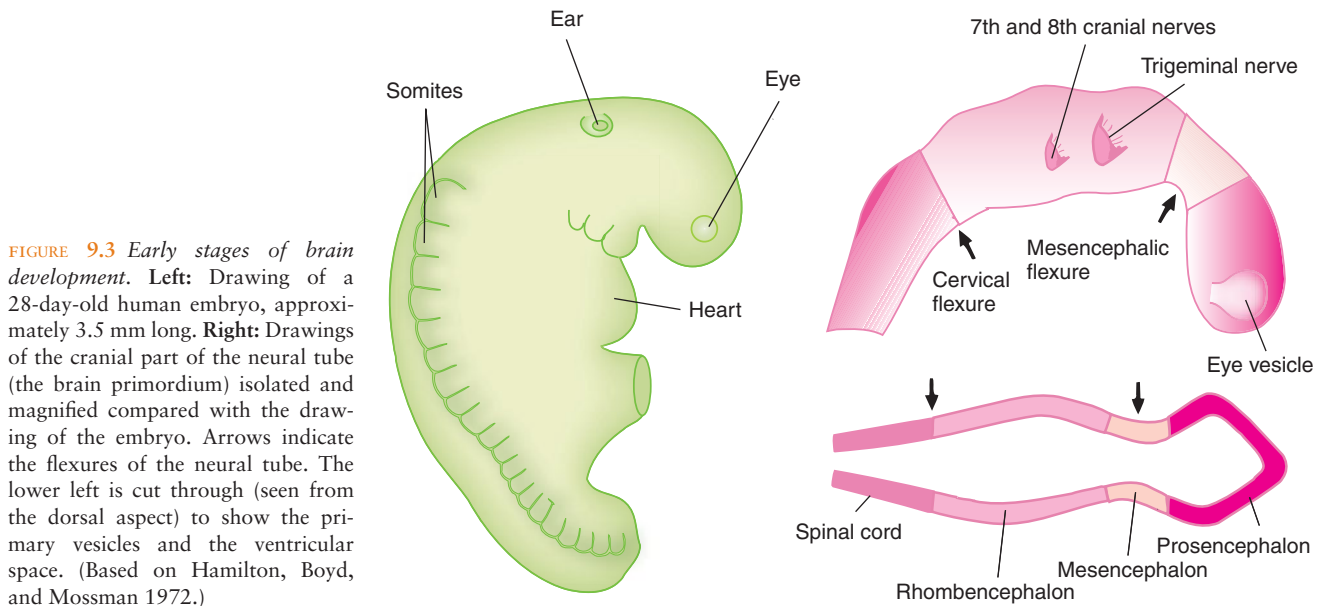


FIGURE 9.3 Early stages of brain development. **Left:** Drawing of a 28-day-old human embryo, approximately 3.5 mm long. **Right:** Drawings of the cranial part of the neural tube (the brain primordium) isolated and magnified compared with the drawing of the embryo. Arrows indicate the flexures of the neural tube. The lower left is cut through (seen from the dorsal aspect) to show the primary vesicles and the ventricular space. (Based on Hamilton, Boyd, and Mossman 1972.)

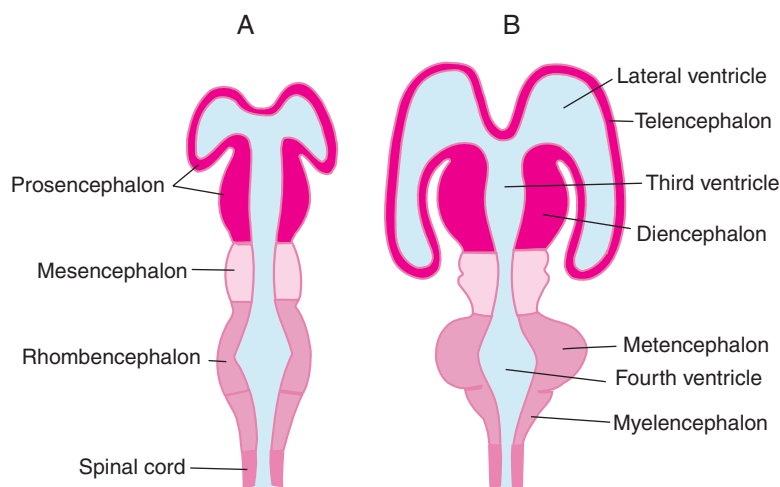


FIGURE 9.4 Different divisions of the brain primordium. Schematic of the cranial part of the neural tube (straightened out and cut open horizontally). **A:** Approximately the same stage as in Fig. 4.3. **B:** The same stage as in Fig. 4.13A. Note the development of the telencephalon (the hemispheres) that gradually covers the diencephalon.

Early Phases in the Development of the Neuroepithelium

Initially, the wall of the neural tube consists of only one layer of cylindrical neuroepithelial cells (Figs. 9.1 and 9.2), bounded externally by the **external limiting membrane** (covered by the pia) and internally toward the cavity by the **internal limiting membrane**. These are basal membranes built of extracellular material, which always develop with surface epithelia. Intense **proliferation** of neuroepithelial cells soon leads to several layers of nuclei. The epithelium does not become truly stratified, however, because all cells retain a thin process reaching the internal limiting membrane (pseudostratified epithelium; Fig. 9.6). The outermost cells move toward the cavity of the neural tube (the future ventricles

and central canal). The innermost layer, the **ventricular zone**, borders the cavity. In the ventricular zone, the cells divide mitotically (Figs. 9.6 and 9.13A). The future neurons, the **neuroblasts**, afterward migrate outward to form the **mantle zone** that later becomes the gray matter. A layer without neurons, the **marginal zone**, forms external to the mantle zone and becomes the white matter. The neurons of the mantle zone send axons into the marginal zone. These axons loop back to the mantle zone, however, to synapse on neurons there (these are the first association connections to arise). Other axons leave the neural tube as motor fibers growing peripherally to contact muscle cells and glands (Fig. 9.7). After neuroblast production ends, various types of glial cells are produced by mitosis of neuroepithelial cells that remain in the ventricular zone. The last to be produced

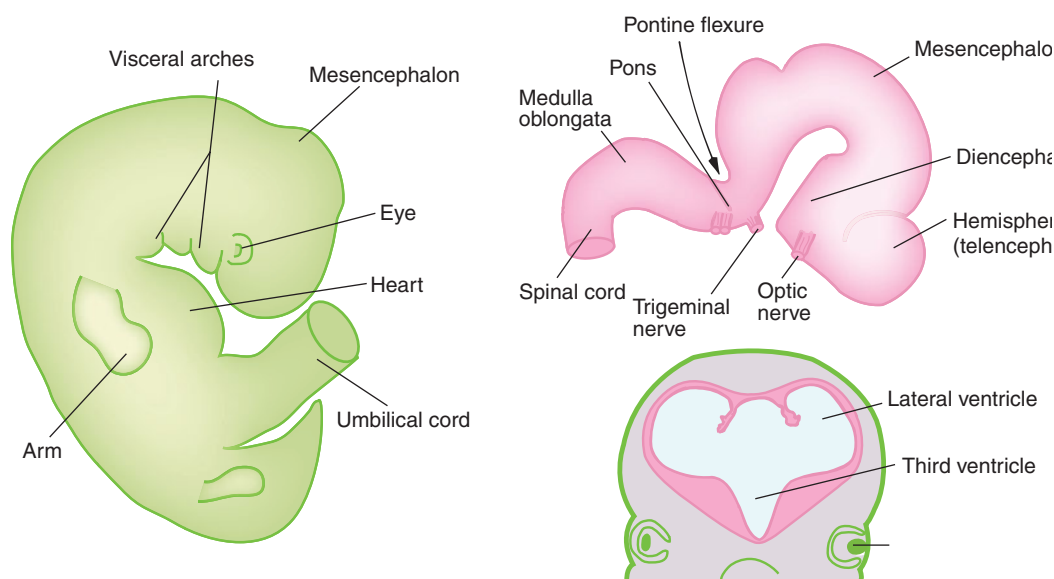


FIGURE 9.5 Early development of the brain. **Left:** Drawing of a 36-day-old human embryo, approximately 11 mm long. **Right:** The

cranial part of the neural tube at the same stage. (Based on Hamilton, Boyd, and Mossman 1972.)

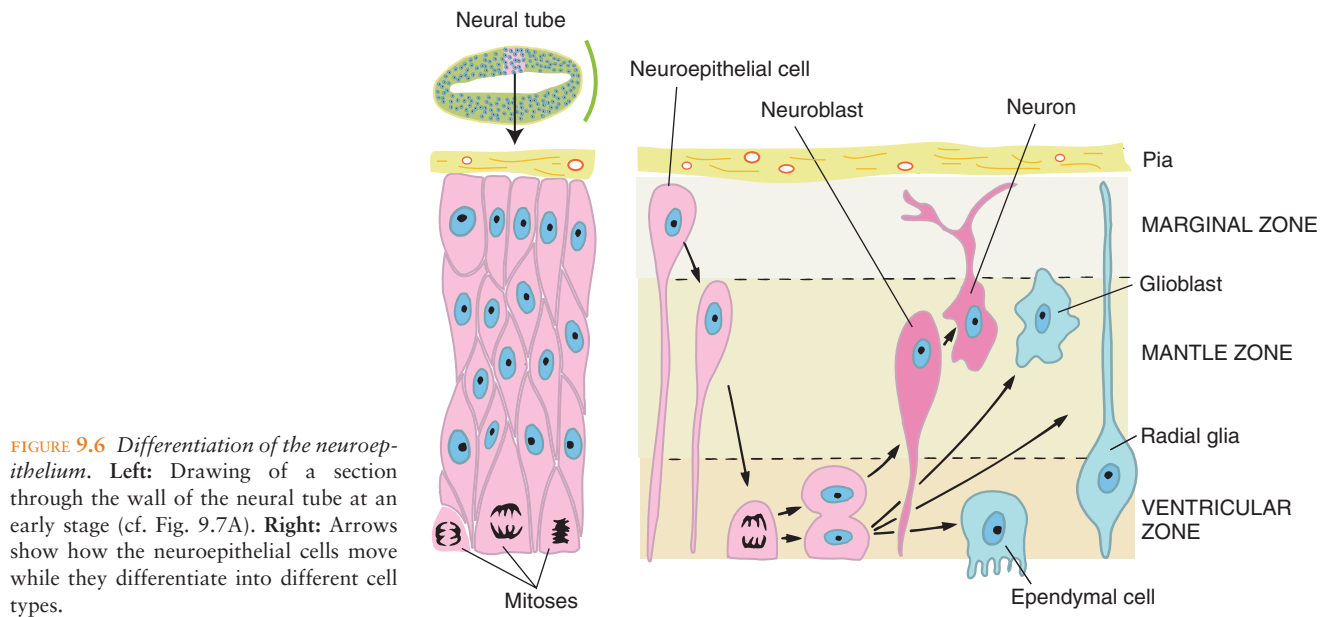


FIGURE 9.6 *Differentiation of the neuroepithelium.* Left: Drawing of a section through the wall of the neural tube at an early stage (cf. Fig. 9.7A). Right: Arrows show how the neuroepithelial cells move while they differentiate into different cell types.

are the **ependymal cells**. These retain their internal position and cover the ventricular face of the neural tube (Fig. 9.6). The cavity is filled with cerebrospinal fluid produced by tufts invaginated from the wall into the cavity (see Fig. 7.6). These tufts—the future **choroid plexus**—are covered by ependymal cells.

The simple layering of the neural tube, with the mantle zone (gray matter) inside and the marginal zone (white matter) outside, is retained with minor changes in the spinal cord. In the cranial part of the neural tube

(the future brain), however, major alterations in the mutual positions of gray and white matter occur. In the developing cerebellar and cerebral cortices, for example, neurons migrate from the mantle zone through the marginal zone and form a layered sheet of gray matter externally, just under the pia.

Neuromeres

We mentioned that the rostral part of the neural tube shows transient, external signs of segmentation—each segment constituting a **neuromere**. Although this phenomenon was observed in the nineteenth century, modern cell biological methods were necessary for closer study of their role in the development of the nervous system. Neuromeres are most convincingly shown in the rhombencephalon, where they are called **rhombomeres**. It is assumed that the mesencephalon consists of two neuromeres, whereas the prosencephalon probably consists of six. The great interest in neuromeres and other external signs of segmentation arose because they provide information about the mechanisms that control the early development from the undifferentiated neural.

Each rhombomere represents a unit of neurons that do not mix with neurons of other rhombomeres during subsequent development. This neuronal specification takes place just when the rhombomere boundaries arise. The rhombomeres arise when adjoining groups of neurons begin to express different surface markers. The neurons of one rhombomere, among other things, have a specified future peripheral target (the neurons of the neural crest in the head region are also specified with regard to their peripheral target before they start to migrate peripherally). The motor trigeminal nucleus,

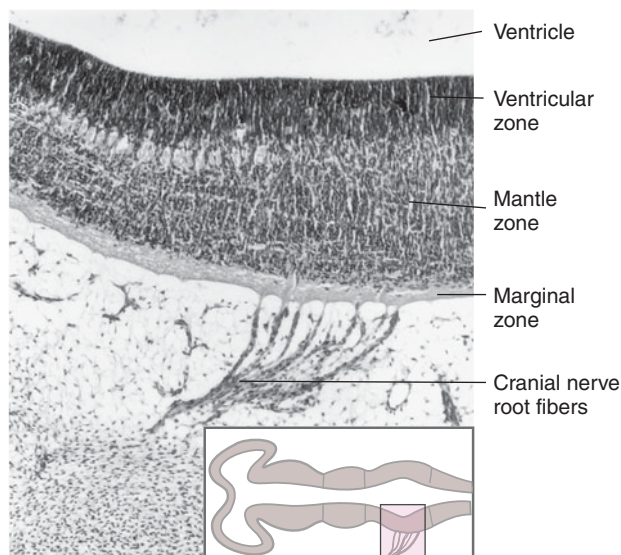


FIGURE 9.7 *Outgrowth of cranial nerves from the rhombencephalon.* Photomicrograph of a horizontal section through a chicken embryo at about the same stage as in Figure 4.3. Thin bundles of axons leave the marginal zone and penetrate the connective tissue that surrounds the brain primordium.

for example, is formed in rhombomeres r2 and r3, whereas the motor facial nucleus develops in r4 and r5. The motor and sensory cranial nerve axons already have a specified target when they start growing out from the neural tube (Fig. 9.7). Thus, when a rhombomere is transplanted to another place in the chicken embryonic brain, it develops as if it were still in its original place and not corresponding to its new location. Such specification must be caused by the switching on of certain genes—which start to express themselves by production of mRNA—while probably other genes are switched off.

Hox Genes

Many so-called *Hox* genes have been identified that are expressed in a pattern corresponding to neuromeric boundaries in vertebrates. Most of these genes code for proteins that act as **transcription factors**. These bind to DNA of other genes and regulate their transcription. Typically, transcription factors are expressed temporarily during specific phases of development. (*Hox* genes not only control regional development of the nervous system but also act in pattern formation in other parts of the body.) When a certain *Hox* gene is switched on, a cascade of changes in the expression of other genes is initiated, producing signal molecules that give the neuronal groups their identity—for example, regarding location and connections. A crucial question is, of course, what controls the regionalized expression of *Hox* genes giving rise to the rhombomeres? One important factor is **retinoic acid (vitamin A)**, which normally occurs with an anteroposterior (rostrocaudal) concentration gradient in the embryo (highest concentration posteriorly or caudally). The retinoic acid seems to stem from the mesoderm adjacent to the neural tube. In low concentrations, retinoic acid acts on *Hox* genes that specify anterior parts of the neural tube, while in

high concentrations it induces differentiation of posterior parts. This explains why adequate dietary levels of vitamin A are necessary for the normal development of the CNS, but also why too much also may cause malformations (as may occur in women treated for acne with retinoic acid in early pregnancy).

Neuromeric borders, identified with genetic markers, are more reliable indicators of future borders between anatomically and functionally different areas than are borders between brain vesicles. For example, the cerebellum, traditionally regarded as arising only from the metencephalon, is also formed by neurons in the adjoining part of the mesencephalon. The rostral border of neurons forming the cerebellum coincides with a border for the expression of the *engrailed 2* gene.

Further Development of the Spinal Cord and the Brain Stem

In the fourth week, the proliferation of neuroblasts in the mantle zone produces a large ventral thickening and a smaller dorsal one on each side of the neural tube. These thickenings are called the **basal plates** and the **alar plates**, respectively (Figs. 9.8, 9.9, and 9.10). A shallow furrow, the **sulcus limitans**, marks the border between them. This remains visible in the lower part of the brain stem in the adult (see Fig. 3.17), while it disappears early in the spinal cord. The basal plate contains neuroblasts that later become motor neurons, whereas many alar plate neuroblasts become sensory neurons. This corresponds to the functional division between the ventral and dorsal horns of the cord. In the adult brain stem, the sulcus limitans marks the border between the motor and the sensory cranial nerve nuclei (Fig. 9.11). In the open part of the rhombencephalon—later to become parts of the medulla and the pons—the motor nuclei lie medially and the sensory nuclei laterally (see Figs. 27.2 and 27.3). This is caused by lateral

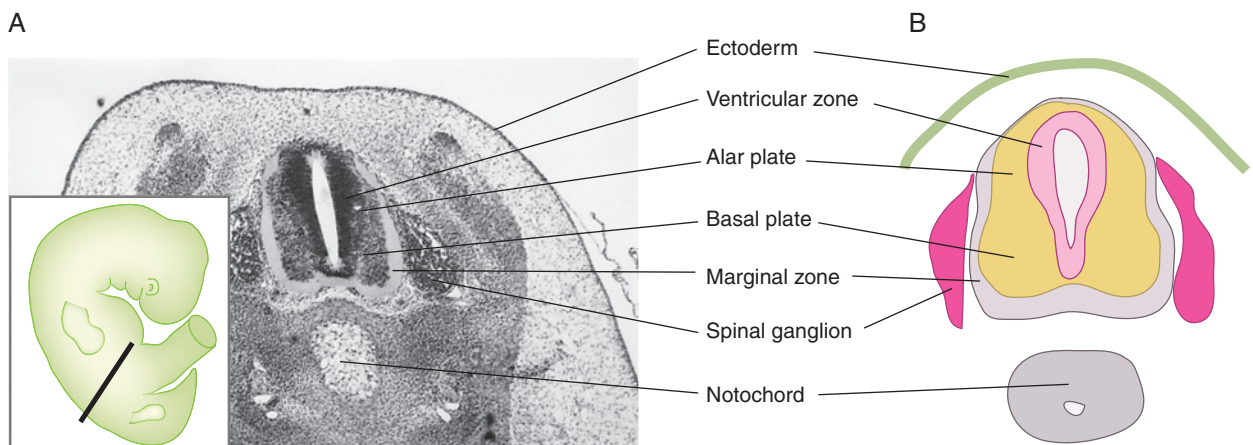


FIGURE 9.8 Early development of the spinal cord. **A:** Photomicrograph of a cross section of a chicken embryo (corresponding approximately to

a 6-week-old human embryo; cf. Fig. 9.5). **B:** Drawing based on a photomicrograph of the human spinal cord at about 7 weeks' gestation.

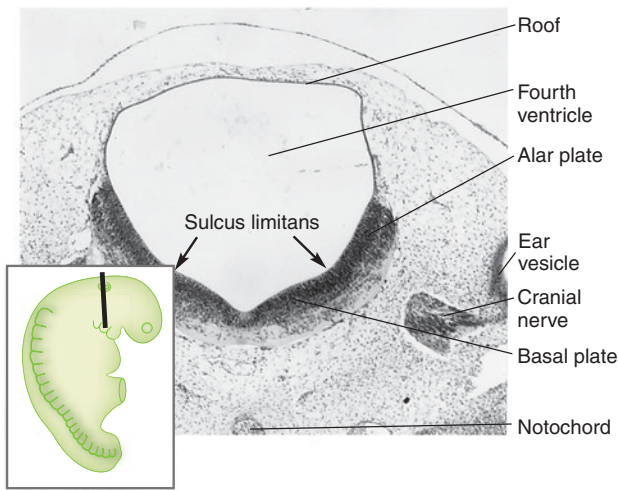


FIGURE 9.9 *The rhombencephalon.* Early stage of development, approximately as in Fig. 9.3. Photomicrograph of a cross section of a chicken embryo. **Inset:** A human embryo at a corresponding stage of development. Note the thin rhombencephalic roof attached to the edge of the alar plate.

bending of the alar plates—away from each other—so that the roof of the rhombencephalon becomes only a thin membrane (Figs. 9.9 and 9.10). At a later stage, the **cerebellum** develops from the margins of the alar plates (the **rhombic lip**).

At an early stage, several neuronal groups in the brain stem **migrate** from their “birthplace” in the alar or basal plates. In the pons, neuroblasts move from the rhombic lip in a ventral direction and form the pontine nuclei (see Fig. 3.18). Similarly, in the medulla the inferior olive (another nucleus projecting to the cerebellum; see Fig. 3.17) is formed by neuroblasts moving ventrally from the rhombic lip. Another example is the **motor**

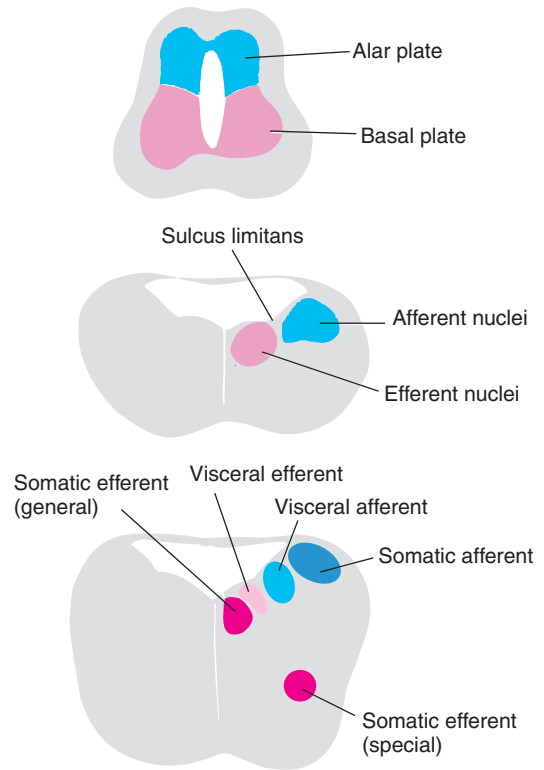


FIGURE 9.10 *Development of the cranial nerve nuclei.* Schematic. Three stages are shown to explain the mutual positions of the nuclei. After their formation, the special somatic efferent nuclei move ventrolaterally. See also Figs. 17.2 and 17.3.

cranial nerve nuclei that innervate visceral (branchial) arch muscles (see Fig. 9.10). These neurons move in a ventral direction during early development after they have started to send out axons. The course of the root fibers in the brain stem therefore shows the path followed by the migrating neurons (see Fig. 27.11).

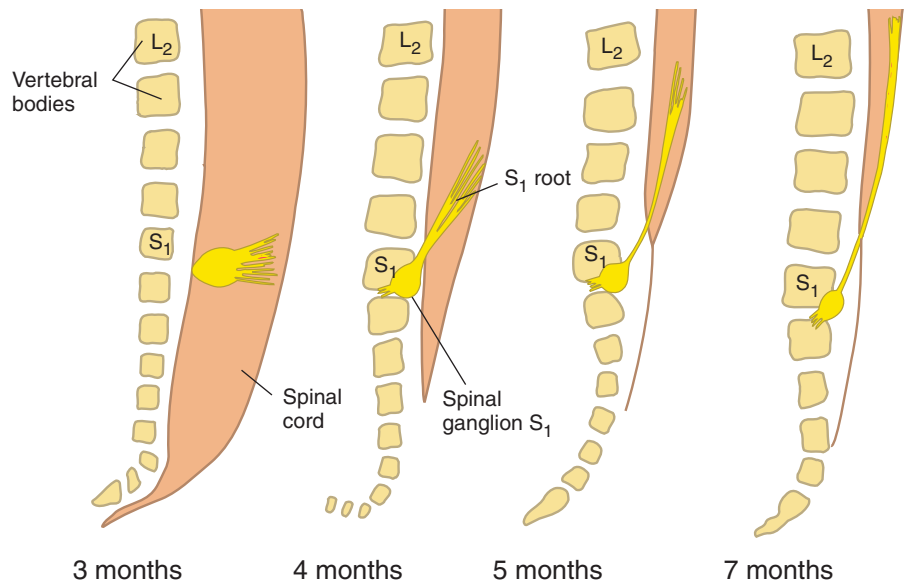


FIGURE 9.11 *Relationship between the lengths of the spinal cord and the vertebral column.* (Based on Hamilton, Boyd, and Mossman 1972.)

Until the eighth to ninth week, the spinal cord extends the full length of the spinal canal of the embryo. After that, however, the vertebral column and the coverings of the cord grow more rapidly than the spinal cord itself, producing a gradually increasing length difference (Fig. 9.11). This explains the oblique course of the **spinal nerves** in the vertebral canal before they leave through the intervertebral foramina. The lower end of the cord is at the level of the third lumbar vertebra at birth, in contrast to the first lumbar vertebra in adults. The dural sac, containing spinal nerve roots (the cauda equina), continues down to the second sacral vertebra (see Fig. 3.3).

Cranial Nerves and Visceral Arches

Cranial nerves 5, 7, 9, and 10 (trigeminal, facial, glossopharyngeal, and vagal; see Figs. 6.15 and 6.19) innervate structures developed from visceral arches (or branchial arches). In fish, the two upper visceral arches are parts of the viscerocranium surrounding the oral cavity. They are not equipped with gills, in contrast to the lower (3–6) arches that are true respiratory organs. In mammals, none of the visceral arches has respiratory functions. Parts of them are used for new tasks, whereas other parts have regressed. There are no signs of gill development at any stage of human embryogenesis. The term “branchial arch” (from Greek *brankhion*, gill) is therefore not quite appropriate.

A visceral arch consists of a skeletal part (cartilage, later replaced by bone in some arches), muscles, skin, mucous membranes, and a nerve of its own. As mentioned, neural crest cells give rise to the skeletal part. In human embryos, some visceral arches appear as ventral bulges in the head and cervical region (Figs. 9.3 and 9.5). The **first visceral arch**—producing the upper and lower jaw with attached masticatory muscles, along with the hammer and the anvil of the middle ear—is innervated by the **fifth cranial nerve** (the trigeminal). The second visceral arch—forming, among other things, the upper part of the hyoid bone, the stirrup of the middle ear, and the facial muscles—is innervated by the **seventh cranial nerve** (the facial). The **third visceral arch**—forming most of the hyoid bone and the posterior part of the tongue—is innervated by the **ninth cranial nerve** (the glossopharyngeal). The **fourth, fifth, and sixth visceral arches** form the cartilaginous skeleton of the larynx and the muscles of the larynx and the pharynx. These structures are innervated for the most part by the **tenth cranial nerve** (the vagus).

Cranial nerves 3, 4, 6, and 12 (the oculomotor, trochlear, abducens, and hypoglossal; see Fig. 3.14) innervate structures that most likely develop from segmentally arranged **somites** (somites are paired cubical masses giving rise to muscles, the axial skeleton, and the dermis of the skin). These nerves are homologous to spinal

ventral roots. As for the **eleventh cranial nerve** (the accessory), it is not settled whether the two muscles it innervates (the sternocleidomastoid and trapezius) develop from somites or from visceral arches. The latter hypothesis is supported by the fact that the accessory nerve root fibers, coming from the upper cervical segments, exit the cord more dorsally than the spinal ventral roots (corresponding to the level where visceral arch nerves leave the brain stem; see Fig. 27.8).

Further Development of the Diencephalon

The diencephalon represents the caudal part of the original prosencephalic vesicle (Figs. 9.4 and 9.5). The lateral wall in this part becomes thicker at an early stage and develops into the **thalamus** (Fig. 9.12A). The floor plate forms the **hypothalamus** and the **posterior pituitary** (see Figs. 6.22 and 6.24). The latter arise as an evagination of the floor plate. The furrow (hypothalamic sulcus; see Fig. 6.23) marking the border between the thalamus and the hypothalamus might be a continuation of the sulcus limitans (Fig. 9.10). It is not settled, however, whether the arrangement with basal and alar plates continues rostrally into the diencephalon. The thin roof plate of the diencephalon forms by invagination the **choroid plexus** of the third ventricle (see Fig. 7.5). Further, the roof plate produces the **pineal body** by evagination (see Fig. 6.23). The **eye vesicles** occur at an earlier stage (before the further differentiation of the prosencephalon; Fig. 9.3) but retain connection with the diencephalon by the future optic nerve (Fig. 9.5).

Further Development of the Telencephalon

In the fifth week, development of the **cerebral hemispheres** starts with the appearance of one vesicle on each side of the prosencephalon (Figs. 9.4 and 9.5). These are called the **telencephalic (cerebral) vesicles**. Their cavities form the lateral ventricles, which initially have wide openings to the third ventricle (interventricular foramen, Fig. 9.12A). The mantle zone of the basal part of the telencephalic vesicle thickens rapidly to form the corpus striatum of the **basal ganglia** (Fig. 9.12A). The thinner overlying part, called the **pallium**, becomes the cerebral cortex. The pallium grows dorsally, rostrally, and caudally in relation to the diencephalon (Fig. 9.4). The caudal and ventral parts of the hemispheres later fuse with the diencephalon (Fig. 9.12B).

The basal ganglia primordium is later divided into two parts—the lateral and medial **corpus striatum**—by descending axons from the cerebral cortex. These descending axons form the **internal capsule** (Fig. 9.12B). As development proceeds, the caudate nucleus and the thalamus come to lie medial to the internal capsule whereas the lentiform nucleus (the putamen and the globus pallidus) lies laterally (see Fig. 13.2).

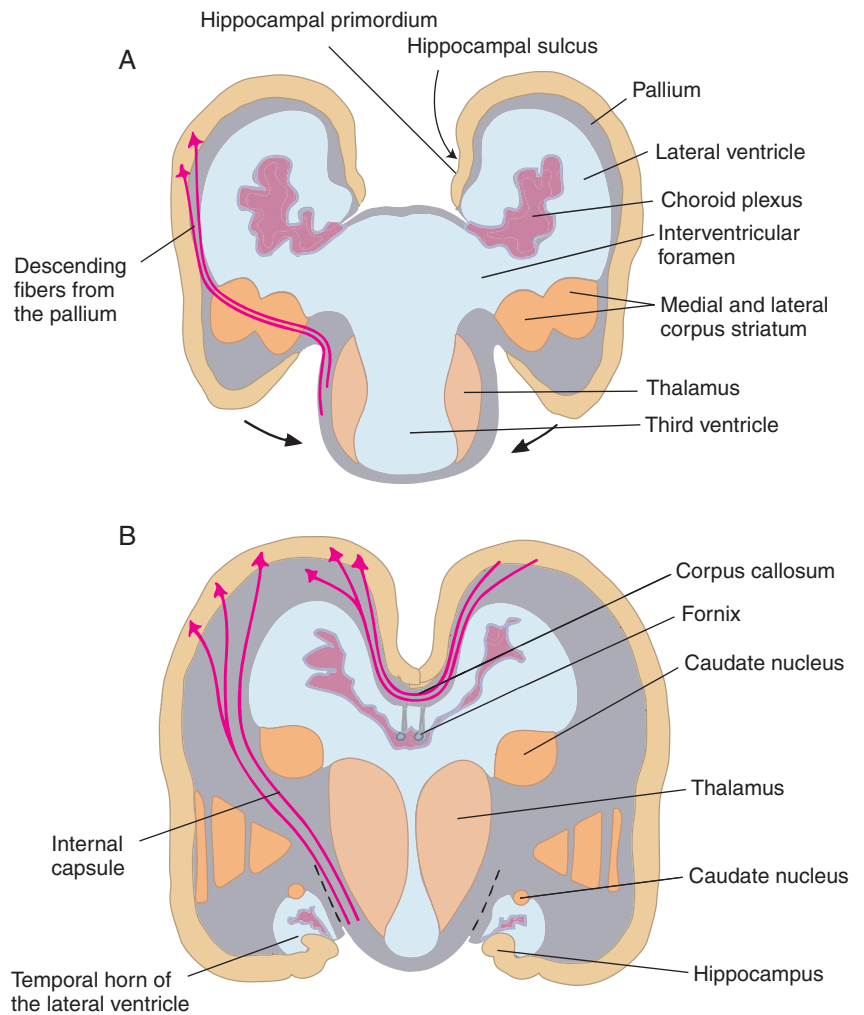


FIGURE 9.12 *Development of the telencephalon.*

A: Early stage, in which the telencephalon is still separated from the diencephalon. Descending fibers have started their growth from the pallium (primordium of the cerebral cortex) to the brain stem and the spinal cord. **B:** The telencephalon is now attached to the diencephalon laterally, and the medial and lateral striatum have been separated by the internal capsule. The hippocampus has changed its position, due to the growth of the hemispheres (cf. Fig. 3.36). (Based on Hamilton, Boyd, and Mossman 1972.)

In the medial wall of the pallium, just above the attachment of the choroid plexus, a thickening arises that bulges into the lateral ventricle (Fig. 9.13A). This is the beginning of the **hippocampus**, which is partly separated from the rest of the pallium by the hippocampal sulcus. As the hemispheres grow, their shape changes so that the temporal lobes come to lie ventrally. This produces the characteristic curved shape of the lateral ventricles and the structures in their wall, such as the hippocampus and the caudate nucleus (see Figs. 7.4 and 31.2). The hippocampus thus moves from the position in Fig. 4.12A to that in Fig. 4.12B.

The **choroid plexus** of the lateral ventricles is formed by invagination of the thin part of the pallium (together with the pia) close to the dorsal aspect of the diencephalon (Fig. 9.12B; see also Fig. 7.6).

Development of the Cerebral Cortex

At first, the telencephalic vesicle consists of only one layer of neuroepithelial cells. These proliferate, however, producing rapid growth of the prospective cerebral

hemisphere. At a later stage, we can differentiate a ventricular zone, mantle zone, and marginal zone, just as in other parts of the neural tube (Fig. 9.13A). At the beginning of the **eighth week**, neuroblasts from the mantle zone **migrate** into the marginal zone and start establishing the **cortical plate** (Fig. 9.13B), which in due course will develop into the mature cerebral cortex by waves of cells migrating toward the cortical surface. The peak of migratory activity probably occurs between the third and fifth months, while migration ends in the third trimester. By the end of the seventh month the cortex has developed six layers, as in the mature cortex (see Figs. 33.1 and 33.2). Synapses begin to occur in the fourth month (earliest in the prospective somatosensory cortex).

The deepest cortical layers are established first; thus, neurons destined for superficial layers have to pass through the deep layers. Neuroepithelial cells in the ventricular zone that have ceased dividing are termed **postmitotic**. The exact time a neuron becomes postmitotic—its **birth date**—appears to decide which cortical layer it will join. Radially oriented glial cells, **radial glia** (Fig. 9.6), with processes extending from the ependyma to the pia,

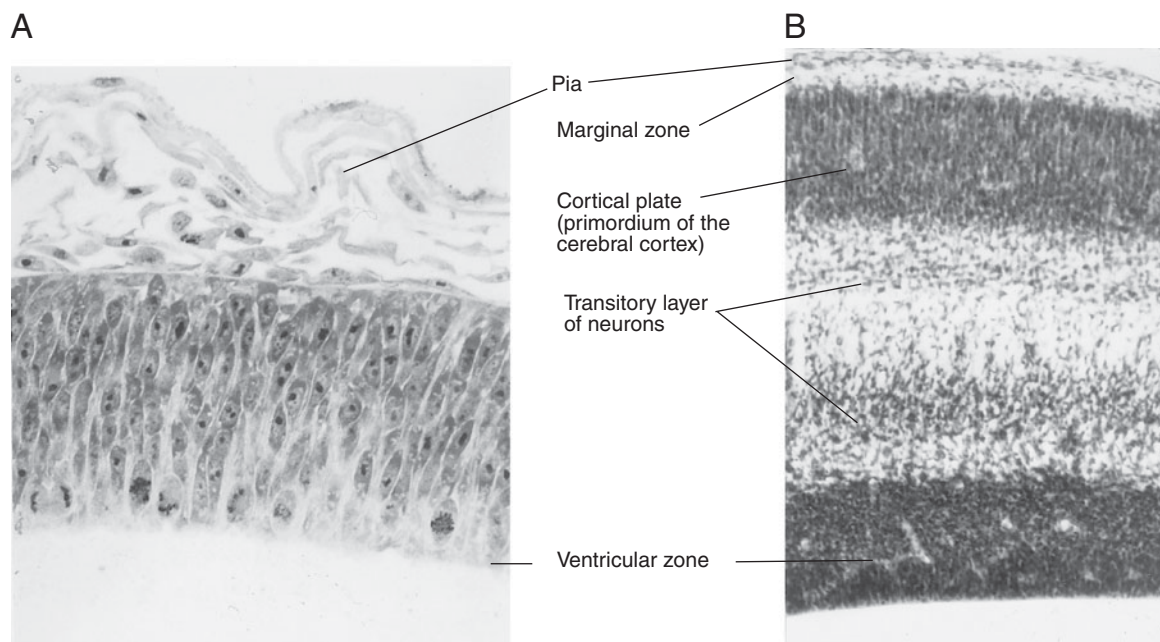


FIGURE 9.13 *Differentiation of the prosencephalic neuroepithelium to the cerebral cortex.* **A:** Photomicrograph of a cross section through the neural tube (prosencephalic part) at an early stage (chicken embryo, corresponding to early fourth week gestation in humans). Mitotic activity occurs in the ventricular zone. **B:** Photomicrograph

of a section through the telencephalon (rat) at a much later stage than in **A**, corresponding approximately to the fourth to fifth month in human development. The ventricular zone is densely packed with neuroblasts that will soon migrate toward the cortical plate. In further development, the cortical plate develops into the adult six-layered cortex.

guide the migration of postmitotic neurons toward the cortex. The phenotype of postmitotic neurons, for example, whether they will develop into interneurons or projection neurons, appears to be specified at the time they start migrating toward their final destination.

Shortly after the establishment of the cortical plate, thalamocortical fibers start to invade the telencephalic wall, although they must “wait” several weeks below the cortical plate before they find their final destination in the developing cortex. The earliest afferent fibers to arrive, however, are **monoaminergic** (at about 7 weeks).

While probably all progenitors of projection neurons arise in the ventricular zone and migrate radially to their final destination, many prospective cortical **interneurons** arise from subcortical sites in the ventral forebrain. They then migrate tangentially along the cortical plate for varying distances before they change course and migrate perpendicularly into the cortex.²

Specification of Cortical Cytoarchitectonic Areas

The adult cerebral cortex consists of many areas that differ structurally and functionally (see Fig. 33.4).

² All cortical interneurons are **GABAergic** but fall into different groups structurally and with regard to whether they co-express the neuropeptide **somatostatin** (SST), **parvalbumin** (PVA), or **calretinin** (CR). The three main groups of GABAergic interneurons seem to be specified at the time they migrate from the so-called ventricular (ganglionic) eminences.

The differentiation of the cortical plate into distinct areas (see Fig. 33.4)—a process called **arealization**—depends on both genetic factors intrinsic to the developing cortical progenitor cells and extrinsic influences. In the beginning, local patterning centers in the periphery of the ventricular zone produces gradients of **morphogens** that define four main, overlapping domains in the cortical plate.³ The morphogens in turn initiate more discrete expressions of transcription factors in cortical progenitor cells. These transcription factors are involved in the further differentiation of the cortical plate into areas with sharp borders (as well as in other aspects of cortical development).

Among **extrinsic influences**, thalamocortical afferents appear to be of particular importance for the mature **cytoarchitectonic** characteristics of an area. For example, the characteristic cytoarchitectonic differences between the primary sensory areas (compare Figs. 33.2 and 33.4) depend on from which specific thalamic nucleus the areas receive their afferents. For example, transplanting a piece of visual cortex to the somatosensory cortex makes the transplanted tissue acquire the cytoarchitectonic features that are typical of the somatosensory cortex. Further, immature projection neurons transplanted from one area to another develop axonal

³ For example, the arealization of the frontal cortex seems to be initiated by two **fibroblast growth factors** (Fgf8 and Fgf17).

ramifications appropriate for the area to which they are transplanted. Thus, the local environment contributes significantly to the neuronal phenotype.

In addition to the genetically determined development described here, proper use of cortical areas is critical for the realization of their functional specialization. This involves **use-dependent** plastic processes, stabilizing useful synaptic connections, and eliminating those that prove to be superfluous or maladaptive.

Migration and Migration Disorders

We know some of the factors governing neuronal migration, such as recognition molecules, adhesion molecules, cytoskeletal components, and others. There is a complex interplay among them, and the different factors must be present in the proper concentration at the proper time and place. For example, the early presence of certain neurotransmitters influences neuronal motility by acting on ion channels that increase the intracellular Ca^{2+} concentration. The glycoprotein **reelin** (among other factors) governs the final positioning of migrating neurons in the cortex. Because reelin is produced by **Cajal-Retzius cells** in the marginal zone (Fig. 9.6), it would seem logical that a concentration gradient of reelin between the marginal and ventricular layers is critical for normal development. The mechanisms behind the effects of reelin appear to be more complex, however. Nevertheless, mice with a mutated gene for reelin exhibit characteristic malformations of layered structures; in the cerebral cortex, the late-arriving neurons do not migrate past those that arrived first to occupy the outer layers. Although the neurons survive and establish apparently normal connections, the brain does not function normally. Another mutation, shown in some humans, affects the migration *before* the neurons enter the cortical plate and is associated with a smooth cortex lacking the normal six-layered structure (**Miller-Dieker lissencephaly**). This is just one example of a large and varied group of malformations in humans—**migration disorders**—caused by delayed or deficient migration of postmitotic neurons.

Migration disorders affect primarily the cerebral cortex and the cerebellum. Many are inherited recessively, although ischemia, radiation, and other influences can cause migration disorders (as shown in animal experiments). In the cerebral cortex, migration disorders typically are associated with defective development of the gyri (Fig. 9.14), which may be lacking (**lissencephaly**), too small (**polymicrogyria**), or show other abnormalities. Usually, such a cortex is called **dysplastic**, and some use the term “dysplastic cortex” as synonymous with migration disorders.

Migration disorders cause a number of syndromes characterized by **cognitive** and behavioral defects (mental retardation is common). Typical of many such syndromes

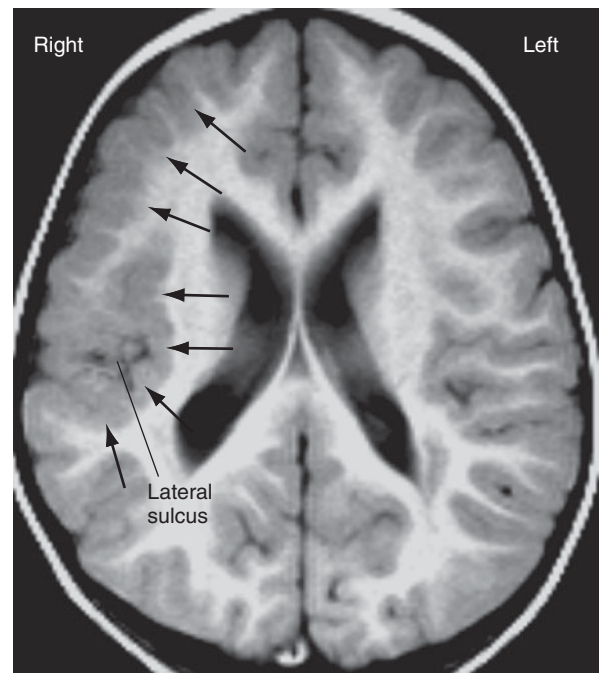


FIGURE 9.14 *Neuronal migration disorder.* Horizontal T1-weighted MRI. The right frontal cortex is dysplastic: it is thicker than normal with obliteration of normal sulci (polymicrogyria). Similar alterations occur around the lateral sulcus. (Courtesy of Dr. S.J. Bakke, Rikshospitalet University Hospital, Norway.)

is **epilepsy** (which is often difficult to treat pharmacologically). This is compatible with the fact that migration disorders also are associated with molecular abnormalities of cortical neurons. Thus, animal experiments suggest that epilepsy can arise in dysplastic cortex because of an imbalance between the expression of excitatory and inhibitory receptors (up-regulation of amino-methylisoxazole propionic acid [AMPA] receptors and down-regulation of GABA_A receptors).

Programmed Cell Death and Competition for Trophic Factors

As previously mentioned, many more neurons are formed during the proliferation phase than are found in the mature nervous system. Cell death usually coincides in time with the period during which the neurons establish synaptic contacts. For the most intensively studied neuronal groups, such as spinal motoneurons and retinal ganglion cells, the amount of elimination depends on **target size**. If the target (e.g., striated muscle cells) is experimentally expanded, more motoneurons survive, whereas reducing the target size increases elimination. This can be explained by the neurons needing a sufficient supply of a growth-promoting substance—a **trophic factor**—to survive. The target cells produce limited amounts of this trophic factor; that is, there is not enough to keep all neurons alive. The neurons

innervating the target **compete** for the trophic factor, and only the winners survive. Because a limited number of synapses can be formed on each neuron, the axons arriving first have an advantage. The trophic factor is taken up by the nerve terminals at the synaptic sites and transported retrogradely to the cell body. The elimination of surplus neurons appears to occur rapidly, perhaps during a few days in some systems. Overall, this kind of cell death is believed to ensure optimal numerical relationships between, for example, motoneurons and muscle cells. Cell death probably also serves to eliminate incorrect synaptic connections.

Programmed cell death is not uniform throughout the nervous system. In the primordial **cerebral cortex** probably about 90% of the neurons die, whereas in the **spinal cord** few interneurons die. About 50% of motoneurons die, and a similar rate holds for a type of interneuron in the **retina** (amacrine cells). We do not know the reasons for such differences, and many questions remain unanswered concerning programmed cell death. For example, the number of cells that die is not always related to the size of the target.

Cell death as a normal developmental process is termed **apoptosis** (in contrast to necrosis, which means cell death caused by abnormal, nonphysiological influences).⁴ What triggers this process? When certain nerve cells die without the presence of a growth factor, does it happen because the cell needs the factor for vital metabolic processes or because the factor inactivates genes that trigger apoptosis? There is much evidence in favor of the latter explanation—that is, the existence of a genetic death program in each cell. For example, mutation of certain genes prevents apoptosis that normally would have occurred at a certain stage of development.

Neurotrophins and Other Growth Factors

The Italian Nobel Prize winner Rita Levi-Montalcini discovered around 1950 a substance that stimulates axonal growth in the peripheral nervous system. This substance—termed **nerve growth factor** (NGF)—is a protein produced by the cells of the target organ. Antibodies against NGF inhibit the growth of axons from autonomic ganglion cells and sensory cells derived from the neural crest. In cell cultures, axons grow into an area containing NGF and retract from areas without it. NGF binds to specific membrane receptors. It is transported retrogradely to the cell body, enabling effects on gene expressions. After the discovery of NGF, several related neuronal growth factors have been identified. Together they form the **neurotrophin family**, consisting of NGF, **brain-derived neurotrophic factor**

(BDNF), and several others (neurotrophin NT-3, NT-4, and so forth). Apart from regulating many aspects of neuronal growth and differentiation during prenatal development, they are involved in synaptic plasticity and neuronal survival in the adult nervous system. The neurotrophins bind to specific **tyrosine-kinase receptors** (Trk) and to an unrelated receptor, **p75NTR** (p75 neurotrophin receptor).⁵ The two kinds of receptor may help to explain further why the effects of neurotrophins are highly complex and not fully understood. Among other differences, the effects mediated via Trk and p75NTR receptors tend to be opposite. Thus, nerve growth factor prevents programmed cell death by binding to TrkA receptors, whereas binding to p75NTR promotes it. Thus, the effects of a neurotrophin on a certain neuronal population depend on the local expression of neurotrophin receptors (e.g., the balance between expression of Trk and p75 receptors). The expression of the neurotrophins and their receptors appear to be dynamically regulated by a complex interaction among intrinsic and extrinsic factors.⁶ Thus, neurotrophin effects would differ among neuronal populations, and on the same population at different points of time.

Among other growth factors acting in the brain are the **fibroblast growth factors** (FGFs), which influence differentiation and survival of several kinds of neurons. Several other growth factors that were initially discovered in the peripheral tissues are also expressed in the brain.

Neurotrophins, Plasticity, and Disease

As mentioned, neurotrophins play a role not only during normal development but also in the adult brain. The finding that expression of neurotrophins can be use dependent has attracted special interest in their role in **synaptic plasticity**. Brain-derived neurotrophic factor (BDNF), for example, is involved in the induction of LTP. In the visual cortex, the expression of the receptor TrkB parallels the critical period in development, and the synthesis increases with increased use of the visual system. Beneficial effects of **physical activity** on cognitive functions and neurogenesis appear at least in part to be mediated by increased expression of BDNF (and perhaps NGF).

5 The Trk receptors constitute a family of three—TrkA, TrkB, and TrkC—each of which can be activated by one or more of the neurotrophins NGF, BDNF, NT-3, and NT-4. Overall, NGF seems to exert its main effects via TrkA receptors, while BDNF acts mainly via TrkB. The p75NTR belongs to the tumor necrosis factor receptor (TNFR) family (tumor necrosis factor is an inflammatory cytokine released from leukocytes, inducing apoptosis via its receptors).

6 For example, while the expression of p75NTR in normal adult brain is minimal, it increases after injury. At the same time, Trk receptor expression may be reduced. This would tend to push the effect of neurotrophin receptor activation from supporting cell survival to inducing cell death.

4 Apoptosis is characterized by the breakdown of DNA to smaller fragments and the subsequent dissolution of the cell.

Neurotrophins are also intensely investigated for their possible beneficial effects after **brain injury**. For example, administration of growth factors at the site of injured neurons might help them survive. In animal experiments, injection of NGF into the ventricular system prevented delayed cell death in the hippocampus after a period without blood supply. The role of neurotrophins in **neurodegenerative diseases** has also attracted interest. One theory proposes that the senile plaques in the brain of Alzheimer patients down-regulate FGF in their environment, thus causing neuronal death. In Parkinson's disease and Huntington's disease as well, the level of FGF is apparently reduced. The connection between such changes and the disease process is not clarified, however.

Prolonged **stress** can cause neuronal death in experimental animals. Decreased production of neurotrophins—BDNF in particular—may mediate the effect of stress. Alterations of growth factors also occur in patients with severe **depression**, and both antidepressant drugs and electroconvulsive therapy have been reported to increase expression of BDNF (among other effects).

Myelination

Myelination of axons starts in the **fourth month** of gestation and is largely completed 2 to 3 years after birth. Although many axons in the CNS remain unmyelinated, the process of myelination is clearly related to functional maturation of neuronal interconnections. Full functional capacity cannot be expected before myelination is completed. As for the individual neuron, myelination starts at the soma and proceeds distally. Different tracts are myelinated at different times. Overall, tracts concerned with basic tasks, necessary for life, are the first to be myelinated (such as sucking, swallowing, retraction from harmful stimuli, emptying of bowel and bladder, and so forth). Such connections are also phylogenetically the oldest.

In the **spinal cord**, myelination starts in the cervical region and proceeds in the caudal direction. First to be myelinated are the propriospinal fibers (interconnecting various spinal segments). Ventral root motor fibers are myelinated earlier than the dorsal root sensory fibers. Myelination of ascending spinal tracts starts in the sixth fetal month, and tracts descending from the brain stem follow (reticulospinal and vestibulospinal tracts). These tracts need to be functioning at birth. In contrast, the **pyramidal tract**, which controls the most precise voluntary movements, is fully myelinated only about 2 years after birth. Connections from the cerebral cortex to the **cerebellum** (see Fig. 24.7) are myelinated at the same time as the pyramidal tract, which seems logical because these connections are important for coordination of voluntary movements.

Myelination of the **cranial nerves** starts in the sixth fetal month, except for the optic nerve (which is a central tract and not a peripheral nerve). Myelination starts shortly before birth in the optic nerve.

In the **cerebral cortex**, myelination begins shortly before birth, first in motor and sensory areas. The association areas are mainly myelinated during the first 4 months after birth, although myelination continues after that period. The last regions to become fully myelinated are the association areas in the frontal lobe (prefrontal cortex).

Although the myelination largely occurs according to the pattern described in the preceding text, longitudinal magnetic resonance imaging (MRI) studies indicate that the relative proportion of white matter in the brain increases (although slightly) until adult age. For example, the cross-section of the corpus callosum increases from 5 to 18 years of age. Presumably, after the age of 2 to 3 years the increase in white matter is caused by increased myelin-sheath thickness of already myelinated axons.

Malformations of the Nervous System

We discussed migration disorders as a cause of malformations, especially of the cerebral cortex. These malformations are special since the development of the cerebral cortex is so protracted, ending late in the prenatal period. Here we consider other kinds of malformations that occur before the gross shape of the CNS has been established.

Malformations of the nervous system, as in other organs of the body, may be caused by genes or environmental factors (or by both). External agents, such as viruses and drugs, are most likely to cause serious malformations or maldevelopment if they act in the period of maximal differentiation—that is, from the **third to eighth week** after fertilization (most organs are formed during the embryonic period from the fourth to eighth week). Harmful influences before this stage usually lead to early death of the embryo.

Among the most common malformations is **defective closure** of the neural tube, which may be caused by various genetic and environmental factors. Normally, the closure is completed by the end of the fourth week. Lack of closure may affect the whole length of the neural tube but is most often restricted to either the cranial or the caudal end. When the neural tube does not develop normally, neither do overlying structures such as the skull, parts of the vertebral column, and the skin. Their normal development depends on induction by diffusible substances (morphogens) from the nervous tissue. With defective closure of the cranial end, the brain does not develop, and the remainder of the neural plate degenerates. This condition is termed **anencephaly**. Such fetuses may sometimes live until birth but always

die shortly after. A defective closure of the spinal cord (most often in the lumbosacral part) is termed **spina bifida** because the vertebral arch and soft tissue dorsal to the cord do not develop normally. This condition may vary in severity. In the most serious cases the vertebral arches, muscles, and skin are absent and there is herniation of the coverings of the cord that contain degenerated nervous tissue (**meningomyelocele**). Less severe cases may involve herniation of the coverings but keep the spinal cord intact, whereas the least affected have only partial lack of the vertebral arch (**spina bifida occulta**). In the most serious cases, the cord does not develop normally, leading to pareses and sensory disturbances of the legs. When the nervous supply is deficient, the muscles do not develop to their normal size and strength.

Reduced drainage of the cerebrospinal fluid caused by, for example, abnormal narrowing or obliteration of the cerebral aqueduct (see Fig. 7.5) leads to **hydrocephalus**. If left untreated, this will cause death or seriously impair brain development (see also Chapter 7, under “Brain Edema, Herniation, and Hydrocephalus”).

MECHANISMS FOR ESTABLISHMENT OF SPECIFIC CONNECTIONS

The main morphologic features of the nervous system—such as its macroscopic form, the positions of major nuclei and their interconnections—arise before birth and shortly after. In a sense, this represents the **hard wiring** of the brain. In this section, we discuss mechanisms important for forming the brain’s “wiring diagrams.” The growth of axons is often surprisingly goal-directed, indicating the existence of guiding mechanisms. We will discuss some mechanisms that can aid axons in selecting their target. The number of known interacting players at the cellular and molecular levels is enormous, and many more are probably yet to be identified. It should therefore not come as a surprise that we cannot fully explain how the amazing connectional specificity of the mature nervous system arises.

Trial and Error

Although trial and error cannot explain the overall development of orderly connections in the nervous system, it nevertheless plays a role at the local level. Indeed, modern imaging methods permitting in vivo observation of growing neurons show that growth and retraction of neuronal processes are highly dynamic processes. Thus, at the same time as growth cones randomly explore their immediate environment and new spines bud from dendrites, errors are corrected continuously by retraction of unsuccessful growth cones and spines. The development of neuronal networks is therefore a

very complex interplay of simultaneous building up and tearing down, resembling the work of a sculptor who adds excess of clay to be able to carve out the fine details.

Distances Are Small in the Early Embryo

Knowing the complexity of the mature nervous system, one might think that most of the human genome must be devoted to specification of neuronal connections. This is not the case of course. We should bear in mind that the whole embryo is only a few millimeters to a couple of centimeters during the stages of most intense axonal outgrowths. Thus, the distances that axons have to grow to reach their targets are usually very short. Further, the topography of the nervous system at these early stages is also much simpler than later, as not all nuclear groups develop simultaneously. Our present knowledge suggests that combined actions of several mechanisms, each of them relatively simple, can explain how the specificity of the nervous system arises.

Time of Neuronal “Birth”

The genetically programmed time of neuronal birth can explain the development of specific connections in many cases (Fig. 9.15). If, at the time of axonal outgrowth from one neuronal group, only certain neurons are present in the direction of growth, the axons will hit their correct target without specific recognition molecules. In addition, programming of neurons for maximal synapse formation during a limited period ensures that synapses are established upon arrival of the proper axons. The time during which neurons readily produce synapses is usually limited. Axons encountering a particular neuronal group at a later stage cannot establish synapses, and therefore they either retract or grow past to other targets.

Trophic Factors

Timing of neuronal birth and maturation cannot explain all aspects of specificity, however. For example, what decides the direction of **axonal outgrowths**? For some neurons, such as pyramidal cells in the cerebral cortex, the initial growth direction is genetically determined. After that, however, signals in the environment of the axons determine the growth direction. Thus, cortical pyramidal cells are occasionally “inverted” with their apical dendrite pointing toward the white matter instead of toward the cortical surface. In such cases, the axons start growing toward the pial surface but soon reverse direction and grow toward the white matter, as do normal pyramidal cells (see Fig. 33.5). Such findings are best explained by the target organs producing growth-promoting substances—**trophic factors**—that

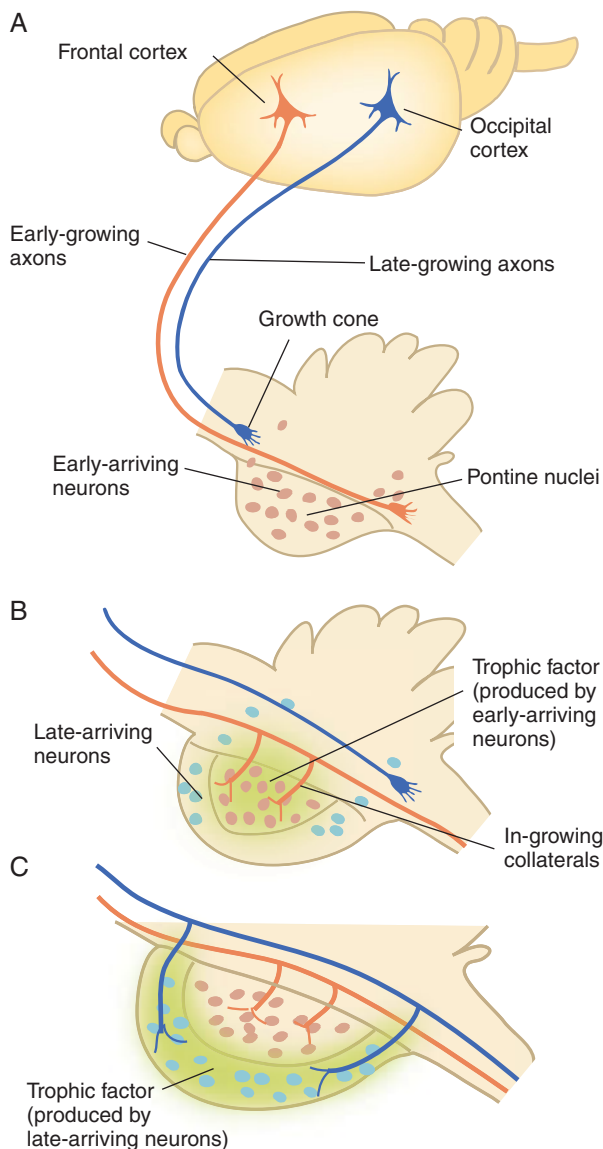


FIGURE 9.15 Development of topographically organized connections: example from the corticopontine projection (cf. Fig. 14. 7). **A:** Because the axons from the frontal lobe start growing earlier than axons from the occipital lobe, they have reached further down in the brain stem. At this stage, neurons migrate into the ventral pons to form the pontine nuclei. **B:** At a later stage, the early-arriving pontine neurons produce a trophic substance that attracts collaterals from the descending axons. However, only the axons from the frontal lobe are sufficiently mature to emit collaterals at this stage (2 days elapse from the time the axons arrive in the pons until they can form collaterals). Immature neurons continue to invade the ventral pons and form a shell around the early-arriving neurons. **C:** Late-arriving pontine neurons are now sufficiently mature to produce the trophic factor, whereas the early-arriving ones have stopped their production. At this later stage, only the axons from the occipital lobe can emit collaterals. Thus, axons from different parts of the cerebral cortex end in different parts of the pontine nuclei, forming the orderly topographic arrangement seen in the adult (see Fig. 24.8). In this example, the topographic arrangement can be explained by genetically programmed differences in the time of birth for neurons in various parts of the cerebral cortex and the pontine nuclei. Other mechanisms may operate as well. (Based on experimental studies in the rat by Leergaard, Lakke, and Bjaalie 1995.)

diffuse in the tissue. Axons then grow in the direction of increasing concentration of the factor, which binds to specific receptors, so that only axons expressing the receptor are attracted. This may explain why axons from a neuronal group in some cases grow toward their correct target even if the neuronal group has been moved to another site before axonal outgrowth.

Cell-Adhesion Molecules and Fasciculation

In other instances, neurons or glia along the route apparently express specific molecules that function as “signposts.” When the first **pioneer axons** have reached their target, the rest of the axons can get there simply by following the pioneers. (The pioneers may have a relatively simple task because, as mentioned, the distances they grow are very short and the “landscape” is simple.) Axons with a common target can express at their surface **neuronal-cell adhesion molecules** (N-CAMs) that make them sticky (N-CAMs are proteins related to the immunoglobulins). Axons expressing a particular kind of N-CAM are then kept together, whereas others are repelled or inhibited in their growth. In this way axons with common targets form bundles, or **fascicles**, and the phenomenon is termed **fasciculation**. The well-defined tracts of the nervous system arise in this way.

Growth Cones and Their Interactions with N-CAMs and Other Molecules

The many trophic factors, “signpost” molecules, recognition molecules, and N-CAMs governing the establishment of specific connections mostly act on the axonal **growth cone**. The growth cone is an expanded part of the tip of a growing axon (Figs. 9.15). It continuously sends out small extensions, or **filopodia**, as if exploring its immediate surroundings (Fig. 9.16). Filopodia encountering the proper molecules in the tissue are stabilized, and this determines the direction of further growth, while other filopodia retract. The stabilization depends on increased number of **actin** molecules in the filopodia and of **microtubules** in the axon. Several N-CAMs influence the growth of the axon when they are present near the growth cone. These N-CAMs can be expressed at the surface of both the growth cone and the nearby cells. Binding of the N-CAM molecules to each other causes stickiness. In other instances, N-CAMs bind to specific receptors in the growth cone membrane and thus activate intracellular second messengers (by influx of Ca^{2+} ions). There are also **extracellular molecules** with actions on the growth cone, such as **laminin**, which is bound to the basal lamina. Laminin binds to receptors in the growth cone membrane. Laminin can occur transiently along the path of growing axons in the peripheral nervous system, guiding the growth

toward the target organ. Although the action of laminin is not by itself specific, its time-specific expression ensures that only axons present at the proper time will grow. This is another example of the importance of **timing**—in this case of axonal outgrowth—during development.

Examples of Axonal Pathfinding

The “inverted” pyramidal cells, previously mentioned, exemplify that gradients of substances in the environment of the axon govern growth direction. Another example concerns the development of descending connections from the cerebral cortex—that is, the outgrowth of axons from **projection neurons** in the cerebral cortex. Many such axons reach the spinal cord and, in addition, emit collaterals to nuclei in the brain stem. Initially, the axons grow toward the cord without sending out collaterals (Fig. 9.15A). After a certain interval or waiting time, however, collaterals grow out and innervate the **pontine nuclei** (Fig. 9.15B). The collaterals to the pontine nuclei arise just when the pontine neurons have reached a certain stage of maturation (postmitotic age). Most likely, the pontine neurons at this stage produce a trophic factor that “attracts” growing axons. Such a mechanism, depending on diffusion, can operate only over short distances. Thus, it is less likely that trophic factors produced in the spinal cord are responsible for the goal-directed growth of corticospinal axons.

When the distance to the target is too long for trophic factors alone to guide axonal growth, we assume that

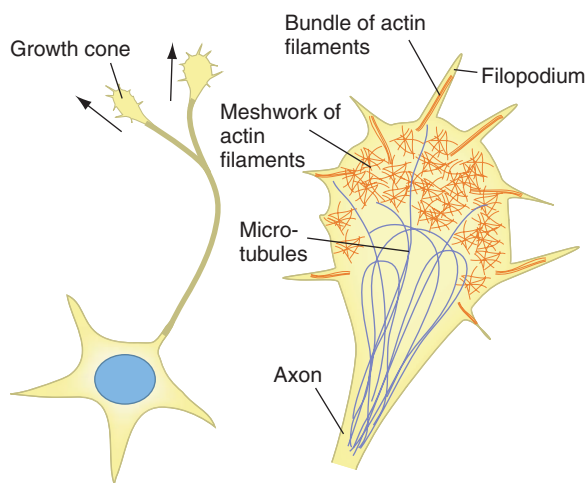


FIGURE 9.16 *The axonal growth cone.* A growth cone contains a central and stable bundle of microtubules. In addition, dynamic microtubules extend toward the filopodia and work together with actin filaments. The filopodia contain bundles of actin filaments providing motility. The filopodia extend or retract depending on the specific molecules they meet in their immediate environment. (Based on Kalil and Dent 2005.)

there are “**signpost**” molecules along the route. One example illustrating this is the growth of axons from the retina through the **optic chiasm**. Figure 16.14 shows the arrangement of the axons from various parts of the retina as they pass through the optic chiasm in the adult. Axons from the nasal retina cross to the other side, whereas axons from the temporal retina pass ipsilaterally—that is, without crossing. In the adult, crossing and ipsilateral axons are segregated as they approach the optic chiasm. During the first outgrowth, however, axons from the nasal and temporal parts of the retina are mixed. Nevertheless, they take the correct path when they reach the region of the chiasm, even when this requires that some have to bend 90 degrees, and that some axons must cross each other. Interaction between axons from the two eyes is not necessary, either. Thus, even if one of the eye primordia is removed before the axons have reached the optic chiasm the axons from the remaining eye still find their correct way through. Neither can trophic factors from the target organ of the axons (a thalamic nucleus) play a decisive role because this nucleus is not yet established at the time the axons grow through the optic chiasm. Therefore, local clues in the region of the optic chiasm must guide the axons—at least during the first pioneering phase. In the next few weeks—when thousands of axons follow the pioneers through the chiasm—cell adhesion molecules and fasciculation are important.

Another example of the importance of local “signpost” cues for axonal pathfinding is the development of skeletal muscle nerve supply by **spinal motoneurons**. Even after removal of the primordial muscle, the axons that would normally supply it still find their way to the site of the (removed) muscle.

Elimination of Axon Collaterals and Synapses

As mentioned, the terminal area of a group of axons is often more extensive initially than after maturation, forming a surplus of synapses. Newborn monkeys, for example, have higher numbers of synapses on each neuron in many parts of the brain than adult monkeys. Thus, many collaterals and synapses are eliminated during further development. When it occurs in early phases of development (prenatally), elimination is probably due mainly to programmed cell death. Later on, competition for growth factors may be more important. In this way, the axonal ramifications of each neuron are pruned, thereby increasing the spatial precision of connections. In other instances, neurons initially send axon collaterals to two (or more) nuclei, but only one collateral survives while the other disappears during further development.

Elimination of axon collaterals is exemplified by the development of descending connections from the **visual cortex**. Initially, the axons grow down to the spinal

cord, sending off collaterals to the pontine nuclei (among other areas) on their way (Fig. 9.15). After the pontine collaterals are established, those to the cord disappear (in adult animals there are no connections from the visual cortex to the cord). We do not know how this happens; perhaps trophic factors from the cord or medulla “trick” the axons to grow beyond their real targets. We may further assume that once in the cord, the axons are not able to establish synapses, perhaps because they lack specific recognition molecules or simply because the spinal neurons are not receptive at the time the axons arrive.

A further example concerns descending axons from the forelimb region of the **motor cortex** (of rodents). Initially, such axons reach both the cervical and the lumbar parts of the cord (innervating the forelimb and the hind limb, respectively). Collaterals enter the spinal gray matter only in the cervical region, however, and the branch to the lumbar cord disappears without having established synapses.

Formation and Elimination of Synapses from Newborn to Adult

While the number of neurons in the human cerebral cortex appears to be fairly constant after the twenty-eighth gestational week, the **synaptic densities** undergo marked changes until the end of puberty. In general, the synaptic density in the cerebral cortex increases steeply from before birth to late in the first postnatal year. Thereafter, the density declines slowly—presumably due to elimination of synapses—until reaching adult values between age 10 and 15. There appears to be large variations among cortical areas, with earlier maximum density and a shorter period of synapse elimination in primary sensory areas than in association areas. In the visual cortex, synaptic density at birth is about 10% of that at 4 to 8 months after birth, when the number of synapses per neuron is estimated to be 15,000. Thereafter, the number declines to about 7500 at 10 to 12 years of age. This number seems then to be stable for many years. Similarly, in the auditory cortex, synaptic density peaks at about 3 months after birth, while synapse elimination ends at around age 12. In the frontal association areas, maximum synaptic density occurs about 15 months postnatally, while elimination ends around age 16.

We can only speculate on the functional meaning of such changes of synaptic numbers and densities. It seems reasonable, however, that a large surplus of synapses is useful in phases with large plastic changes. Presumably, there is a large pool of labile (perhaps “silent”) synapses, of which only some become stabilized by proper use. It may be significant that the postnatal overproduction of synapses is particularly marked in the human cerebral cortex, as compared with animals

with relatively less developed cerebral cortex. Most likely, this is related to the enormous human potential for learning and adaptation: evolution has favored flexibility and learning capacity at the expense of “secure” genetic preprogramming of brain synaptic interconnections.

The Brain Changes during Adolescence

For some unknown reason, the rate of synapse elimination appears to be particularly high just before and during puberty. In the monkey visual cortex, for example, the maximum number of synapses occurs around the third postnatal month. Thereafter, the number is fairly constant until puberty, when a marked reduction occurs (40% loss) to obtain stable adult values. Longitudinal MRI studies in children and adolescents indicate that the ratio of gray and white matter changes, especially around puberty—in spite of unaltered total brain volume. First, there is an increase of gray matter that is most marked in the frontal lobes. After puberty, gray matter declines while white matter increases. These changes appear to continue into the late 20s. Although their functional meaning is not clear, such data strengthen the impression that considerable plastic changes occur in the brain during adolescence.

Establishment of Topographic Maps

When axons arrive at their target, they usually do not establish synapses at random among the neurons but, rather, in a restricted part of a neuronal group. Thus, the incoming axons together form a topographic map so that, for example, different body parts (see Fig. 14.7) or parts of the visual field (see Fig. 16.19) are represented by spatially separate neurons. Most connections in the brain exhibit some degree of topographic organization, and this is not restricted to systems that convey sensory information and motor commands. Some topographic maps are simple, while others may appear highly complex.

In some parts of the brain—for example, in the **superior colliculus** where axons from the retina form a map of the visual field—there exist **gradients** of specific receptors along different axes during development.⁷

Genetic programming of **neuronal time of birth** is another important mechanism. (Time of birth also decides the scheduling of axonal outgrowth and the

7 In the early development of the superior colliculus, neurons express membrane-bound **ephrins** (ephrin A [EphA] and B [EphB]) that form mediolateral and anteroposterior gradients. Axons arising in different parts of the retina differ with regard to the kind of ephrin (EphA or EphB) receptor they express and are thereby specified to establish synapses in certain parts of the colliculus. Activation of the Eph receptor requires contact between the axon and the target neuron. In general, it appears that activation of EphA receptors leads to axon repulsion—that is, the growth cone collapses—while activation of EphB leads to attraction with stabilization of the growth cone (partly by actin polymerization). Ephrins are also involved in neuronal migration and synapse formation.

ability to form and receive synapses, as discussed above.) Figure 9.15 illustrates how topographic maps in the **pontine nuclei** may arise. In the adult, axons from the cerebral cortex end in a precise topographic pattern in the pontine nuclei. Functionally different parts of the cerebral cortex connect with different parts of the nuclei (Fig. 24.9). This pattern appears to arise because axons from different parts of the cortex start growing at different times and because subgroups of pontine neurons are born at different times.

Although the basic pattern of topography is genetically determined, proper use of the system is necessary to obtain maximum precision of spatial arrangements. We return to this next in considering the environmental effects of experience on the development of the nervous system.

Prenatal Development of a Neuronal Network

We use as an example the development of the **spinal network** that controls **locomotion**. This network arises long before locomotor movements are of any use. In human fetuses, rhythmic movements of the extremities (although uncoordinated) occur as early as 10 weeks after fertilization. At first, the spinal network initiates movements without any sensory information or commands from higher levels of the CNS. Gradually the coordination improves to complete locomotor patterns (improvement continues after birth when the system is used in a goal-directed manner). The prenatal improvement is probably due to several factors: altered electric properties of the neurons, expression of novel transmitters and receptors, and development of the connections among the network neurons. These alterations depend on the development of descending connections from the

brain stem (especially important might be those that contain serotonin). Maturation of the locomotor function starts with the forelimb and proceeds caudally to the hind limb, in parallel with the growth of descending axons.

Postnatal Growth of the Brain

The weight of the human brain triples during the first year of life (from 300 to 900 g), and at the same time major changes of synaptic density take place, as discussed in the preceding text. After the first year, the weight increases more slowly to reach adult values around the age of 5 to 7 (1400 g for men and 1250 g for women). This weight gain is caused by the growth of existing neurons and their processes, the myelination of axons, and the proliferation of glial cells. The neuronal growth mainly involves expansion of the dendritic trees and formation of axon collaterals with nerve terminals. The growth of dendritic arbors is especially marked in the human cerebral cortex during the first 2 years of life (Fig. 9.17). There is good evidence that increased dendritic ramifications relate to an increased number of synapses. Myelination of corticocortical connections takes place during the same period and is a sign of functional maturation. Increased conduction velocity enhances both the precision and the capacity of the neural networks; that is, their potential for information processing increases.

Human Brain Weights

Typical values of average adult brain weight from large autopsy studies are about 1400 g in men and 1250 g in women. Usually, such studies are based on brains of

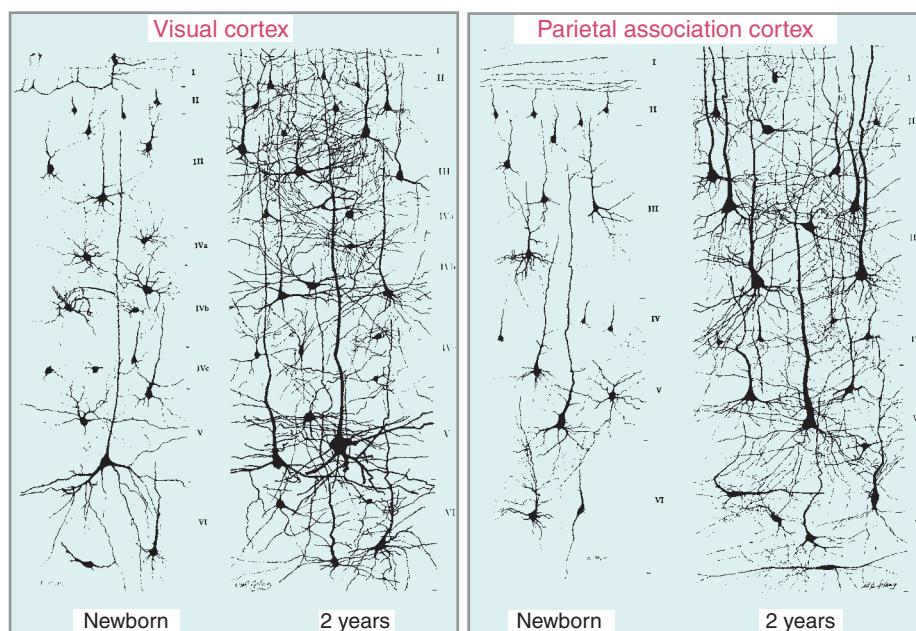


FIGURE 9.17 Postnatal development of the human cerebral cortex. Drawings based on Golgi sections from the visual cortex showing the increase in dendritic arborizations from birth and until the age of 2 years. The increase in dendritic arborization is most likely related to the increase in the number of synapses per neuron. (From Conel 1939.)

adults between the ages of 20 and 40 years. More interesting than average weight is the range of variation among normal individuals. An investigation of brains from 200 persons 17 to 40 years of age reported variations from 1120 to 1780 g in men and from 1070 to 1550 g in women. Older autopsy studies indicate that adult brain weight (volume) is reached between ages 5 and 10. An MRI study of 85 normal children, aged 5 to 17, did not find any volume increase after the age of 5. MRI studies show a weak correlation between brain volume and IQ, both among children and adults.

THE ROLE OF THE ENVIRONMENT IN DEVELOPMENT OF THE NERVOUS SYSTEM

Growth and Use-Dependent Plasticity

Although the brain's postnatal weight gain is largely genetically determined, the final functional state of the various neural systems and networks depends critically on their proper use during a certain period: they must be used at the proper time and in a meaningful way. The ensuing **use-dependent** structural changes occur mainly at the synaptic level, determining the final number of synapses, their precise distribution, and postsynaptic effects. Both establishment and elimination of synapses occur at high rates during the development toward a fully functioning system. As we have discussed, an enormous surplus of synapses arises in infancy, with subsequent synaptic elimination as the neuronal networks mature. Nevertheless, networks that are properly used appear to end up with a higher number of synapses than networks not subjected to normal challenges.⁸

"Enriched Environments" and Synaptic Plasticity

Numerous animal experiments have shown robust effects of environmental conditions on brain structure and biochemistry, as well as on behavior. Early experiments showed, for example, that dendritic arborizations were more extensive in the cerebral cortex of rats raised in an **enriched environment** (a simulated natural environment with ample space and access to toys) than in rats raised in standard laboratory cages.⁹ Brains of wild and tame animals represent a naturally occurring

analog of experiments with enriched environments. Indeed, wild animals have somewhat larger brains than tame animals of the same species. The responsible influence must occur quite early, because animals born in the wild and later tamed have the same brain size as wild animals. The difference is not genetic, however, because individuals of the first generation born in captivity have smaller brains than their wild relatives have. Experimental enrichment in the early postnatal period of rats induced increased synaptic density in the **hippocampus** and improved performance in certain learning and memory tasks. Results from experiments on the **motor system** further support the association between synaptic plasticity and learning: Among young adult rats doing different motor tasks, some developed more synapses per neuron in the cerebellar cortex than others did. The decisive factor was not the amount of motor activity but that the activity implied learning of new motor behaviors. Other experiments show that learning of specific skills is associated with synaptic changes in cortical areas involved in the task. For example, as monkeys gradually improved their performance in distinguishing tones of different frequencies, the part of the **auditory cortex** representing the particular frequencies increased in size. Similar changes occur in the motor cortex during learning of motor skills.

The preceding examples strongly suggest that formation and modification of synapses are closely linked with **learning**. It is plausible, for example, that life in the wild (or in an artificially enriched environment) requires the acquisition of a broader repertoire of adaptive behaviors than life in the cage or the bin. In general, there is good reason to assert that task-specific networks become operational during childhood because of learning processes driven by active interactions between the individual and his or her environment.

Information Must Be Meaningful

Nerve impulses are not by themselves sufficient for normal development, as shown in many animal experiments. For example, in goldfish exposed only to diffuse light (devoid of information) at the time retinal axons form synapses in the optic tectum, the normal ordered map of the visual environment does not develop properly. Another example concerns the development of connections from the retina to the visual cortex. At first, neurons in the visual cortex are influenced with equal strength from each eye. Soon after birth, however, neurons in the cortex segregate into groups, with a dominant input from one eye and a weaker input from the other. This phenomenon is called **eye dominance**. When all impulse traffic from the retina is blocked in kittens, eye dominance does not develop: each neuron continues to respond equally well to signals from either eye. Artificial electric stimulation of the

8 Kittens that do not use their vision during the phase of maximal synapse production (sensitive period), end up with about two-third of the normal number of synapses in the visual cortex. Generalizing from this observation, one-third of synapses in the cerebral cortex may depend on proper use of the functional system in which they participate.

9 It should be emphasized that so-called enriched environments are enriched only as compared with standard laboratory conditions. The latter is a situation of **deprivation** rather than of normality. Both the standard and the enriched-conditions are therefore highly artificial. Nevertheless, when interpreted with caution, experiments with enriched environments give robust evidence of the role of environmental factors in brain development and function.

optic nerve can nevertheless induce eye dominance even though the animal is blind, but only if the signals from the two eyes arrive with a minute time difference (corresponding to what occurs under natural conditions with light falling on the two retinas). With natural use of the system during development, axon terminals conveying signals from the two eyes compete for the available synaptic sites on each neuron.

Both of the aforementioned examples from the visual system show that, in order to induce the normal synaptic pattern and connectivity, nerve impulses must convey **meaningful information**—that is, information that helps the animal adapt to its environment.

Early Social Experience Alters Brain Structure and Function

Human experiences and animal experiments strongly suggest that social conditions in early childhood can influence adult emotional and cognitive behavior, and that this is associated with alterations of brain structure. For example, rat pups of mothers spending much time licking and grooming them develop higher synaptic density in the **hippocampus** than do pups with mothers paying less attention to them. In agreement with alterations of the hippocampus, the pups of “high-licking” mothers also show enhanced spatial learning and memory. Further, as adults, the offspring of “high-licking” mothers show a different response to stress than offspring of “low-licking” mothers.¹⁰ Another example concerns the effects of exposing rat pups to traumatic emotional experiences, for example by separating them from their mothers. Such pups show synaptic alterations in the **anterior cingulate gyrus** (see Fig. 6.26), a region that is involved in emotional processing. Further, separation was associated with altered development of neural networks that control emotional responses mediated by the **autonomic nervous system**.

Even though early experiences seem capable of producing life-long alterations of brain structure, improved environmental conditions at a later stage can reverse the changes. For example, exposure to an enriched environment around puberty was found to normalize the structural changes of the rat hippocampus induced by early traumatic experiences. As summarized by the U.S. neuroscientist Robert M. Sapolsky (2004, p. 792): “Thus, early experience can have lifelong consequences

ranging from the molecular to the behavioral level. . . . But, to the great relief of many of us, early experience is not necessarily destiny. . . .”

Sensitive (Critical) Periods

The term **sensitive** or **critical period** pertains to the development of a particular function of the nervous system and refers to the period when the system responsible for the function is maximally plastic—that is, its capacity for structural and functional adaptations is at its highest.¹¹ Further, the full development of a particular function requires proper use of the relevant neuronal networks during the sensitive period. For example, the ability of the mature brain to process sensory information depends on use of the particular sensory system during periods in early postnatal development. The same holds for the development of many skills. Further, we know that later use of a system cannot fully compensate for the lack of use during the sensitive period.

Sensitive periods occur at different times and vary in duration for different systems and behaviors. The **opening** of a sensitive period coincides with increased plasticity and with intense use of the neural networks responsible for the particular behavior. In normal children, for example, intense training of walking starts at about 1 year of age, and the vocabulary develops almost explosively between ages 2 and 3. We can only speculate what drives—or **motivates**—a normal child to train so intensely just during the phases of maximum plasticity.

The phases of system-specific, maximum plasticity must coincide with **meaningful use** of the systems to ensure optimal development. For example, infant monkeys prevented from using their vision until the age of 3 to 6 months needed several months after regaining vision to learn to distinguish a circle from a square. A normal infant monkey learns this simple task in a few days. Monkeys deprived of vision during the sensitive period do not develop the proper synaptic connections in the cortex that enable them to extract the meaning of visual information. Further, they do not develop the ability to integrate vision and the other senses (see Chapter 34, under “The Parietal Lobe and the Development of the Ability to Integrate Somatosensory and Visual Information”). There is much evidence that later use and extra training cannot fully compensate for the lack of use during the sensitive period. For example, although language may be acquired even if training starts later than normally, development of the full potential requires that training start during the first few years of life.

10 This influence appears to be mediated by (at least) two intracellular pathways. First, a **high-licking** mother, as compared with a **low-licking** one, induces higher levels of serotonin in the pup’s brain. This in turn leads to increased NGF-expression in the hippocampus. In addition, the glucocorticoid receptor gene in the hippocampus is demethylated. This makes the gene permanently more accessible to activation by NGF, resulting in permanent high levels of glucocorticoid receptors in the hippocampus. The level of glucocorticoid receptors relates to behavioral and endocrine responses to stress in the adult animal.

11 Many researchers now prefer the term “sensitive” period because “critical” period may give a false impression of an all-or-nothing phenomenon. Thus, even though the sensitive period undoubtedly is of special importance, most systems remain plastic also after the end of the sensitive period.

Although the period of intense synaptic proliferation is brief and well defined, the sensitive period for functional development is usually much longer and need not have a clear-cut endpoint. Moreover, sensitive periods in **humans** are generally much longer than in animals with a shorter period of postnatal development. In humans the sensitive period for vision, for example, probably lasts for the first 2 to 3 years, although the development is most rapid during the first year. Various aspects of vision have different sensitive periods, and at the cellular level, neurons in different layers of the visual cortex develop their characteristic properties at different times. More complex behaviors have, as one would expect, later sensitive periods than simple behaviors. For example, in the visual system the sensitive periods for development of binocular vision ends long before the sensitive period for the analysis of complex objects.

Compared with other phases of development, a sensitive period is characterized by both increased plasticity and increased **vulnerability** of the nervous structures involved. These factors are mutually dependent: one cannot have the one without the other. Thus, lack of proper stimulation, lack of opportunities for practice, or exposure to harmful environments has effects that are more serious during sensitive periods than at other times.

Cellular Mechanisms and Sensitive Periods

We do not fully understand the **cellular mechanisms** underlying the enhanced plasticity during sensitive periods. We do know, however, that sensitive periods in experimental animals correlate with the ease of LTP induction (long-term potentiation). We also know that **monoamines** play an important role. Thus, a sufficient level of these transmitters must be present in the cerebral cortex at the time of synaptic proliferation. Monoaminergic fibers are the first to grow into the cerebral cortex during development, perhaps to prepare the ground for the sensitive period. Through their modulatory actions, monoamines may ensure the necessary responsiveness of neurons to new synapses. Several growth factors and **neurotrophins**, known to influence neuronal plasticity, are expressed during sensitive periods.

Start and End of Sensitive Periods

We do not fully understand which mechanisms start and end a sensitive period. In general terms, it seems likely that a sensitive period **starts** when a genetically determined development of relevant neuronal networks reaches a certain stage, so that, as it were, experience has a substrate to work on. We know that the start of a sensitive period coincides with increased plasticity. In addition, experience in itself seems to boost plasticity,

as witnessed by delay of the plasticity increase if use of the system is prevented. In the visual system, the earliest sensitive period seems to start with a genetically determined proliferation of synapses in a particular neuronal network. Figure 9.18 shows the increase of synapses per neuron from birth to adulthood in the cat visual cortex. The largest increase starts at the time the kittens open their eyes at about 8 days and ends after about 1 month. Further experiments showed that part of this increase takes place only if the visual system is used. In one group of kittens, the eyelids were sutured in the first postnatal week, whereas in another group the optic nerves were cut. Both groups had about 30% fewer synapses per neuron in the visual cortex than controls. It is worth noting that the effect was the same regardless of whether the visual cortex was completely cut off from afferent signals from the retina or only lacked a meaningful sensory input (with sutured eyelids, action potentials still travel in the optic nerve).

As to what **ends** a sensitive period, reduced levels of monoamines probably contributes. Another factor appears to be signal molecules from myelin that reduce plasticity. Further, the development of GABAergic inhibition seems to correlate in time with the end of sensitive periods. Indeed, a proper balance between excitation and inhibition is a prerequisite for normal functioning of neuronal networks. In functional terms, the sensitive period would seem to end when the neuronal networks have attained the level of structural refinement that enables them to perform the tasks demanded of them. If a system is not used at the right time, the start of the sensitive period may be delayed for some time: the

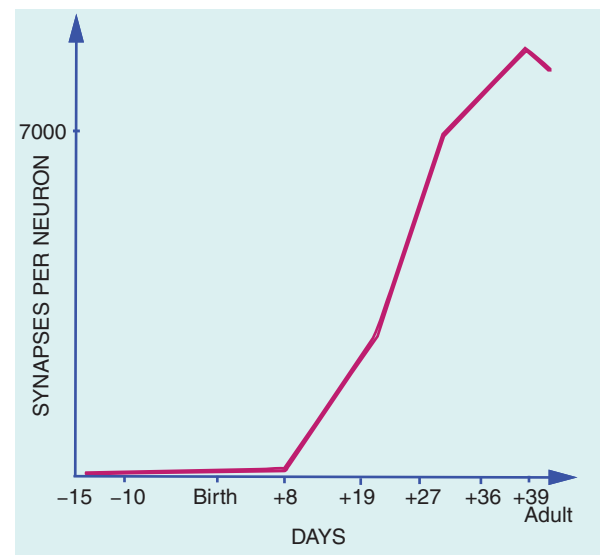


FIGURE 9.18 Postnatal increase in number of synapses during a sensitive period. The graph shows the change in the number of synapses per neuron in the visual cortex of kittens. The number increases steeply when the kittens open their eyes 8 days after birth. (Based on Cragg 1972.)

system “waits” for the right signals. However, if time passes without proper use, the networks seem to be taken over by other systems. In this way, they become less and less accessible for their proper inputs. One example concerns children that are born blind: as adults, their visual areas are activated by somatosensory stimuli (e.g., during Braille reading).

Examples of Sensitive Periods in Humans

Infants born with opaque lenses in their eyes—**cataracts**—can develop normal vision if the lens is removed at a very early stage. The longer the time before operation, the smaller the chances are that the child will attain normal vision. Persons attaining their sight after puberty (for example, by removal of an opaque lens) have grave difficulties using their sight. The “new” sense may cause trouble rather than being the expected blessing. Some choose to return to life as a blind person. A boy operated on at the age of 8 illustrates this problem. Several months of patient training were needed before he could recognize objects by sight (objects he was familiar with from the use of other senses). The surgeon who treated the boy concluded afterward: “To give back his sight to a congenitally blind patient is more the work of an educationalist than of a surgeon” (cited by Zeki, 1993, 216). The main reason for the problems encountered by this boy (and others in his situation) is most likely that he had not

established the cortical networks needed for integrating visual information with other sensory modalities. Thus, his brain was not capable of using the wealth of information provided by his eyes. In contrast, if the visual system has been used normally during the sensitive period in infancy and early childhood, even many years of temporary blindness have no serious effects on visual capacities.

Another example concerns children who are born **deaf** and later receive a **cochlear implant** that supplies the brain with information about sounds of different frequencies. Experience shows that such children can develop useful language and hearing behavior if the implant is provided early—that is, during the first 2 to 3 years of life. As with vision, access to auditory information during the sensitive period is necessary for proper development of the hearing system. To use sounds as basis for development of language, for example, numerous specific connections must be formed in the brain between the auditory cortex and other areas of the cerebral cortex. Such connections cannot be properly formed after the sensitive period. At least in part, this is due to other systems taking over the parts of the cerebral cortex that are normally engaged in auditory functions. Thus, deaf children activate the auditory cortex when using sign language (this is called cross-modal plasticity). Cochlear implants in such children do not restore sound-activation of the auditory cortex.

10 | The Nervous System and Aging

OVERVIEW

Many presumably harmful changes occur in the brain as we grow older. There are, for example, reductions of average brain **weight**, **synaptic numbers**, and **neurotransmitters**. There is also a loss of myelinated nerve fibers, and together all these changes would be expected to cause less efficient communication in neuronal networks. Age-related changes appear to affect especially networks of the **prefrontal cortex** and the **medial temporal lobe**—regions critically involved in memory and other cognitive functions. Nevertheless, most healthy elderly persons function remarkably well in spite of biologic alterations—with only a minor reduction of recent memory and some slowing of movements and mental processes. This is believed to result from compensatory, use-dependent plasticity initiated in response to the biologic changes. Thus, elderly individuals seem to activate larger parts of the brain when solving mental tasks (often both hemispheres are activated in contrast to strictly unilateral activation in younger people). Aging also entails a fairly marked loss of **peripheral receptors**, which may compromise vision, hearing, and balance. Such loss of peripheral receptors is often more bothersome than the changes occurring in the brain.

In **neurodegenerative diseases**, loss of neurons and their processes usually proceeds for many years before symptoms occur. This is at least partly due to compensatory processes going on in parallel with neuronal loss. The emerging symptoms depend mainly on which parts of the brain are subject to the neuronal loss. In spite of being caused by different mutations, **misfolding** of intracellular proteins occurs in many neurodegenerative diseases. Misfolding appears to initiate cytotoxicity. Neurodegenerative diseases with massive loss of neurons in the cerebral cortex lead to **dementia**, with Alzheimer's disease as the most common. This disease is characterized by—in addition to cortical neuronal loss—degeneration of the cholinergic **basal nucleus** (of Meynert) and marked loss of acetylcholine in the cortex.

AGE-RELATED CHANGES IN THE NORMAL BRAIN AND THEIR CONSEQUENCES

Biologic Changes and Their Interpretations

Comparisons of the brains of young and old persons have revealed several biologic differences. These include

a reduction of average brain **weight**, enlargement of ventricles (indicative of tissue atrophy), the appearance of **degenerative patches**, and **neuronal shrinkage** (that occurs primarily in very old people). **Dendritic trees** and **synaptic numbers** appear to be reduced in the cerebral cortex. Reduced **blood flow** of the whole brain or certain regions has been found in several studies. Several other changes, particularly in **neurotransmitters** and their **receptors**, have been reported. With imaging techniques, several alterations have been reported, notably changes of both **gray** and **white matter**. The relationship between biologic changes and impairments of brain performance is still debated, however. This is partly because of conflicting evidence on several important points and uncertain interpretations of findings. Interpretations are hampered, for example, because the biologic differences usually represent small group averages, while the individual differences within each age group may be much larger. Because detailed information of a person's behavioral and cognitive performance is seldom available when examining the brain after death, correlations must be tentative. In animal experiments, formal testing can be performed before examining the brain, enabling more affirmative conclusions regarding the correlation between behavior and age-related brain alterations. Unfortunately, some biologic alterations that are well documented in aged rats and monkeys are controversial in humans.

Nonuniform Distribution of Changes

In spite of the uncertainties just mentioned, one important point seems clear: Alterations of brain structures with advancing age are not uniformly distributed but, rather, are concentrated in specific parts of the brain. Thus, changes in a small part of the cortex may cause functional deficits without significantly affecting overall numbers of neuronal elements. Most consistently, changes have been observed in the **hippocampal formation** and the **prefrontal cortex**. In general, psychological tests show that functions of the brain that are served by these altered parts of the brain also show the most marked reductions by normal aging.

Elderly People Differ Widely with Regard to Brain Functions

It should be emphasized that individual differences with regard to brain functions are as marked among the

elderly as among young people. When we characterize persons in general terms according to their age, we often forget that persons of the same age—be it 25 or 80 years—differ widely not only physically but also emotionally and cognitively. For example, our personalities are quite stable over the adult life span. While certain traits are characteristic of elderly people as a group (as for adolescents and for middle aged people) the individual differences are actually more prominent. This even concerns **memory** for recent events, which is probably the most constant cognitive sign of aging. Investigations of aged rats, monkeys, and humans with standard tests for recent memory reveal that a relatively large minority does as well as young controls. Indeed, a study of more than 1600 persons with a simple memory test (repeat a list of 20 words just presented) showed that the performance of the oldest group (88 years) overlapped that of the youngest (25 years) by 50%. Another characteristic of elderly people is a reduction in **psychomotor** speed. They may speak more slowly, use more time to consider a question or to recall a name, and motor responses are less brisk than in young people. For many, however, this reduction is apparent only when the performance demand is high.

Loss of Neurons Does Not Explain Cognitive Decline in Normal Aging

Reduced mental capacities in older people are often explained simply by referring to neuronal death (as we know, lost neurons are generally not replaced). However, even though a modest loss of neurons may occur, there seems to be agreement that this can explain neither cognitive decline nor the reduction of brain weight reported in normal aging.

From a theoretical viewpoint, a modest diffuse neuronal loss would not necessarily cause clear reductions in the performance of the nervous system. As discussed earlier, large **populations** of neurons share responsibility for most specific tasks, especially higher mental functions such as abstract thinking, language, and memory. These are the products of **distributed networks** that connect neuronal groups in many parts of the brain. A diffusely distributed reduction of neurons in these networks would not be expected to cause functional deficits, except perhaps with maximal demands on performance.

Animal studies support that neuronal death is not the cause of normal age-related cognitive decline. Thus, old rats and monkeys showing cognitive impairments do not have fewer neurons than young animals in the hippocampal region and the prefrontal cortex.

How Many Neurons Are Lost?

Early studies of postmortem human brains suggested that as much as 20% to 50% of neurons are lost

during aging. More reliable methods later showed these numbers to be far too high (an important source of error being different shrinkage—due to different water content—of young and old brains). Indeed, recent studies with stereological methods find little or no neuronal loss in the human cerebral cortex. One study of 90 persons of both sexes suggested a loss of about 10% of cortical neurons from 20 to 90 years, while other studies did not find any significant neuronal loss in selected parts of the cortex with advancing age. Relatively few brains examined, and different sex and age distribution may probably explain some of the differences among studies. The total number of neurons in the cerebral cortex of humans may vary by 100% among individuals, according to studies with stereological methods. As noted by Alan Peters and coworkers (1998, p. 297): “Such large variations make it virtually impossible to accurately determine if there is a significant loss of neurons from an individual brain, and raises doubt about the significance of a loss up to 10% . . .”

Loss of Gray and White Matter

Even though few neurons are lost during aging, the brain nevertheless shrinks, as witnessed by data of brain weights after death and brain volumes during life (with magnetic resonance imaging [MRI]). MRI studies show that as we grow older, we lose both gray and white matter and the cerebral ventricles expand correspondingly. The reduction of **gray matter** is most likely due to shrinkage of cell bodies, loss of dendritic branches (especially the smaller ones and spines), loss of thin axonal branches and nerve terminals, and loss of water (as in all tissues of the body, the brain's water content decreases with age). Shrinkage of cell bodies seems to occur especially after 80 years. While the exact distribution of age-related cortical thinning (that is, gray matter loss) differs somewhat among studies, two findings appear to be consistent: first, the distribution is patchy with regions with marked thinning alternating with regions with no thinning, and second, alterations are most marked the **prefrontal cortex**.

There is good evidence from animal studies that many **synapses** are lost during aging.¹ For example, in the prefrontal cortex of old monkeys, there are on average 30% fewer synapses in some cortical layers than in young individuals. Further, there is a clear correlation between the magnitude of the synaptic loss and degree of cognitive decline in the old monkeys. Interestingly, the loss is particularly marked in layers 1, 2, and 3, giving off and receiving the bulk of corticocortical (association) connections. Corresponding to the

1 Whether corresponding age-dependent loss of synapses occur in the human cortex is not clear, however, owing to conflicting data from studies using comparable methods.

anatomic data, spontaneous excitatory postsynaptic potentials were reduced in the same layers in aged monkeys, suggesting reduced excitability of cortical neurons with aging.

Age-dependent loss of **white matter** of the cerebral hemispheres occurs in humans and in experimental animals. This appears to be due to loss of (mostly thin) myelinated fibers and alterations of myelin structure, presumably leading to degraded corticocortical connectivity. Corticocortical connections that link the prefrontal cortex with parietal and temporal cortical areas are of special importance for cognitive functions. Accordingly, neuropsychological tests of cognitive functions suggest a relationship between loss of white matter—especially in the frontal lobes—and cognitive decline.

In conclusion, it seems that the aging cortex is characterized by loss of synaptic contacts and corticocortical connectivity, affecting the prefrontal cortex to a larger degree than other parts of the brain. These alterations would be expected to slow down communication among cortical neurons, and might help to explain the typical slowing of mental processes in elderly people.

How Much Does the Brain Shrink during Aging?

A large study comparing average brain weights at 25 and 80 years of age found 1400 and 1300 g for men respectively, and 1250 and 1150 g for women. Thus, it would appear that the brain is on average some less than 10% lighter at age 80 than at age 25. For methodological reasons, this is probably an underestimate, and in vivo MRI studies indicate a volume loss of 20% to 30% from adolescence to 80 years of age. However, the shrinkage is not evenly distributed in the brain. A longitudinal study with MRI scanning of each person with an interval of 5 years suggested that the most marked cortical shrinkage affects association areas in the frontal, parietal, and temporal lobes. Marked shrinkage affected also the cerebellar hemispheres, the hippocampus, and the caudate nucleus. Other longitudinal studies have come to largely the same conclusions (although there are differences with regard to specific regions).

Neurotransmitters and Receptors

Reduced age-related levels of several neurotransmitters and their receptors—including glutamate, acetylcholine, dopamine, and norepinephrine—have been reported. However, most of these findings are difficult to evaluate with regard to their contributions to decline of brain functions. For example, alterations of glutamate transmission in prefrontal cortex might simply be the result of loss of excitatory synapses. In other instances, changes in transmitters and receptors may be the result of compensatory mechanisms rather than the cause of functional deficits. Further, compensatory changes of

other transmitter systems may prevent that a small change in density of a certain receptor causes a loss of function.

Recently, age-related change of **dopamine** receptors has attracted special interest. Dopamine plays a role in regulating attention (among other actions), and loss of dopaminergic receptors—especially in the prefrontal cortex—has been suggested to contribute to the cognitive deficits observed in the elderly. Loss of dopamine receptors probably reduces the **signal-to-noise ratio** for cortical neurons, making them less specific and more prone to erroneous responses. Indeed, PET studies have shown a relationship among density of dopamine receptors, cognition, and age. However, we do not know whether reduced dopamine transmission is a major factor in cognitive aging.

Why Is Psychomotor Speed Reduced in the Elderly?

Most likely, the reduction in psychomotor speed mainly reflects longer time needed for signal transfer and information processing in the old brain, caused by slower axonal conduction in myelinated fibers and loss of excitatory synapses. A reduced number of excitatory synapses on a neuron would prolong the time needed to reach the threshold for eliciting action potentials. Reduced amounts of available neurotransmitters and receptors would have the same effect. Such alterations might also explain that the **reaction time** is longer in older than in younger persons.

Memory Impairment in the Elderly

Solid evidence links age-dependent memory loss to changes in specific areas of the brain. Studies of aged rats indicate that their memory impairments are due to anatomic and physiological changes in the **hippocampal region**. The hippocampus and surrounding parts of the temporal lobe are necessary for the storage and retrieval of events, faces, names, and so forth (see Chapter 32). The changes occurring in the hippocampus are quite specific and affect only certain kinds of neurons and synapses. Some changes are presumably caused by compensatory mechanisms. For example, one kind of neuron receives fewer synapses, but each nerve terminal releases more transmitter. Long-term potentiation (LTP) is more difficult to produce and does not last as long in aged rats as in young ones. The memory impairment in aged monkeys has the same characteristics as in young monkeys after removal of the medial temporal lobe. Thus, it seems likely that age-dependent loss of recent memory is caused primarily by specific changes in the hippocampus and surrounding parts of the medial temporal lobe. In addition, changes of the dorsolateral parts of the prefrontal cortex are associated with reduced working memory (the ability to hold a number of items temporarily in memory).

Loss of Peripheral Sensory Receptors with Age

For everyday problems of the elderly, changes in the CNS may be less important than loss of sensory cells and neurons of the peripheral sensory organs. These losses are of sight and hearing in particular, but joint sense and sense of equilibrium also deteriorate with advancing age. For example, animal experiments show that muscle stretch receptors (muscle spindles) lose dynamic sensitivity. This might result in slower reflex responses to unexpected, sudden movements of the limbs. In general, thick myelinated, fast conducting fibers are more vulnerable during aging than thin fibers. Especially, impaired proprioception and cutaneous sensation in the lower extremities may contribute to increased risk of falling among the elderly.

Transmission of sensory signals at lower levels of the CNS is serial—that is, the different links in the chain are coupled one after another (see Fig. 14.1). If one link is completely broken, all transmission stops, and the brain cannot compensate for the loss of sensory information. This contrasts with the high degree of parallel processing taking place at higher levels, particularly in the cerebral cortex.

Dizziness and Loss of Vestibular Receptors

Studies of eye movements show that the **vestibuloocular reflex (VOR)** is less accurate in persons older than the age 75 than in younger persons. This reflex ensures that the gaze is kept fixed at one point in the environment when the head rotates (to keep the retinal image stable). Further, **optokinetic eye movements** become more sluggish in old age. Such eye movements are elicited when the environment moves—for example, when looking out of a car—so that the gaze is kept fixed. Suppression of the VOR, which is necessary when the head rotates in the same direction as the visual scene, is also impaired in many elderly persons. All these changes may cause difficulties with vision and orientation while moving, especially if the movement is rapid. This may be an important factor in the dizziness bothering many elderly people. At the cellular level, a major cause of impaired equilibrium and eye movements may be loss of sensory cells in the vestibular apparatus. A 40% loss has been reported in persons 75 years of age.

Plasticity May Compensate for Age-Dependent Losses

As we grow older, potentially harmful changes occur in the brain, as discussed in the preceding text. Yet, most elderly people manage remarkably well and show only minor functional deficits, which often become apparent only in situations with high demands. As said by Denise Park and Patricia Reuter-Lorenz (2008, p. 183): “The puzzle for cognitive neuroscientists is not so much in

explaining age-related decline, but rather in understanding the high level of cognitive success that can be maintained by older adults in the face of such significant neurobiological changes.” There is now much evidence that this seeming paradox arises because the nervous system remains plastic throughout life (even though the plasticity decreases with advancing age). Thus, even in old age we can learn and thereby uphold functions that are threatened by loss of neuronal elements. Probably, this process is not principally different from what takes place at any age when the brain is challenged—be it by damage or disease, need for new skills, or novel environments.

Interestingly, brain-imaging studies show that elderly and young people have different patterns of **cortical activation** during cognitive tasks, even when performance is equal. In particular, processes that are strongly lateralized in the young are more evenly divided between the two hemispheres in the elderly (Fig. 10.1). Much evidence supports that this activation pattern in the elderly is a sign of functional compensation rather than of faulty processing. For example, the tendency to use both hemispheres is associated with higher performance compared with other elderly persons using mainly one hemisphere. Elderly persons show less hippocampal activation than young people on certain **memory** tests, presumably because of age-related alterations in the hippocampal region. Among the elderly, those showing increased prefrontal activation (compared with young persons and age-matched controls) performed better on memory tests than those with less prefrontal activation. This suggests that increased prefrontal activation in the

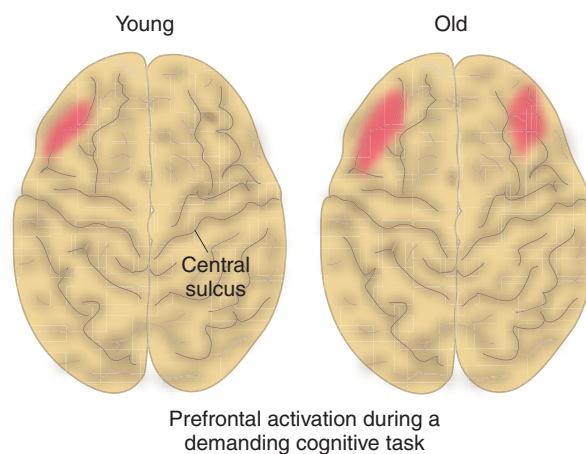


FIGURE 10.1 Altered brain activation patterns in the elderly. Comparisons with functional MRI of young and elderly people (older than 65 years) during execution of various cognitive tasks shows clear differences. This figure shows (in a very simplified form) that during a demanding verb-generation task, elderly persons activate the prefrontal cortex on both sides, whereas young people activate only one side (differences are apparent also in other parts of the brain). This extra activation most likely is due to compensatory (plastic) processes. It would mean that elderly people, while solving the task as well as young people, must allocate more resources to the task.

elderly compensates for reduced performance of the hippocampal region. Disrupting cortical processing by transcranial magnetic stimulation (TMS) supports that the bilateral frontal activation in the elderly is of functional significance. Thus, in the young, TMS of the left prefrontal region influenced memory performance most severely, while in older adults application to either the left or the right hemisphere reduced their performance equally. Also for successful **motor performance**, larger parts of the cortex and the cerebellum are recruited in elderly than in young persons.

Together, these and other observations strongly suggest that plastic changes occur in the aging brain, and that they counteract the detrimental effects of age-related loss of neural elements.

The Benefit of Experience

Normal aging does not affect everyday activities significantly. Indeed, it appears that all activities (motor or intellectual) that have become highly automated by long practice are quite resistant to age-dependent decline. Both speed and precision can then be maintained into advancing age. We need only think of musicians, such as Rubinstein and Horowitz, performing with excellence after 80. Preserved superior spatial memory in old taxi drivers provides another example of the effect of experience. Further, a study comparing young and old bridge and chess players concluded that there was no clear age-related decline in performance, with prior experience being more important than the player's age. A generally reduced short-term memory in the older players is presumably compensated by superior card recognition and specific memory for cards. Thus, old experienced players remembered more cards after a 1-second exposure than young untrained ones did, and this difference persisted even if the younger players were given more time. Another example of the beneficial effect of experience comes from a study comparing old and young bank employees. Although the older employees scored on average less on reasoning tests than the younger ones did, this did not correlate with poorer job performance. In conclusion, specific experience and continuous training appears to be more important for performance than normal age-related reductions in cognitive functions.

Presumably, continuous use of neural systems makes them more resistant to age-related impairment. One reason may be that more neurons are included in task-specific networks.² This might make the networks more robust and less vulnerable to a loss of synapses and

other elements. Further, age-related changes may be slowed by use-dependent production of growth factors and neurotrophins. Finally, by continuously challenging the systems, compensation by recruiting additional neuronal groups would counteract the inevitable negative effects of aging on the brain.

Physical Exercise May Protect the Aging Brain

There is evidence from both experimental animals and humans that physical training (especially cardiovascular conditioning) can improve cognition. This has also been examined specifically with regard to the aging population. For example, a study including more than 5000 women older than 65 years showed that those with the highest level of physical activity were less likely to develop cognitive decline during the next 6 to 8 years. Other studies indicate that **aerobic capacity**—as expressed in maximal oxygen uptake—is the factor most clearly linked with beneficial effects on cognitive functions. Especially executive cognitive processes—such as planning, task coordination, and working memory—seem to benefit from aerobic training. Increased production of **BDNF** (brain derived neurotrophic factor)—known to enhance neuronal plasticity—may mediate some of the effects of physical training on the brain.

NEURODEGENERATIVE DISEASES AND DEMENTIA

Dementia

Dementia can be defined as an acquired, global impairment of intellect, reason, and personality, but without impairment of consciousness. It is uncommon before the age of 60 but occurs with increasing frequency, especially after the age of 75. One-third of people older than 85 years of age may exhibit signs of dementia; although with varying severity. Dementia can have different causes, but all cases share extensive damage to neurons and connections of the cerebral cortex. One cause of dementia is loss of brain tissue due to ischemic brain lesions. Usually, the patient has suffered from repeated small infarctions of the white matter, causing **vascular cognitive impairment** (VCI).³ Dementia can also be caused by **intoxications** (alcohol, solvents, and carbon monoxide), **large tumors**, or **infections** (for example, HIV).

Neurodegenerative diseases leading to progressive neuronal loss, however, cause most cases of dementia. Most people in the latter group have **Alzheimer's disease** (AD). This disease was described just after 1900 as

2 London taxi drivers were found to have larger hippocampi than controls, and the difference increased with increasing experience (Maguire et al. 2000). Presumably, this represents a learning effect caused by their long-term engagement in spatial navigation.

3 Vascular cognitive impairment was formerly termed vascular dementia, and that term is still widely used. Cerebrovascular lesions may be the main cause in as many as 15% of all patients with dementia.

a distinct kind of dementia, which characteristically developed before the age of 60 (**presenile dementia**). Around 1970 it was realized, however, that the majority of cases of dementia beginning after age 60 also are of the Alzheimer kind. Despite all the research since then, so far, no one theory can explain the complex biochemical and pathological aspects of this devastating disease. **Frontotemporal dementia** (FTD) is less common than Alzheimer's disease and affects different parts of the cerebral cortex. This explains why the symptoms differ in these two forms of dementia. In many patients, multiple cerebrovascular lesions may coexist with Alzheimer pathology, aggravating the cognitive deficits (mixed type dementia).

Common Molecular Mechanisms in Neurodegenerative Diseases

Alzheimer's disease, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis (ALS), and other neurodegenerative diseases have in common that neurons slowly die in a long period preceding the debut of neurologic symptoms. These diseases affect different parts of the nervous system, occur in different age groups, and show different heritability. They therefore present as separate disease entities. They may nevertheless share basic cellular malfunctions that eventually cause neuronal death. One common finding is that neurons accumulate large amounts of **misfolded** proteins, which thereby induce cytotoxicity. Various factors, such as gene mutations, environmental influences, and aging may induce misfolding of proteins. Most likely, a modest amount of misfolded proteins occurs normally, and cellular processes are disturbed only when the amount of misfolded proteins exceeds the cell's capacity to remove them. Disturbed cellular processes comprise transcription, energy handling, axonal transport, synaptic function, and apoptosis. Apparently, different mutations can produce similar clinical pictures, the crucial point being which neuronal populations are affected. Thus, dementia will ensue if sufficient numbers of neurons in certain parts of the cerebral cortex die, regardless of the cause of cellular death. Neurodegenerative diseases are characterized by clumps of misfolded proteins—**inclusion bodies**—in the neuronal cytoplasm. Most likely, damaging effects of the misfolded proteins is exerted before they are collected in clumps, with the latter representing just the end stage in a cascade of molecular events.

Alzheimer's Disease

There is now little doubt that Alzheimer's disease (AD) represents a distinct disease entity, not just an exaggerated form of normal aging. The disease manifests itself by gradual decline of mental functions. Impaired recent

memory is an early sign: the person forgets appointments and names, repeats questions, and may appear confused and helpless. There is typically a loss of interest and initiative, and a neglect of daily activities. Initial symptoms are usually vague and hard to distinguish from normal aging. In some patients, the early signs of dementia may be misdiagnosed as depression. If such patients receive antidepressant drugs, their condition can deteriorate dramatically with confusion, loss of bladder control, and other symptoms. This may probably be explained, at least in part, by the anticholinergic effect of such drugs.

Microscopically, **senile plaques** and **neurofibrillary tangles** in the cerebral cortex characterize AD. The latter consist of thickened, intraneuronal fibrils. The senile plaques are patchy, degenerative changes with extracellular deposits of amyloid and other proteins, in which **amyloid β -protein** ($A\beta$) is a major component. The plaques also contain many abnormal axon terminals. $A\beta$ is formed from a large membrane protein called **amyloid precursor protein** (APP). We do not know the function of APP, but it is present normally in most neurons and is transported anterogradely in the axons to the terminals. $A\beta$ is normally secreted from neurons, and we do not know why it precipitates in AD. The **neurofibrillary tangles** consist of a specific kind of filament (**paired helical filaments**; PHF). A major component of PHF is an abnormal kind of the microtubule-associated protein **tau**. The protein lacks its normal ability to attach to microtubules, however, and this may explain why tau aggregates as abnormal filaments.

There is a marked loss of **synapses** in AD, as shown via electron microscopic observations. Most notable are losses in the layers two and three of the cerebral cortex (see Fig. 33.1) that send out and receive association connections. Figure 10.2 shows the typical distribution of early changes in the form of cortical atrophy. The pathologic changes are more severe in the **association areas** of the frontal and temporal lobes than in the primary sensory and motor areas. The earliest changes appear to occur regularly in the **entorhinal cortex** (part of the hippocampal region in the medial temporal lobe). It appears that severity of dementia shows a closer correlation with the degree of synaptic loss than with the density of senile plaques and neurofibrillary tangles. What causes the cell loss (and loss of synapses) is not quite clear, although both $A\beta$ and tau are involved in the process. Neither is the relationship between these two actors clear. The so-called **amyloid-cascade hypothesis** implies that overproduction of $A\beta$ leads to amyloid precipitation that induces formation of neurofibrillary tangles, which then cause neuronal death.

Alzheimer patients have a severe neuronal loss in a diffusely distributed cell group at the base of the cerebral hemispheres, called the **basal nucleus** (of Meynert) (Fig. 10.3). Many neurons in this nucleus are cholinergic and send

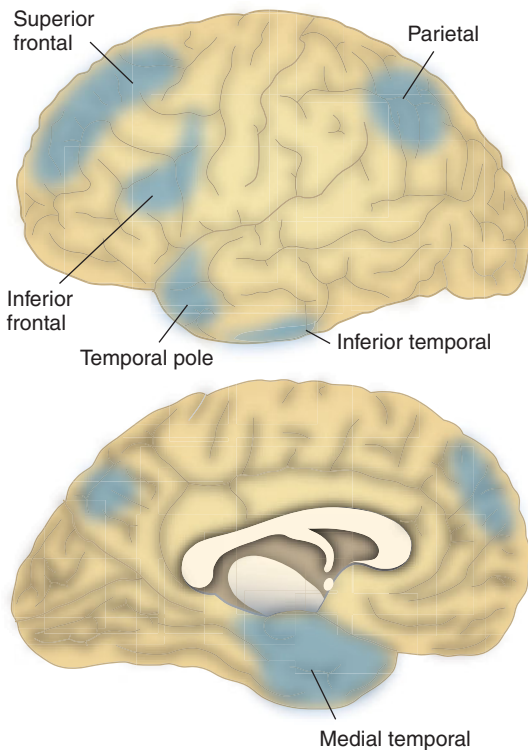


FIGURE 10.2 *The distribution of cortical atrophy in early stages of Alzheimer's disease.* The blue patches show approximate locations of regions with the most marked thinning of the cortex, based on quantitative analysis of MRI data. More extensive parts of the cortex become atrophic as the disease progresses. (Based on data of Dickerson et al. 2009.)

their axons to the cerebral cortex (see Chapter 31, under “Cholinergic Neurons Projecting to the Cerebral Cortex”). Loss of neurons in the basal nucleus may therefore explain why Alzheimer patients have severely reduced amounts of **acetylcholine** in the cortex. Acetylcholine raises the

excitability and improves the signal-to-noise ratio of cortical neurons, enhancing their ability to handle information rapidly and accurately. These facts form the basis of the first pharmacological treatment of AD, which aimed at increasing the amount of acetylcholine available in the cerebral cortex.

The relationship between the loss of basal nucleus neurons and the pathological changes in the cerebral cortex is not clear, however. Thus, not all aspects of the functional impairment in AD can be induced by destruction of the basal nucleus in experimental animals. Further, transmitters other than acetylcholine are also affected by AD. Many monoaminergic neurons die in the locus coeruleus and the raphe nuclei, which may explain why the levels of **norepinephrine** and **serotonin** in the cortex are markedly reduced. Depleting the cerebral cortex of acetylcholine, norepinephrine, or serotonin (rats) triggers a rapid rise of APP synthesis. Such transmitter depletion can be produced by destroying the basal nucleus, the raphe nuclei, or the locus coeruleus. These findings suggest, first, that APP synthesis is normally controlled by synaptic actions and, second, that precipitation of amyloid might be caused by loss of transmitter actions.

Even though patients with AD are quite similar with regard to brain pathology and clinical manifestations, they differ in terms of causal factors. **Heritability** plays a variable role but is especially important in patients with an early onset of symptoms (before 60 years). Many in the latter group inherit the disease dominantly. These patients have mutations of three genes: two genes coding for **presenilins** (1 and 2) and the **APP** gene. All three mutations increase A β synthesis. The much more common late-onset AD shows less obvious familial accumulation of cases. A large number of genes have been implicated as possible risk factors, however.

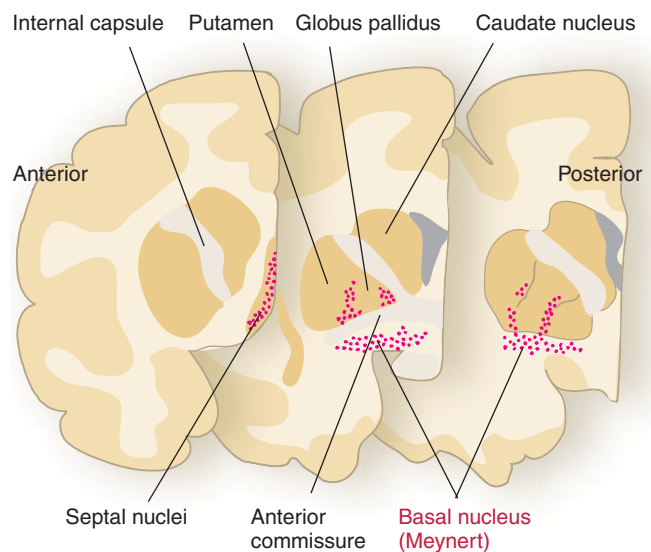


FIGURE 10.3 *The basal nucleus (of Meynert).* Frontal section through the left hemisphere (monkey). The cholinergic neurons of the basal nucleus are indicated with red dots. The basal nucleus is continuous anteriorly with the septal nuclei and other cholinergic cell groups of the forebrain. (Redrawn from Richardson and DeLong 1988.)

For example, persons with the *ApoE4* gene (coding for a apolipoprotein) run a much higher risk of AD than do persons with genes for another variety of apolipoprotein. Apolipoproteins normally transport cholesterol in the blood. ApoE4, however, also binds to amyloid β -protein and is present in senile plaques and neurofibrillary tangles. Therefore, it may facilitate the precipitation of amyloid.

Frontotemporal Dementia

Perhaps as many of 15% of all patients with dementia have **frontotemporal dementia** (FTD), formerly called

Pick's disease. It can be distinguished from AD by the distribution of degenerative changes, which affect primarily frontal and temporal parts of the cerebral cortex. Further, the initial symptoms are behavioral aberrations, typical for lesions of the affected regions, rather than loss of memory (patients with FTD are often misdiagnosed as suffering from a psychiatric disease). Both FTD and AD, however, are associated with abnormal phosphorylation of the microtubule-associated protein **tau**.

Restitution of Function after Brain Damage

OVERVIEW

Whatever the nature of the insult, brain damage ultimately leads to loss of neurons and their interconnections. Although new neurons are produced in limited parts of the adult brain, there is no effective replacement of lost neurons after brain damage. Reduced or abolished blood supply—**ischemia**—initiates a cascade of cellular events ending in cell death. Excessive release of **glutamate** and ensuing **excitation** appear to be crucial steps in a vicious cycle. In a region with focal ischemia (e.g., caused by occlusion of an artery) there is central zone where all cells will die and a peripheral zone—the **penumbra**—with diminished supply, in which neurons may silenced but do not die. They may be rescued if circulation is restored within a certain time (e.g., by thrombolytic drugs or reduction in edema).

The nervous system retains its **plasticity** throughout life, and this property is a prerequisite for recovery of function after damage of nervous tissue. Yet what changes do occur in the brain when a patient recovers functions after a stroke or other acute brain insults? Judging from animal experiments and brain imaging studies of human stroke patients, recovery is due to a learning process in which remaining parts of the brain are modified. We call this process **substitution**. The modification is driven by continuous efforts to perform tasks that were made difficult or impossible by the brain damage. Crucial factors determining the degree of progress are **motivation**, **focused attention**, and **amount of training**. Stroke patients who have recovered activate much larger parts of the cortex than normal persons when performing simple movements. In addition, plastic changes may be **compensatory**, in the sense that they do not restore the lost function but reduces disturbing symptoms or enable goal attainment by other means than those enabled by the damaged parts of the brain.

Infants with **early brain damage** may in some cases recover remarkably well, especially if lesions (even very large ones) are confined to one hemisphere. In other instances, with bilateral or lesions affecting deeper parts of the brain (basal ganglia, brain stem), complex functional impairments become stable and may even worsen during the first years of life. This may at least partly

be due to a disturbance of ongoing developmental processes.

BRAIN INJURIES AND POSSIBLE REPARATIVE PROCESSES

Ischemic Cell Damage

Abolished or reduced blood supply to an organ is called **ischemia**. Regardless of cause, even a few minutes of severe brain ischemia can produce massive neuronal death, but, surprisingly, the neurons often do not die until several hours after the ischemic episode. This is so because the lack of oxygen (hypoxia) and glucose triggers a cascade of events, in which each step takes some time but results eventually in neuronal death. Although many factors are implied, **glutamate** and glutamate receptors are under particularly strong suspicion as the main villains of this drama. Thus, ischemia leads to excessive release of glutamate (see later, “The Glutamate Hypothesis,” and Chapter 5, under “NMDA Receptors: Mediators of Both Learning and Neuronal Damage”). In an ischemic region, whether it is due to edema, bleeding (e.g., after head injuries), or vascular occlusion, there is usually a central zone where the ischemia is so severe that all neurons die.¹ Outside this region, however, there is a zone—the **penumbra** (from Latin *paene* = almost, and *umbra* = shadow)—characterized by neurons that are nonfunctional (unexcitable) but still viable.

The discovery of the delay between an ischemic episode and irreversible cell damage initiated an enormous research activity to find drugs that prevent or reduce the brain damage. In clinical situations, **focal ischemia** due to occlusion of a brain artery by thrombosis or embolus is more common than **global ischemia** (shutting down the blood supply to the whole brain, as can occur in cardiac arrest). In cases of focal ischemia hopes were raised to reduce the size of the brain infarct by

¹ Cell death in the central region completely devoid of blood supply is due to necrosis, as judged from the electron microscopic appearance. There is some evidence that apoptosis may cause cell death in the penumbra.

keeping neurons in the penumbra alive until the circulation improves (improvement may occur spontaneously or by use of thrombolytic drugs that dissolve the vascular obstruction). So far, however, the results with **glutamate-receptor blockers** (especially of the *N*-methyl-D-aspartate [NMDA] receptor) have been disappointing in humans in spite of convincing results in animal experiments. One problem has been intolerable side effects with the doses that are necessary to obtain protection. This might not be surprising considering that glutamatergic transmission participates in virtually every neuronal network. Another reason for the failure of numerous attempts to achieve neuroprotection after stroke may be that although the activation of NMDA receptors undoubtedly is harmful in the early stages of a stroke, it may be necessary for recovery in the delayed phase. Further, it appears that the cellular events evolving in the penumbra are much more complex than assumed—with regard to both the number of substances involved and the temporal profile of cellular changes.

Excessive release of glutamate cannot readily explain *all* aspects of ischemic cell death, however. For example, after global ischemia, the cell death is not diffusely distributed. Especially **vulnerable** are the neurons in the CA1 field of the hippocampus (see Fig. 32.10). Regional differences in glutamate release or glutamate receptors are probably not the reason. More likely explanations concern differences among regions regarding the presence and regulation of neurotrophic factors.

The Glutamate Hypothesis of Ischemic Cell Damage

According to the glutamate hypothesis, the sequence of events after a temporary stop of blood flow may be as follows (Fig. 11.1):

1. The loss of energy supply rapidly reduces the activity of the sodium–potassium pump, leading to (among other things)
2. Increased extracellular K^+ concentration that depolarizes the neurons so they fire bursts of action potentials, which leads to
3. Steadily worsening imbalances of ion concentrations across the cell membrane that make the neuron incapable of firing action potentials (electrically silent), while at the same time
4. The extracellular glutamate concentration rises steeply (because, among other things, the ionic imbalances reverse the pumping of glutamate by high-affinity transporters), leading to
5. Enormous activation of glutamate receptors, among them NMDA receptors; this in turn is believed to induce
6. Cell damage, probably because of an uncontrolled rise in intracellular Ca^{2+} concentration. This may harm the cell in various ways, for example, by activating

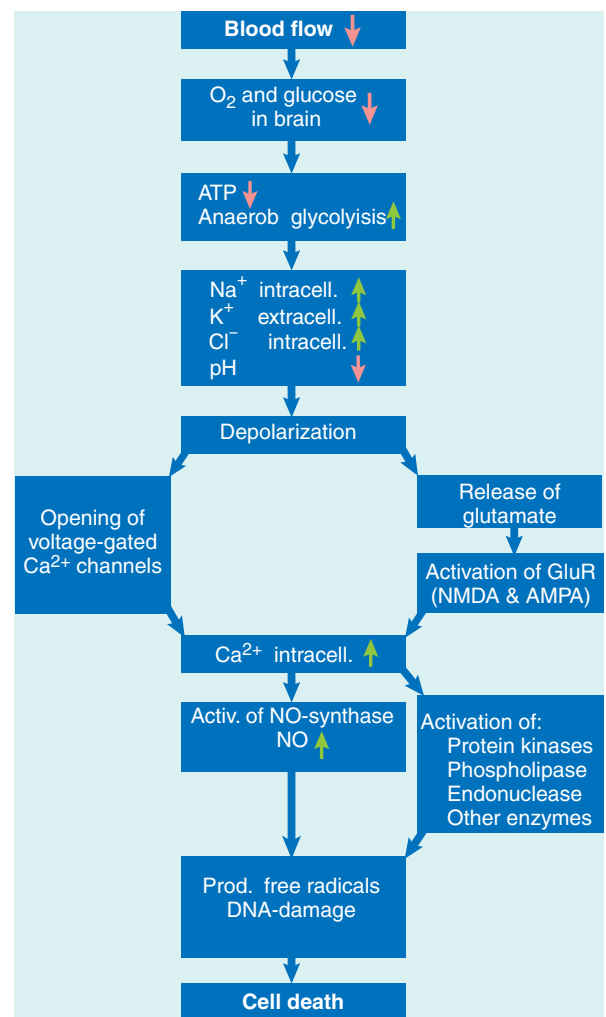


FIGURE 11.1 Sequence of cellular changes after focal ischemia leading to cell death (hypothetical). (Based on Samdani et al. 1997.)

enzymes that degrade proteins and nucleic acids and by activating nitric oxide (NO) synthases that leads to production of free radicals. In turn this destroys other vital enzymes.

7. Before the strong inflow of calcium there is an inflow of Na^+ accompanied by Cl^- and water, which causes neurons and glia to swell.

That **NMDA receptors** are involved in ischemic cell death is indirectly supported by animal experiments involving NMDA-receptor blockers. If such drugs are given even a few hours after an ischemic episode, they reduce or prevent the cell damage. Also, blockers of AMPA receptors provide protection against ischemic cell death in such experiments.

Neurogenesis: Production of New Neurons in the Adult Brain

In adult mammals, new neurons are produced continuously in parts of the subventricular zone and the dentate

gyrus of the hippocampal formation (see Fig. 32.10). The neurons produced in the subventricular zone migrate into the olfactory bulb (claims that some also populate the cerebral cortex have not been substantiated). Although only a minority of the newly formed neurons appears to survive, some are incorporated into existing networks. The finding that the number of surviving neurons is use-dependent further strengthens the assumption that the new neurons are of functional significance. Neurogenesis thus seems to represent an additional but spatially restricted form of plasticity, supplementing the ubiquitous synaptic plasticity discussed earlier in this book. It remains to be determined, however, how much and in what way neurogenesis contributes to the role of hippocampus in memory formation.

Nevertheless, adult neurogenesis has attracted much interest and raised hopes that it may be induced in regions with neuronal loss due to injury or disease. Indeed, increased neurogenesis occurred in the subventricular zone after ischemic brain injury in rats, and some neurons were incorporated in adjacent parts of the striatum. The clinical relevance of such observations is not yet known, however.

Why Is Neurogenesis and Regeneration So Restricted in the Human Brain?

Because neurogenesis—contrary to earlier beliefs—*does* occur in adult mammals, one may ask why it is so limited in distribution. Thus, in reptiles and birds neurogenesis occurs in large parts of the nervous system. One reason may be that new neurons might be disturbing rather than beneficial if not properly integrated with existing networks, and that the increasing complexity of the mammalian brain has rendered successful integration of new neurons less likely. Mutations that diminished neurogenesis may therefore have given an evolutionary advantage. The situation in the dentate gyrus may be special because of its specific role in memory acquisition—perhaps “ordinary” synaptic changes are insufficient or less efficacious than addition of new neurons.

Further, it is probably not coincidental that the mature CNS produces substances that **inhibit axonal growth** (such substances are not present during early development). An uncontrolled axonal growth in the mature brain might cause harm rather than help restitution. Correspondingly, although supply of growth promoting factors after brain damage might rescue sick neurons and promote axonal growth, it might also be expected to disturb the connections and functions of the healthy neurons.

Axonal Regeneration in the CNS—Hope for Spinal Cord Repair?

Why do severed axons regenerate in the peripheral nervous system but not in the CNS? This question, puzzling

neuroscientists for a century, has recently been answered at least partly. Thus, in the adult brain, the **growth cone** of an axon trying to regenerate quickly collapses due to the presence of inhibitory substances. Among several such substances, **myelin-associated proteins** produced by oligodendroglia have been most thoroughly studied. The proteins bind to receptors in the membrane of the growth cone and activate intracellular pathways that quickly down-regulate protein synthesis. In addition, scar formation at the site of injury also inhibits axonal regeneration.

Central axons may regenerate under certain conditions, however. Thus, the proximal stump of a cut central axon can grow into a piece of peripheral nerve if it contains viable **Schwann cells**. Indeed, growth of central axons for several centimeters has been demonstrated in nerve transplants in experimental animals. Another special kind of glial cell—**olfactory ensheathing cells (OECs)**—can also induce growth of central axons and has been subject to clinical trials. These cells cover the olfactory-receptor cell axons that extend from the nasal mucosa to the olfactory bulb. Interestingly, the olfactory receptor cells are renewed throughout life, and the OECs provide a permissive environment enabling the axons to grow into the CNS. Local supply of substances that block the growth-inhibiting proteins combined with transplantation of OECs is now used experimentally in patients with transverse lesions of the cord in the hope of inducing axonal regeneration across the lesion. So far, however—in spite of positive results in experimental animals—such procedures have not provided convincing functional improvement in patients. Animal studies (rodents) suggest that a combination of regeneration-promoting local measures and active rehabilitation might give the best results.

Collateral Sprouting Can Aid Restitution

Neurons that have lost their afferents may be supplied with new ones from normal axons in the vicinity. This is called **collateral sprouting**. After cutting afferent axons to a neuronal group (deafferentation), the axons degenerate and glial cells remove their nerve terminals. In a relatively short time, however, new nerve terminals fill the vacant synaptic sites. This probably occurs because a trophic substance becomes available from the deafferented neurons. The trophic substance stimulates nearby normal axons to send out sprouts. Obviously, collateral sprouting can only act locally in restoration of neuronal activity. There is no evidence that it can restore connections between more distant cell groups. Nor do all systems seem to exhibit collateral sprouting. Neither can collateral sprouting restore the original pattern of innervation because the neurons responsible for that pattern are gone. Thus, there will always be a loss of specificity of connections, compared with the

normal situation. Indeed, restitution of function seen after brain damage usually entails a loss of precision; for example, recovery of movement force is usually better than recovery of the ability to carry out delicate movements.

Although collateral sprouting probably aids recovery in many situations, it may not always be beneficial. For example, after a stroke that damages the motor pathways descending from the cerebral cortex, axon collaterals from sensory neurons may fill the vacant synaptic sites on motor neurons in the cord. This would not help to improve the voluntary control of the muscles but might, rather, contribute to the abnormally increased reactivity of the muscles to sensory stimuli that is characteristic of such cases (as in **spasticity**).

Can We Help Restitution by Neurotrophic Factors, Drugs, or Transplantation?

The well-established effects of **neurotrophins** and **growth factors** on plasticity and neuronal survival raise the question of whether they can be used therapeutically. Thus, one might expect that raising the level of such substances in the damaged brain might rescue neurons and stimulate the plastic changes necessary for restitution. The application in the CNS of such substances meets difficulties, however. Because they are proteins, the substances do not pass the blood–brain barrier readily, and even if they are delivered directly into the cerebrospinal fluid, their tissue penetration is restricted. Also, long-term supply of proteins to the brain may have uncontrollable actions and perhaps dangerous side effects. Therefore, the hope is to develop drugs that can stimulate the synthesis of specific neurotrophic factors in the brain itself or to improve their penetration of the blood–brain barrier.

Drugs that increase brain levels of **monoamines** (e.g., amphetamine) help recovery after brain lesions in experimental animals, apparently by increasing plasticity. From a theoretical point of view, it seems likely that the same would apply to humans. Indeed, some studies suggest that amphetamine speeds up recovery in stroke patients. Presumably, the effect would be best when combined with function-specific training.

Another means of helping restitution might be to replace lost neurons by **transplantation** of neural precursor cells—**neuroblasts**—into damaged regions. Embryonic neurons can still send out axons, provided the embryo is so young that the neuroblasts have not yet sent out axons. Besides numerous animal experiments, this approach has been tried in a limited number of patients with Parkinson's disease, giving promising results. Animal experiments show convincingly that neuroblasts can survive and send out axons after transplantation to an adult brain. If they are transplanted to a site where the adult neurons have been destroyed, the axons may even reach the normal targets of

the destroyed neurons. Functional restoration has also been demonstrated in some studies. Embryonic neuroblasts are obviously not affected by the myelin-associated factors that normally inhibit axonal growth in the mature CNS.

Many problems complicate the transplantations of embryonic tissue, however. One is the ethical issue of using brains of early abortions for therapy. Neuroblasts grown in culture may replace embryonic cells, however. Such cells may also be genetically modified to, for example, improve survival in the host. Other problems with neural transplantations are technical, such as the necessity for growth over long distances in the human brain (as compared with the rat) and survival of a sufficient number of neuroblasts. The chances of success are best with restoring diffusely organized, fairly short connections, whereas chances are presumably remote with precisely organized, long-ranging connections in the human brain.

BRAIN PROCESSES UNDERLYING RECOVERY OF FUNCTION

Substitution and Compensation

Differentiated neurons are unable to divide mitotically, and the brain does not possess a store of undifferentiated neurons that can multiply and replace lost ones (with some possible exceptions, as discussed earlier). Thus, as a rule, dead neurons are not replaced. We also know that cut axons do not regenerate in the CNS (the latter in contrast to the peripheral nervous system). Therefore, a reparative process in which the damaged structures regenerate or are replaced by new ones cannot explain the recovery of function after brain damage. Rather, recovery must be due to changes among the remaining undamaged neurons and glial cells. Indeed, there is much evidence to suggest that neural circuits **reorganize** to adapt to a novel situation. We discuss two kinds of adaptation next, both of which can probably best be regarded as **learning processes**.

In one type of adaptation—which we call **substitution**—intact neuronal groups take over and substitute for the damaged parts. The neuronal groups responsible for the substitution normally carry out tasks similar to those of the damaged ones. Thus, after a stroke destroys the cell groups that are responsible for initiating certain kinds of movement, the cells responsible for other kinds of movements may take over to some extent. For example, other cortical areas may partially substitute for functions lost when a lesion destroys the **primary motor cortex**, and neuronal groups that normally control only one side of the body may expand their activity to the other side as well. Substitution can seldom restore a system to its premorbid functional level, however. After all, an

amateur, who still has to take care of his former duties, has replaced a specialist!

The other kind of adaptation, called **compensation**, implies that any remaining structures change their normal activity to diminish disturbing symptoms produced by the injury. This is relevant when the brain damage not only leads to loss of functions, such as muscular weakness or sensory loss, but also produces disturbing phenomena such as involuntary movements or sensory confusion. For example, unilateral destruction of the **vestibular apparatus** in the inner ear initially causes severe dizziness and disturbances of posture and eye movements. This is due to a mismatch between the vestibular information reaching the brain from the two sides of the body. The symptoms usually subside quickly, because the vestibular system compensates by altering its sensitivity and information handling. Although this does not normalize the sense of equilibrium, it reduces disturbing symptoms. By substitution, the brain can learn to rely on other sources of information to control posture and movements. Thus, the postural adjustments necessary for the upright position gradually improve because visual information substitutes for vestibular. When denied the use of vision (as in the dark), the person becomes very unsteady.

Some patients do not recover useful function of a body part after brain injury. To some extent goals can still be achieved by using other movement strategies, for example, using the (normal) left arm instead of the paralyzed right, using a stick to keep balance, using both hands instead of one, and so forth. Also in such cases, the degree of success depends on a learning process, even though the level of performance as a rule will be severely reduced compared with the normal situation. This may also be regarded as a form of compensation, and should be distinguished from substitution.

Restitution after a Stroke Can Be Divided into Two Phases

After a person has a stroke, there is usually a **first phase** of rapid improvement lasting from days to weeks, followed by a **second phase** of slower progress lasting from months to years. Acute damage to neural tissue is usually caused by head injuries with crushing of neural tissue and bleeding or by vascular occlusion caused by thrombosis or an embolus. Secondary changes occur in the penumbra (the tissue surrounding the damaged area) such as edema (tissue swelling) and disturbed local circulation. If the edema subsides quickly and the circulation improves in the penumbra, neurons will regain their normal activity (compare with the transient weakness and loss of cutaneous sensation produced by pressure on a peripheral nerve). This is probably why there is often a marked improvement of the patient's condition during the first week or two after the accident.

For example, an arm that was totally paralyzed the day after a stroke may in a matter of days be only slightly weaker and clumsier than before the stroke. However, when the condition deteriorates rapidly after the initial insult, the reason is usually that the edema worsens and compromises the blood supply to more and more of the brain.

Certainly, plastic changes occur shortly after an injury (hours, days), as witnessed by animal experiments. Thus, there is probably no sharp distinction between the rapid and the slow phases of recovery—the nervous system starts its adaptation to the new circumstances immediately.

Methods to Study Neuronal Activity and Connectivity in the Living Brain

Throughout this book, references are made to studies using brain imaging techniques, particularly those that enable determination of regional cerebral blood flow (see Chapter 8, under “Regional Cerebral Blood Flow and Neuronal Activity”). Because of the link between a change of neuronal activity and local cerebral blood flow, the flow of blood through specific parts of the brain can be correlated with certain stimuli or specific sensory, motor, or mental activities. Computer tomography (CT) can be used to visualize the distribution of a radioactive substance in the living brain. This method, called **positron emission tomography (PET)**, is based on the use of isotopes that emit positrons. Positrons fuse immediately with electrons, producing two gamma rays going in opposite directions, thus permitting the identification of their origin in the brain with the aid of a powerful computer. PET produces images that show the distribution of an inhaled or injected radioactively labeled substance at a given time. By labeling substances of biological interest, one can determine their distribution in the body. In the case of **blood flow** measurement, radioactively labeled water is injected into the bloodstream. **Functional MRI (fMRI)** is the other main method to visualize dynamic changes in the brain. This method is based on the fact that the magnetic properties of hemoglobin depend on whether or not it is oxygenated, and small differences in blood oxyhemoglobin concentration can be detected with MRI. For unknown reasons, the oxygen uptake of nerve cells does not increase simultaneously with increased activity. Thus, it appears that neurons work anaerobically during a brief period of increased activity, despite a sufficient oxygen supply. The glucose uptake increases, however, and so does the blood flow. When the blood flow increases without increased oxygen uptake, more oxygen remains in the blood after passing the capillaries—that is, the arteriovenous O_2 difference is reduced. MRI can detect this change, and in this way the regions of increased or reduced blood flow can be visualized. An advantage

over PET is that the picture of blood flow changes can be compared with the precise MRI picture of the same person's brain. This permits localization of blood flow changes to anatomic structures with a spatial resolution down to less than 2 mm.

A drawback of blood flow measurement as an indicator of neuronal activity is its low time resolution compared with the time scale for signal transmission in the brain. In this respect, recording the electrical activity of the brain is superior. This is usually done by placing many electrodes on the head and is called **electroencephalography** (EEG, see Chapter 26, under "Electroencephalography"). EEG has been developed to enable study of topographic patterns of cerebral activation in relation to the performance of specific tasks, and coherence of EEG activity in different cortical areas suggests that the areas are functionally interconnected.

A drawback with EEG is its low spatial resolution, which means that only a crude correlation is possible between electrical activity and its origin in the brain. A more recent technique, **magnetic encephalography** (MEG), records the magnetic fields produced by the electric activity of the brain. The spatial resolution is much better than with EEG—at best, down to a few millimeters. (For both methods, however, the spatial resolution becomes poorer the deeper in the brain the source of activity is located.) A further advantage with the MEG technique is that it can be combined with MRI. This enables precise localization of the brain areas participating in a task, at the same time as the temporal aspects are analyzed (such as the sequence in which various cell groups are activated).

Electric stimulation of the brain through the skull (transcranially) requires an intensity that causes pain, and is therefore seldom used. With **transcranial magnetic stimulation** (TMS; also termed magnetic brain stimulation), however, neurons in the cerebral cortex can be activated painlessly with a short, intense magnetic pulse applied to the head. The magnetic field penetrates the skull and creates an electric current in the brain, primarily in the outer parts of the cerebral cortex. Magnetic brain stimulation also enables study of how disruption of normal activity in a specific part of the cortex affects behavior and cognitive functions.

Connectivity in the living brain can be studied with TMS and with **diffusion-weighted imaging** (DWI). TMS has been used to study the connections between the motor cortex and the spinal cord in healthy persons and in persons with motor dysfunction (e.g., multiple sclerosis). It can also be used to study whether parts of the brain are interconnected, and in particular, whether connectivity changes in the course of therapeutic interventions (such as rehabilitative training programs for stroke patients). DWI (and further developments of this method) enables visualization of major pathways in the brain, such as connections among various cortical areas,

and between the cerebral cortex and subcortical nuclei. The method cannot determine the polarity of connections and the spatial resolution is limited. Provided the results are critically evaluated in conjunction with experimental tracing data from primates, the method gives important information.

Studies of Recovery after a Stroke in Humans

Many stroke patients have muscular weakness (pareses) in the opposite side of the body as a dominating symptom. This is called **hemiplegia** (*hemi*, half; *plegia*, from Greek *plegē*, stroke). The symptoms are caused by destruction (due to occlusion of an artery or a bleeding) of motor pathways from the cerebral cortex to the brain stem and the cord (see Fig. 22.3). Usually, the injury occurs in the internal capsule (see Fig. 22.13) where the descending fibers from the cerebral cortex are collected. When such injuries cause hemiplegia, we use the term **capsular hemiplegia**. Because patients with this clinical condition are so common, and because the site of their injury is usually well localized, they have been the subjects for many studies of restitution. Some examples elucidating mechanisms behind restitution in humans are presented here.

Regional cerebral blood flow is closely linked with neuronal activity. Thus, when we find altered blood flow—using PET or fMRI brain imaging—in a specific part of the brain after a stroke, it is taken as evidence of altered activity caused by the stroke. One group of patients with capsular hemiplegia was tested 3 months after the stroke. They were then completely or substantially recovered; for example, they could all perform opposition movements with the fingers. The test was to touch the thumb sequentially with the fingers, in a rhythm determined by a metronome. In normal control persons, this task is associated with increased blood flow primarily in the opposite **motor cortex** (and in subcortical motor regions like the cerebellum and the basal ganglia). This fits with the fact that the pyramidal tract, which is necessary for precise finger movements, originates in the motor cortex and is crossed. The patients differed from the controls by showing increased activity in cortical areas that are not normally activated in this kind of simple, routine movement, notably in parts of the, **insula**, **posterior parietal cortex**, and **cingulate gyrus**. In controls, these areas show increased activity with complex movements and problem-solving tasks that require extra attention. The regions have in common that they send association connections to the **premotor area** (PMA; see Fig. 22.10) and by this route can influence voluntary movements. Indeed, increased activity is also present in the premotor area. In addition, some of the patients differed from the controls by showing increased activity in motor cortical areas on the same (ipsilateral) side as the affected hand. Such findings

suggest that functional recovery after hemiplegia is related to the patients learning to use larger parts of the cerebral cortex for control of simple finger movements than before the damage, and their using motor areas on the same side as the pareses. A large number of studies confirm the above findings, although the exact distribution of activation may vary somewhat among studies. Use of **transcranial magnetic stimulation (TMS)** in restituted stroke patients has furthermore showed that not only are cortical activation differently distributed but there is also evidence of altered connectivity, as assessed in a resting situation. Further support for **structural changes** during recovery comes from use of MRI-based morphometry, suggesting that an intense rehabilitative training program is associated with increased cortical gray matter in the regions most activated by movements.

The involvement of larger parts of the cerebral cortex may explain why simple, previously effortless movements require so much more **attention** and **mental energy** than before the stroke. There is evidence that descending fibers from various cortical motor areas, such as the primary motor cortex (MI), the supplementary motor area (SMA), and premotor area (PMA) occupy different parts of the internal capsule. Thus, when a capsular stroke damages the most powerful and direct pathway from the cortex to the motor neurons—arising in MI—other, parallel, descending pathways may be “trained” to activate motor neurons more powerfully than before the stroke.

The Contribution of the Undamaged Hemisphere to Recovery

Although many observations suggest that the undamaged hemisphere participates in recovery, the responsible mechanism is not clear. In particular, results are conflicting as to whether or not descending motor pathways from the undamaged hemisphere contribute—the other possibility being actions via the lesioned hemisphere and its remaining descending connections. One should note that most studies involve a small number of patients with somewhat differently placed lesions, making it difficult to draw general conclusions. The weight of evidence suggests that different mechanisms may operate during recovery in patients with a similar functional and clinical picture.

Some observations support that use of **uncrossed motor pathways** contribute to recovery. For example, a group of patients with strokes affecting the pyramidal tract in the posterior part of the internal capsule (as verified with MRI) had initially severe paresis of the hand contralateral to the stroke, but they gradually recovered the ability to perform fractionate finger movements. In these patients, transcranial magnetic stimulation of the motor cortex of the undamaged hemisphere evoked movements of both hands—not just

of the contralateral hand, as in normal persons. Further, in some recovered stroke patients, involuntary movements of the unaffected fingers—**mirror movements**—accompany voluntary movements of the paretic fingers. Finally, a peculiar experiment of nature strongly suggests that the undamaged hemisphere participates in recovery in some patients. Thus, two patients with purely motor symptoms who were in good recovery from capsular hemiplegia suffered a second stroke in the internal capsule of the other hemisphere. As expected, their previously normal side now became paretic, but so did also the recovered side. This phenomenon is difficult to explain if we do not assume that the restitution had involved the use of motor pathways from the normal hemisphere to ipsilateral muscles.

Most patients do not exhibit mirror movements, however, and several studies found no evidence of contribution of descending pathways from the undamaged hemisphere (e.g., with the use TMS).

Examples of Substitution from Animal Experiments

Substitution implies that neuronal groups and pathways that normally participate only marginally gradually take over the responsibility for the execution of a task. Clinical recovery after brain damage resembles a long-term learning process, involving strengthening of specific synaptic connections by repeated use.

Some redirecting of impulse traffic can occur even immediately after the damage, however. When the arm region of the **motor cortex** in monkeys doing specific wrist movements is cooled, the movements immediately become slower and weaker. This is expected because the cooling inactivates many pyramidal tract neurons. Activity in the somatosensory cortex increases simultaneously, however, as if this region were prepared to take over immediately. If the somatosensory cortex is then also cooled, the monkey becomes paralytic and is unable to do the movements at all. Cooling of the somatosensory cortex alone has almost no effect on the movements and is not accompanied by increased activity in the motor cortex. These data suggest that the impulse traffic can be switched very rapidly from one path to another, so that command signals for movements are redirected to the somatosensory cortex. If the motor cortex were permanently inactivated, we might expect that establishment of new synapses would gradually strengthen the new impulse routes. Further experiments indicate that this is indeed what happens. After the first link of the disynaptic pathway from the **cerebellum** to the motor cortex is severed in monkeys (Fig. 11.2A), voluntary movements become jerky and uncoordinated. In 2 to 3 weeks, however, most symptoms disappear, and movements are carried out almost as before (Fig. 11.2B). Thus, considerable restitution has occurred, despite permanent damage to the connections.

If the somatosensory cortex is then destroyed, however, the original symptoms recur with full strength (Fig. 11.2C). After this second operation, recovery takes much longer and is incomplete. Obviously, during the first phase of recovery, the somatosensory cortex substituted the cerebellum regarding information to the motor cortex. This information is presumably carried by the numerous association connections from the somatosensory to the motor cortex. This assumption is supported by the electron microscopic demonstration of increased **synaptic density** in the motor cortex after recovery. These experiments thus suggest that new synapses can be established in the adult after brain damage, even in a neuronal group far removed from the site of injury.

Restitution Is a Learning Process

Regardless of whether long-term restitution is caused by other neuronal groups taking over a function or by collateral sprouting (or both), there is good reason to believe that the improvement is a result of a learning process, subject to the same rules that underlie learning in an intact nervous system. All learning is likely to involve changes in the properties of existing synapses, formation of new ones, and removal of inappropriate ones. Such use-dependent plasticity continues throughout life and is the nervous system's means of adapting to new and changing conditions, in both the body itself and the environment. The process is bound to take time and is probably slower in a brain deprived of many neurons. We know, from our own experience and from cognitive psychology, the importance of motivation and **focused, selective attention** for effective learning. An example of the importance of focused attention concerns infants and children with a squint (strabismus). Because the retinal images are different in the two eyes, the child attends to signals from one eye only to avoid

contradictory information (double vision). Some children solve the conflict by constantly using one eye and ignoring information from the other, whereas other children use the two eyes interchangeably. It turns out that in those always using the same eye, vision eventually becomes severely reduced in the eye that is not used. Those using their eyes interchangeably, however, retain normal vision in both eyes. The images reaching the cortex from the two eyes—even with a squint—are almost identical. The crucial factor is whether or not the information receives attention. In anesthetized animals, cells in the visual cortex can change their properties after exposure to light stimuli if they are coupled with stimulation of cell groups that are normally active during focused attention. Thus, the learning effect appears to require that two kinds of synaptic input converge at a neuron: one kind provides specific information; the other “tells” the neuron that this is relevant information that should be stored (see Fig. 4.10). As discussed earlier, the release of **monoamines** is probably a factor in such plastic changes.

During restitution, skills are re-learned more or less completely. Of course, the degree of recovery depends on the localization and size of the damaged region and on whether the neuronal loss occurs acutely or chronically (due to a degenerative process). With otherwise identical lesions, however, the recovery depends on well-known factors that contribute to successful learning in normal people. Among these, **motivation** and **amount of training** are of fundamental importance. Other factors, such as encouragement, feedback, and the learner's perception of progress, are crucial for maintaining a high motivation and would be especially important in patients with brain damage. Further, the prospects of recovery seem to be better when training starts very early after a stroke. This agrees with animal experiments showing that the plasticity is highest in a

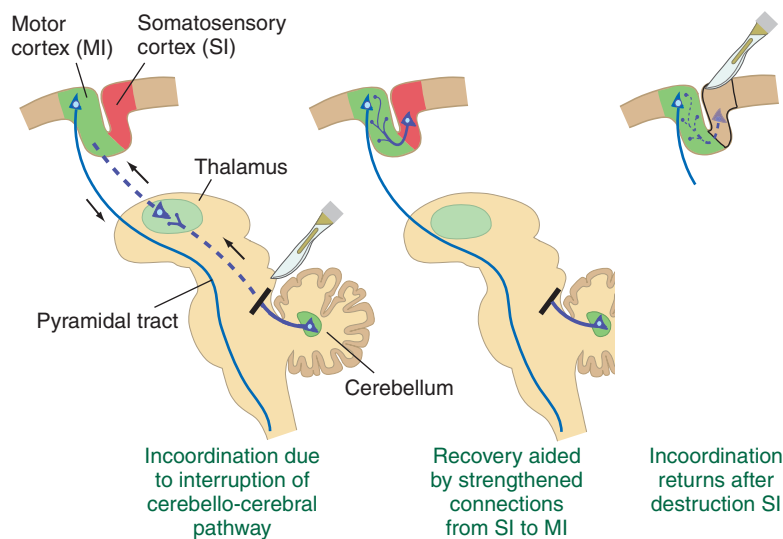


FIGURE 11.2 Example of recovery due to substitution. Recovery of coordinated movements after loss of connections from the cerebellum to the cerebral cortex is aided by strengthening of connections from the somatosensory cortex to the motor cortex. (Based on experiments in monkeys and cats by Mackel 1987, and Keller and coworkers 1990.)

limited period after the infliction of a brain lesion. Finally, although the learning curve (rate of recovery) is usually steepest during the first few months after a stroke, improvement can continue for several years if training is continued. One study even showed that a specific training program started 4 years after a stroke led to rapid improvement in hand function in patients with a deficit that had been stable for several years. These results were obtained by combining carefully planned training of the disabled arm with preventing use of the normal arm (by a sling). This method, called **constraint-induced movement therapy** (CI therapy), is based on results from experiments with constraining the normal arm in monkeys with unilateral lesions of the motor system. Apart from constraining the normal arm, the crucial factors for success seem to be a highly motivated patient, gradually increasing complexity of the training tasks so that the patient experiences progress regularly, and several hours of daily training.

RESTITUTION AFTER DAMAGE IN EARLY CHILDHOOD

The brain is certainly more plastic prenatally and during infancy than in adulthood. Thus, one might expect restitution to be more complete after early than after late brain damage. This holds only for some cases of early brain damage, however. In other cases, the consequences are in fact more severe.

Successful Restitution after Large Lesions of One Hemisphere

The most clear-cut examples of **successful recovery** after early brain damage concern infants with lesions—even very large ones—of one hemisphere.² In such cases, although the child remains hemiplegic, cognitive development may be virtually normal. The reason restitution is so successful is probably only partly that the individual neurons are more apt to send out new collaterals and remove others. Another factor could be as important: because in the young brain various regions, particularly of the cerebral cortex, are not yet fully engaged in the tasks for which they are destined, they are recruited more easily for other purposes if necessary. Thus, the right hemisphere of the brain may take over the processing of **language** if the left hemisphere (which normally does this) is damaged at an early age. In addition, prenatally and in infancy many connections are more widespread than later. During further development, synapses in some targets strengthen while synapses in other targets weaken or disappear. This is at least partly a use-dependent process, and in case of brain damage, persistence of the

widespread connections might probably aid restitution. As long as the damage is restricted to one hemisphere, size of the lesion does not seem to matter for success of restitution. The most crucial negative factor in children with such lesions is the occurrence of **seizures**, which inhibits cognitive development. Presumably, ongoing epileptic activity disturbs the use-dependent development of brain networks.

The Limits of Plasticity: Cerebral Palsy

Although recovery is remarkably good in some cases of early unilateral brain damage, this appears not to be the most common outcome after early brain damage. Indeed, prenatal and perinatal (at the time of birth) brain damage often gives more serious functional deficits than similar injuries acquired later in life. Thus, the child with early brain damage often presents with a mixture of symptoms, such as pareses, postural abnormalities, and involuntary movements. This is the case for children with **cerebral palsy**,³ suffering from the effects of lesions—such as birth-associated hypoxia, intrauterine infections, or genetic aberrations—that may have occurred at different developmental stages. Some children with cerebral palsy have additional cognitive impairments. We have discussed that during sensitive periods of development the brain is not only more plastic but also more vulnerable than later. Thus, a lesion acquired pre- or perinatally might interfere with a number of developmental processes. In addition, collateral sprouting at early stages may not necessarily be helpful. Thus, sprouts from nearby neurons may fill vacant synaptic sites, regardless of whether this is functionally meaningful or not. For example, in the cord sprouts from sensory fibers may innervate motor neurons that have lost their input from the cerebral cortex. Such “blind” sprouting in various parts of the brain and cord may perhaps contribute to the involuntary movements typical of many children with cerebral palsy. Further, long tracts that are already established are apparently not replaced if damaged. Thus, if the left motor cortex (sending fibers to the right spinal cord) is removed in monkeys shortly after birth, these monkeys never learn (with their right hand) the kinds of movements that depend on direct connections from the cerebral cortex to the cord. This is because these connections are already established at birth in monkeys (and in humans). Reports of apparent reestablishment of long fiber tracts after early injury are probably based on experiments in which the injury was inflicted before the axons from the relevant cell groups had reached their targets. In such instances, the outgrowing axons may

2 The situation appears to be less favorable when the lesions affect both hemispheres (even when the lesions are relatively small).

3 Cerebral palsy is used as an umbrella term for a number of clinically different neurological disorders that appear in infancy or early childhood and are dominated by disturbances of bodily movements and posture.

also innervate other cell groups than they normally do. This may be due to a lack of normal competition among outgrowing fiber tracts.

Finally, we will mention one other possible factor—although speculative—that may help to explain unsuccessful restitution after early brain damage. Prenatal brain damage occurs in a brain that is not yet geared to motivated, goal-directed behavior; establishment of functional networks is still mainly under genetic control. Thus, when emerging neuronal networks are disturbed there is presumably little pressure on functional restitution, in contrast to what is usually the case when damage occurs postnatally. This may allow haphazard plastic changes that are not tempered by a pressure to re-establish connectivity that is necessary for certain behaviors.

Early Damage of the Corticospinal (Pyramidal) Tract

Direct connections from the cerebral cortex to the spinal cord are necessary for the acquisition and performance of most skilled movements. Although corticospinal fibers reach the cord at the seventh gestational month, the development of synaptic connections in the cord and myelination of the corticospinal fibers go on at least until the child is 2 to 3 years old. Some connections are strengthened and others are removed during this period. Presumably, this represents a sensitive period for establishment of specific corticospinal connections. Initially, connections from each hemisphere are bilateral, but at the age of 2 the hand is controlled exclusively from the contralateral hemisphere; that is, the

relevant corticospinal fibers do all cross (in the medulla). This development is at least partly use-dependent and governed by the child's efforts to learn new skills.

A frequent kind of cerebral palsy is **hemiplegia** caused by a perinatal stroke. Typically, disturbed movement control and posture in these children appear not immediately but during the first 3 years of life, and often skills acquired early are later lost. This is probably due to plastic processes that are detrimental to normal motor development. Thus, it appears that any remaining corticospinal fibers from the damaged hemisphere are prevented from establishing synaptic connections in the cord by competition from the much stronger connections of the undamaged hemisphere. Indeed, the undamaged hemisphere retains and strengthens its ipsilateral connections to the cord, rather than being weakened as would normally occur (this is shown in animal experiments and indirectly by use of TMS in human infants). Children with cerebral palsy doing **mirror movements** express this: that is, voluntary movements with one hand are always accompanied by the same movement with the other hand. Why this abnormal innervation pattern is associated with poor function is not so obvious, however. In any case, animal experiments show that stimulation of the damaged hemisphere can rescue the remaining fibers and prevent the poor functional outcome. Further, it appears that in infants with cerebral palsy early specific training (constraint-induced movement therapy) of the impaired arm can improve the functional outcome. Probably, in such cases use-dependent plasticity serves to prevent the gradual loss of connections from the damaged hemisphere.

III

SENSORY SYSTEMS

THE first chapter in this part covers the basic features that are common to all sensory receptors, or sense organs. Such knowledge will make reading the following chapters easier. The next three chapters treat various aspects of the **somatosensory system**, which is primarily concerned with sensory information from the skin, joints, and muscles. Chapter 13 deals with the peripheral parts of the somatosensory system; that is, the sense organs and their connections into the central nervous system. Chapter 14 deals with the central pathways and neuronal groups concerned with processing of somatosensory information, while Chapter 15 discusses pain in some more depth. (Sensory information from the internal organs—visceral sensation—is dealt with mainly in Chapter 29.) In Chapter 16, we describe the **visual system**, from the peripheral receptors in the eye to the higher association areas of the cerebral cortex. Chapter 17 deals with the **auditory system** (hearing); Chapter 18 with the **vestibular system** (sense of equilibrium); and finally Chapter 19 discusses **olfaction** and **taste**.

We use the term **system** to indicate that each of these senses is mediated, at least in part, in different regions of the nervous system. In neurobiology, a system usually means a set of interconnected neuronal groups that share either one specific task, or several closely related tasks. Sensory systems are designed to capture, transmit, and process information about events in the body and in the environment, such as light hitting the eye or the

fullness of the bladder. We also use the term “system” for the parts of the nervous system that deal with tasks other than sensory ones. The motor system, for example, initiates appropriate movements and maintains bodily postures.

We use the term “system” rather loosely here: we do not imply that each system can be understood independently of the rest of the nervous system. Further, it is often arbitrary as to which system a particular neuronal group is said to belong. Thus, usually a cell group is used in the operations of several systems. Especially at higher levels of the brain, information from various systems converges. Cortical neuronal groups, for example, may be devoted to both sensory processing and motor preparation. **Convergence** of information from several sensory systems seems necessary for us to perceive that the different kinds of information actually concern the same phenomenon, whether it is in our own body or in the environment. How do we know, for example, that the round object with a rough surface we hold in the hand is the same as the orange we see with our eyes?

Sometimes the term system is misused, so that it confuses rather than clarifies; this happens when we lump structures about which we know too little, or cell groups that have such widespread connections that they do not belong to a particular system. The wish to simplify a complex reality—and the nervous system is indeed overwhelmingly complex—can sometimes become too strong.

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12 | Sensory Receptors in General

OVERVIEW

Sensory signals from nearly all parts of the body are transmitted to the central nervous system (CNS), bringing information about conditions in the various tissues and organs and in our surroundings. The structures where sensory signals originate are called **sense organs**, or **receptors**. The receptors are formed either by the terminal branches of an axon (the skin, joints, muscles, and internal organs) or by specialized sensory cells (the retina, taste buds, and inner ear) that transmit the message to nerve terminals. We are using the word “receptor” differently here than in earlier chapters where we applied it to molecules with specific binding properties. A **sensory unit** is a sensory neuron with all its ramifications in the periphery and in the CNS. The **receptive field** of the sensory neuron is the part of the body or the environment from which it samples information. These terms are applied to neurons at all levels of the sensory pathways, from the spinal ganglia to the cerebral cortex. Because of **convergence**, the receptive fields are larger at higher than at lower levels of the CNS.

Sensory receptors are built to respond preferentially to certain kinds of stimulus energy, which is called their **adequate stimulus** (mechanical, chemical, electromagnetic, and so forth). Regardless of the nature of the stimulus, it is translated into electric discharges conducted in axons; that is, the language of the nervous system. This process is called **transduction** and involves direct or indirect activation of specific cation channels. Activation produces a **receptor potential**, which is a graded depolarization of the receptor membrane (similar to a synaptic potential). We **classify** receptors by their adequate stimulus (e.g., mechanoreceptor, photoreceptor, and so forth), by their degree of adaptation, and by the site from which they collect information. For example, many receptors are rapidly adapting, meaning that they respond mainly to start and stop of a stimulus. Others are slowly adapting, and continue to signal a stationary stimulus. Receptors in the muscles and connective tissue are termed **proprioceptors**, while **exteroceptors** capture information from our surroundings (e.g., light, something touching the skin, and so on). **Enteroceptors** (interoceptors) inform about stimuli arising in the visceral organs. In general, contrasts and brief, unexpected stimuli are perceived much more easily than steady and familiar ones. This is partly because the majority of

receptors **adapt** rapidly, and partly because sensory information is heavily **censored** on its way to the cerebral cortex.

SENSORY UNITS AND THEIR RECEPTIVE FIELDS

Regardless of the structure of the receptor, a sensory neuron transmits the signal to the CNS. We use the term **sensory unit** for a sensory neuron with all its ramifications in the periphery and in the CNS (Figs. 12.1 and 12.2). Such primary sensory neurons have their cell bodies in ganglia close to the cord or the brain stem (similar neurons are found in the retina). The **receptive field** of the sensory neuron is the part of the body or the environment from which it samples information. The receptive field of a cutaneous sensory unit is a spot on the skin (Fig. 12.1; see also Fig. 13.10). For a sensory neuron in the retina, the receptive field is a particular spot on the retina, receiving light from a corresponding part of the visual field. We can determine the receptive field of a sensory unit by recording with a thin electrode the action potentials produced in an axon or a cell body, and then we can systematically explore the area from which it can be activated.

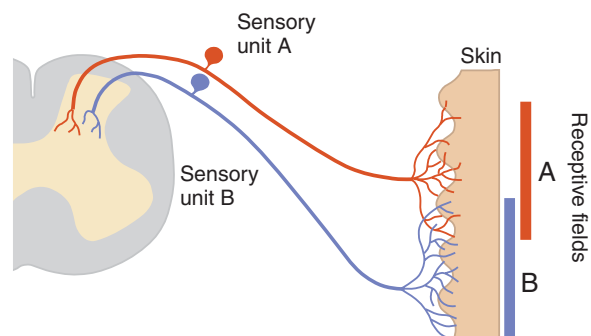


FIGURE 12.1 *Sensory units and receptive fields.* Simplified representation of two sensory units (A and B), which in this case are spinal ganglion cells. The peripheral process of each unit (fibers) ramifies in an area of the skin, which is the receptive field of this particular unit, and the receptive fields of the two units overlap. The density of terminal nerve fiber branches—endowed with receptor properties—is highest in the central part of the receptive field. Therefore, the threshold for eliciting action potentials is lower centrally than peripherally in the receptive field.

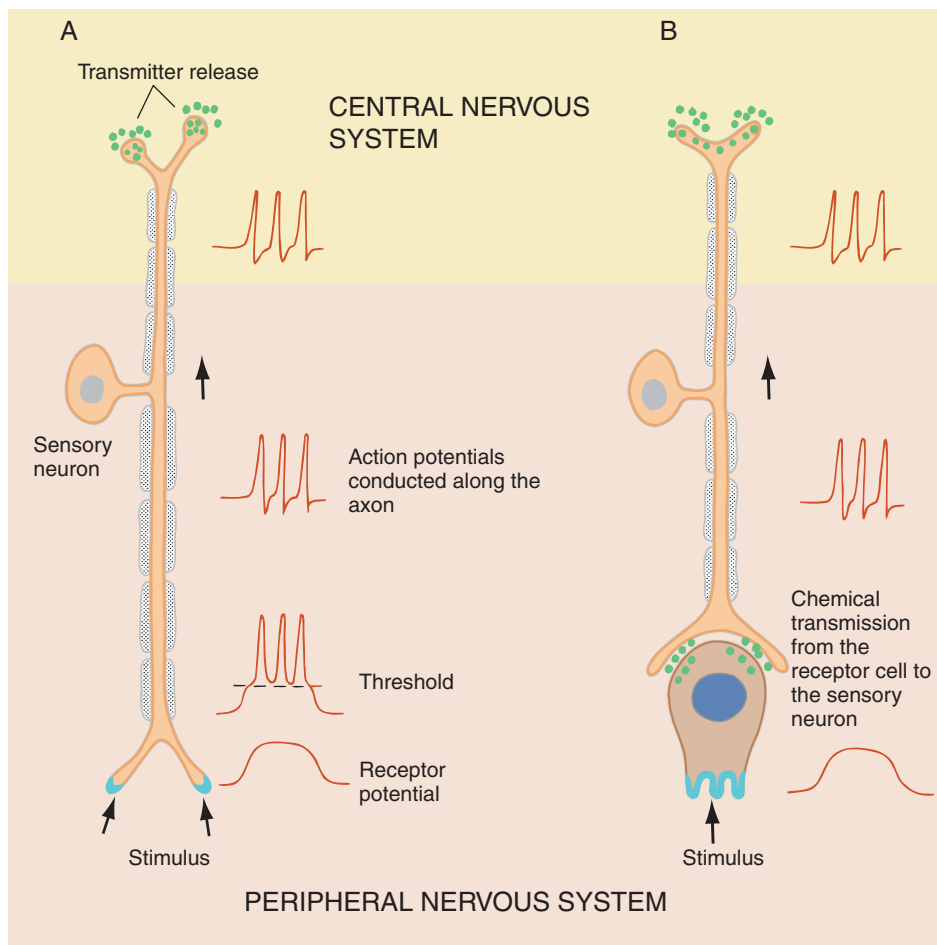


FIGURE 12.2 Two main kinds of sensory receptor. **Left:** The most common kind, found in most parts of the body. The receptor properties are in the terminal ramifications of a sensory axon that belongs to a pseudounipolar ganglion cell. **Right:** The kind of receptor found in the sense organs for taste, equilibrium, and hearing. The receptor properties are located on a sensory cell that transmits the signal to a ganglion cell. In both kinds of receptor, a stimulus produces a graded receptor potential. An action potential results in the ganglion cell to the left if the graded potential reaches the threshold. The sensory cell to the right does not produce an action potential but releases a chemical substance that depolarizes the terminals of the ganglion cell. In the CNS the signal is transmitted chemically to central sensory neurons.

Convergence Makes Receptive Fields Larger

We will come back to the sensory unit and receptive field and exemplify them in the following chapters. Here it is important to note that these terms apply to sensory neurons at any level of the pathways that lead from the receptors to the cerebral cortex. A neuron in the visual cortex, for example, reacts to light hitting only one particular part of the retina, which represents the receptive field of the neuron (see Fig. 7.8). At the cortical level the receptive field depends on convergence from several sensory units at lower levels (retina and thalamus, for example). Thus, receptive fields are usually larger for cortical sensory units than for those closer to the sense organ.

Receptive Fields Are Dynamic and Context-Dependent

One might perhaps expect that the receptive field of a sensory neuron would be a constant, hard-wired property. This is not the case, however; there is ample experimental evidence that receptive fields are **dynamic**. The size of a receptive field depends on the **context** of a stimulus; for example, focused **attention** reduces the receptive fields of neurons in the visual cortex. Such

rapid changes of properties are largely due to specific activation of inhibitory interneurons (see Fig. 13.3). Further, receptive fields can change because of specific **training** (learning) and of lesions that alter the sensory inputs to the CNS.

TRANSDUCTION: THE TRANSLATION OF STIMULI TO ACTION POTENTIALS

The task of the receptors is to respond to stimuli. Regardless of the nature of the stimulus, the receptor “translates” the stimulus to the language spoken by the nervous system—that is, electrical signals in the form of action potentials. We discuss in Chapter 4 how summation of many synaptic inputs evokes an action potential (depolarization to threshold) that is propagated along the axon.

Receptor Potentials

In receptors, the action potential arises near the terminal branches of the sensory neuron. This is true regardless of whether the stimulus acts directly on the terminal

branches (as in the skin; Fig. 12.1; see also Fig. 13.1) or indirectly via receptor cells (as in the retina and the inner ear; see Figs. 16.4 and 17.5). The stimulus alters the permeability of the receptor membrane, thus depolarizing the receptor. The effect of the stimulus on ion channel openings is **graded** like a synaptic potential (see Fig. 4.4). This graded change in the membrane potential is called a **receptor potential** (Fig. 12.2). The receptor potential that arises in the terminal branches spreads electrotonically in the proximal direction (toward the CNS, as in Fig. 12.2; see also Fig. 13.2) to the site where an action potential arises by opening voltage-gated Na^+ channels (provided the receptor potential is strong enough). The process is similar to synaptic activation of a neuron (in which the action potential usually arises where the axon leaves the cell body). In sensory cells, the stimulus does not evoke an action potential but leads to graded release of a neurotransmitter that depolarizes the terminals of the sensory cell (Fig. 12.2, right).

Transduction and TRP Channels

The mechanisms behind “translation,” or transduction, of a stimulus to a receptor potential are only partly known.

Although dozens of receptors and ion channels are involved, receptor potentials in a variety of receptors depend on activation of members of a large family of cation channels, collectively termed **transient receptor potential (TRP) channels** (mammals have more than 30 genes coding for TRP channels). Together, TRP channels exhibit a great variety of activation mechanisms and kinds of stimuli to which they respond.¹

Many receptors are depolarized by binding of a specific **chemical substance** to the membrane. The chemical acts either **directly** on TRP channel proteins (or other channels) or **indirectly** via **G protein-coupled receptors**. The latter can then alter the opening state of ion channels via intracellular second messengers (such as cyclic AMP or GMP). For example, light hitting the photoreceptors in the retina changes the photopigment, which leads to breakdown of cyclic GMP. Cyclic GMP keeps Na^+ channels open, so this breakdown leads to closure of Na^+ channels. Thus, unlike other receptors, the photoreceptors are hyperpolarized by their proper stimulus.

Many receptors are most easily activated by **mechanical forces**. Typical examples are receptors detecting pressure on the skin or rotation of a joint. Mechanical forces can most likely evoke a receptor potential in different ways. Stretching of the cell membrane may open TRP channels directly (Fig. 12.3A). More commonly, the mechanical force transmits to membrane-receptor

proteins via the rigid cytoskeleton (or similar structures outside the nerve terminal), which in turn either opens ion channels mechanically (Fig. 12.3B) or chemically via intracellular second messengers (Fig. 12.3C). Actions of external mechanical forces on membrane proteins have been particularly well elucidated in the sense organs for hearing and equilibrium (see Fig. 17.8).

Other receptors are particularly sensitive to the **temperature** in their surroundings. Change in temperature appears to alter the voltage sensitivity of specific subsets of TRP channels, thereby producing a receptor potential. Substances producing a cooling sensation when applied to the skin, like **menthol**, activate a TRP channel expressed in cold-sensitive receptors.

Finally, it should be noted that many receptors are sensitive to several kinds of stimuli; for example, mechanoreceptors may be sensitive to temperature.² Many receptors evoking pain when activated express various kinds of TRP channels, thus combining sensitivities to mechanical, thermal, and chemical stimuli.

PROPERTIES AND CLASSIFICATION OF RECEPTORS

Modality and Quality of Sensation

When a particular kind of receptor is stimulated with sufficient intensity to cause a consciously perceived sensation, we always get the same kind or **modality** of sensory experience—for example, light, touch, pressure, warmth, pain, or sound. The words modality and quality are both used to describe aspects of sensation but unfortunately somewhat differently by various authors. We use **quality** here to describe further the nature of a sensory modality; for example, pain is a sensory modality that may have a burning quality. That stimulation of a particular receptor always evokes a sensation of the same modality does not mean that the receptor necessarily has been subjected to its adequate stimulus. Thus, most receptors can respond to stimuli of other kinds (inadequate stimuli), although the **threshold** then is much higher for evoking a response. As an example, we can mention the perception of light upon a blow to the eye (mechanical stimulus of the photoreceptors rather than the normal light stimulus), and perception of sound upon chemical, rather than the normal mechanical, irritation of the receptors for hearing. The kind of perceived sensation—the modality or quality of sensation—is thus characteristic for each type of receptor (Müller’s law of specific nerve energies). Irritation of the **axon** leading from the receptor will also evoke the same sensory

¹ TRP channels are not restricted to sensory neurons but are expressed in virtually all tissues of the body.

² This is most likely the explanation why a cold object feels heavier than a warm (The “Weber deception” or “silver Thaler illusion”). Thus, some mechanosensitive skin receptors increase their firing when the temperature drops. These receptors appear to express both mechanosensitive and thermosensitive TRP channels. The biologic meaning—if any—of this dual sensitivity is unknown.

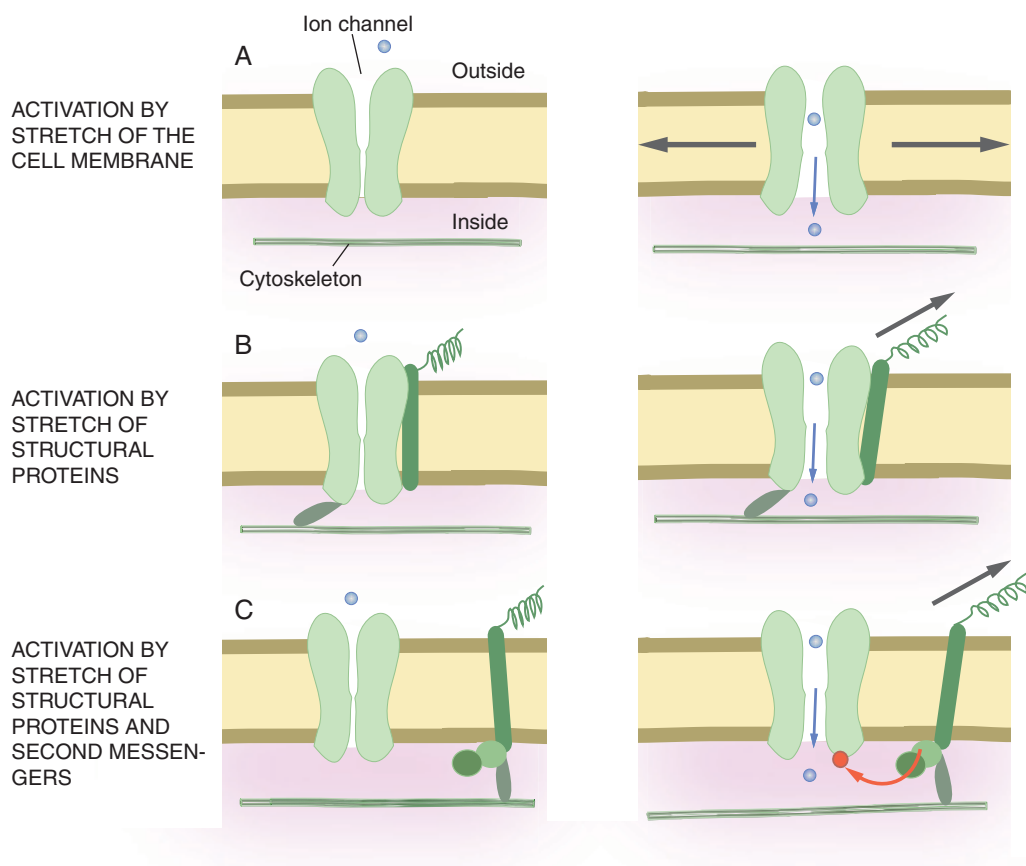


FIGURE 12.3 Possible means for mechanical stimuli to evoke receptor potentials via TRP ion channels. (Based on Christensen and Corey 2007.)

modality as when the receptor is stimulated by its adequate stimulus.

Adequate Stimulus

Most receptors are built to respond only or preferably to one kind of stimulus energy: mechanical, chemical, thermal, and so forth. The kind of stimulus to which the receptor responds most easily—that is, with the lowest threshold—is called the **adequate stimulus** for the receptor. We also say that the receptor is **specific** for this type of stimulus, whether it is mechanical, chemical, electromagnetic (light), or thermal (warmth, cold). As we shall see, each of these broad groups of stimuli is registered by receptors with different properties.

Receptors are classified according to the nature of their adequate stimulus, that is, their stimulus specificity. A large group of receptors, the **mechanoreceptors**, responds primarily to distortion of the tissue in which they lie and thus inform the CNS about mechanical stimuli. Such receptors are numerous in the skin, in deep tissues such as muscles and joint capsules, and in internal organs. Another large group of receptors,

chemoreceptors, responds primarily to certain chemical substances in the interstitial fluid surrounding the receptor. Many chemoreceptors respond to substances produced by or released from cells as a result of tissue damage and inflammation, regardless of the cause—mechanical trauma, burns, infection, and so forth. Other kinds of chemoreceptors are the receptors for taste and smell. Receptors in the retina responding to visible light are called **photoreceptors**. **Thermoreceptors** respond most easily to warming or cooling of the tissue in which they lie.

Threshold

Even when stimulated by their adequate stimulus, receptors vary enormously with regard to the strength of the stimulus needed to activate them; that is, they have different **thresholds** for activation. For example, in the retina, the rods are much more sensitive (have lower threshold) to light than the cones. Another example concerns mechanoreceptors, many of which have a **low threshold** and send signals even on the slightest touch of the skin or a just barely perceptible movement

of a joint. Other mechanoreceptors have a **high threshold** and require very strong stimulation to respond; we usually perceive such stimuli as painful.

Adaptation

Receptors differ in other ways, too. Many receptors send action potentials only when a stimulus starts (or stops). If the stimulus is continuous, this kind of receptor ceases to respond and thus provides information about changes in stimulation only. Such receptors are called **rapidly adapting**. When after a short time we cease noticing that something touches the skin, this is partly because so many of the receptors in the skin are rapidly adapting. Other receptors, however, continue responding (and thus sending action potentials) as long as the stimulus continues. Such receptors are called **slowly adapting** (or nonadapting). Receptors responsible for the sensation of pain exemplify slowly adapting receptors. It would not be appropriate if the body were to adapt to painful stimuli because these usually signal danger and threat of tissue damage. Receptors that signal the position of the body in space and the position of our bodily parts in relation to each other must also be slowly adapting; if not, we would lose this kind of information after a few seconds if no movement took place. That adaptation is a property of the receptors themselves can be verified by recording the action potentials from the sensory fibers supplying various kinds of receptors. For example, afferent fibers from receptors excited by warming of the skin stop sending signals if the same stimulus is maintained for some time, whereas afferent nerve fibers from sense organs in a muscle continue to send signals as long as the muscle is held in a stretched position.

Dynamic and Static Sensitivity

Many receptors respond more vigorously to rapid changes in the stimulus than to slow ones (they adapt rapidly). For example, rapid stretch of a muscle produces much higher firing frequency in the sensory nerve fibers leading from the muscle than when the same stretch is applied slowly (see Fig. 13.8). Such a receptor can therefore inform about the velocity of stretching, not just its magnitude. This property of a receptor is called **dynamic sensitivity**. A receptor that continues to produce action potentials with a constant frequency as long as the stimulus is constant (slowly adapting or nonadapting) is said to have **static sensitivity**. Often, one receptor has both kinds of sensitivity (see Figs. 13.8 and 16.8). The majority of receptors, however, have dynamic sensitivity and are rapidly adapting. This helps to explain the everyday experience that we are much more aware of changing stimuli than those that are constant—such as an insect moving on the skin or a moving light, as opposed to stationary ones.

Origin of Stimulus

Receptor classification can be based on the origin of the stimuli the receptors capture. For example, mechanoreceptors may—depending on their location—give information of very different events (even though all are mechanical). A low-threshold mechanoreceptor in the wall of the urinary bladder provides information about its degree of distension, mechanoreceptors in the inner ear inform us about sound (movement of air molecules), whereas mechanoreceptors around the root of a hair respond to the slightest bending of the hair.

We usually distinguish between **exteroceptors**, **proprioceptors**, and **enteroceptors**. **Exteroceptive** signals reach the body from the outside, from our environment. Most exteroceptors are located in the skin, whereas the receptors in the eye and the internal ear represent important special kinds of exteroceptors that respond to teleceptive signals. **Proprioceptive** signals originate in the body itself. The term is, however, restricted mostly to signals arising in the musculoskeletal system, including the joints. **Enteroceptive** signals arise from the internal (visceral) organs, such as the intestinal tract, lungs, and the heart.

Comprehensive Classification of a Receptor

By the descriptions discussed in the preceding text, we can classify receptors by the characteristics we want to emphasize. We can also give a complete description of the receptor by mentioning all the characteristic properties: for example, a proprioceptive, fast-adapting, low-threshold mechanoreceptor (informing about joint rotation), or an enteroceptive, slowly adapting chemoreceptor (in the wall of an artery informing about the oxygen tension of the blood).

RECEPTORS AND SUBJECTIVE SENSORY EXPERIENCE

Receptor Type and Quality of Sensation: An Uncertain Connection

The kind of conscious sensory experience it evokes is perhaps the most interesting among a receptor's many characteristics. How do I describe what I feel when a particular kind of receptor sends signals to the brain? We use the term pain receptor or **nociceptor** to describe receptors that, when stimulated, produce pain. Stimulation of **cold receptors** causes a feeling of coldness, stimulation of **warm receptors** gives a feeling of warmth, and so on. Only exceptionally is it possible, however, to know whether a sensory experience is evoked by stimulation of one receptor type only, or by the simultaneous activation of several kinds. The connection between the conscious sensory experience and the responsible receptors is therefore often uncertain. Because only human beings can

inform the observer directly of what they feel, animal experiments alone cannot resolve the question of the relationship between receptors and conscious sensations. Moreover, the very same receptors that can be examined physiologically cannot be examined anatomically in human beings. Important insight has nevertheless emerged from correlation of observations obtained in animals with psychophysical observation in humans (see Chapter 13, under “Microneurographic Studies of Human Skin Receptors”).

In most cases, our conscious sensory experiences are due to signals from several kinds of receptor. The brain interprets a barrage of signals and the context in which they arise, and provides us with a unitary sensation. The sensation of taste is an example: it depends not only on signals from taste receptors (on the tongue) but also on signals from olfactory receptors (with some contribution from mechanoreceptors and thermoreceptors in the mouth).

The Brain Does Not Receive “True” Information

Not all signals reaching the CNS from the receptors are consciously perceived. In particular, interoceptive signals are mostly processed only at a subconscious level. For signals from all kinds of receptors, however, a considerable selection and suppression of signals take place at all levels of the sensory pathways in the spinal cord and the brain, to leave out “irrelevant” information. At the same time, “relevant” signals are usually enhanced in the CNS (see also Chapter 13, under “Inhibitory Interneurons Improve the Discriminative Sensation”). We mentioned that receptive fields of sensory cells expand or shrink in a context-dependent manner, and that most receptors respond more strongly to changing than to constant stimuli. Therefore, sensory information is censored and does not provide the brain with an objective representation of the physical world (see Fig. 16.9). Usually, such weighting of sensory signals is advantageous because it ensures that the most behaviorally relevant information is prioritized while irrelevant or less important signals are suppressed. An example concerns our need to be able to distinguish sensory signals that we produce ourselves by voluntary

movements from signals that arise from external perturbations. As rule, our self-produced sensory signals are inhibited while the unforeseen external ones are enhanced.

Central Analysis

We cannot explain how it happens that action potentials, which are of the same kind in all nerve fibers, evoke entirely different conscious sensations, depending on where the stimulus arises. One prerequisite is that different kinds of sensory information are analyzed by different neurons (i.e., neurons located in anatomically distinct parts of the CNS). A complete mixing of information from different receptor types would not be compatible with our discriminative abilities. It is equally clear, however, that there must be ways in which the brain can bring the different kinds of information together, making meaning and unity out of the innumerable bits and pieces of sensory information. Such a synthesis may not necessarily be due to convergence of all relevant information onto a single cell group. Rather, we now believe the final processing to depend on extensive interconnections among cortical areas dealing with different aspects of sensory information (these aspects are treated further in Chapter 16, under “How Are Data from Different Visual Areas Integrated?” and Chapter 34, under “Parietal Association Areas”).

In an elegant study, de Lafuente and Romo (2005) trained monkeys to respond to cutaneous stimuli that were barely detectable, while at the same time recording neuronal activity in the cerebral cortex. It turned out that whenever the monkey responded to the stimulus (as a sign of perception), there was activity among neurons in the premotor cortex of frontal lobe as well as in the somatosensory cortex. Activity in the somatosensory cortex alone was apparently not sufficient for perception. On the other hand, there was frontal activation in instances when the monkey (erroneously) responded without the stimulus being strong enough to evoke activity in the somatosensory cortex. This example serves further to emphasize that our conscious sensations are not hard-wired to the signals from sensory receptors, but depends critically on central analysis.

Peripheral Parts of the Somatosensory System

OVERVIEW

The term **somatosensory** includes, strictly speaking, all sensations pertaining to the body (*soma*). It is therefore more comprehensive than the significance commonly assigned to it—namely, sensory information specifically from **skin**, **joints**, and **muscles** only. In this chapter, we discuss receptors (sense organs) located in the skin and in the musculoskeletal system and the neurons leading from them into the central nervous system (CNS). Receptors in the skin and in deep tissues are either **free** or **encapsulated**. The terminal ramifications in encapsulated receptors are surrounded by connective tissue cells, serving as a filter ensuring that only certain kinds of stimuli reach the axon terminal. We distinguish between **exteroceptors**, located in the skin, and **proprioceptors** in muscles and connective tissue around the joints. Regardless of their location, somatosensory receptors can be classified by their adequate stimulus as **mechanoreceptors**, **thermoreceptors**, and **chemoreceptors**.

High-threshold receptors usually signal impending tissue damage, and are termed **nociceptors**. They are all free nerve endings, but differ with regard to stimulus specificity. Some respond only to intense mechanical stimuli, others to chemical (inflammatory) substances, while others detect extreme heat or coldness. Many nociceptors respond to several kinds of stimuli (**polymodal nociceptors**), while others do not respond to “normal” pain-provoking stimuli but require prolonged stimulation to become active (**silent nociceptors**). In addition to signaling impending acute tissue damage, nociceptors probably play an important role in monitoring the composition of the tissue fluid and thus contribute to bodily **homeostasis**.

Thermoreceptors respond to even very small changes of the temperature of their surroundings (especially in the skin). They are much less sensitive to steady-state temperature. Cold receptors detect cooling below the normal skin temperature, while warm receptors detect a rise in temperature.

Low-threshold mechanoreceptors in the skin and in the deep tissues share many physiological properties. In general, they signal deformation of the tissue in which they reside. Their capsular elements and relation to surrounding tissue components—such as collagen fibrils

or muscle fibers—determine the kind of stimulus they respond to with the lowest threshold. Further, capsular elements and membrane receptor proteins determine whether the receptor is **rapidly** or **slowly adapting**.

Cutaneous low-threshold mechanoreceptors are of four types: **Meissner corpuscles** reside in the dermal papillae of glabrous skin (e.g., on fingertips) and respond to the slightest impression on the skin. They are rapidly adapting, thus signaling only when something touches (or leaves) the skin—not when the skin remains impressed. Like Meissner corpuscles, **Merkel disks** respond to small impressions of the skin but are slowly adapting. **Ruffini corpuscles** are located in the dermis and respond to stretching of the skin. They are slowly adapting, and probably signal the steady tension (important, e.g., when holding objects). **Pacinian corpuscles** are large, lamellate structures located on the transition between the dermis and the subcutaneous tissue. They are extremely rapidly adapting and are well suited to signal vibration (>100 Hz). The receptive fields of Meissner corpuscles and Merkel disks are small (especially on distal parts of the extremities), making these receptors of crucial importance for **discriminative sensation**.

Among **proprioceptive** low-threshold mechanoreceptors, the **muscle spindles** are the most elaborate. They consist of a small bundle of thin, so-called **intrafusal** muscle fibers encircled by sensory nerve endings. Although each muscle spindle is much shorter than the muscle in which it lies, it is connected with connective tissue strands to both tendons of the muscle. Thus, when the muscle elongates, the muscle spindle is stretched and the sensory endings are depolarized. The muscle-spindle sensory endings have both dynamic and static sensitivity, making them suited to record the actual muscle length at any time, length change, and direction and velocity of change. Signals from muscle spindles contribute to reflex control of movements and to **kinesthesia**—that is, our conscious perception of joint positions and movements. **Tendon organs** are located at the musculotendinous junction, and measure the force of muscle contraction. **Joint receptors** are located in the fibrous joint capsule and in ligaments around the joints. They can signal movements and steady positions but their contributions to movement control and kinesthesia are not well understood.

Dorsal root fibers conducting from somatosensory receptors are classified according to **conduction velocity** into myelinated A fibers and unmyelinated C fibers. The thickest A fibers conduct from low-threshold mechanoreceptors, whereas thin A fibers (A δ) and C fibers conduct from nociceptors and thermoreceptors. The thin (A δ and C) and thick dorsal root fibers end almost completely separated in the cord. Probably all primary sensory neurons release a classical transmitter with fast synaptic actions in the spinal cord. Probably all primary sensory neurons release a classical **neurotransmitter** with fast synaptic actions in the spinal cord. In addition, many contain several **neuropeptides**, such as substance P and others. Apparently, release of neuropeptides from the peripheral branches of sensory neurons contributes to local inflammation in several diseases (e.g., arthritis, asthma, and migraine).

The ventral branches of the spinal nerves form **plexuses** supplying the arms and the legs. Each nerve emerging from these plexuses contains sensory and motor fibers arising from several segments of the spinal cord. In the peripheral distribution of the fibers, however, the **segmental** origin of the fibers is retained. Thus, sensory fibers of one dorsal root supply a distinct part of the skin. The area of the skin supplied with sensory fibers from one spinal segment is called a **dermatome**.

EXTEROCEPTORS: CUTANEOUS SENSATION

The skin and the subcutaneous tissue contain receptors that respond to stimuli from our surroundings. Among the almost indefinitely varied sensory experiences that can be evoked from the skin, we usually distinguish only a few, regarded as the basic modalities: **touch**, **pressure**, **heat**, **cold**, and **pain**. There is indirect evidence that each of these sensations can be evoked by stimulation of one receptor type. Yet it is important to realize that terms such as “pain,” “cold,” and “touch” refer to the quality of our subjective sensory experience evoked by certain kinds of stimuli. Thus, a classification of sensations on this basis does not enable us to conclude that for each subjective quality of sensation there exists one particular kind of receptor.

Sensations—such as itch, tickle, dampness, or dryness; the texture of surfaces; and the firmness or softness of objects—are thought to arise from the simultaneous stimulation of several kinds of receptors in the skin and also in deeper tissues. As discussed in Chapter 12, our conscious sensory experience is the result of an interpretation in the brain of information from a multitude of receptors.

Functionally, skin receptors—like receptors in other parts of the body—can be classified by their adequate stimulus as **mechanoreceptors**, **thermoreceptors**, and **chemoreceptors**.

Free and Encapsulated Receptors

We conveniently subdivide receptors in the skin (and in deep tissues) on a structural basis into **free** and **encapsulated**, although there are numerous transitional forms. In encapsulated receptors, the terminal ramifications of the sensory axon are surrounded by a specialized capsule-like structure of connective tissue cells (Fig. 13.1A, C, and D), whereas such a structure is lacking around the free receptors or endings (Fig. 13.1E). The encapsulation serves as a **filter**, so that only certain kinds of stimuli reach the axon terminal inside. **Schwann cells** cover the axonal ramifications of the free receptors, except at their tips (Fig. 13.1E), where their receptor properties presumably reside. Free receptors are the most numerous and widespread, being present in virtually all parts of the body. In the skin, free receptors are particularly numerous in the upper parts of the dermis, and they even extend for a short distance between the cells of the deeper layers of the epidermis.

Free receptors that are structurally indistinguishable can nevertheless have different adequate stimuli. The **cornea** of the eye, for example, contains only free nerve endings but functionally there are at least four different receptors. Thus, anatomically identical receptors can differ functionally as a result of their expression of different repertoires of membrane channels and receptor molecules.

Nociceptors

Receptors that on activation evoke a sensation of pain are termed nociceptors. A more precise, **physiological definition** would be: a receptor that is activated by stimuli that produce tissue damage, or would do so if the stimulus continued. In Chapter 15, we discuss the relationship between nociceptor stimulation and the experience of pain. In the skin and, as far as we know, in all other tissues from which painful sensations can be evoked, nociceptors are **free nerve endings**. It is characteristic that most stimuli we experience as painful are so intense that they produce tissue damage or will do so if the stimulus is continued. Functionally, skin nociceptors are of three main types. One type responds to intense mechanical stimulation only (such as pinching, cutting, and stretching), and is therefore termed a **high-threshold mechanoreceptor**. The other is also activated by intense mechanical stimuli but in addition, by intense **warming** of the skin (above 45°C)¹ and by **chemical**

¹ Heat activates nociceptors by opening the nonselective cation channel TRPV1 (and apparently also other varieties of the TRPV channel). Interestingly, **capsaicin**, responsible for the pungency of hot peppers, activates the heat-sensitive channel. On continued presence, however, capsaicin desensitizes the receptor thus alleviating ongoing pain. Drugs that selectively block the TRPV1 channel have been tested in animals and found effective in alleviating inflammatory pain but also causing a rise in body temperature. Thus, the TRPV1 channel apparently contributes not only to thermal and chemical nociception but also to control of body temperature.

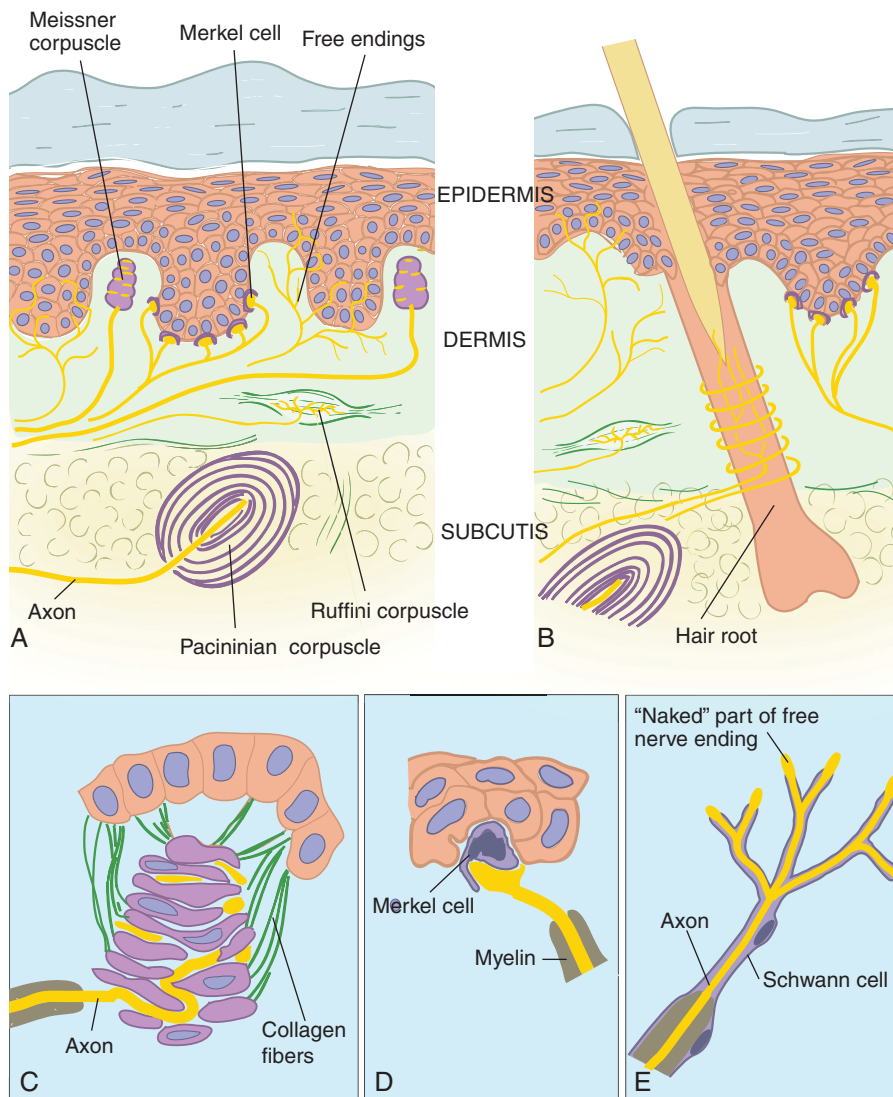


FIGURE 13.1 *Cutaneous receptors.* Schematic of receptors as they appear in sections through the skin. Sensory nerve fibers and receptors in (A) glabrous skin (palms of the hands and soles of the feet) and (B) hairy skin. Nerve endings in hairy skin wind around the hair follicles and are activated by the slightest bending of the hair. C: Meissner corpuscle (from glabrous skin). The axon (red) follows

a tortuous course between flat, specialized connective tissue cells. The whole sense organ is anchored to the epidermis with thin collagen fibers. D: A disk of Merkel (present in both glabrous and hairy skin). The axonal terminal is closely apposed to a Merkel cell in the epidermal cell layer. E: Free nerve endings are covered by Schwann cells except at their tips, where the receptor properties reside.

substances that are liberated by tissue damage and inflammation. Because such receptors can be activated by different sensory modalities, they are termed **poly-modal nociceptors**. In addition, recent studies indicate that many nociceptors are purely sensitive to chemical substances released in inflamed tissue. Since such receptors are unresponsive to most nociceptive stimuli used in animal experiments, they are termed **silent nociceptors**. Silent nociceptors typically require stimulation for 10 to 20 minutes to become active; thereafter, however, they may continue firing for hours. They are present in skin, muscles, and visceral organs and may constitute about one-third of all nociceptors. They appear particularly

suited to communicate about disease processes in the tissues, and are probably important for communication between the immune system and the brain.

Many substances can excite nociceptors, and specific membrane receptors and ion channels have been identified for some. **ATP**, for example, excites nociceptors by binding to purinoceptors (Fig. 13.2). Because ATP is normally present only intracellularly, its extracellular occurrence is an unequivocal sign of cellular damage. The peptide **bradykinin**, which is produced by the release of proteolytic enzymes from damaged cells, acts on specific membrane receptors in nociceptors. Several other mediators of **inflammation**—such as prostaglandins,

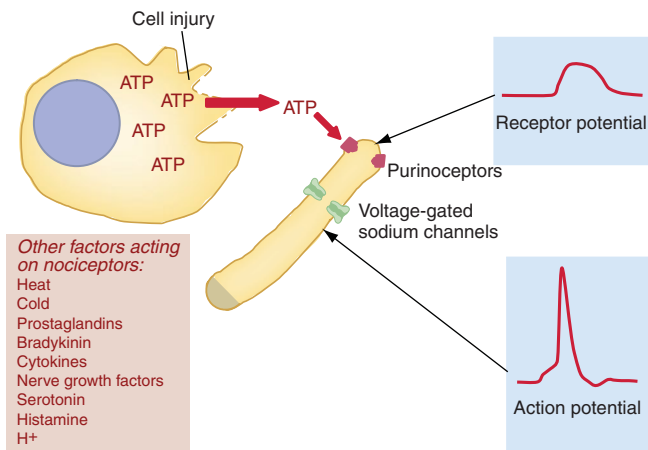


FIGURE 13.2 *Nociceptor activation.* ATP is one among many substances that activate nociceptors. Because the extracellular ATP concentration is extremely low normally, ATP is a very sensitive signal of impending cell damage. Binding of ATP to specific receptors depolarizes the nerve terminal (receptor potential). If the receptor potential reaches threshold an action potential arises due to opening of voltage-gated sodium channels. The action potential is conducted to the CNS. Some other means of activating nociceptors are listed.

histamine, serotonin, substance P, and adenosine—may contribute to nociceptor activation and sensitization. H^+ ions (pH < 6) activate nociceptors effectively and, in addition, seem to increase their responses to inflammatory substances.

A characteristic feature of nociceptors is their tendency to be **sensitized** by prolonged stimulation (this is the opposite phenomenon of adaptation). Sensitization is due to accumulation of inflammatory products that up-regulate voltage-gated Na^+ channels and transient receptor potential (TRP) channels expressed in sensory neurons. Sensitization partly explains why even normally non-noxious stimuli (such as touching the skin) may be felt as painful when the skin is inflamed. A condition of abnormal intensity of pain compared to the strength of the stimulus is called **hyperalgesia**. As discussed later in this chapter, hyperalgesia may also be caused by altered properties of neurons in the CNS, especially in the spinal dorsal horn. The term **allodynia** is used when innocuous stimuli, such as light touch, evoke intense pain.

Signals from nociceptors are conducted in thin myelinated (A δ) and unmyelinated (C) fibers. This is further discussed later in this chapter, under “Classification of Dorsal Root Fibers in Accordance with Their Thickness.”

Nociceptors, Voltage-Gated Sodium Channels, and Inherited Channelopathies

Sensory neurons express a multitude of ion channels and receptors, among them several kinds of voltage-gated sodium (Na^+) channels. Such channels are expressed in all neurons and are necessary for excitability and impulse

conduction. Ten voltage-gated sodium channels have been identified, sharing a basic structure but with somewhat different properties and distribution. Sensory neurons express several kinds, and some of these occur primarily in small nociceptive spinal ganglion cells. Nerve injury as well as inflammation cause rapid changes in the expression of several voltage-gated sodium channels, thus rendering the ganglion cells hyperexcitable (as mentioned earlier in relation to hyperalgesia).

One channel— $Na_v1.7$ —appears to be particularly important for the generation of action potentials in axons leading from nociceptors. Rare inherited diseases caused by mutations of the gene (*SCN9A*) coding for $Na_v1.7$ shed light on the function of this channel. Two gain-of-function mutations cause attacks of intense pain but with different mechanisms. Patients with **paroxysmal extreme pain disorder** suffer from attacks of pain in the eyes, jaw, and rectum, apparently caused by incomplete inactivation of the $Na_v1.7$ channel. Patients with **erythromelalgia** (primary erythermalgia) experience burning pain and redness in the hands and feet. The mutation in the latter patients cause the channel to open too easily (lowered threshold). A third variety of inherited $Na_v1.7$ channelopathy was quite recently found among members of a Pakistani family. Those afflicted show complete absence of pain sensitivity and suffer serious injuries as they grow up. The mutation in these patients appears to cause a loss of function of the $Na_v1.7$ channel, resulting in abolished impulse traffic from nociceptors.

Other voltage-gated sodium channels, expressed in nociceptive ganglion cells, also appear to be involved in clinical pain conditions. Great efforts are now being made to develop drugs that can block specific voltage-gated sodium channels. Nonspecific blockers, such as **lidocaine**, have long been used for local anesthesia.

Silent Nociceptors, the Immune System, Homeostasis, and Sickness Behavior

Since mediators released in inflamed tissues activate them, silent nociceptors most likely contribute to the pain that follows tissue damage. Nevertheless, in contrast to other nociceptors they may not function primarily as a warning system for impending tissue damage but, rather, play a long-term role in evaluating the status of the tissue microenvironment. Thus, they would play a role in bodily **homeostasis**. Signals from silent nociceptors may be particularly important to modulate the activity of the **immune system**. Thus, we know that the nervous system can influence the properties of the cells of the immune system, as we shall return to in Chapter 30. To do this, the nervous system needs information from the “battle-field” of the immune system. Silent nociceptors seem well suited to carry out this task, as substances released from leukocytes activate them. We know, for example,

that cytokines released in the gastric mucosa stimulate nerve fibers in the **vagus nerve**. This produces **sickness behavior** in rats: they move less around, lose appetite, are uninterested in their surroundings, and so forth. In addition, cytokines in the bloodstream can reach neurons in certain parts of the brain, either by passing the blood–brain barrier or by acting on places devoid of a blood–brain barrier. It seems likely that signals from silent nociceptors—not only in the skin but also in deep tissues and visceral organs—contribute to “how we feel,” or the subjective feeling of the physiological state of the body.

Thermoreceptors

Free endings of thin sensory nerve fibers are responsible for the perception of heat and cold. Although warm and cold receptors look the same, they express different kinds of TRP channels. **Cold receptors** respond with an increase of firing frequency to cooling of the skin below the normal temperature (about 32°C). They stop responding, however, at very low skin temperatures. Surprisingly, cold receptors can also be excited by skin temperatures above 45°C. This explains why a hot shower may feel cold at the beginning (until we feel pain, as mentioned previously, temperatures above 45°C excite polymodal nociceptors). This phenomenon is called **paradoxical cold** (sensation). Signals from cold receptors are conducted in thin myelinated (A δ) fibers. **Warm receptors** respond to warming of the skin above the normal temperature up to about 45°C. The signals are conducted in unmyelinated (C) fibers.

The **adequate stimulus** for thermoreceptors is the temperature of the tissue surrounding them or, rather, changes in the temperature. The receptors send action potentials with a relatively low frequency at a steady temperature, whereas a small change in the temperature elicits a marked change in the firing frequency. A heat receptor, for example, fires at constant room temperature with a low frequency, but warming the skin even slightly increases in the firing rate. The response is particularly brisk if the warming happens rapidly (thus, we perceive lukewarm water as hot if the hand is cold when put into it). A change in skin temperature of 0.2°C is sufficient to cause a marked change in firing rate from a thermoreceptor. Thus, the thermoreceptors do not give an objective measure of the actual skin temperature but, rather, signal **changes** that may be significant in our adjustment to the environment (to keep the body temperature constant). Thus, they are important for the maintenance of bodily **homeostasis**.

Mechanoreceptors of the Skin

The study of receptors for pain and temperature sensation is more difficult than the study of mechanoreceptors. These are particularly well studied, therefore, and among

them, we know most about the low-threshold mechanoreceptors. The following account mainly deals with low-threshold mechanoreceptors of the skin. High-threshold mechanoreceptors in the skin are nociceptors and do not differ significantly from such receptors in muscles and around joints. As far as we know, they are always free nerve endings (Fig. 13.1E).

There are several kinds of **low-threshold mechanoreceptors** in the skin, ranging from free receptors to those with an elaborate capsule. Some adapt slowly or not at all; others adapt very rapidly. For example, receptors found close to the roots of **hairs** (Fig. 13.1B) are rapidly adapting. They are activated by even the slightest bending of a hair, as can easily be verified by touching the hairs on the back of one’s own hand. If the hair is held in the new position, however, the sensation disappears immediately. Unmyelinated afferent fibers from hairy skin that signal light touch may be of special importance for mediating **emotional** aspects of touch (rather than precise, discriminative information).²

Although the thick, **glabrous skin** on the palm of the hand and on the sole of the foot lacks hair, the elaborate encapsulated receptors are particularly abundant at these locations and are obviously related to the superior sensory abilities of these parts—the fingers, in particular (Table 13.1). One such receptor is the **Meissner corpuscle**, which mediates information about touch. Meissner corpuscles are small oval bodies located in the dermal papillae just beneath the epidermis—in fact, as close to the surface of the skin as possible without being directly exposed (Fig. 13.1A and C). Several axons approach the corpuscle and follow a tortuous course inside the capsule between the lamellae formed by connective tissue cells.³ Meissner corpuscles respond by sending action potentials even when indenting only a few micrometers the skin overlying the receptor. If the skin is kept indented, however, the receptor stops sending action potentials. On release of the pressure, a few action potentials are again elicited. Meissner corpuscles are thus **rapidly adapting** and obviously have a low threshold for their adequate stimulus. These corpuscles are presumably well suited to, among other things, signal **direction** and **velocity** of objects moving on the skin.

2 A woman with selective loss of large-diameter myelinated sensory fibers provided an opportunity to study the properties of **low-threshold C-fibers** (Olausson et al. 2002). Light touch (stroking with a soft brush) applied to the back of the hand was felt as a very faint and diffuse but pleasant pressure (no sensation was evoked by brushing the palm of the hand, corresponding to the lack of unmyelinated low-threshold afferents from glabrous skin). Interestingly, functional magnetic resonance imaging (fMRI) showed activation of the insula known to be related to affective aspects of sensation but not in SI that is responsible for the discriminative aspects.

3 In addition to a thick myelinated axon, the Meissner corpuscle receives thin unmyelinated axons. The latter contain neuropeptides and express receptors typical of nociceptors. So far, however, there is no direct evidence of a contribution from Meissner corpuscles in nociceptive signaling.

TABLE 13.1

RECEPTIVE FIELDS	ADAPTATION	
	Rapid	Slow
Small	Meissner corpuscles: Glabrous skin, in dermal papillae Touch: moving stimuli Vibration < 100 Hz	Merkel disks: Close to the epithelium Touch (judging form & surface of objects)
Large	Pacinian corpuscles: Border dermis-subcutis; Vibration > 100 Hz	Ruffini corpuscles: Dermis, parallel to the skin surface; stretch of the skin (friction)

Ruffini corpuscles are also low-threshold mechanoreceptors and are **slowly adapting**. They consist of a bundle of collagen fibrils with a sensory axon branching between the fibrils (Fig. 13.1A). The collagen fibrils connect with those in the dermis, and stretching of the skin in the direction of the fibrils is the adequate stimulus for the receptor. Stretching the skin tightens the fibrils, which, in turn, leads to deformation and depolarization of the axonal ramifications, thus producing action potentials in the afferent fiber. It is therefore assumed that Ruffini corpuscles function as low-threshold **stretch receptors** of the skin, informing us about the magnitude and direction of stretch.

Another kind of **slowly adapting**, low-threshold mechanoreceptor in the skin is the **Merkel disk** (Fig. 13.1A and D), present particularly on the distal parts of the extremities, the lips, and the external genitals. An axon ends in close contact with a large epithelial cell in the basal layer of the epidermis. Even after several minutes of constant pressure on the skin overlying the Merkel disks, the receptor continues to send action potentials at about the same rate.

A final type of low-threshold mechanoreceptor, **Pacinian corpuscles** (Fig. 13.1A), are found at the junction between the dermis and the subcutaneous layer; they are also present at other locations, such as in the mesenteries, vessel walls, joint capsules, and in the periosteum. Pacinian corpuscles are large (up to 4 mm long) ovoid bodies, which can be seen macroscopically at dissection. A thick axon is surrounded by numerous lamellae, which are formed by a special kind of connective tissue cell. Between the cellular lamellae there are fluid-filled spaces. Pacinian corpuscles are very **rapidly adapting**, eliciting only one or two action potentials in the afferent fiber at the onset of indentation of the skin. The adequate stimulus is therefore extremely rapid indentation of the skin. In practice, this is achieved by **vibration** with a frequency of 100 to 400 Hz. If a vibrating probe

is put in contact with the skin, the frequency of action potentials in the afferent fiber follows closely the frequency of vibration. Vibration with a frequency below 100 Hz appears to be signaled by Meissner corpuscles.

Itch and Tickle

Itching is a peculiar, unpleasant sensation evoked by stimulation of free nerve endings in superficial parts of the skin and mucous membranes. The signals are conducted in unmyelinated sensory fibers to the spinal cord. From the cord to higher levels, the signals follow the pathways for pain, because cutting these also abolishes the sensation of itching. (In fact, we know that weakly painful stimuli, such as scratching the skin, suppress the sensation of itching.) Further, liberation of **histamine** in the skin, for example, from mast cells in allergic reactions, evokes itching. The leaves of stinging nettle (*Urtica dioica*) provoke intense itching because they contain histamine. Microneurographic studies in humans have identified a subgroup of peripheral sensory units (with unmyelinated fibers) that react vigorously to histamine and produce itching when stimulated. These units have been regarded as itch-specific (even though capsaicin and inflammatory substances such as prostaglandin E and bradykinin also activate them). Further, some units in the dorsal horn (lamina 1) respond primarily when histamine is applied in their receptive fields. Histamine appears to evoke activity in the same parts of the cerebral cortex as painful stimuli do, as shown via functional magnetic resonance imaging (fMRI). Nevertheless, animal experiments suggest that itch involves neuronal groups—from the cord to the cortex—that are at least partly separate from those treating noxious stimuli. Thus, the sensation of itch may be served by a specific subdivision of the neural system for pain. Itching is a symptom in several diseases (cancer, metabolic disorders, skin diseases, and others). Histamine does not seem to be involved in such cases, however.

Tickling is another peculiar sensory phenomenon. It is evoked by stimulation of low-threshold mechanoreceptors, but we do not know which subgroup is involved. The sensation of tickling is strongly influenced by the **context** in which it is evoked: the same stimulus may be experienced as tickling in one situation and merely as light touch in another. For example, we know that we are unable to tickle ourselves, probably because the brain can easily distinguish sensory signals produced by our own actions and those coming from other sources. We also know that our emotional state influences whether or not a stimulus is felt as tickling. This example may serve to emphasize a point we will return to repeatedly: the sensory messages sent from the receptors are subject to extensive processing before a conscious sensation is produced.

What Information Is Signaled by Low-Threshold Cutaneous Mechanoreceptors?

Together, the four types of low-threshold mechanoreceptors described in the preceding text are thought to mediate the different qualities of our sense of touch and pressure, which are so well developed in glabrous skin (fingers, toes, and lips). One important aspect is the ability to judge the speed and direction of a moving object in contact with the skin, as well as the friction between them. Thus, we may perceive quickly that an object is slipping from our grip and judge from the friction the force needed to stop the movement. Two of the receptor types, Merkel disks and Ruffini corpuscles, are slowly adapting and, as long as the stimulus lasts, continue to provide information about slight pressure and stretching of the skin, respectively (Table 13.1). The other two, Meissner and Pacinian corpuscles, are rapidly adapting and signal only the start and stop of stimuli. It seems likely that Meissner corpuscles would be particularly well suited to signal the direction and speed of a moving stimulus. It should be noted that low-threshold mechanoreceptors in the skin also contribute to joint sense (see later), and control of posture and movements (see Chapter 18, under “Receptors of Importance for Upright Posture,” and Chapter 21, under “Cutaneous Receptors and the Precision Grip”).

Microneurographic Studies of Human Skin Receptors

Studies with techniques that enable **recording** from and **stimulation** of peripheral nerves in conscious human beings have provided important information with regard to the functional properties of receptors (Fig. 13.3). The Swedish neurophysiologists Hagbarth and Vallbo pioneered this technique around 1970. With the use of very thin needle electrodes, one records the activity of single sensory axons within a nerve, such as the median nerve at the forearm. Thus, it is possible to determine the receptive field of this particular sensory unit, along with its adequate stimulus. In glabrous skin of the fingers and palms, four types of low-threshold mechanoreceptors have been so characterized. Most likely, they correspond to the four encapsulated types described earlier (Table 13.1). Thus, there are two types of rapidly adapting sensory units: one with a small receptive field (most likely the Meissner corpuscle), and the other with a large and indistinct receptive field (Pacinian corpuscle). The two other types of sensory units are slowly adapting; again, one has a small receptive field (Merkel disk) and the other a large but direction-specific receptive field (most likely the Ruffini corpuscle).

Stimulation of the axons of the sensory units that have just been recorded enables correlations to be made between the conscious sensory experiences evoked by stimulation of only one sensory unit. Stimulation of single sensory

units that most likely end in Meissner corpuscles produces a feeling of light touch, like a tap on the skin with the point of a pencil. As a rule, the person localizes the feeling to exactly the point on the skin previously found to be the receptive field of the sensory unit. Activating a sensory unit that presumably leads off from Merkel disks evokes a sensation of light, steady pressure (as long as the stimulus lasts). Stimulating axons that appear to end in Pacinian corpuscles gives a feeling of vibration.

Receptive Fields of Cutaneous Sensory Units

It has been known for a long time that cutaneous sensation is punctate; that is, there are distinct tiny spots on the skin that are sensitive to different sensory modalities. We therefore use the terms “cold,” “warm,” “touch,” and “pain” spots. Cold spots are most easily demonstrated. Between the spots sensitive to cooling of the skin, there are others where contact with a cold object is felt only as pressure. This is so because each sensory unit distributes all its peripheral ramifications within a limited area of the skin. Thus, the sensory unit can be activated only from this part of the skin, which constitutes its **receptive field** (Fig. 13.3; see Fig. 12.1). Certain parts of the skin lack ramifications belonging to “cold” sensory units; consequently, sensations of cold cannot be evoked from such areas. Correspondingly, many small spots on proximal parts of the body—for example, on the abdomen and the upper arm—lack nociceptors; therefore, insertion of a sharp needle at such places is felt only as touch. The receptive fields of nociceptive sensory units are so closely spaced, however, that one has to make a thorough search to find painless spots.

The **size** of the receptive field depends on the area of the skin receiving axonal branches from the sensory neuron. In general, the **density** of sensory units—that is, the number of units innervating; for example, 1 cm² of the skin—is highest in distal parts of the body (fingers, toes, and lips), and the receptive fields are smaller distally than proximally (Fig. 13.3). This explains why the stimulus threshold is lower and the ability to localize a stimulus is more precise in the palm of the hand than at, for example, the upper arm.

Discriminative Sensation

The punctate arrangement of the cutaneous sensation is important for our ability to **localize** stimuli. Being able to determine that *two* pointed objects (such as the legs of a compass) touch the skin rather than one must mean that separate units innervate the two spots. Not surprisingly, the distance between two points on the skin that, when touched, can be identified as two is shortest where the density of sensory units is highest, and the receptive fields are smallest—that is, on the fingertips (Fig. 13.3A and B). Determining this distance gives a measure of what

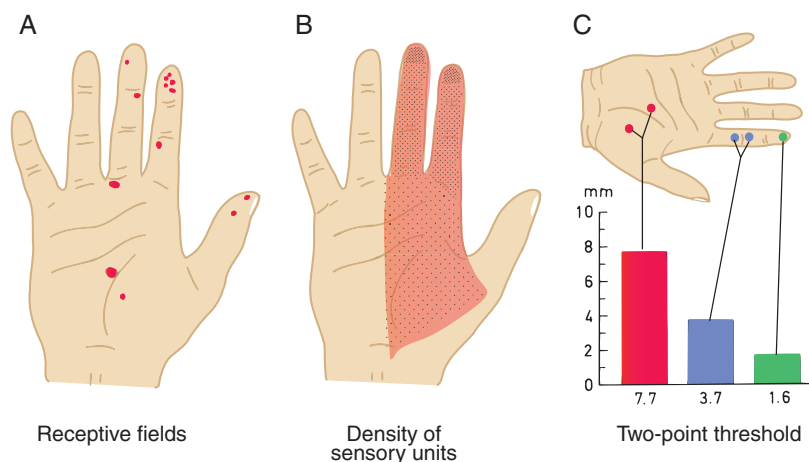


FIGURE 13.3 *Receptive fields.* **A:** Size and location of the receptive fields of 15 sensory units, determined by recording from the median nerve. All of these sensory units were rapidly adapting and were most likely conducting from Meissner corpuscles. Within each receptive field there are many Meissner corpuscles, all supplied by the same axon. **B:** Relative density of sensory units conducting from Meissner corpuscles (i.e., number of sensory units supplying 1 cm²). The density increases distally and is highest at the volar aspect of the fingertips. **C:** Two-point discrimination. The numbers give the shortest distance between two points touching the skin that can be identified by the experimental subject as two (reducing the distance further makes the person experience only one point touching the skin). Average of 10 subjects. (Based on microneurographic studies by Vallbo and Johansson 1978.)

we call **two-point discrimination** and is often used clinically. The smallest distance at which two stimuli can be discriminated is the **two-point threshold** (Fig. 13.3C). A useful test for this kind of discriminative sensation is the writing of letters or figures on the skin (with the subject's eyes closed). The figures that can be interpreted are quite small on the fingertips, somewhat larger on the palms, much larger on the upper arm, and even larger on the trunk. As one might expect, the pathways conducting the sensory signals from the spots on the skin are arranged topographically so that signals from different parts of the skin are kept separate at all levels up to the cerebral cortex.

Lateral Inhibition: Inhibitory Interneurons Improve the Discriminative Sensation

The two-point threshold (Fig. 13.3C) does not depend solely on the size and density of the cutaneous receptive fields. Inhibitory interneurons in the cord (and at higher levels) restrict the signal traffic from the periphery of a stimulated spot, thereby improving the discriminative ability compared with what might be expected from the anatomic arrangement of receptive fields (Fig. 13.4). The cutaneous sensory units (neurons) send off collaterals in the CNS that activate inhibitory interneurons that, in turn, inhibit sensory neurons in the vicinity. This phenomenon is called **lateral inhibition**, and occurs at all levels of the sensory pathways and in all sensory systems. Figure 13.4 shows an example with a pencil pressed lightly onto the skin. The sensory units with receptive fields in the center of the stimulated spot receive the most intense stimulation. Therefore, they excite the inhibitory interneurons strongly, with consequent strong inhibition of sensory units leading from the periphery of the spot. Sensory units with receptive fields in the periphery are less strongly stimulated but receive strong inhibition. Thus, they cannot inhibit the transmission of signals from the center of the spot. Together, the impulse traffic from the periphery of the

stimulated area is reduced, and we perceive the stimulated area as smaller and more sharply delimited than it really is. In this way, the CNS receives distorted sensory information.

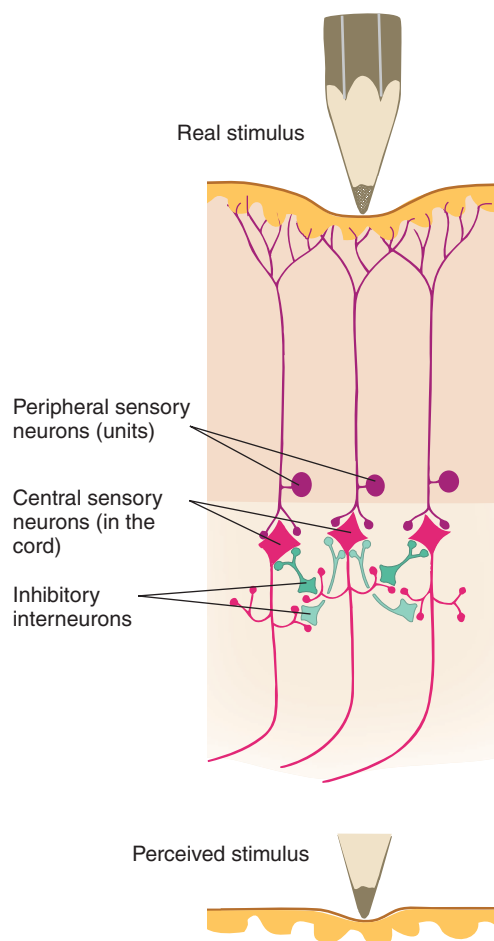


FIGURE 13.4 *Lateral inhibition.* Simplified presentation of how inhibitory interneurons in the CNS can improve the precision of the sensory information reaching consciousness.

PROPRIOCEPTORS: DEEP SENSATION

As mentioned, the term “proprioceptive” is used for sensations pertaining to the **musculoskeletal system**—the muscles, tendons, joint capsules, and ligaments. There are many similarities between proprioceptors and cutaneous receptors. For example, numerous free receptors (belonging to thin myelinated and unmyelinated axons) occur in the muscles, the muscle fascia, and the dense connective tissue of joint capsules and ligaments. Many of these are nociceptors as in the skin.

Here we are dealing primarily with specialized sense organs in muscles and around joints, which are of crucial importance for control of posture and goal-directed movements. These are low-threshold mechanoreceptors, and the signals from them are conducted centrally in thick, myelinated axons. The adequate stimulus of these receptors is stretching of the tissue in which they lie. Whether the receptors are located in a muscle or in a joint capsule, joint movement is the natural stimulus that leads to their activation.

We first discuss specialized sense organs in muscles—muscle spindles and tendon organs—and then discuss receptors in joint capsules and ligaments. In Chapters 18 and 21, we treat the role of proprioceptors in control of movements.

Classification of Muscle Sensory Fibers

Muscle afferents, that is, sensory fibers leading from muscles, are classified according to size into groups I to IV (size or thickness is closely related to conduction velocity). **Group I** muscle afferents contain fast conducting, thick myelinated fibers, while **group II** contain medium-sized myelinated fibers, and **group III** comprises the thinnest myelinated fibers. **Group IV** contains the slowly conducting unmyelinated fibers. Group I is further divided into **Ia** and **Ib** fibers, with Ib fibers conducting slightly more slowly than Ia fibers. Signals from low-threshold mechanoreceptors in muscles—that is, from muscle spindles—are conducted in group I and II fibers, while the tendon organs are supplied with Ib fibers. Unfortunately, other terms are used for classification of cutaneous sensory fibers and efferent (motor) fibers. Broadly, group I and II fibers correspond with regard to conduction velocity to A α and A β fibers, while group III and IV correspond to A δ and C fibers, respectively (see later in this chapter under “Classification of Dorsal Root Fibers in Accordance with Their Thickness”).

Nociceptors in Muscle and Tendon

As mentioned, muscles are supplied with numerous free nerve endings of thin myelinated and unmyelinated axons (i.e., the most slowly conducting fibers). Microscopic examination of muscle nerves in experimental animals

shows that almost 40% of all the axons are either thin myelinated or unmyelinated sensory fibers, terminating in free endings (Fig. 13.5). Many—probably the majority—of the unmyelinated and thin myelinated axons lead from nociceptors in the muscle. This has been shown by recording the activity of single sensory units that innervate a muscle while systematically exploring their adequate stimuli. Such units were excited by both strong mechanical stimuli and substances liberated in inflamed tissue (such as bradykinin, known to provoke pain in humans). Inflammation also **sensitizes** the nociceptive sensory units; making them respond to normal movements (this may partly explain the muscle soreness after heavy exercise). Like cutaneous nociceptors, those in muscle are also sensitized by prolonged stimulation, and some sensory units are activated by **ischemia**. For example, muscle ischemia caused by a thrombotic artery, produces pain in humans.⁴

Microneurographic studies have identified sensory units in human muscle nerves that have nociceptor-like properties similar to those described in the preceding text in experimental animals. Thus, human units are slowly adapting, have a high threshold for mechanical stimuli, and can be activated by inflammatory substances. Electrical stimulation of small nerve fascicles in human muscle nerves produces pain as the only sensory experience. The pain is felt to be deep (not in the skin) and has a cramp-like quality. The subject localizes the pain to the muscle supplied by the nerve, not to the site of nerve stimulation that is at a distance from the muscle (recording and stimulation of single sensory fibers have confirmed these findings). If the stimulation continues, or its intensity is increased, the pain radiates to other regions than the muscle itself (this phenomenon is called **referred pain**; see later, “Spinothalamic Cells Receive Signals from Both Somatic and Visceral Structures: Referred Pain”).

A small number of axons from nociceptors in a muscle **tendon** have been investigated with microneurography. Stimulation of such a sensory unit produced a sharp pain, different from the muscle pain described above. The subject localized the pain to the tendon.

Ergoreceptors and Other Kinds of Free Receptors in Muscles

Not all of the thin, slowly conducting sensory fibers in a muscle nerve are nociceptors. Thus, in experimental animals, some of these fibers have a low mechanical threshold; that is, ordinary muscle contraction or gentle

⁴ Animal experiments using cultured dorsal root ganglion cells indicate that the adequate stimulus for many muscle nociceptors is a combination of protons, ATP, and lactate, acting on ASICs (acid sensing ion channels), P2X (purinergic type 2), and TRPV receptors, respectively. Increased levels of these metabolites would presumably mediate the pain provoked by heavy muscular exercise (causing ischemia).

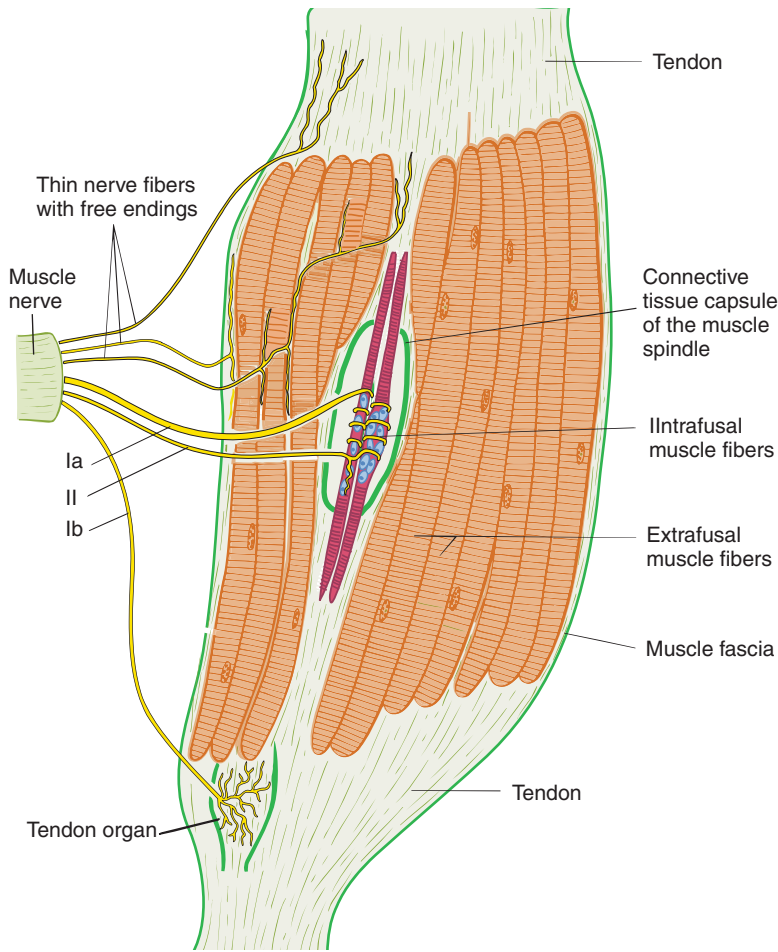


FIGURE 13.5 *Sensory innervation of skeletal muscles.* The size of the receptors relative to the muscle is exaggerated. Because the muscle spindle attaches to the tendons via connective tissue fibers, the muscle spindle is stretched whenever the whole muscle is stretched. Many of the free nerve endings are nociceptors.

stretching of the muscle activates them. Their maximal firing rate is reached by stimuli much weaker than the stimuli producing pain. Such receptors may be responsible for circulatory and respiratory reflex effects known to occur at the start of muscular activity. Thus, there is a slight increase of pulse and breathing rate that occurs too early to be caused by a rise in blood CO_2 or lowered pH. Although the receptors responsible for such reflex effects have not been identified with certainty, they are called **ergoreceptors** (ergoceptors).

The Structure and Innervation of the Muscle Spindle

The name muscle spindle is derived from the oblong shape of this sense organ. The muscle spindles are located within the muscle, among the striated muscle cells, and consist of a few (2–14) specialized muscle cells enclosed in a connective tissue capsule. The capsule is approximately 0.2 to 0.3 mm in diameter and up to 5 mm long. The muscle fibers (or muscle cells) of the spindle are called **intrafusal** and are much thinner and shorter (7–10 mm long) than the ordinary, **extrafusal**, muscle fibers. In contrast to the extrafusal muscle fibers, the

intrafusal fibers show cross-striation only at their ends. This means that they are able to contract only these parts and not their middle portions. There are two main types of intrafusal fibers (Fig. 13.6). One type is called the **nuclear bag fiber** because the nuclei are all collected in the middle part of the muscle fiber. In the other type, the **nuclear chain fiber**, the nuclei are evenly distributed along the muscle fiber.

The **nerve supply** of the muscle spindles is highly complex, and only the main features will be treated here (Figs. 13.5–13.7). A thick afferent fiber ends with a spiraling course around the middle portion of the nuclear bag and to a lesser extent nuclear chain fibers, forming the **primary sensory ending** of the muscle spindle. In addition, a thinner afferent fiber ends mainly in relation to the nuclear chain fibers, forming the **secondary sensory ending**. Afferent nerve fibers from muscles are classified with regard to thickness (and thus to conduction velocity) in groups I to IV, with group I fibers comprising thick myelinated axons, and group IV unmyelinated ones. Group I is further divided into Ia and Ib, the former being the thickest. The primary sensory ending belongs to a **group Ia** afferent fiber, whereas the

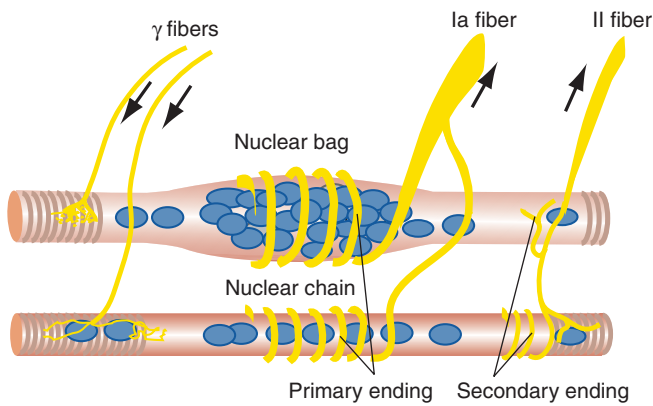


FIGURE 13.6 *The muscle spindle.* Schematic of the two kinds of intrafusal muscle fibers and their innervation. (Modified from Matthews 1964.)

secondary sensory ending originates from a **group II** afferent fiber (Figs. 13.7A and B). Both types of sensory neuron have their cell bodies in the spinal ganglia. The **adequate stimulus** for the primary and secondary sensory endings is stretching of the intrafusal muscle fibers. This deforms the spiraling axonal branches and thus elicits depolarization and (if the depolarization is strong enough) action potentials in the group Ia and II sensory fibers.

The muscle spindle is also supplied with motor axons, called **fusimotor**, or γ , fibers, coming from γ **motoneurons** located in the ventral horn (Fig. 13.7C). The γ fibers end in the distal cross-striated parts of the intrafusal fibers and make them contract (Fig. 13.6). The contraction leads to stretching of the noncontractile middle part of the intrafusal fibers, where the sensory endings are located, increasing the sensitivity of the intrafusal fibers to stretch. It should be emphasized that because the intrafusal fibers are so few and thin, their contraction does not contribute to the tension or shortening of the whole muscle.

Number and Density of Muscle Spindles

The number of muscle spindles has been determined in approximately 200 human muscles, and varies greatly from muscle to muscle. The functional significance of such variations is not clear, and has been subject to different interpretations. In large muscles such as the latissimus dorsi the **total number** of muscle spindles is about 400, whereas the small abductor pollicis brevis contains about 80. The largest absolute number, close to 1300, was found in the quadriceps muscle. The soleus muscle contains approximately 400 muscle spindles, the medial gastrocnemius 150, and the extensor digitorum longus muscle (EDL) 190. Although the absolute number is low in small muscles, the **density** of muscle spindles may be very high. For example, the tiny lumbrical muscles

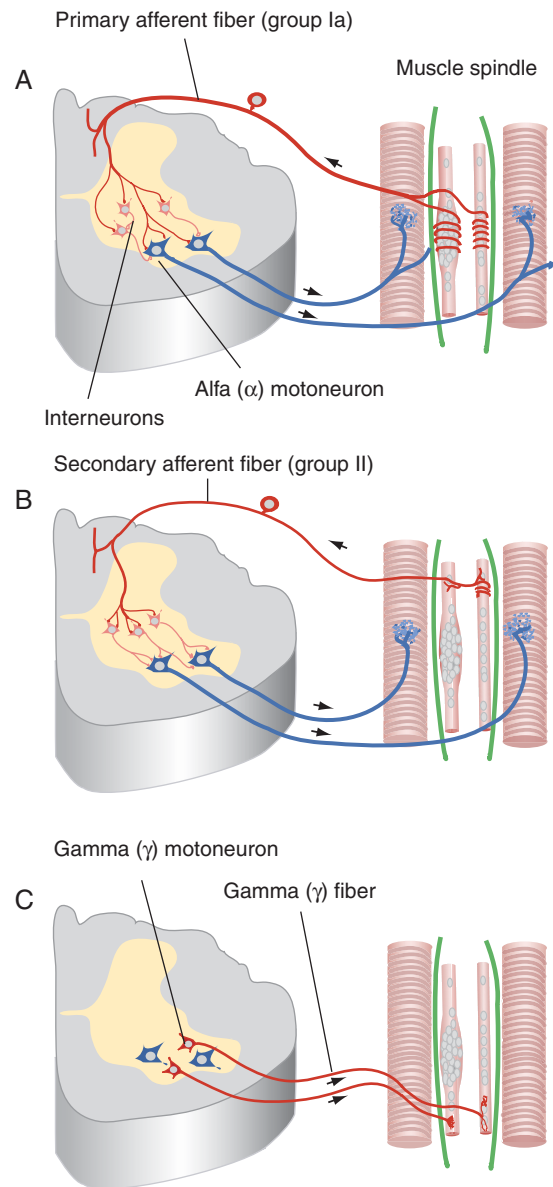


FIGURE 13.7 *Innervation of the muscle spindle.* **A:** The fast-conducting group Ia fibers end monosynaptically on α motoneurons in the ventral horn. This is the reflex arc that mediates the brief muscle contraction elicited by a tendon tap (monosynaptic or phasic stretch reflex). In addition, Ia afferents end on interneurons and send ascending and descending collaterals into the dorsal columns. **B:** The group II fibers end on interneurons in the intermediate zone and the ventral horn. **C:** The small γ motoneurons lie scattered among the larger α motoneurons in the ventral horn.

of the hand contain 12 spindles per gram of muscle with a total count of only 18 muscle spindles. The corresponding numbers for the soleus and the EDL are 0.4 and 3.7, respectively.

When comparing the number of muscle spindles with the number of **motor units** of the muscle, however, the ratio is not necessarily higher in small than in large muscles. (A motor unit is a motoneuron and all the extrafusal muscle fibers it supplies, and is the basic functional unit of

a muscle. Motor units are larger; that is, there are more muscle fibers per motoneuron, in large than in small muscles; see Chapter 21.) Thus, the ratio for the first dorsal interosseus muscle is 0.29, the first lumbrical muscle 0.54, the biceps muscle 0.41, and the tibialis anterior muscle 0.64.

Another relationship exists between density of muscle spindles and kind of extrafusal muscle fibers. In general, it appears that muscle spindles are most abundant in “red” parts of a muscle (**Type 1 muscle fibers**, see Chapter 21).

Functional Properties of the Muscle Spindle

To understand how the muscle spindle functions, one must know that it is arranged in **parallel** with the extrafusal muscle fibers. Thus, both ends of the spindle are attached to the connective tissue within the muscle and are thereby indirectly anchored to the muscle tendons (Fig. 13.5). From this structure one may deduce that when the whole muscle shortens as a result of contraction of the extrafusal fibers, the intrafusal fibers will be shortened passively. Conversely, stretching of the whole muscle will stretch the intrafusal muscle fibers. The rate of shortening or lengthening will be the same for the muscle spindle as for the whole muscle.

Action potentials can be recorded from single group Ia and II afferent fibers in the dorsal roots of anesthetized animals. It is then possible to study how the primary and secondary sensory endings behave in response to various stimuli. As expected from the anatomic facts described in the preceding text, both types of afferent fibers increase their firing rate (i.e., the frequency of

action potentials) as the length of the muscle increases (Fig. 13.8). If the muscle shortens, the firing rate decreases (if the muscle is sufficiently shortened, no action potentials can be recorded; see Fig. 13.9). When the length of the muscle is kept constant, the firing rate is also constant (static phase in Fig. 13.8); the muscle spindle afferents are thus **slowly adapting**. This property of the muscle spindle is called **static sensitivity**. Because the firing rate of both group Ia and group II fibers depends on the length of the muscle, both inform the CNS about the **length of the muscle** at any time (or the static length).

During the phase in which the muscle length is changed, however, group Ia and group II afferent fibers behave differently (dynamic phase in Fig. 13.8). The firing rate of the group Ia fiber is much higher during stretching than when the length is kept stationary in the stretched position, but the group II fiber does not show this same change in firing rate. During the shortening phase, the Ia fiber becomes completely “silent.” Although not shown in Fig. 13.8, the firing rate of the group Ia fiber also depends on the velocity of the length change. Thus, the Ia fiber signals that the length of the muscle is changing, as well as the **velocity** with which it is occurring. This property is called **dynamic sensitivity**.

These facts indicate that the **primary sensory ending** of the muscle spindle has both static and dynamic sensitivity: this ending is capable of informing about the actual length of the muscle (position of a joint), whether the length is constant or changing (joint movement), and the velocity of change (velocity of the movement).

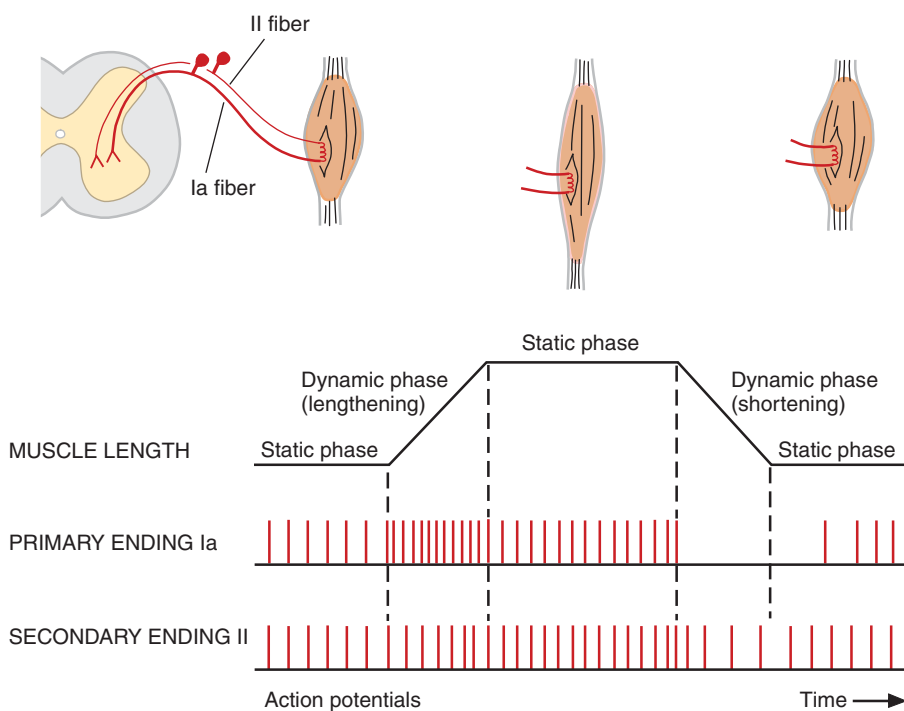


FIGURE 13.8 Functional properties of the muscle spindle. Both the primary and the secondary sensory endings signal the static length of the muscle (static sensitivity), whereas only the primary ending signals the length changes (movements) and their velocity (dynamic sensitivity). The diagram is based on recordings from single dorsal root fibers of anesthetized cats. The change of firing frequency of group Ia and group II fibers can then be related to static muscle length (static phase) and to both stretching and shortening of the muscle (dynamic phases). The density of the vertical lines on the two lower rows indicates the frequency of action potentials in the dorsal root fibers. (The muscle spindle is not under the influence of γ motoneurons in this experiment.)

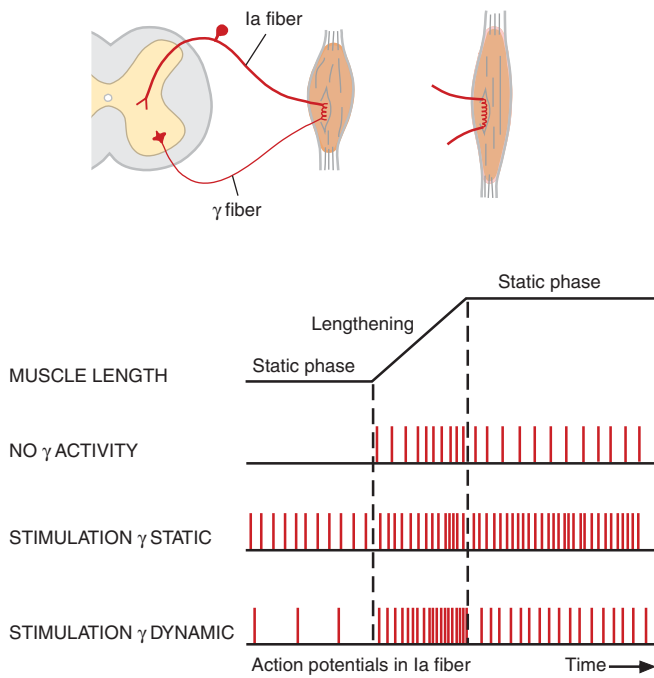


FIGURE 13.9 Action of γ motoneurons on the muscle spindle. The experimental setup is as in Fig. 13.8, except that in addition to recording the activity of group Ia fibers in the dorsal root, γ axons are isolated in the ventral roots so they can be electrically stimulated. In this example, there is no firing of the Ia fiber at the resting length of the muscle when the γ fibers are not stimulated. Stimulation of a static γ fiber (innervating the same spindle that the Ia fiber conducts from) makes the Ia fiber fire even at the static resting length; stretching the muscle to a new static length increases the firing frequency to a new stable level. Stimulation of a dynamic γ fiber increases the firing frequency of the Ia fiber mainly during the stretching phase.

Because the **secondary sensory ending** almost totally lacks dynamic sensitivity, it should be able to inform primarily about the static length of the muscle.

There is much evidence to suggest that the **nuclear bag fibers** are responsible for the dynamic sensitivity of the primary sensory ending, whereas the **nuclear chain fibers** are responsible for the static sensitivity of both the primary and secondary sensory endings. That the primary and secondary sensory endings (and their afferent nerve fibers) have different properties is most likely due to differences in viscoelastic properties of the nuclear bag and nuclear chain intrafusal muscle fibers.⁵

The preceding description of the properties of the muscle spindle derives from experiments in which there was no impulse traffic in the **fusimotor γ axons** (because the ventral roots were cut before recording from the dorsal roots). As discussed in the next section, however,

the properties of the muscle spindle are markedly influenced by the activity of the γ motoneurons. To understand the functioning of the muscle spindle in an intact organism, we must therefore also know the actions of the γ innervation.

Effects of γ Innervation on the Properties of the Muscle Spindle

As mentioned, signals in γ fibers elicit contraction of the distal, cross-striated parts of the intrafusal muscle fibers. This stretches the midportion of the intrafusal fibers with the sensory endings (Fig. 13.6). In addition, it also alters the **stiffness** of the intrafusal fibers so that their reaction to stretch is altered. In general, the γ motoneurons and their γ fibers enable the brain to control the **sensitivity** of the muscle spindle to length and changes in length.

In animal experiments, single γ fibers in the ventral roots have been stimulated while, at the same time, the activity of group Ia and group II afferent fibers in the dorsal roots were recorded. It has thus been shown that there are **two types of γ motoneurons** (Fig. 13.9). One type increases the **dynamic sensitivity** of the muscle spindle and is therefore called γ_d (**gamma dynamic**). On a fairly rapid stretch of the muscle, the firing rate of a group Ia fiber increases more when the muscle spindle receives signals from γ_d motoneurons than without such influence, but the firing rate during static length is not significantly altered (Fig. 13.9). The muscle spindle's increased sensitivity to stretch enables the CNS to react more rapidly and forcefully to any unwanted change in muscle length (imposed, e.g., by external forces that upset body balance or ongoing movements).

Signals from the other type of γ motoneurons increase the **static sensitivity** of the muscle spindle and are therefore called γ_s (**gamma static**). The activity of γ_s motoneurons increases the firing rate of muscle spindle afferent fibers during constant length, as compared with a situation without γ activity. Although not shown in Fig. 13.9, the firing rate of both group Ia and group II afferents increases. This influence of the γ system may be important to prevent the muscle spindles from becoming "silent"—that is, sending no action potentials—during the shortening of the muscle. In other words, the **length sensitivity** of the muscle spindle increases. Thus, the muscle spindle may signal the length of the muscle in its entire range of movements, which is for precise movement control and for our awareness of joint positions.⁶ (Figure 13.7 shows that stimulation of γ_s in fact *reduces*

⁵ In reality, the conditions are even more complex. Among other things, there are two types of nuclear bag fibers that differ ultrastructurally and histochemically: Only one of them, called **bag₁**, appears to be responsible for the dynamic sensitivity of the primary sensory ending. The other one, called **bag₂**, behaves more like a nuclear chain fiber and contributes presumably only to the static sensitivity.

⁶ The γ innervation poses a problem for the brain when judging joint positions and movements, however, since the signals from muscle spindles depend not only on the absolute length of the muscle but also on the degree of γ activity. Since muscle spindles are crucial for our perception of joint positions and movements, the brain must in some way be able to account for the γ activity and end up with a true measure of muscle length.

the dynamic sensitivity of the primary sensory ending, since there is no extra increase in firing rate of the Ia fiber during the stretch phase. Under the influence of γ , the primary sensory ending behaves more like a secondary one.)⁷

Even though this description of the properties of muscle spindles is based on experiments in animals, there is evidence that it applies to the human muscle spindle. Certainly, however, results from anesthetized animals, often with the spinal cord isolated from the rest of the brain, do not enable us to draw conclusions as to the functions of the muscle spindle in intact organisms, for example, in human voluntary movements and in proprioception. It may seem paradoxical that the muscle spindle is among the most studied sense organs, yet its functional roles remain only partially understood.

Muscle Spindles in Humans and α - γ Coactivation

The activity of group Ia afferent fibers in the nerves of the arm and the leg has been recorded in conscious human subjects via **microneurographic** techniques. It appears (unexpectedly, based on animal experiments) that in a resting muscle there is little or no impulse traffic from the muscle spindle. Indirectly, this shows that there is no fusimotor (γ) activity, either. But if the muscle contracts isometrically (i.e., without change of length), there is a sharp increase of the firing frequency of Ia fibers, which must be caused by increased fusimotor activity that occurs simultaneously with the increase in α -motoneuron activity (which evokes the contraction of the extrafusal fibers). This phenomenon of simultaneous activation of α and γ motoneurons is called **α - γ coactivation**. This ensures that the sensitivity of the muscle spindle is increased whenever the muscle is being used. In fact, the firing rate of the Ia fiber is maintained or increased even if the muscle is shortened during active contraction. This must mean that the fusimotor activity (firing rate of the γ motoneurons) increases during active shortening of the muscle.

The preceding example of α - γ coactivation does not mean that the γ motoneurons are activated only in conjunction with the α motoneurons, even though direct proof of separate activation is scarce. There are situations in which it would be desirable to have increased sensitivity of the muscle spindle without simultaneous muscle contraction. One piece of indirect evidence comes from studies of stretch reflexes during **mental imagery**. In such a situation—when the subject imagines the performance of a movement—the stretch reflex response of the relevant muscles is increased without concomitant increase of α -motoneuron excitability, so

it appears that the sensitivity of the muscle spindle has been increased by **selective activation** of γ motoneurons. It is also difficult to understand why the elaborate γ system has developed if its activity were always to reflect that of the α system. In fact, there are collaterals of α axons, so-called **β axons**, that innervate some intrafusal muscle fibers, and in submammalian species (e.g., in the frog) there are only β fibers. During evolution, the β system appeared first.

Signals from Muscle Spindles Reach Consciousness

There is good evidence that signals from muscle spindles contribute to our conscious awareness of joint angle and movement (discussed further later in this chapter). Signals from a single muscle spindle are not sufficient to produce a conscious sensation, however (in contrast to some cutaneous and joint low-threshold mechanoreceptors), so that microneurographic stimulation of single spindle afferents from human intrinsic hand muscles does not evoke any sensation. Many muscle spindles must obviously be activated simultaneously for the signals to be consciously perceived, and presumably this always happens when a muscle is stretched. Therefore, it seems unnecessary and perhaps also disturbing (“noise”) to the brain if every muscle spindle were to evoke a sensation on its own.

Especially in weak contractions, different parts of a muscle undergo different length changes; consequently, the muscle spindles would fire with different frequencies. In this way the signals from single muscle spindles may provide useful feedback to the motor control machinery in the cord about smaller parts of the muscle. The cerebral cortex is presumably concerned only with the muscle as a whole and extracts necessary information by integrating the signals from the numerous muscle spindles.

The Tendon Organ

The other kind of proprioceptive receptor we describe here is the tendon organ, also called the **Golgi tendon organ**. It is built more simply than the muscle spindle and consists of a sensory nerve fiber that follows a convoluted course among collagen fibrils of the tendon, close to the musculotendinous junction (Fig. 13.5). The number of tendon organs in a muscle appears to be only slightly lower than the number of muscle spindles. The thick, myelinated fiber leading from the tendon organ belongs to group I and is called a **group Ib** fiber. There is no efferent innervation of the tendon organ (in contrast to the muscle spindle): its sensitivity cannot be controlled from the CNS.

The **adequate stimulus** of the tendon organ is stretching the part of the tendon in which it lies. Stretching tightens the collagen fibers, and thus the axonal branches

⁷ Because the bag fibers appear to be solely responsible for the dynamic sensitivity of the muscle spindle, it has been assumed that dynamic γ fibers end on bag fibers and static γ fibers end on chain fibers. This is not fully clarified, however.

between them are deformed (probably stretched, similar to the group Ia afferents). This depolarizes the receptor and, if the stimulus is of sufficient intensity, evokes action potentials in the afferent Ib fiber. Recording the activity of Ib fibers shows that the receptor is **slowly adapting**. It is important to realize that the tendon organ, in contrast to the muscle spindle, is coupled in **series** with the extrafusal muscle fibers. Both passive stretch and active contraction of the muscle increase the tension of the tendon and thus activate the tendon organ receptor. The tendon organ, consequently, can inform the CNS about the **muscle tension**. In contrast, the activity of the muscle spindle depends on the muscle length and not on the tension.

Recording from single group Ib fibers in the dorsal root of anesthetized cats (Fig. 13.10) confirms what was expected on the basis of the structure of the tendon organ. In addition, however, such experiments have shown that the tendon organ is much more sensitive to tension produced by **active contraction** than to that produced by passive stretch. The tendon organ therefore appears to be primarily concerned with signaling how hard the muscle is contracting rather than with how hard it is passively stretched.

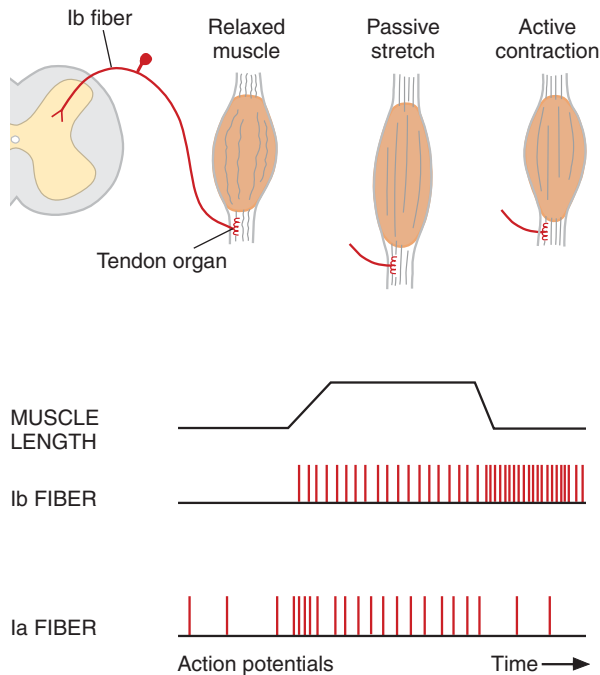


FIGURE 13.10 *Functional properties of the tendon organ.* The experimental setup is as in Fig. 13.8. Action potentials are recorded from isolated Ib fibers in the dorsal roots. Vertical lines on the lower rows indicate the firing frequency. Both passive stretching and active contraction of the muscle increase the firing frequency of the Ib fiber, but active contraction produces the greatest increase. The firing frequency of a Ia fiber during the same experiment is shown for comparison.

Why Tendon Organs Are More Sensitive to Contraction than to Passive Stretch

Structural details may explain why the tendon organ is more sensitive to active contraction than to passive stretch. Each tendon organ is directly attached to a small bundle of extrafusal muscle fibers. If one or a few of these contract, the tension set up in this particular small part of the tendon is much higher than the tension measured for the whole muscle. To obtain the same tension in this particular tendon organ by passive stretch, higher overall tension of the muscle would have to be produced. The muscle fibers attached to one tendon organ appear to belong to several motor units (see Chapter 21, under “Motor Units”). Because each tendon organ probably monitors the tension produced by only a few motor units, the CNS is informed not only of the overall tension produced by the muscle but also of how the workload is distributed among the different motor units.

The Actions of Ib Afferents on Spinal Motoneurons

Activation of the group Ib afferents from tendon organs was shown a long time ago to inhibit (via interneurons) motoneurons of the muscle in which the tendon organs lie (homonymous inhibition). Although these experiments were performed on anesthetized animals, and therefore should be interpreted with caution, the task of the tendon organ was said to be to prevent contractions from being too strong. More recent studies have shown that the effects of the Ib afferents in the cord are not limited to homonymous inhibition. In awake animals, the effect on the motoneurons depends on the locomotor phase, and the effect is reversed from inhibition to excitation when moving from the swing phase to the standing phase. Thus, tendon organs in hind limb extensors excite extensor motoneurons when the leg is in the standing phase (via excitatory interneurons). In this case the signals from the tendon organ serve to amplify the contraction of the extensor muscles that keep the upright position. This is an example of how higher motor centers (in the brain stem and cerebral cortex) can switch the impulse traffic from one route to another in the cord, depending on the motor task. Further examples of this phenomenon are provided in Chapter 21.

Perception of Muscle Force

Our conscious perception of how hard the muscles contract may depend on two sources of information. One is the total activity of neurons in the motor cortex that send commands to the muscles that contract. This requires that other parts of the brain—for example, the somatosensory cortex—receive a copy of the motor commands sent to the muscles. This is called **efferece copy**, or **corollary discharge**. Based on previous experience, the

somatosensory cortex (in cooperation with other cortical regions) may estimate the muscle force that corresponds to the motor command. The other source of information is the **proprioceptors** that inform about the tension in the muscles themselves and in connective tissue that is stretched by muscle contraction. As discussed, **tendon organs** are particularly suited to informing about the tension in a contracting muscle, but because muscles often insert in the joint capsule, **joint receptors** may also contribute to communication of information.

That the motor cortex output plays a part is witnessed by the fact that persons with pareses due to muscle disease judge objects to be heavier than they actually are. To compensate for the weakened muscle, the motor cortex output is presumably higher than normal while the proprioceptive feedback is correct (informing about the real muscle tension). Nevertheless, the proprioceptive information seems to be necessary also, since patients with neuropathies may experience difficulties with holding a steady force and judging the weight of objects.

Low-Threshold Mechanoreceptors around the Joints

Not only receptors in muscles and tendons but also receptors in the connective tissue around the joints provide information important for our awareness of movements and for motor control. While the relative importance of information from joint and muscle receptors is not clear, the prevailing view is that the contribution from joint receptors to proprioception is less important than that of the muscle spindles.

Many sensory nerve fibers end in the joint capsules and in the ligaments around the joints (Fig. 13.11). Many are **free-ending receptors**; others are **encapsulated** endings that correspond anatomically and with regard to response properties to encapsulated receptors in

glabrous skin. The encapsulated joint receptors are **low-threshold mechanoreceptors** and have been divided into four groups. The **type 1 joint receptor** resembles the Ruffini corpuscle in the dermis (Fig. 13.1A). A myelinated axon ramifies among collagen fibrils, within a thin connective tissue capsule. They are found almost exclusively in the fibrous part of the joint capsules. The **adequate stimulus** of these Ruffini-like receptors is increased tension in the part of the capsule in which they lie. The higher the capsular tension is, the higher is the firing rate in the afferent sensory fiber from the receptor. Like the Ruffini corpuscle in the skin, this joint receptor is **slowly adapting**. Because the tension in various parts of the capsule depends on the joint position, type 1 receptors would appear suited to signal the position of the joint. For example, receptors in the posterior part of the elbow joint capsule would be highly active in a flexed position of the joint, which stretches the capsule, and less active in an extended position, which relaxes the capsule. The receptor also has **dynamic sensitivity**, giving a stronger response (higher firing rate) to a rapid movement than to a slow one. The type 1 or Ruffini-like receptor thus seems capable of signaling static joint position, joint movements, and direction and speed of movements. As discussed later, however, the ability of the type 1 receptor to signal static joint position appears to be limited.

The **type 2 joint receptor** structurally and functionally resembles the Pacinian corpuscle but is considerably smaller (it is also called “Paciniform receptor”). Type 2 receptors are present only in the fibrous part of the joint capsules. They are **rapidly adapting**, and their **adequate stimulus** is stretching of the part of the capsule in which they lie. Owing to their rapid adaptation, they can inform only on joint movements, not of static position. They appear particularly suited to signal **movement velocity** and have also been called acceleration receptors.

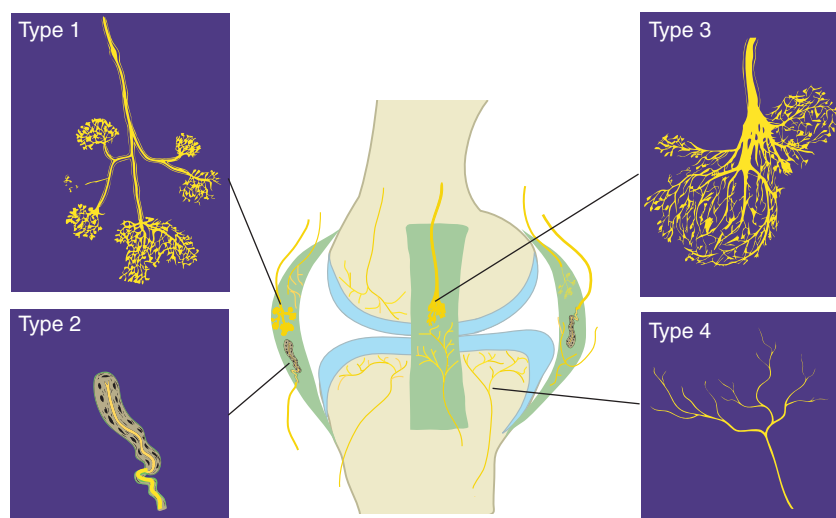


FIGURE 13.11 *Joint innervation.* Drawing of a knee joint, showing the distribution of the various kinds of joint receptors. The morphology of the four main receptor types is shown in more detail in the side panels. Type 1, 2, and 3 are low-threshold mechanoreceptors responding to small changes in joint-capsule tension. Type 4 are free nerve endings (most of them nociceptors).

A third kind of encapsulated receptor (**type 3**) resembles the tendon organ and is present in **ligaments** only. It is slowly adapting (like the tendon organ), but its functional role is unknown. A protective role in signaling overstretching of joints has been proposed but has not gained experimental support.

Microneurographic studies in humans with stimulation of single afferent fibers from finger joints show that signals from low-threshold joint receptors can reach consciousness. Thus, when a fiber leading from a presumed type 1 receptor was stimulated, the subject reported the feeling of a movement of the relevant joint (the location and adequate stimulus of the receptor were determined before stimulation). In other instances of single-unit stimulation, the subject reported only a nonpainful, punctate feeling at the joint.

Joint Nociceptors

The fibrous part of the joint capsule and the ligaments are richly supplied with thin myelinated and unmyelinated axons ending in free receptors. These have been termed **type 4** joint receptors. Many of these, like free endings in other tissues, are nociceptors. The joint capsules and ligaments contain **high-threshold mechanoreceptors**, **polymodal nociceptors**, and “**silent**” **nociceptors**. Stimulation of the fibrous capsule and the ligaments during surgery of the knee joint confirms that pain can be evoked from both the capsule and ligaments. The sensation is punctate—as in the skin—so that certain spots when probed with a needle evoke pain while other spots evoke a feeling of pressure or no sensation at all.

Many of the thin fibers ending in the joint capsule contain **neuropeptides**, such as substance P and calcitonin gene-related peptide (CGRP). Release of such peptides is believed to have a role in the development of inflammatory **arthritis** in humans (especially rheumatoid arthritis). Thus, the severity of induced inflammatory arthritis in experimental animals is reduced after depleting the joint sensory fibers of substance P (see later in this chapter under “Inflammatory diseases and release of neuropeptides from peripheral branches of sensory neurons”).

Other free joint receptors appear to be **ergoreceptors**—that is, they play a part in circulatory and respiratory reflexes that are elicited by passive joint movements.

Innervation of the Synovial Membrane

While autonomic efferent fibers have been traced to the synovial membrane, innervation by **sensory fibers** has been a matter of dispute. Thus, such fibers were not identified by use of heavy metal impregnation, which reliably showed nerve fibers in the fibrous capsule. Nevertheless, recent immunocytochemical studies in humans showed the presence of thin nerve fibers that end freely in the synovial membrane. These fibers contain **neuropeptides**

typical of primary sensory neurons (substance P, CGRP, and others). Due to the low density of such fibers, doubt nevertheless exists as to their functional significance. Studies in humans using various ways of stimulating the synovial membrane (during surgery under local anesthesia) have given conflicting evidence: some report that pain sensations were evoked from the synovial membrane, others not. In probably the most careful study, however, pain (or any other sensation) was only occasionally evoked from the synovial membrane of the knee joint with mechanical and chemical stimuli. This contrasted with the ease with which pain was evoked from the fibrous capsule and ligaments, as described earlier.

In contrast to the sparse sensory innervation, the synovial vessels possess a rich **efferent (autonomic) innervation**. These fibers contain norepinephrine and usually also neuropeptide Y (NPY). Release of norepinephrine can induce the release of substance P from sensory nerve fibers in the vicinity, and substance P may be involved in producing inflammation in **arthritis**. Indeed, by removing the autonomic innervation of the joint, arthritic inflammation can be reduced in animal models of arthritis. It should be emphasized, however, that norepinephrine and substance P are present also in the fibrous capsule. Therefore, sensory fibers in the synovial membrane may not necessarily be responsible for the inflammatory response.

Kinesthesia

The term **kinesthesia** (sometimes used synonymously with joint sensation and proprioceptive sensation) is commonly used to refer to the **perception of joint position, joint movements, and the direction and velocity of joint movements**. Strictly speaking, the word “kinesthesia” (Greek: *kinesis*, movement) encompasses only the dynamic and not the static aspect of sensation. The term “joint sensation” is not a good alternative, however, because it may give the false impression that kinesthesia depends only on joint receptors. Some also include in the term “kinesthesia” the perception of force, effort, and heaviness in relation to muscle contractions.

Our ability to judge the position of a joint (without seeing it), even after a long period without movement, is usually fairly precise, as can be verified by trying to match the position of the joint to be tested with the joint of the other side (e.g., finger or elbow joints). This is the **static** part of kinesthesia. Nevertheless, the precision of our judgment is increased considerably if movements are allowed, particularly active movements.

The static aspect of kinesthesia depends on slowly adapting receptors that change their firing rate with changing joint position. The **dynamic** aspect of kinesthesia—the ability to perceive that a movement is taking place and to judge the direction and velocity of the movement—depends on receptors with dynamic sensitivity, many of

them rapidly adapting. Receptors with such properties, which are influenced by joint movements, are found in muscles, around the joints, and in the skin. The accumulated evidence today indicates that **muscle spindles**, **joint receptors**, and **skin receptors** all contribute to kinesthesia. Muscle spindles appear to contribute most importantly to kinesthesia with regard to large joints, such as the **hip** and knee joints, whereas joint receptors and skin receptors may have more significant contributions with regard to **finger** and toe joints.

Significance of Various Receptors for Kinesthesia

The views on which receptor types are responsible for kinesthesia have undergone considerable changes. At the beginning of the past century, the newly discovered muscle spindle was held solely responsible but during the 1950s and early 1960s investigations indicated that joint receptors had the necessary properties to signal all the information needed for kinesthesia. It was also argued that the muscle spindle cannot give the necessary information, since the firing rate of its afferent nerve fibers depends not only on the actual position and movements of a joint but also on whether the γ motoneurons are active. While it was held that signals from the muscle spindles do not reach consciousness, it has now been convincingly demonstrated that signals from the muscle spindles can reach consciousness and that they contribute to our kinesthetic sense. A simple demonstration to this effect was performed by vibrating the biceps muscle in a normal subject. **Vibration** is known to stimulate the primary sensory endings of muscle spindles (the stimulus consists of brief stretches of the muscle). The subject, who is blindfolded, feels that the forearm is moving downward even though no such movement is occurring—that is, there is an illusory extension movement at the elbow joint. This corresponds to a lengthening of the biceps muscle, and under normal circumstances would be the normal cause of an increased firing rate in muscle spindle afferents.

Reexamination of the properties of joint receptors showed a striking paucity of slowly adapting joint receptors (type 1) that are active in midrange positions of the joint, the range in which the precision of kinesthesia is best. Because most type 1 receptors appear to reach their maximal firing rate only toward extreme joint positions, it seems unlikely that joint receptors alone can provide all the necessary information. Further, examination of patients with **artificial joints** who lack joint capsules (and thus, presumably, most of their joint receptors) show that their kinesthesia is only slightly reduced, at least with regard to the hip joint and the metacarpophalangeal joints. Presumably, muscle spindles (not tendon organs) are responsible for the remaining kinesthesia in such cases, even though skin receptors may contribute as well (particularly for the metacarpophalangeal joints). For the knee joint, however, elimination of

presumably all afferent signals from the joint capsule and the overlying skin by local anesthesia does not impair kinesthesia appreciably. Local anesthesia of finger joint capsules and the skin of the fingers provides more marked reduction of kinesthesia, but even in such cases the loss of kinesthesia is not complete.

Proprioceptors, Balance, and Voluntary Movements

Patients with neuropathies causing loss of thick myelinated fibers in their peripheral nerves have problems with voluntary movements, especially when they cannot see the moving parts. They also depend on visual information to keep upright and perceiving the position of their body parts. Their problems must be caused by the loss of information from low-threshold mechanoreceptors, presumably mainly proprioceptors. Most disturbed are movements requiring **coordination** of several joints, such as slicing bread, hitting a nail with a hammer, unlocking a door, and so forth. The regulation of muscular force is inaccurate, and they have particular problems with maintaining a **constant force** for some time. Presumably, information from proprioceptors is necessary for a continuous upgrading of the central motor program, so that the commands issued to the muscles are adapted to the actual position of the body parts. Thus, the influence of gravity changes continuously during a movement, and so do the mutual forces exerted by the parts (e.g., between the arm and the forearm). The movements become inaccurate and insecure if the force of muscle contraction is not adapted to these changes, which may differ slightly every time we, for example, raise a glass of water. Such **proprioceptive feedback** is particularly important for small, precise movements, when unforeseen disturbances occur during the movement, and when we learn new movements. Vision can only partially compensate for the loss of proprioceptive information.

As mentioned, **kinesthesia** does not depend solely on information from proprioceptors; cutaneous receptors also contribute. Patients with peripheral **neuropathies** and loss of proprioceptive information usually also have reduced cutaneous sensation that may contribute to their disturbances of posture. Thus, specifying the individual contribution of muscle spindles, tendon organs, joint receptors, and cutaneous receptors to the control of voluntary movements is difficult. Further, when information of one kind is reduced, the patient will learn to rely more on information from other kinds of receptors. In addition, the analysis is complicated by the fact that the various proprioceptors differ regarding their contribution to kinesthesia in different joints, and this probably is true for control of voluntary movements as well. The brain extracts what it needs for motor control from the collective information provided by all these receptors. Nevertheless, it is striking that loss of proprioceptive information is poorly compensated: the person becomes permanently dependent upon the cumbersome use of

visual information to control posture and voluntary movements.

We return to proprioceptors and their central actions in Chapter 14, and discuss their relation to motor control in Chapters 18 and 21.

Clinical Examples of Loss of Somatosensory Information

In his book *The Man Who Mistook His Wife for a Hat*, the neurologist Oliver Sacks gives a vivid description of a young woman, Christina, who completely lost kinesthetic sensation. A sensory neuropathy of unknown origin deprived her suddenly of virtually all kinds of proprioceptive information. Her cutaneous sensation was only slightly reduced, and motor axons were essentially spared. Nevertheless, at first she could not stand without continuously watching her feet. She could not hold anything in her hands, and they wandered around without her awareness. When stretching out to grasp an object she usually missed it—the movement stopped too soon or too late. “Something awful’s happened, I can’t feel my body. I feel weird—disembodied,” she said, and “I may ‘lose’ my arms. I think they’re one place and I find they’re another.” After having proprioception explained, she said: “This ‘proprioception’ is like the eyes of the body, the way the body sees itself. And if it goes, as it’s gone with me, it’s like the body is blind . . . so I have to watch it—be its eyes. Right?”

Another example concerns a 36-year-old man who gradually lost both cutaneous and kinesthetic sensation of the extremities due to a sensory neuropathy (described by Rothwell and coworkers 1982). His muscle power was hardly reduced, and he did surprisingly well on several routine tests of motor function. He performed, for example, various finger movements that require cooperation between muscles in the forearm and the hand. He could move his thumb with fair precision over three different distances and with three different velocities, and he could judge reasonably well the resistance to a movement. In spite of this, his hands were almost useless in daily life. He could not hold a cup with one hand, hold a pen and write, or button his shirt. Most likely, this can be explained by lack of automatic adjustment of ongoing movements and by an inability to maintain constant muscle force for more than a few seconds (without seeing the part). The problems seemed to arise also because he was unable to do longer sequences of simple movements without constantly watching what he was doing.

THE SENSORY FIBERS AND THE DORSAL ROOTS

Afferent (sensory) fibers from the receptors follow the peripheral nerves toward the CNS. Close to the spinal cord, the sensory fibers are collected in the **dorsal roots** and enter the cord through these (see Fig. 6.5 and 13.12).

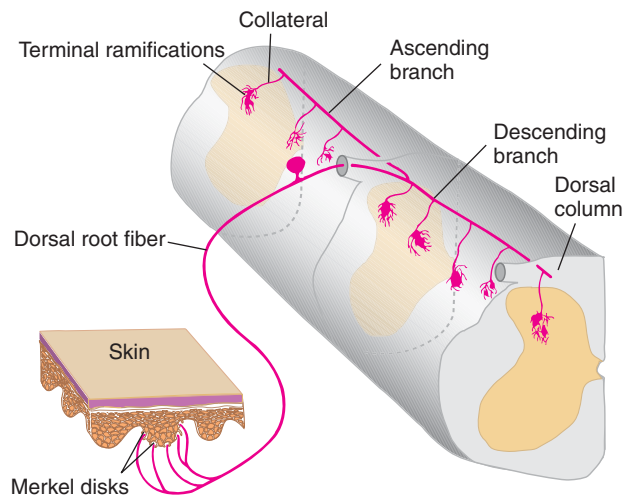


FIGURE 13.12 *Terminal pattern of a dorsal root fiber.* A dorsal root fiber (in this case conducting from Merkel disks) divides into an ascending and a descending branch after entering the cord. These branches give off several collaterals that end in the dorsal horn. The piece of the cord shown is about 1 cm long, but the axon continues beyond this in both directions. Corresponding reconstructions have been made for sensory units leading from several other kinds of receptors, and each sensory unit has a characteristic terminal pattern in the dorsal horn. (Based on Brown 1981.)

The sensory fibers of the spinal nerves have their cell bodies in the dorsal root **ganglia** (see Figs. 6.5 and 6.9). Likewise, the sensory fibers in the cranial nerves have their cell bodies in ganglia close to the brain stem (see Fig. 27.5).

As mentioned, the spinal ganglion cells are pseudounipolar (see Figs. 1.5 and 12.2) and send one long process peripherally, ending freely or in encapsulated sense organs. Functionally and structurally, both the peripheral and the central processes are axons. The **central process** enters the cord and then divides into an ascending and a descending branch (Fig. 13.12). These branches give off several collaterals ventrally to the gray matter of the cord. One sensory neuron, entering the cord through one dorsal root, can therefore influence spinal neurons at several segmental levels of the cord.

Classification of Dorsal Root Fibers in Accordance with Their Thickness

The dorsal root fibers vary in thickness, from the thickest myelinated ones, with a diameter of 20 μm and conduction velocity of 120 m/sec, to the thinnest unmyelinated fibers, with a diameter of less than 1 μm and conduction velocity of less than 1 m/sec. The thick fibers belong to the ganglion cells with large cell bodies, and the thin fibers belong to those with small cell bodies (Fig. 13.13). We have previously in this chapter described classification of sensory axons from muscle by their thickness (conduction velocity) into groups I to IV (see under “Classification of Muscle Sensory Fibers”).

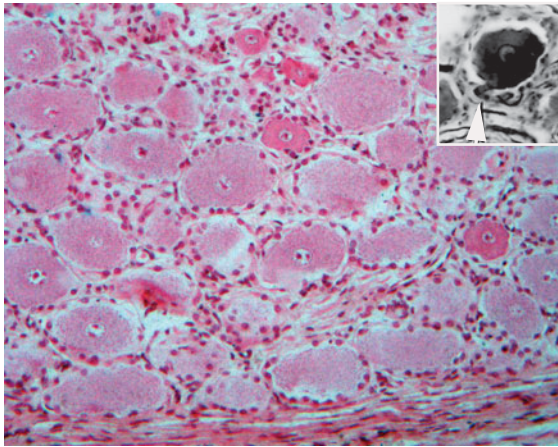


FIGURE 13.13 *Spinal ganglion.* Photomicrograph of a section from a human spinal ganglion. The cell bodies of the pseudounipolar neurons are of different size. The cell bodies are surrounded by satellite cells. **Inset:** Photomicrograph of section treated with a silver impregnation method showing axons. Arrow points to the axon leaving the cell body of the ganglion cell.

Another classification of sensory fibers is used mainly for cutaneous afferents. Myelinated fibers fall within **group A**, and **group C** contains the unmyelinated fibers. In the A group, the fastest conducting (thickest) fibers are termed $A\alpha$, somewhat slower conducting fibers are $A\beta$, and the thinnest of the myelinated fibers are called $A\delta$.

The different kinds of sensory receptors are supplied with axons of characteristic thickness. Signals from **low-threshold mechanoreceptors** are conducted in thick myelinated fibers ($A\alpha$ and $A\beta$); signals from **cold receptors** are conducted in thin myelinated fibers ($A\delta$), while unmyelinated (C) fibers conduct from **heat receptors**. Signals from **nociceptors** are conducted in $A\delta$ and C fibers. In the spinal cord, the terminations of the $A\delta$ and C fibers are almost completely separated from those of the $A\alpha$ and $A\beta$ fibers (Fig. 13.16).

Are There Sensory Fibers in the Ventral Roots?

We now know that there are exceptions to the rule that sensory fibers enter the cord through the dorsal roots and that motor fibers enter through the ventral roots (the law of Magendie). The fact that cutting the dorsal roots—dorsal **rhizotomy**—does not always abolish pain indicates that not all sensory information passes through the dorsal roots. Electron microscopic investigations have proved the presence of many **unmyelinated fibers** in the ventral roots of several species, including humans. Many of these are efferent, preganglionic autonomic fibers, but physiological data show that others (30% in the rat) are sensory and react to stimulation of **nociceptors**. However, few, if any, of these unmyelinated fibers have been proved actually to enter the cord through the ventral root.

With retrograde transport methods, their cell bodies have been shown to lie in the **dorsal root ganglia**. Some ventral root sensory fibers leave the ventral root to innervate the pia at the ventral aspect of the cord; others reverse direction before they enter the cord. Some of these then enter the cord through the dorsal root, whereas the further course of the others is unknown. Neither do we know whether these peculiar arrangements have a special biological meaning.

Fiber Categories and Conscious Sensations

Relationships between signals conducted in sensory fibers of various size (thickness) and conscious sensations have been investigated primarily via graded electrical stimulation of peripheral nerves and selective blockage of axonal conduction. By electrical stimulation of peripheral nerves, the weakest stimulation evokes activity only in the thickest myelinated fibers, and with increasing intensity, the thinner fibers are recruited progressively. Thus, the thickest fibers have the lowest electrical threshold for activation. In human subjects, **pain** is evoked by such stimulation only if the stimulus is strong enough to activate **$A\delta$ fibers**. The person then typically reports that the pain is of a sharp, pricking quality. If the stimulus strength is increased to recruit **C fibers** as well, the person experiences an intense, often burning pain that continues after the stimulus stops. These experiments are in agreement with the common experience that one usually can distinguish two phases of pain after an acute injury. The first phase, or **fast pain**, is experienced immediately after the stimulus, is well localized, and not very intense; the second phase, or **slow pain**, occurs with a longer latency and is more unpleasant, is not well localized, and usually continues after the end of the stimulus. The slow pain is delayed because it depends on being conducted in C fibers with a conduction velocity of less than 1 m/sec. The difference in conduction velocity between fibers that give rise to fast and slow pain is most easily observed when something hits the foot hard enough to cause pain—for example, when a toe is bumped against a hard object. The pathway from the toe to the cerebral cortex is longer than from any other part of the body, so the time lag between the signals conducted in thick and thin fibers is greatest. In fact, the very first sensation is only that something touched the foot, due to activation of low-threshold mechanoreceptors. Almost simultaneously, the sharp and well-localized fast pain is perceived, and we then know that the pain will soon be worse—the diffuse, burning, and intensely unpleasant slow pain continuing for some time.

Injection of **local anesthetics** around a peripheral nerve blocks the thinnest (C) fibers first and the thickest myelinated ones last. Accordingly, with local anesthetics the pain disappears first, whereas the tactile sensation remains considerably longer, and some sensations may

remain throughout the period of anesthesia. When peripheral nerves are subjected to **pressure**, conduction in thick fibers is blocked first, and, accordingly, there first occurs a reduction of the ability to perceive light touch and to judge the position of joints, but pain perception is still present.⁸

The Segmental Innervation: The Dermatomes

The **ventral branches** (rami) of the spinal nerves form plexuses supplying the arms (see Fig. 21.1) and the legs. Each nerve emerging from these plexuses contains sensory and motor fibers arising from several segments. In the peripheral distribution of the fibers, however, the segmental origin of the fibers is retained. Thus, sensory fibers of one spinal segment—that is, of one dorsal root—supply a distinct part of the skin. The area of the skin

supplied with sensory fibers from one spinal segment is called a **dermatome**. In the thorax and abdomen, dermatomes form circular belts; in the extremities, however, conditions are more complicated (Figs. 13.14 and 6.15). Knowledge of the segmental innervation of the skin (and also of the segmental innervation of muscles and viscera, dealt with in Chapters 21 and 27) is of great practical value in clinical neurology. For example, if a dorsal root is interrupted, the skin sensation is reduced or abolished in the corresponding dermatome.

Because the dermatomes **overlap**, each spot on the skin is innervated by sensory fibers from at least two dorsal roots.⁹ Interruption of a single dorsal root therefore may not produce a clear-cut sensory deficit. Nevertheless, careful examination usually shows a narrow zone (centrally in the dermatome) where the cutaneous sensation

8 For local anesthetics, which act by blocking voltage-gated Na⁺ channels in the axonal membrane, access to the channels is more direct in unmyelinated fibers than in thick, myelinated ones. Pressure affects primarily thick myelinated fibers, presumably because of their high oxygen demand that make them more vulnerable than unmyelinated fibers to ischemia.

9 Animal experiments indicate that the dermatomes are about twice as large as what appears from ordinary testing conditions, and the overlap is correspondingly larger. Inhibitory propriospinal fibers in the tract of Lissauer (Fig. 6.15) seem to inhibit weak sensory inputs from the peripheral parts of the dermatome, so that only signals from its central parts evoke a sensation. Under pathological conditions, however, the spinal neurons may become hyperexcitable and therefore respond to sensory inputs that normally are too weak to activate them.

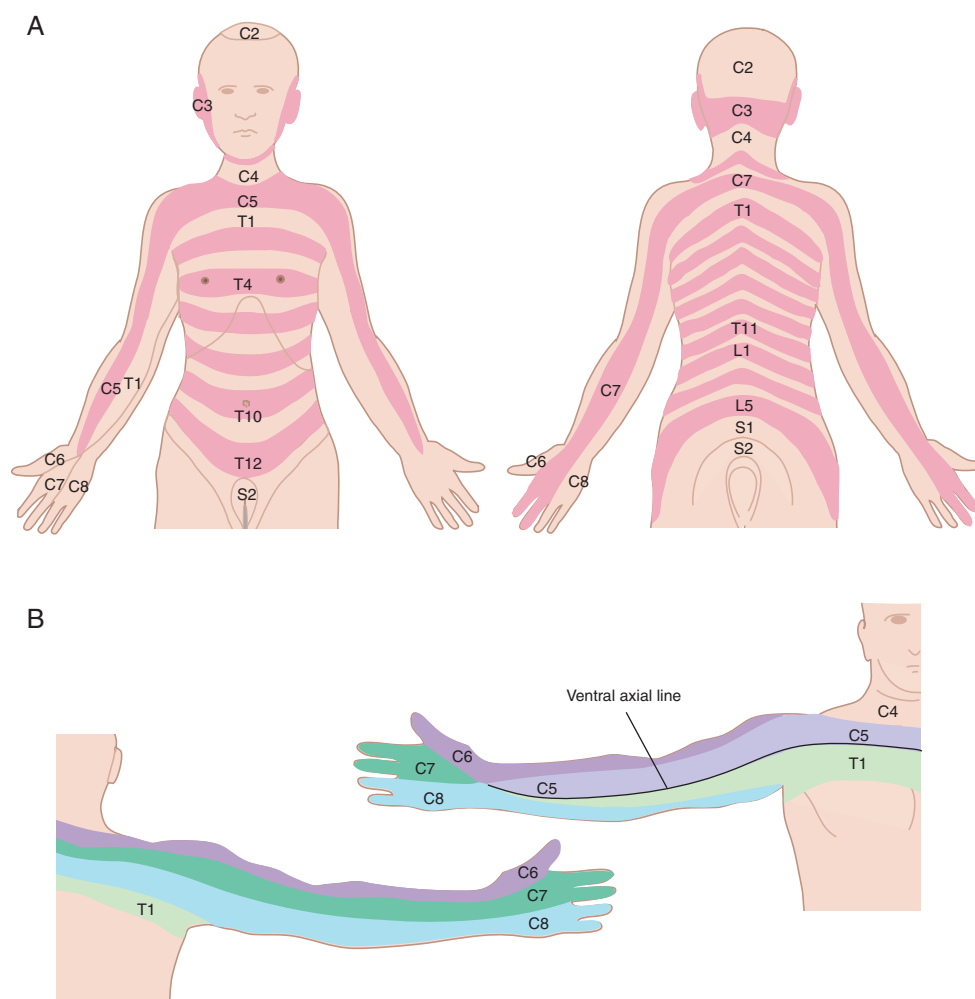


FIGURE 13.14 Dermatomes of the trunk and the upper extremity. Dermatomes not supplied by neighboring spinal segments meet at the ventral axial line (C₅ and T₁). The map gives a false impression of the borders between the dermatomes; in reality neighboring dermatomes overlap considerably. (From Keegan and Garrett 1948.)

to touch is slightly reduced and that for pain is abolished (analgesia). The usually more marked reduction in pain than in touch sensation is due to less extensive overlapping of fibers coming from nociceptors than of fibers coming from low-threshold mechanoreceptors.

When a dorsal root is subjected to **irritation**, as may occur by compression or stretching in connection with growth of an intraspinal tumor or protrusion of an intervertebral disk, this can cause pain and other sensory phenomena (numbness, pricking, tingling, and so forth) in the vicinity of the dermatome. Often the symptoms are felt only in smaller parts of the dermatome. With a protruding (herniated) intervertebral disk in the lumbar spine, for example, most often the roots of the fifth lumbar or first sacral nerves are affected, and the pain is felt in the leg (**sciatica**).

All dermatomal maps are composites of many single observations; no more than one or a few dermatomes have been determined in any single person. For this reason, all maps showing dermatomes for the whole body are approximations, not taking into account, for example, the considerable individual variations that exist. This, together with the fact that different methods have been used, probably explains why the dermatomal maps of different authors vary so much. For the student the main emphasis should therefore be on learning the main features of the dermatomal distribution rather than the artificial (and falsely defined) borders indicated on the maps.

How the Dermatomes Have Been Determined

The oldest method for determining the dermatome is to follow the distribution of the nerves by dissection. To follow the course of fibers from a root through the plexuses is, of course, far from easy. Certain diseases may affect single dorsal roots and produce changes restricted to the dermatome. **Shingles** (herpes zoster), for example, is a viral infection of the spinal ganglion cells that produces skin eruptions in the dermatome of the affected dorsal roots. Examination of many patients with this disease served as a basis for maps showing the dermatome (Head 1920). **Electrical stimulation** of dorsal roots during operations (Foerster 1933) and comparison of observations during operations for herniated intervertebral disks with the information given previously by the patient of where the pain and sensory loss were localized also help determine the location of dermatomes. **Local anesthesia** of single or several dorsal roots in healthy volunteers has also been of value. The best method is to eliminate impulse conduction in several dorsal roots on each side of one that is left intact (**method of remaining sensibility**). Sherrington (1898) did this experimentally in monkeys but the results are not directly applicable to humans. The German neurosurgeon Foerster (1933) made more

sporadic observations based on the method of remaining sensibility in patients in whom the dorsal roots were cut to relieve pain.

The dermatomal map presented here (Figs. 13.14 and 13.15; reproduced from Keegan and Garrett 1948) is based on observations of a large number of patients with **root compressions** (usually due to a herniated intervertebral disk) and, in addition, on examination of the distribution of reduced sensation in volunteers who had been subjected to local anesthesia of dorsal roots. The skin regions with reduced sensation (**hypoesthesia**) were carefully mapped out before operation and during operation it was determined which root was affected. Local anesthesia of a dorsal root also produces a sensory loss that is much less extensive than the total distribution of sensory fibers of the root. Thus, the borders between dermatomes as presented in Figs. 13.14 and 13.15 are imaginary. They ignore, for example, the great overlap between neighboring dermatomes, as well as the fact that the dermatomes are much wider than the zones of hypoesthesia occurring after damage to one dorsal root.

Fibers from Different Receptors End in Different Parts of the Dorsal Horn

The thin ($A\delta$ and C) and thick ($A\alpha$ and $A\beta$) dorsal root fibers end almost completely separated in the cord (Fig. 13.16). This indicates that afferents from nociceptors and low-threshold mechanoreceptors make monosynaptic contacts on different neuronal populations, as confirmed by recordings from single units in the dorsal horn. Nevertheless, the extensive dendrites of spinal neurons (see Fig. 6.12) and numerous interneurons enable convergence of signals from different receptor types. Thus, in the dorsal horn, some neurons are **modality specific**, whereas others integrate signals from different kinds of receptors.

Thin $A\delta$ and C fibers conducting signals from **nociceptors** end almost exclusively in the dorsalmost parts of the dorsal horn, in **laminae I and II** (**substantia gelatinosa**), but to some extent the $A\delta$ fibers also terminate in lamina V. Low-threshold receptors around the joints send signals mainly to lamina VI. Signals from cutaneous **low-threshold mechanoreceptors** activate mostly neurons in deep parts of the dorsal horn—that is, in **laminae III to V**.¹⁰ Signals from low-threshold mechanoreceptors

10 Combination of physiological and anatomic techniques verified the differential terminal patterns of dorsal-root fibers. Further, such experiments made it possible to study in detail the termination of individual sensory units, as exemplified in Fig. 13.12. Single axons in the dorsal root were penetrated with thin glass microelectrodes (pipettes). After determination of the receptive field and adequate stimulus of the sensory unit, a tracer substance was injected in the axon with same pipette. The axon and its ramifications were subsequently traced in serial sections of the spinal cord. A remarkable degree of specificity exists in the pattern of termination of fibers that belong to functionally different receptors.

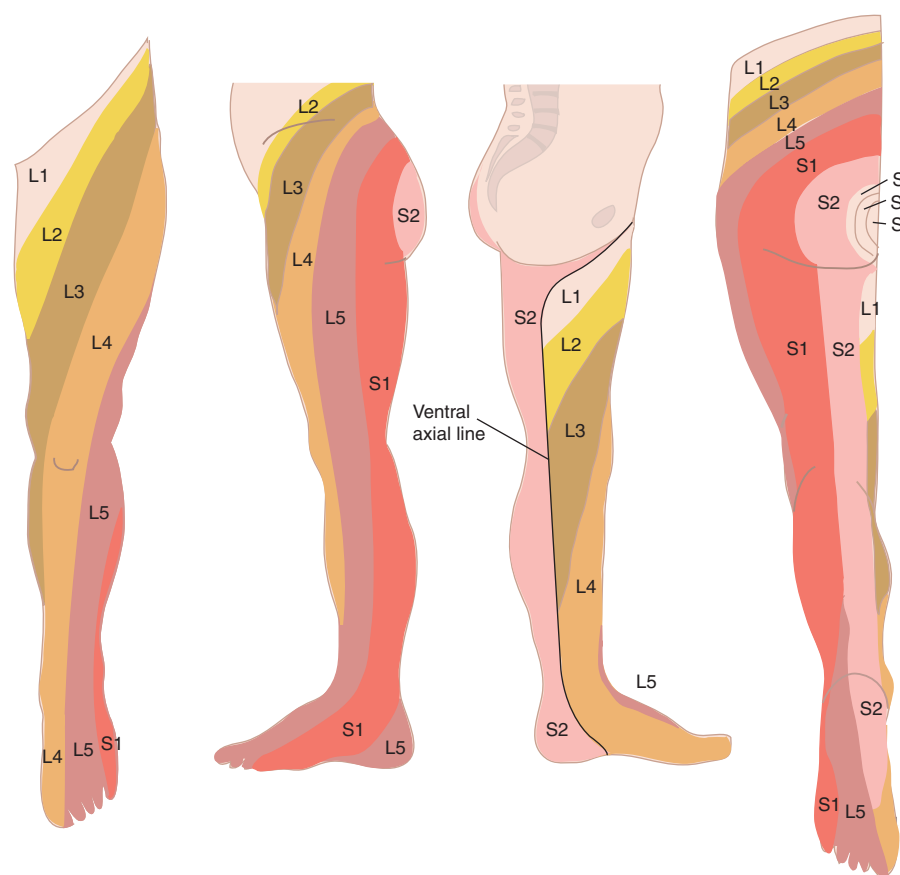


FIGURE 13.15 *Dermatomes of the lower extremity.* (From Keegan and Garrett 1948.)

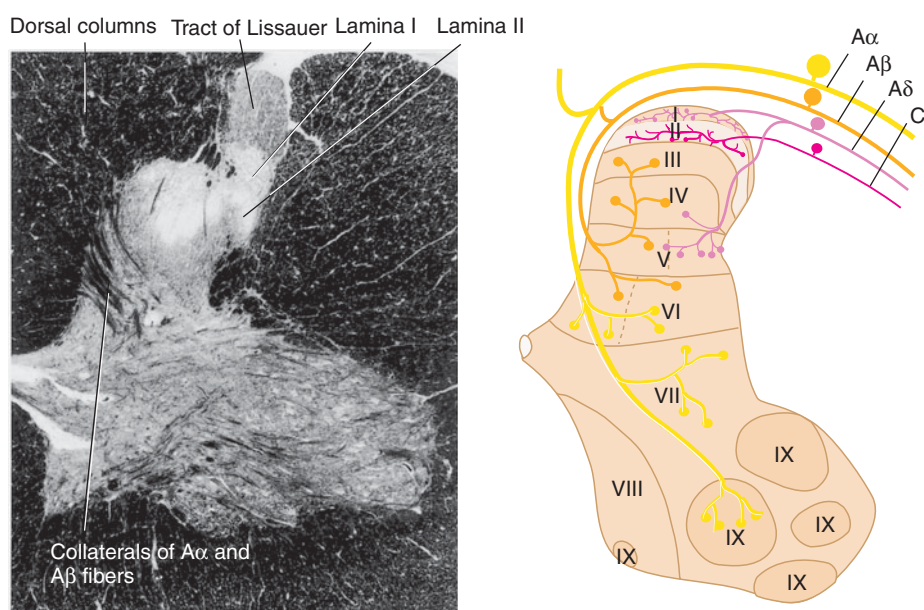


FIGURE 13.16 *Terminal regions of the dorsal root fibers in the cord.* **Left:** Photomicrograph of myelin stained transverse section through the cervical cord. **Right:** Schematic drawing. The thickest myelinated fibers ($A\alpha$, from muscle spindles and tendon organs) end in the deep parts of the dorsal horn and partly also in the ventral horn. Thick, myelinated fibers from cutaneous mechanoreceptors ($A\beta$) end in laminae III–VI. The thinnest myelinated and unmyelinated dorsal root fibers ($A\delta$ and C)—many of them leading from nociceptors—end in laminae I, II, and parts of V. Based on experiments with axonal transport of tracer substances.

and from nociceptors (and thermoreceptors) follow different routes from the dorsal horn to the cerebral cortex, as we will discuss in Chapter 14.

Thin Sensory Fibers from Muscles, Joints, and Viscera: Nociception and Homeostasis

Whereas the thin afferent fibers (the majority coming from nociceptors and thermoreceptors) from the **skin** end mainly in **laminae I and II**, corresponding fibers from the **viscera** appear to end almost exclusively in **laminae I** (and, to some extent, lamina V), thus avoiding the substantia gelatinosa. Thin **muscle** and **joint** afferents appear to terminate in the same parts of the dorsal horn as the fibers from viscera (although there are some conflicting data). Another feature of afferents from muscles and joints is their extensive rostrocaudal distribution in the cord. For example, afferent fibers from a single **facet joint** of the back terminate in seven to eight segments of the cord (cat). It is a common experience that pain of visceral origin has different qualities than pain evoked from the skin; visceral pain is much more diffuse and difficult to localize. In addition, pain that arises in muscles and joints is less precisely localized than cutaneous pain and radiates out from the site of the noxious stimulus. We also mentioned that muscle pain has a cramp-like quality. Presumably, the anatomic arrangements in the dorsal horn may contribute to such differences.

Lamina I of the dorsal horn has attracted interest because of its possible role in **homeostasis**. Due to the unique convergence of thin sensory fibers (especially C fibers) from virtually all tissues of the body—many of them chemoreceptors—lamina I neurons may monitor the metabolic status of the organism (such as the concentration of lactic acid and other metabolites produced in working muscles). At higher levels of the CNS, information from lamina I neurons most likely elicits appropriate autonomic and endocrine adjustments, while increasing afferent activity is perceived as discomfort and pain. The latter feelings are, of course, strong “recommendations” to the brain to change behavior (so that homeostasis is reestablished).

Primary Sensory Fibers and Neurotransmitters

Probably all primary sensory neurons release a classical transmitter with fast synaptic actions in the spinal cord. Among other evidence, this is based on the observation that boutons originating from dorsal root afferents contain small, clear vesicles, which have been shown in other parts of the nervous system to contain this kind of neurotransmitter. It has been estimated that at least 70% of all dorsal root fibers release an excitatory amino acid transmitter. Both ionotropic and metabotropic **glutamate receptors** are present in the dorsal horn, with

the highest concentration in the dorsalmost laminae (I and especially II). **NMDA** receptors have attracted much interest because of their possible role in development of central sensitization in chronic pain. There is also evidence that some primary sensory fibers release **ATP** (probably together with glutamate), exerting fast, excitatory synaptic actions.

Several **neuropeptides** are present in the central and peripheral terminals and in the cell bodies of spinal ganglion cells, as shown with immunocytochemical techniques. These include **substance P (SP)**, **vasoactive intestinal polypeptide (VIP)**, **cholecystokinin (CCK)**, **somatostatin**, **calcitonin gene-related peptide (CGRP)**, **galanin**, and others. Many ganglion cells contain more than one neuropeptide; for example, 80% of all SP-containing cells contain CGRP as well. The neuropeptides probably always colocalize with a classical transmitter with a fast, excitatory action. The peptides appear to mediate slow, modulatory synaptic actions in the dorsal horn, probably largely by acting on **extrasynaptic receptors** (see Fig. 5.1). When applied locally in the dorsal horn, SP and CGRP increase the release of glutamate. Further, SP receptors (neurokinin receptors) and *N*-methyl-D-aspartate (NMDA) receptors (glutamate) interact, making the NMDA receptors more sensitive to glutamate. Release of SP in the dorsal horn might therefore enhance and prolong the excitation produced by incoming signals from sensory receptors. We return in Chapter 15 to how this phenomenon may relate to increased pain sensitivity.

A relationship seems to exist between the kind of **tissue** a neuron innervates and its neuropeptide content. For example, many more of the ganglion cells innervating viscera contain SP and CGRP than do those cells innervating the skin. About two-thirds of the ganglion cells supplying joints contain SP and CGRP. The evidence so far is too limited, however, to draw conclusions regarding relations between neuropeptides, sensory modalities, and tissue or organ specificity.

Neuropeptides in Spinal Ganglion Cells and Nociceptors

Most of the neuropeptides are found only in **small ganglion cells** (Fig. 13.13) which have thin axons; these are probably mainly C fibers but also some A δ fibers. Substance P, for example, is present in most of the small ganglion cells (in about 20% of all, large and small together). The **neuropeptide receptors** are concentrated in **laminae I and II** (with the exception of CGRP receptors). These data suggest that the neuropeptides are particularly involved in transmission from **nociceptors** and **thermoreceptors**. Correspondingly, several neuropeptides and their receptors in the dorsal horn are up- or down-regulated in conjunction with **inflammation**, **nerve injury**, and **enduring pain**. Especially SP and its receptors

(**neurokinin 1**, or **NK-1**) are likely to be involved in processing of signals from nociceptors. Thus, SP is released from dorsal root fibers in the dorsal horn on nociceptor activation, and microinjection of SP in the dorsal horn make neurons more susceptible to sensory stimulation. Blocking NK-1 receptors prevents this effect.¹¹

Inflammatory Diseases and Release of Neuropeptides from Peripheral Branches of Sensory Neurons

The function of neuropeptides present in the peripheral ramifications of primary sensory neurons is less understood than their functions in the central terminals. We do not know, for example, whether these peptides are released under normal circumstances and take part in the normal homeostatic control. We know, however, that the peptides can be released in the peripheral tissue by **noxious stimuli** and by **antidromic activation** of the axon (see Chapter 29, under “Antidrome Signals and the Axonal Reflex”). Several of these peptides, when released in the tissues, have profound effects on vessels, as shown in the skin and mucous membranes. **SP** and **VIP** both produce vasodilation, and thereby increased blood flow and extravasation of fluid from the capillaries leading to edema. Furthermore, SP can activate cells of the immune system, resulting in phagocytosis and release of inflammatory mediators. Inhalation of irritating gases may provoke release of SP from peripheral sensory fibers in the **airways**, and the same takes place in the **skin** upon strong mechanical stimulation, such as scratching. The liberation of SP in such cases is probably due to an axonal reflex because afferent signals from the receptors are transmitted not only toward the spinal cord but also distally in branches of the sensory fibers (that is, distally in the branches that were not stimulated).

Injury to the nerve or the innervated tissue changes the neuropeptide content of the ganglion cells. For example, **experimental arthritis** leads to a marked increase of SP and CGRP in the cell bodies of the ganglion cells that innervate the affected joint (the arthritis is produced in animals by injecting the joint with a local irritant). SP enters the synovial fluid, and induces release of such substances as prostaglandins and collagenase

from leukocytes. Such substances may contribute to the damage of the joint cartilage in arthritis. Blocking the SP receptors (NK-1) or depleting the nerves of SP with **capsaicin** reduces the inflammatory reaction. Therefore, peripheral release of neuropeptides is now believed to play a role in several human diseases, including **rheumatoid arthritis**, **asthma**, **inflammatory bowel disease**, and **migraine**.

Sensory Fibers Are Links in Reflex Arcs: Spinal Interneurons

Sensory information reaching the spinal cord through the dorsal roots is further conveyed to higher levels of the CNS. In addition, many of the spinal neurons that are contacted by dorsal root fibers are not links in ascending sensory pathways but have axons that ramify within the cord—that is, they are **spinal interneurons**. The axons of these interneurons establish synaptic contacts with other spinal neurons, among them **motoneurons** (see Fig. 6.6) and **sympathetic neurons** in the intermediolateral cell column (see Fig. 3.8), giving origin to efferent fibers to smooth muscles and glands. In this manner, **reflex arcs** (see Fig. 21.9) for several important somatic (skeletal muscle) and autonomic (visceral) reflexes are established. Most, if not all, spinal interneurons also establish connections between neurons at different segmental levels (propriospinal fibers). Each spinal interneuron thus establishes synaptic contacts with a large number of other neurons in the spinal cord. Signals entering the cord through one dorsal root may influence neurons at several segmental levels, by both their own ascending and descending collaterals and their influence on interneurons with propriospinal collaterals (Fig. 13.12; see also Fig. 21.10).

How far the signals from one dorsal root fiber spread from interneuron to interneuron depends on the other synaptic influences these interneurons receive. For example, **descending connections** from the brain can selectively facilitate or inhibit spinal interneurons. This enables the impulse traffic from dorsal root fibers to be directed so that certain reflex arcs are used, whereas others are “switched off,” in accordance with the need of the organism as a whole. **Presynaptic inhibition** is an important mechanism in this respect (see Fig. 4.7). For example, separate groups of interneurons mediate presynaptic inhibition of group I muscle afferents, group II muscle afferents, and group Ib tendon organ afferents. Spinal reflexes are treated in more detail in Chapter 21.

11 Although SP is clearly associated with nociception, the correlation is not absolute (as judged from studies combining physiological and immunocytochemical characterization of single spinal ganglion cells). Thus, an SP-containing ganglion cell is not necessarily nociceptive, and many nociceptive neurons do not contain SP.

14 | Central Parts of the Somatosensory System

OVERVIEW

There are two major somatosensory pathways, both consisting of **three neurons** forming a chain from the receptors to the cerebral cortex (Fig. 14.1). The first, the **primary sensory neuron**, has its cell body in a spinal ganglion or in a cranial nerve ganglion; the next, the **secondary sensory neuron**, has its cell body in the gray matter of the spinal cord or in the brain stem; and the third, the **tertiary sensory neuron**, has its cell body in the thalamus. Both somatosensory pathways are **crossed**, so that signals from one side of the body are brought to the cerebral hemisphere of the other side. The actual crossing over takes place at different levels for the two pathways, however (Fig. 14.1). Another important point is that the pathways are **somatotopically organized**, which implies that neurons that conduct signals from different parts of the body are kept separate.

Whereas axons conducting from different kinds of receptors lie intermingled in the peripheral nerves and the dorsal roots, they are grouped according to their thickness as soon as they enter the spinal cord. The thick dorsal root fibers ($A\alpha$ and $A\beta$) pass medially, whereas the thin ones ($A\delta$ and C) follow a more lateral course into the dorsal horn. Largely, then, neurons conveying signals related to low-threshold mechanoreceptors and nociceptors (and thermoreceptors) are kept separate in the spinal cord, as previously discussed. This segregation is maintained in the pathways that lead from the cord to higher levels.

The medially located, **thick dorsal root fibers** continue without synaptic interruption rostrally in the **dorsal columns** (dorsal funiculus), without synaptic interruption in the cord. The first synaptic interruption occurs in the **dorsal column nuclei**, which contain the cell bodies of the secondary neurons in this pathway. The secondary axons cross in the medulla to end in the **thalamus** on the opposite side, forming the so-called **medial lemniscus**. From the thalamus, the tertiary neurons send their axons to the **primary somatosensory cortex (SI)** in the postcentral gyrus. Together, these three links constitute the so-called **dorsal column–medial lemniscus pathway**. From the prevalence of thick fibers conducting from low-threshold mechanoreceptors, it follows that the dorsal column–medial lemniscus pathway is important

for perception of **touch, pressure, vibration, and kinesthesia**, but it is of primary importance for the **discriminatory aspects** of sensation—that is, the ability to distinguish differently placed and different kinds of stimuli. The pathway appears not to be necessary for the mere perception of, for example, light touch or movement of a joint.

The central pathway followed by signals conducted in the **thin dorsal root fibers** differs from that followed by the thick fibers. The thin fibers make synaptic contacts in the gray matter of the dorsal horn, where most of the secondary sensory neurons of this pathway are located. The axons of the secondary neurons cross to the other side of the spinal cord and form the **spinothalamic tract**. As the name implies, the fibers of this tract terminate in the thalamus. The spinothalamic tract is primarily important for the perception of **pain and temperature**, which is consistent with the observation that it transmits information mainly from $A\delta$ and C dorsal root afferents. (There are also some other, less specialized, pathways capable of transmitting somatosensory information.)

As mentioned, the sensory signals conducted in the medial lemniscus and the spinothalamic tract finally reach the **somatosensory cortex (SI)**. Here, the body is represented **somatotopically** with the face most laterally on the convexity and the foot on the medial side of the hemisphere. In addition, the spinothalamic tract sends signals to several other parts of the cerebral cortex (notably the anterior cingulate gyrus and the insula). Sensory information reaching the SI is subject to some processing before being forwarded to the **posterior parietal cortex** for further analysis and integration with other sensory modalities. In addition, many fibers pass from SI to the precentral motor cortex, contributing to control of voluntary movements.

CENTRAL SOMATOSENSORY PATHWAYS

Chapter 13 dealt with the peripheral parts of the somatosensory system—the receptors and the primary sensory neurons. We now turn to the tracts and nuclei conveying and processing somatosensory information within the central nervous system (CNS). The term “somatosensory pathways” is not entirely appropriate, however, because these pathways transmit signals not only from somatic

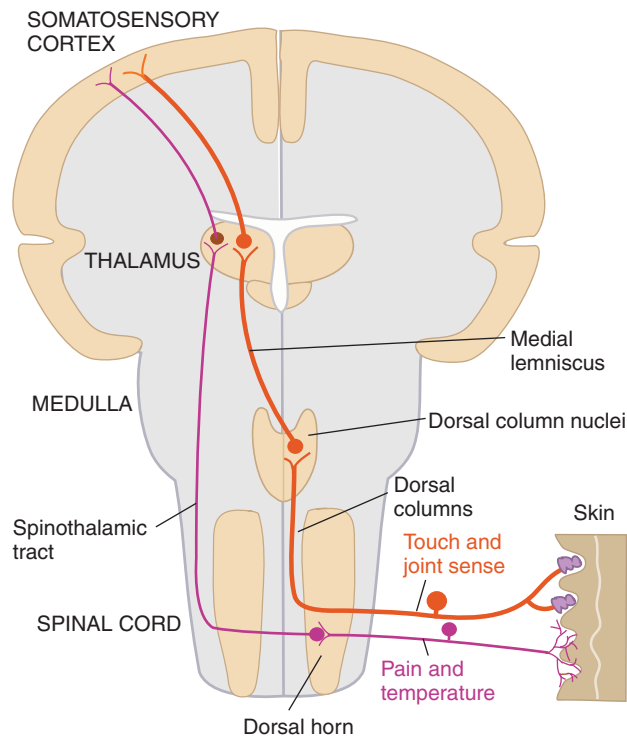


FIGURE 14.1 *Somatosensory pathways.* Highly simplified to show the main features of the two major pathways: the medial lemniscus–dorsal column pathway and the spinothalamic tract. The two pathways cross at different levels and differ in the sensory modalities they mediate.

structures, such as skin, muscles, and joints, but also from internal (visceral) organs. Most signals from internal organs are not consciously perceived, and visceral sensory processes have been less intensively investigated than somatosensory ones. We treat sensory information from the internal organs in Chapter 29.

Before we describe the somatosensory pathways in more depth, a few basic features of the thalamus need to be emphasized.

Lesion and Stimulation Experiments Are Used to Determine the Functions of Cell Groups

A fundamental approach to the study of the function of specific parts of the nervous system is to make circumscribed lesions in animals and to observe the functional disturbances that ensue. Such a lesion may constitute interruption of fiber tracts, destruction of neurons within a nucleus, or removal of large parts, such as a whole lobe of the cerebral hemisphere. It is also possible to **cool circumscribed regions** reversibly, so that neurons are “silenced” only temporarily. Corresponding reversible effects can be obtained with the use of local anesthetics. To study the functional and behavioral changes, highly sophisticated test methods may be necessary. Also, control experiments with lesions of other parts are usually crucial.

Examples of **lesion experiments** in animals and observations in **humans with brain damage** are mentioned throughout this book. The interpretation of the association between the normal function of the structures and the symptoms that ensue after lesions is often far from straightforward, however. For example, a lesion may destroy not only a certain group of neurons but also fibers passing through the area. In such cases, dysfunction of neuronal groups distant from the lesion may produce the symptoms. The interpretation is generally least problematic when lesions are confined to large, well-delimited tracts, whereas symptoms after lesions of the cerebral cortex may be much more difficult to interpret. A fundamental problem remains, however, in all such experiments: to what extent can normal function of a part of the brain be deduced from the deficits and disturbances resulting from its removal? In many instances, the symptoms occurring are generally a consequence of dysfunction and compensations of cell groups not damaged by the lesion.

Electrical stimulation of tracts and specific cell groups can elucidate their functions, assuming that the physiological and behavioral effects are closely related to their normal function. One example is electrical stimulation of the motor cortex in the precentral gyrus, which elicits more or less isolated muscle contractions in the opposite side of the body. This example also illustrates the limitations of such methods: the experiments tell us that the motor cortex is important for the start of movements, but they tell us very little about how the complicated pattern of activity in many muscles—which is characteristic of our voluntary movements—comes about. The difficulties of interpretation become much greater when regions with multifarious connections with other parts of the brain are stimulated (this may be particularly obvious for stimulation experiments of the so-called limbic system; see Chapter 31, under “Behavioral Effects after Damage or Stimulation of the Amygdala”).

Stimulation experiments often have to be performed on anesthetized animals, so this further limits the conclusions about normal cell function. Many cell groups are much less excitable during general anesthesia. This problem can be overcome by the use of chronically **implanted electrodes**, which enable stimulation to be performed in conscious animals. Such electrodes are inserted in the brain under general anesthesia and are fixed to the skull to remain in place (the electrodes cause no pain because the nervous tissue is devoid of nociceptors). The behavioral effects of stimulation of specific parts can then be observed repeatedly. After the experiments, the exact location of the electrode can be verified histologically. Such experiments also entail problems of interpretation. After all, the evoked activity is artificial, and one usually cannot know whether identical patterns of activity occur naturally.

The Thalamus: Relay Station for Sensory Pathways

All pathways conducting sensory information from the receptors to the cerebral cortex (except the olfactory pathways) are synaptically interrupted in the thalamus. In addition, the thalamus has a decisive influence on the general level of neuronal activity of the cerebral cortex and thus on the level of consciousness and attention. The macroscopic appearance of the thalamus is described and illustrated in Chapter 6 (see Figs. 6.22, 6.24, and 6.27). Three major subdivisions, delimited by the Y-shaped **internal medullary lamina**, can be identified macroscopically (Fig. 14.6; see also Fig. 6.24): an **anterior** nuclear group (or complex), a **medial** nuclear group, and a **lateral region** or part made up of a **dorsal** and a **ventral** nuclear group (Fig. 14.6; see also Fig. 33.7). Within and close to the internal medullary laminae are several less clearly defined groups of neurons, called the **intralaminar thalamic nuclei**.¹ These are of particular interest because of their relation to the thalamic influence on consciousness and sleep (this is further described in Chapter 26). The intralaminar nuclei are probably also important for the perception of pain.

Each of the three major thalamic subdivisions can be further subdivided into smaller nuclei based on cytoarchitectural differences. These are called the **specific thalamic nuclei**, because most of them are relays in precisely organized, major pathways that reach only certain parts of the cerebral cortex. The various specific nuclei have different functional tasks, and they receive fiber connections from the somatosensory nuclei, the retina, the nuclei of the auditory pathways, the cerebellum, the basal ganglia, and some other cell groups. As a rule, each nucleus receives afferents from only one of these sources. The somatosensory pathways terminate in the ventral nuclear group, as is dealt with in more detail later in this chapter.

The Dorsal Columns and the Medial Lemniscus

The thick, myelinated fibers in the medial portion of the dorsal roots curve rostrally within the **dorsal columns** (funiculi) just after entering the cord. Many of these fibers ascend to the dorsal column nuclei in the medulla (see Figs. 6.16 and 6.19), where they terminate and establish synaptic contacts (Fig. 14.1). As the fibers ascend in the dorsal columns, they send off collaterals ventrally to the spinal gray matter (see Fig. 13.12). Most of these collaterals establish synaptic contact with interneurons, but some reach as far as the ventral horn

and contact α motoneurons (primary afferents from muscle spindles).

The fibers occupying the medial part of the dorsal columns—the **gracile fascicle**—conduct signals from the lower part of the trunk and the legs. These fibers end in the **gracile nucleus** (Fig. 14.2). Signals from the upper part of the trunk and the arms are conducted in the lateral part of the dorsal columns, the **cuneate fascicle**. The fibers of the cuneate fascicle terminate in the **cuneate nucleus**. Why the longest fibers of the dorsal columns lie most medially is explained by the simple fact that they enter the cord at the lowermost level, where no other long ascending fibers are present. At higher levels, fibers entering from the dorsal root occupy positions lateral to those that have entered at more caudal levels. Initially the fibers of the dorsal columns are arranged **segmentally**, but as they ascend, the fibers rearrange themselves so that they are organized **somatotopically**—that is,

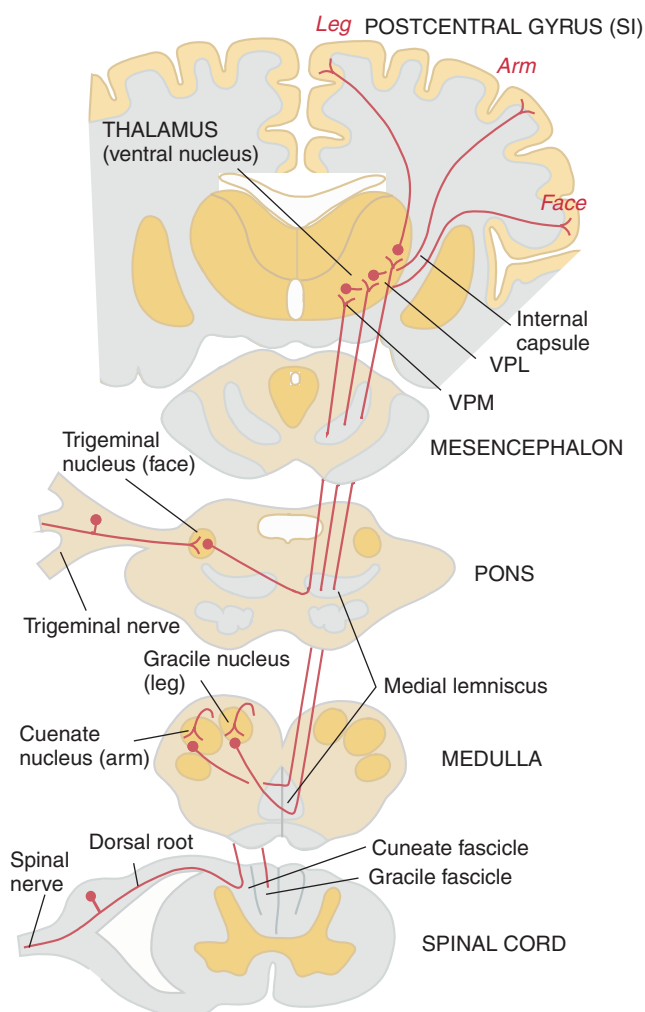


FIGURE 14.2 The dorsal column–medial lemniscus pathway. This is the main pathway for transmission of signals from low-threshold mechanoreceptors. Fibers leading signals from mechanoreceptors in the face join the medial lemniscus in the brain stem.

¹ The intralaminar thalamic nuclei were formerly called the “unspecific thalamic nuclei” because their connections with the cerebral cortex were thought to be diffusely distributed, in contrast to the connections of the specific thalamic nuclei. More recent studies have questioned the nonspecific nature of the intralaminar nuclei, however.

fibers conducting from the hand lie together, separated from those of the forearm, and so on (Fig. 14.2). Thus, fibers from different dorsal roots are mixed at higher levels of the dorsal columns.

The primary afferent fibers ascending in the dorsal columns end in a particular cytoarchitectonic subdivision—the **cluster region**—of the dorsal column nuclei. The morphology and arrangement of the neurons in the cluster regions ensure a particularly precise topographic arrangement of the afferent and efferent connections. The neurons of the cluster regions of the dorsal column nuclei send their axons rostrally to the thalamus, forming the **medial lemniscus**. The fibers first course anteriorly and cross the midline to occupy a position just dorsal to the pyramid (Fig. 14.3; see also Fig. 6.17). In the pons and the mesencephalon, the medial lemniscus is placed more laterally (Fig. 14.3; see also Figs. 6.18 and 6.20).

The medial lemniscus ends in the **ventral posterolateral nucleus, VPL** (Figs. 14.2 and 14.6). The fibers of the medial lemniscus are **somatotopically** organized,² and this pattern is maintained as the fibers terminate in the VPL. Fibers from the gracile nucleus (sensory signals from the leg) terminate most laterally, with fibers from the cuneate nucleus (arm) terminating more medially. Most medially, in a separate nucleus, the **ventral posteromedial nucleus (VPM)** ends the fibers from the sensory trigeminal nucleus (relaying signals from the face; Fig. 14.6).

Neurons of the dorsal column nuclei that send their axons in the medial lemniscus most likely use **glutamate** as transmitter (like most other long, precisely organized tracts in the CNS).

The Dorsal Columns Contain More than Primary Afferent Fibers

Many of the **primary afferent** fibers in the dorsal columns do not reach the dorsal column nuclei, even though they may pass through many segments—for example, from the lumbar to the cervical levels. Further, not all fibers of the dorsal columns are primary afferents. Thus, the dorsal columns contain a large number of axons from neurons located in the spinal gray matter. Most of these are **propriospinal** fibers extending just for one or a few segments to end in the cord (propriospinal fibers). Others, however, end in the dorsal column nuclei and are called **postsynaptic dorsal column neurons**. The possible relationship of these fibers to transmission of **nociceptive** signals is discussed later in this chapter (“Single-Unit Properties in the Dorsal Column–Medial

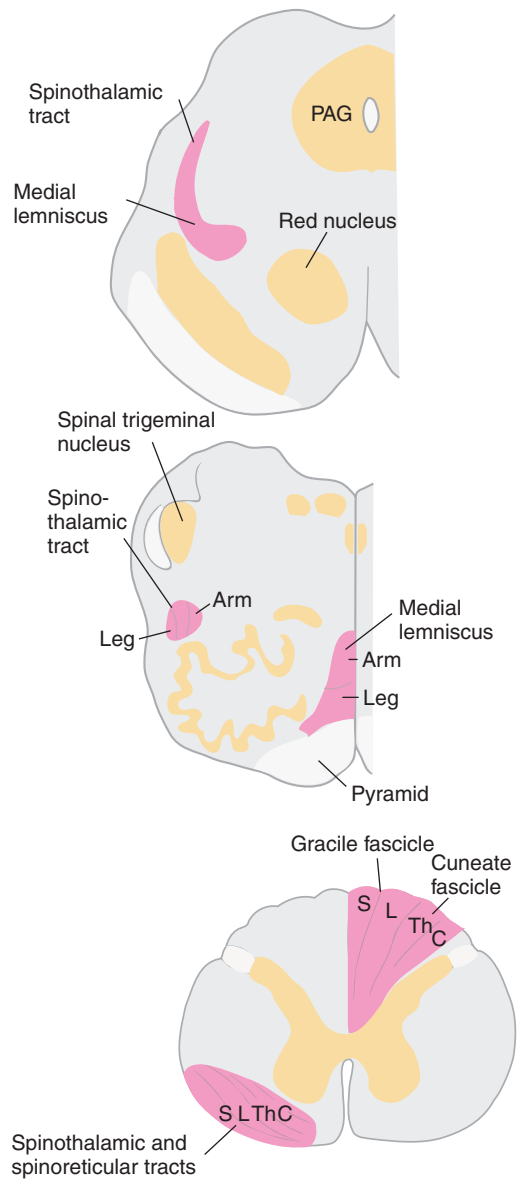


FIGURE 14.3 Somatosensory pathways. Position and segmental arrangement in the cord, medulla, and mesencephalon. The spinal cord is disproportionately large. Termination of collaterals of spinothalamic fibers in the brain stem are not shown (see Fig. 15.2).

Lemniscus System” and “Additional Pathways from Nociceptors”).

The dorsal columns also contain some **descending** axons making synaptic contacts in the dorsal horn (especially in lamina V, containing neurons excited by signals from nociceptors). These descending fibers come from neurons in a subdivision of the dorsal column nuclei (the reticular zone) that receives afferents from the cerebral cortex and the reticular formation. Thus, the dorsal column nuclei are part of a neuronal network that—by way of descending connections—control the flow of sensory information from the cord (see later in

² That fibers carrying signals from the leg are located more anteriorly within the medial lemniscus than are those related to the arm (Fig. 14.3) is simply because the gracile nucleus is situated more caudally than the cuneate nucleus. When fibers from the cuneate nucleus join those from the gracile nucleus, the anterior position is already occupied.

this chapter under “Transmission of Sensory Signals Is Controlled from the Brain”).

Thalamocortical Pathway to SI and SII

The neurons of the VPL and the VPM send their axons into the internal capsule (Fig. 14.2) and further through this to the **postcentral gyrus**. This part of the cortex, made up of cytoarchitectonic fields 3, 1, and 2 (after Brodmann; see Fig. 34.3), constitutes the **primary somatosensory area, SI** (Figs. 14.2 and 14.6). In addition, some fibers from the VPL and the VPM end in the **secondary somatosensory area, SII**, situated in the upper wall of the lateral cerebral fissure (Fig. 14.6). On **electrical stimulation** of SI or SII, conscious human subjects report sensory phenomena such as tingling, itching, numbness, and so forth. Just as the somatosensory pathways are **somatotopically** organized, this is also the case within SI and SII (Figs. 14.2 and 14.7). Fibers conducting signals from the leg end most medially within the postcentral gyrus, then follow fibers conveying signals from the trunk, arm, and face successively in the lateral direction.

Epileptic Seizures Demonstrate the Cortical Somatotopic Pattern

On irritation of the cortex within the postcentral gyrus—for example, by a chip of bone from a skull fracture—the patient may experience fits of abnormal sensations. In the same person, the fits have the same characteristic pattern each time: The sensations are felt in one particular part of the body and then spread gradually to other parts. The spreading follows the known somatotopic pattern within SI (see Fig. 14.8). For example, the patient may first experience a tingling sensation in the thumb; then it moves to the index finger and the other fingers; then to the forearm, upper arm, shoulder, and even further. Such epileptic seizures are called **Jacksonian fits** (after the famous British neurologist, Hughlings Jackson). They signify the presence of a local disease process of the brain, and the starting point of the abnormal sensations indicates the focus of the disease. Often the sensory phenomena are followed by muscle spasms (convulsions) due to spreading of the abnormal cortical electrical activity to the motor cortex of the precentral gyrus.

Single-Unit Properties in the Dorsal Column–Medial Lemniscus System

As mentioned, the dorsal columns contain primarily fibers coming from **low-threshold mechanoreceptors** in the skin, muscles, and joints. Recording the activity of single units in the dorsal columns has confirmed this and has shown that there is a predominance of **rapidly adapting** sensory units; relatively few are slowly adapting. They have small **receptive fields**, mostly at the **distal**

parts of the extremities. Recordings from neurons in the cluster regions of the dorsal column nuclei show that many neurons are activated by only one kind of receptor. Some are activated only by joint movements, others only by light touch of the skin, others only by vibration, and so forth. These neurons are called **modality specific** (because they only react to one kind of stimulus) and **place specific** (because they are activated only from one restricted part of the body).

Neurons in the VPL and SI also have the same characteristic response properties as those described for the neurons of the dorsal column nuclei, even though an increasing number of neurons are activated by more than one kind of receptor. In addition, the receptive field tends to be somewhat larger for neurons in SI than, for example, in the dorsal column nuclei.

Some of the axons that end in the dorsal column nuclei do not belong to primary sensory units but to neurons with their cell bodies in the dorsal horn (**post-synaptic dorsal column neurons**). Unexpectedly, these postsynaptic neurons are activated not only from cutaneous low-threshold receptors but also from **visceral nociceptors**. We return later in this Chapter to these special dorsal-column sensory units and their possible role in nociception (see under “Additional Pathways from Nociceptors”).

Functions of the Dorsal Column–Medial Lemniscus System

Most of the axons at all levels of the dorsal column–medial lemniscus system are thick and rapidly conducting. This, together with the data from single-unit recordings mentioned above, enable us to conclude that the dorsal column–medial lemniscus system is particularly well suited to bring fast and precise information from the skin and musculoskeletal system about the type of stimulus, the exact site of the stimulus, and when the stimulus starts and stops. Thus, it provides information about the **sensory quality** and the **spatial and temporal characteristics** of any stimulus of low intensity (“what,” “where,” and “when”). The next question is then: how is this information used by the CNS? Unfortunately, on this point conclusions are largely based on the deficits observed when the system is not working. Further, because of the adaptive changes taking place after an injury, we have to distinguish between the acute and long-term functional deficits. Other problems of interpretation arise because with incomplete lesions, functional deficits may be revealed only by tests that require full use of the system.³

3 For example, experiments in cats show that sparing as little as 10% of the fibers of the dorsal columns—as compared with a lesion comprising all fibers—markedly reduces the sensory deficits with regard to, for example, the discrimination of surfaces with different roughness.

Most studies (with some exceptions) indicate that in monkeys, as in humans, **acute damage** to the dorsal columns produces severe **ataxia** (insecure and incoordinate movements), which recedes partly or completely within weeks to months after the damage. In some patients, the ataxia may be so severe that they cannot walk without support. Observations some time after the damage indicate, however, that the dorsal column–medial lemniscus system is not necessary for all aspects of cutaneous sensation and kinesthesia. First, temperature and pain perception are unaltered by lesioning the dorsal columns; second, light touch of the skin can easily be felt, as can passive joint movements. Two-point discrimination may not be appreciably reduced, and some reports even indicate that the ability to recognize objects by manipulation may be retained (clinical observations do not support the latter point, however). What appears to be consistently impaired is the ability to solve tasks that require spatially and, in particular, temporally very accurate sensory information. Thus, a coin pressed into the palm of the hand may perhaps be recognized, but the patient is unable to decide which is the larger of two coins. The patient may also correctly identify that something is moving on the skin, but not the direction of the movement. To ask the patient to identify figures written on the skin, for example, is one sensitive test of the function of the dorsal column–medial lemniscus system. Further, some careful clinical observations indicate that the perception of joint position and movement is abnormal after lesions of the dorsal columns.

The above-described sensory deficits occurring after lesions of the dorsal columns have in common that they concern spatial and temporal comparisons of stimuli, or what we call **discriminative sensation**. Such sensory information is crucially important for the performance of many **voluntary movements**; indeed, disturbances of voluntary movements are characteristic of the lesions that affect the dorsal column–medial lemniscus system. After the acute phase, the movement deficits first concern movements that require fast and reliable feedback information from the moving parts. For example, the ability to adjust the grip when an object is slipping is clearly reduced. Delicate movements, such as writing and buttoning, are performed only with difficulty after lesions of the dorsal columns. It is not possible to throw an object accurately or to perform a precise jump, presumably because such activities require feedback information from skin receptors to judge the pressure exerted on the hand by the object or by the ground against the sole of the foot.

In conclusion, the dorsal column–medial lemniscus system is of primary importance for complex sensory tasks, such as determination (comparison) of direction and speed of moving stimuli. Further, many precise voluntary movements—especially of the hand—depend on the fast sensory feedback provided only by this system.

In fact, after damage to the system, the motor deficits may be more disturbing than the purely sensory ones.

The Dorsal Columns and Kinesthesia

Observations in humans have provided conflicting results as to whether lesions of the dorsal column–medial lemniscus system give impaired kinesthesia. A thorough clinical study by Nathan and coworkers (1986, p. 1032), however, concluded that complete lesions of the dorsal columns do produce clear-cut and enduring kinesthetic deficits. But they emphasize that: “routine examination of tactile sensibility does not show up these defects as well as everyday activities of living. The further one gets away from this testing situation, the easier it is to see the effects of these disturbances of sensibility.” One example may illustrate this point: a patient with damage to the dorsal columns was aware of a toe being passively moved by the examiner; nevertheless, his shoe would easily slip off his foot without his noticing, and he was unable to roll over in bed because he did not realize that one leg was hanging off the bed.

In monkeys, Vierck and Cooper (1998) described deficient kinesthetic sensation of the hands after cutting the cuneate fasciculus, although only specific and detailed testing revealed the problems. Thus, the perception of passive finger movements was impaired only if the movements were small or slow. This is indeed what you would expect when eliminating a system devoted to very precise sensory information. It is furthermore worth noticing that the monkeys had more obvious problems with precise hand movements than with kinesthesia.⁴

Clinical Examination of the Dorsal Column–Medial Lemniscus System

Many of the deficits that occur after damage to the dorsal column–medial lemniscus system may not be revealed by a routine neurological examination. Nevertheless, they may render the patient severely handicapped in daily life. We described earlier in this chapter similar symptoms occurring after loss of thick myelinated nerve fibers in the peripheral nerves. This is not surprising because many of these fibers continue into the dorsal columns. The deficits may be more severe with peripheral loss, however, because reflex effects from low-threshold

4 In the legs, however, no defect of joint sense occurred in monkeys after lesions of the dorsal columns at thoracic or cervical levels. This can perhaps be explained by less rigorous testing of the legs than of the hands. Another explanation is possible however. Thus, primary afferent fibers conveying signals from slowly adapting low-threshold mechanoreceptors in leg muscles and joints leave the gracile fasciculus at the low thoracic level (in monkeys). Then they enter the dorsal horn, where they synapse on second-order sensory neurons. The axons of the latter continue in the dorsal part of the lateral funiculus, not in the dorsal columns. Thus, a lesion of the dorsal columns at cervical levels would not interrupt signals from leg proprioceptors. We do not know, however, whether this arrangement pertains also to humans.

mechanoreceptors (among other things) are also lost. Based on these considerations, a routine examination of kinesthesia (asking the patient to indicate the direction of movement in a joint being passively moved) would not provide definite information because such a test may be negative (i.e., normal performance) in spite of damage to the dorsal columns or the medial lemniscus. Neither is testing the sense of vibration a reliable source of information; several clinical studies show that vibration is not always reduced after damage to the dorsal columns. A routine testing of cutaneous sensation would not necessarily reveal the lesion. The most reliable information is presumably obtained by testing the ability to judge the direction of a stimulus to the skin and to examine the patient's ability to identify numbers written on the skin.

The Spinothalamic Tract

As mentioned, the fibers in the lateral portion of the dorsal roots are predominantly thin (A δ and C fibers). Immediately after entering the cord, such fibers divide into short ascending and descending branches, passing only for one or two segments in each direction. The collaterals and terminal branches of these fibers pass ventrally to end in the gray substance of the dorsal horn, where they establish synaptic contacts with other neurons (Figs. 14.4 and 14.5). The thinnest (largely C fibers) among the ascending and descending branches of the dorsal root fibers form a small bundle immediately dorsal to the dorsal horn, called the **tract of Lissauer**, or the *zona terminalis* (see Fig. 13.16).⁵

Many neurons in the dorsal horn send off long ascending axons. Most of these first course almost horizontally and somewhat ventrally across the midline through the gray substance (Figs. 14.1 and 14.4). After having entered the lateral funiculus on the opposite side, the fibers curve sharply in the rostral direction. As mentioned, some of these fibers ascend without interruption as far as the thalamus, thus forming the **spinothalamic tract**. Many of the spinothalamic fibers give off **collaterals** in the brain stem to nuclei involved in autonomic control (circulation, respiration) and in behavioral responses to signals from nociceptors. The majority of spinothalamic tract fibers are located in the **anterolateral quadrant** (funiculus), that is, anteriorly within the lateral funiculus (Fig. 14.3). Here the spinothalamic fibers intermingle with numerous propriospinal fibers and

⁵ There has been a long-standing controversy with regard to the fiber composition of the tract of Lissauer, largely because of difficulties in tracing unmyelinated fibers. Investigations with improved methods indicate that about 80% of the fibers of the tract of Lissauer are primary afferent (dorsal root) fibers. The other fibers are propriospinal fibers ensuring cooperation between neighboring segments. This concerns in particular Rexed's laminae I and II, which also receive most of the thin primary afferents.

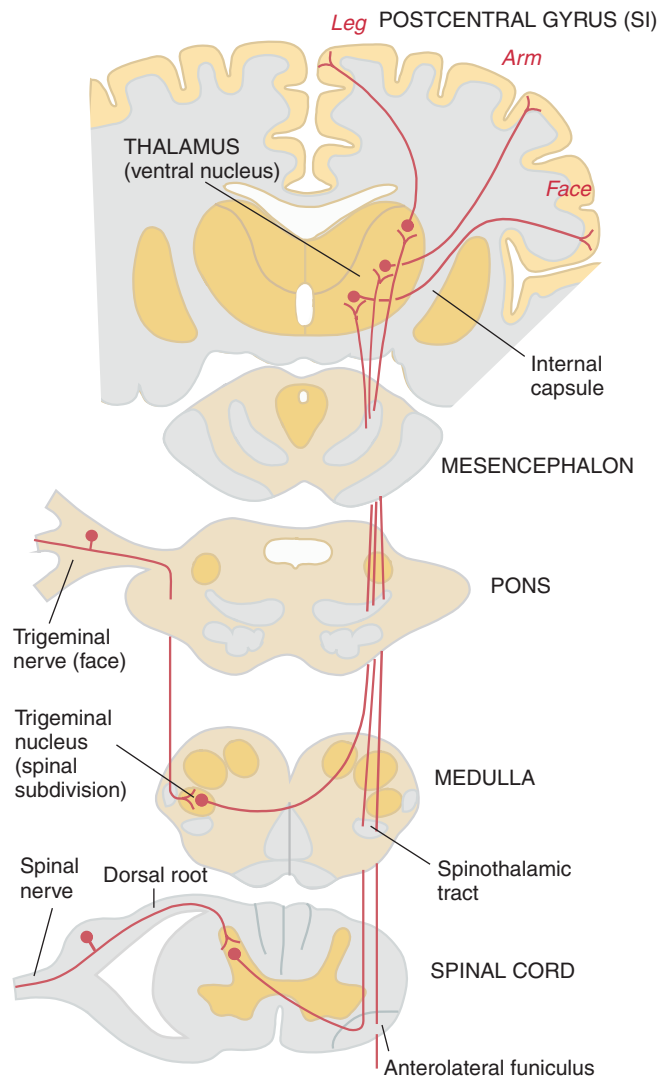


FIGURE 14.4 *The spinothalamic tract.* This is the main pathway for transmission of signals from nociceptors and thermoreceptors. Nociceptive and thermoceptive signals from the face join the spinothalamic tract in the brain stem (the spinal trigeminal nucleus).

fibers terminating in the brain stem. In the brain stem, the spinothalamic tract lies laterally and fairly superficially (Fig. 14.3).

Synaptic Coupling between Dorsal Root Fibers and Spinothalamic Neurons

The location and properties of spinothalamic neurons have been determined by use of axonal transport methods combined with physiological studies of anatomically identified neurons (Fig. 14.5). Particularly, conditions in monkeys may be expected to correspond closely to those in humans. The major group of spinothalamic cells is located in **lamina V**, particularly in those parts

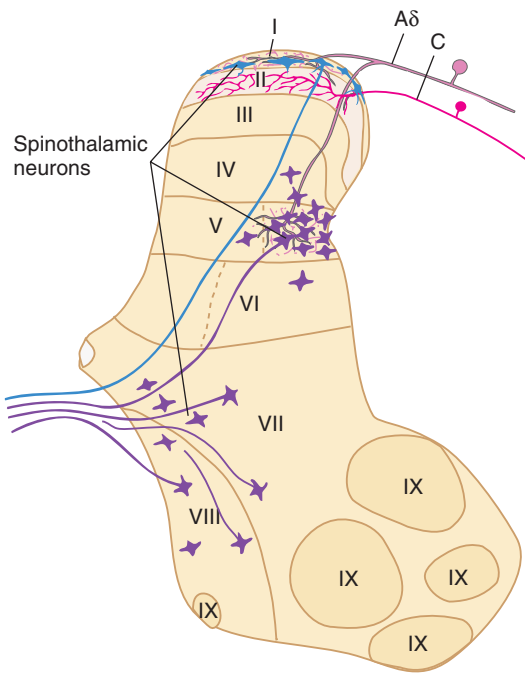


FIGURE 14.5 *Signal transmission from nociceptors in the spinal cord.* The figure shows the terminal regions of thin dorsal root fibers leading from nociceptors and the position of neurons sending their axons to the opposite thalamus (spinothalamic neurons). Lamina II (substantia gelatinosa) receives many C fibers, but the neurons of this lamina do not send their axons to the thalamus.

that receive A δ fibers from the dorsal roots. The other major population of spinothalamic cells are located in **lamina I**. When we recall that most of the A δ and C fibers establish synapses in lamina I and V (Fig. 14.5), the relationship between the spinothalamic tract and perception of pain becomes understandable. In addition, scattered spinothalamic cells are found more ventrally in **laminae VII and VIII** (Fig. 14.5).

The anatomic data indicate that many thin dorsal root fibers do not synapse directly (monosynaptically) onto the spinothalamic cells but influence them indirectly via spinal interneurons. The numerous small neurons in **lamina II**, which are contacted by dorsal root C fibers, send their axons in part dorsally into lamina I—where spinothalamic cells can be influenced—and in part ventrally in the dorsal horn to contact other interneurons, which in their turn form synapses on spinothalamic cells in laminae V, VII, and VIII. In addition, many of the neurons in lamina II send axon collaterals to spinal segments above and below their own (passing in the tract of Lissauer).

The synaptic arrangements underlying further transfer of signals transmitted in thin dorsal root afferents, many coming from nociceptors, must be complex. The large number of very small **interneurons** and the numerous neuroactive substances (e.g., numerous neuropeptides) in the dorsal horn make it a difficult task to clarify how

the signals from nociceptors are processed. For example, hardly any of the neurons in lamina II (substantia gelatinosa) send fibers into the spinothalamic tract, yet they receive a large proportion of the dorsal root C fibers. Generally, the interneurons of the dorsal horn—and those of substantia gelatinosa in particular—will have a decisive influence on whether the signals from nociceptors will be transmitted to higher levels of the nervous system. These interneurons can both inhibit and facilitate the signal traffic that will ascend in the spinothalamic tract. Thus, they will strongly influence how intense the experience of pain will be on a certain stimulation of nociceptors. As we discuss later in this chapter, the CNS can control signal transmission from receptors to higher levels by descending fibers acting on the dorsal horn interneurons.

Properties of Spinothalamic Tract Units

Physiological studies of single spinothalamic cells and of thalamic cells in the terminal regions of spinothalamic fibers are in accord with their preferential activation from thin (A δ and C) dorsal root fibers.

The spinothalamic cells in the cord are classified by their response properties:

1. **Low-threshold units**—neurons that react only to light mechanical stimuli (e.g., light touch of the skin)
2. **Wide dynamic range units (WDR)**—neurons that react to stimuli of high intensity (activating nociceptors) and to light stimuli; the impulse frequency of these cells increases with increasing stimulus intensity
3. **High-threshold units (HT)**—neurons that respond only to stimuli of an intensity sufficient to activate nociceptors
4. **Thermosensitive units**—neurons that respond only to either warming or cooling of the skin, indicating that they are activated by thermoreceptors.

These properties, determined in animal experiments, agree well with clinical observations in patients who have been subjected to therapeutic interruption of the antero-lateral funiculus (containing the spinothalamic tract and other fibers). This operation, called **cordotomy**, is sometimes applied to alleviate intense pain that cannot be overcome in any other way. After such an operation, sensations of pain and temperature are almost totally abolished in the opposite side of the body in the parts supplied by sensory fibers from the spinal segments below the level of interruption (although only temporarily; see under “Additional Pathways from Nociceptors”).

Although both lamina I and lamina V contain a mixture of WDR and HT neurons, it appears that lamina I contains a majority of HT (nociceptive specific) neurons, whereas WDR-neurons dominate in the deeper parts of the dorsal horn.

Terminations of the Spinothalamic Tract and Further Projections to the Cortex

The **thalamic** termination site of the spinothalamic tract is more extensive than that of the medial lemniscus, and the same holds for the further signal transmission to the cerebral cortex. Many of the spinothalamic fibers end in the **VPL** with a **somatotopic** pattern (Fig. 14.4), but not in exactly the same parts as the fibers of the medial lemniscus (corresponding fibers from the spinal trigeminal nucleus end in the **VPM**). The terminal ramifications are also different for the two pathways, and nociceptor-activation of VPL neurons requires more summation than activation from low-threshold mechanoreceptors. In addition, many of the spinothalamic fibers arising in lamina I end more posteriorly in the **ventromedial nucleus** (VM). Spinothalamic fibers also end in parts of the **intralaminar nuclei** (e.g., the central lateral nucleus, CL), in the **mediodorsal nucleus** (MD), and some other nuclei.

The multiple terminations of spinothalamic fibers probably explain how signals from nociceptors, by way of thalamocortical fibers, can reach several regions of the cortex in addition to SI and SII. Recordings of single-unit activity in monkey SI indicate that neurons activated by high-intensity (presumably noxious) cutaneous stimuli are concentrated in a narrow zone at the transition between Brodmann's areas 3 and 1 (Fig. 14.6). This may primarily concern signals relayed through the VPL, whereas signals from other parts of the thalamus receiving spinothalamic fibers may directly and indirectly influence other cortical areas, such as the **anterior cingulate gyrus** and the **insula** (this is further treated later in this chapter,

under "Which Parts of the Cerebral Cortex Process Nociceptive Information?").

Does the Spinothalamic Tract Consist of Distinct Discriminative and Affective Parts?

The spinothalamic tract has been proposed to consist of two anatomically and functionally different components. One part—ending in the lateral thalamus (mainly in VPL and VPM) with further transmission to SI—would be responsible for the discriminative aspects of pain perception; that is, our ability to localize a painful stimulus and to judge its quality (sticking, burning, cramp-like, and so forth). The other part—consisting of fibers ending in the medial thalamus (especially in the intralaminar nuclei) and projecting on to the **insula**—would be responsible for the affective, emotional aspects of pain. There is much evidence, however, that this division is at least an oversimplification. For example, many spinothalamic fibers send collaterals to both lateral and medial thalamic nuclei (and to parts of the reticular formation in the brain stem). Furthermore, microstimulation of the lateral thalamus in humans (presumably in the ventromedial nucleus) evoked pain that could be precisely localized *and* at the same time evoked strong emotions (anxiety, discomfort).

Spinothalamic Cells Receive Signals from Both Somatic and Visceral Structures

Recording from spinothalamic cells in the spinal cord has shown that many can be activated by nociceptive stimuli applied to **visceral organs** and to the **skin**. Signals from

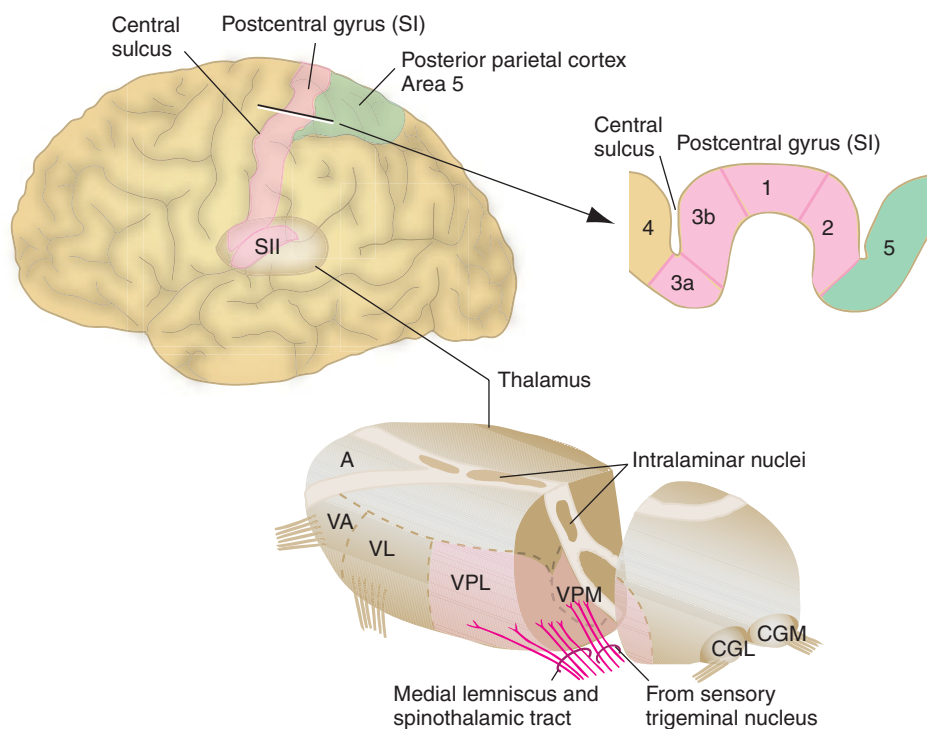


FIGURE 14.6 The somatosensory cortex (SI) and its thalamic afferent nucleus. **Upper left:** The location of the SI and SII and parts of the posterior parietal cortex in the left hemisphere. **Upper right:** The extension of the various cytoarchitectonic areas of the central region. **Below:** The VPL—supplying the SI and SII with somatosensory signals—is marked on a schematic drawing of the left thalamus. The main afferent pathways to the VPL are also indicated. A, anterior thalamic nucleus; Lg, lateral geniculate body; MG, medial geniculate body; VA, ventral anterior nucleus; VL, ventral lateral nucleus; VPM, ventral posteromedial nucleus; VPL, ventral posterolateral nucleus.

the skin and viscera converge onto the same neuron, which then conveys the information to the thalamus. Nociceptive signals from **muscles** and skin can also converge onto the same spinothalamic neuron. Sensory convergence of this kind pertains to regions of the skin and deep tissues and to visceral organs that receive sensory innervation from the same segments of the cord. The primary sensory fibers can activate the spinothalamic cells monosynaptically or through one or more interneurons (polysynaptically). In addition, in some spinal ganglion cells the peripheral process divides, with one branch innervating the skin and the other a visceral organ or a muscle.

The observations of convergence on spinothalamic cells and branching of sensory fibers may help explain why pain arising from a visceral organ is often felt as if it comes from the skin. This phenomenon is called **referred pain** and is discussed further in Chapter 29.

What Tracts in the Anterolateral Funiculus Can Mediate Alone: A Case History

A woman with a knife-stab partly severing the cord at the Th₃ level illustrates what the anterolateral funiculus alone can mediate of somatic sensations (Danzinger and coworkers 1996). The lesion, which was verified during open surgery, severed the cord completely except for the anterolateral fascicle and adjoining parts of the lateral funicle on the left side. Shortly after the accident, with some difficulty, she was able to perceive and localize light touch on both sides of the body below the lesion. She was able to perceive passive joint movements on the left side, but not vibration. Eighteen years after the lesion her sensibility was virtually unchanged. The lesion did not abolish control of the bladder and the rectum, and she retained some voluntary control of movements of the left leg (whereas the right leg was paralytic, she was, for example, able to lift the left leg 30° in the supine position).

Additional Pathways from Nociceptors

Other pathways than the spinothalamic tract may transmit signals from nociceptors to consciousness, even if their contribution may be minor under normal conditions. It should be emphasized that nociceptive signals reach numerous subcortical nuclei (in the rat apparently more than 20), many of which have ascending projections. Concerning transmission from **visceral nociceptors**, postsynaptic fibers in the **dorsal columns** seem to contribute in addition to the spinothalamic tract. There is also evidence that signals in the dorsal columns may contribute to chronic pain in some instances (this is further discussed in Chapter 29, under “Visceral Pain”). Further, many nociceptor-activated spinal neurons send their axons to brain stem nuclei that mediate automatic responses to noxious stimuli, such as changes of

blood pressure, heart rate, sweating, breathing, muscle tone, bladder and bowel function, and so forth. Parts of the **reticular formation**, the **PAG** (periaqueductal gray), and the **parabrachial nucleus** all receive signals from nociceptors and participate in automatic behavioral responses to nociceptor activation. Among these, the parabrachial nucleus is special since it receives spinal afferents mainly from **lamina I**, which contains many neurons specifically activated from nociceptors.⁶ In addition, these brain stem nuclei send fibers rostrally to end in the **amygdala**, **hypothalamus**, and **thalamus**. Such connections are probably involved in the varied affective, autonomic, endocrine, and behavioral responses to noxious stimuli but may also gain access to the cortical network responsible for pain perception. Whereas the spinothalamic tract is mainly crossed, these other pathways are bilateral. Thus, unilateral stimuli would reach the thalamus (and cerebral cortex) of both sides.

We know that nociceptive neurons are easily **sensitized** by prolonged stimulation. For example, recordings during brain surgery in humans with **chronic pain syndromes** revealed neurons in the reticular formation and thalamus that responded to noxious stimuli; while such responses were not found in the same regions in pain-free controls (the patients were undergoing brain surgery for movement disorders). These and other data strongly suggest that various cell groups and pathways—in addition to the spinothalamic tract—are recruited under special, poorly understood circumstances.

After interruption of the spinothalamic tract by **cordotomy**, pain usually returns in some months to a year. This probably depends, at least partly, on transmission of nociceptive signals in pathways other than the spinothalamic tract (nociceptor-activated postsynaptic units in the dorsal columns, spino-parabrachial-amygdala connections, and others). Presumably, after severance of the dominant spinothalamic pathway, other minor pathways, which normally do not contribute significantly to our pain perception, take over.

Homeostatic Surveillance: A Task of Ascending Tracts from Lamina I?

As mentioned, perception of pain and temperature are particularly dependent on the spinothalamic tract, although, as witnessed by clinical observations, this tract can also mediate a relatively crude sense of touch and pressure. In a broad perspective, pathways leading from nociceptors constitute an **alarm** system: they report that something is wrong in the body and a change

⁶ In the cat, the majority of lamina I neurons with an ascending axon terminate in the parabrachial nucleus and the PAG (twice as many in the parabrachial nucleus as in PAG). Only about one sixth appears to terminate in the thalamus. If this is so also in humans, it would underscore the role of lamina I in homeostatic control.

of behavior is required to avoid damage. Especially many neurons in **lamina I**—responding to a variety of potentially harmful tissue substances and to heat and cold—seem suited for this task. The system notifies impending cell damage regardless of cause (metabolites, inflammatory mediators, extreme temperatures, or mechanical forces), and is an integral part of the defense systems of the body. Pain elicits a **stress response** consisting of autonomic, somatic, and endocrine components aiming at restoring homeostasis. Activation of brain stem nuclei (parabrachial nucleus, PAG), the hypothalamus, and the amygdala elicits the stress response, either by ascending tracts carrying signals from nociceptors or by descending signals from the cortical pain network (or both). In daily life, most of the signals conveyed through A δ and C fibers do not reach consciousness: small adjustments prevent the sum of stimulation to reach a level sufficient to cause pain. We alter position, change clothing, or protect ourselves in other ways without needing to pay attention. Persons born without the ability to feel pain illustrate the importance of the alarm system: they incur frequent and serious injuries of various kinds such as burns, wounds, infections, overstretched joints, and fractures.

The Brain Controls the Transmission of Sensory Signals

Descending fibers from the cerebral cortex and the brain stem end in the relay nuclei of the somatosensory pathways. One important group of such connections arises in **SI** (the primary somatosensory area) and terminates in the **thalamus** (VPL and other nuclei), the **dorsal column nuclei**, the **sensory trigeminal nucleus**, and the **dorsal horn** of the cord (see Fig. 22.8). These connections are somatotopically organized and enable selective control of sensory signal transmission from particular parts of the body and from particular receptor types. Physiological studies indicate that descending connections from the SI sometimes can **facilitate** signal transmission through the sensory relay nuclei, but **inhibitory** effects appear to be most common. The latter effects are mediated via inhibitory interneurons. Generally, it appears that signals from somatosensory receptors are continuously regulated to adapt to changing needs—for example, whether sensory information is received passively or actively sought, whether it is needed for movement control, or just arise as a trivial result of self-initiated movements (see later, “Why We Cannot Tickle Ourselves”).

Recordings from single units of the medial lemniscus in conscious monkeys have shown reduced impulse traffic from cutaneous receptors immediately before a voluntary movement. There is indirect evidence of the same phenomenon in humans: the threshold for perceiving a vibratory stimulus is elevated immediately before a voluntary movement. Perhaps this happens because proprioceptive signals are of greater importance than cutaneous ones

in this particular situation. During walking, signals from mechanoreceptors in the sole of the foot are let through to the motor cortex immediately before heel strike, while the traffic is inhibited during most of the stance phase.

We know from daily life that we have the ability to leave out sensory signals that are irrelevant at the moment. Without such filtering mechanisms, we would be flooded by sensory information. The sensory information finally reaching the cerebral cortex is therefore “censored” and distorted compared with the stimuli received by the receptors.

Later in this chapter we deal specifically with descending control of transmission in the “pain pathways.”

Why We Cannot Tickle Ourselves

The fact that we are unable to tickle ourselves is an interesting special case of sensory-information control. Identical signals from cutaneous low-threshold mechanoreceptors can be perceived as tickling if they are caused by another person stroking our skin but as mere touch if a self-initiated movement causes them. In the latter situation, the activation of SI is reduced. Apparently, only signals that are unexpected or unpredictable are let through to SI without suppression. Presumably, experience has provided us with (subconscious) knowledge of the sensory signals that a certain motor command will produce. The cerebellum may have a role in such situations by informing the cerebral cortex about which somatosensory signals might be expected. Probably, the cerebellum receives a copy of the motor command issued from the motor cortex (**efference copy**).

THE SOMATOSENSORY CORTICAL REGIONS

As mentioned, the sensory signals conducted in the medial lemniscus finally reach the two somatosensory areas, SI and SII. In addition, the spinothalamic tract sends signals to several other parts of the cerebral cortex. Both SI and SII receive somatotopically-organized projections from the VPL and VPM (Figs. 14.7 and 14.8), transmitting signals primarily from low-threshold mechanoreceptors and, to a lesser extent, from nociceptors. Somatosensory signals also reach other cortical regions, however, such as the **motor cortex** (MI) in the precentral gyrus. Not unexpectedly, primarily signals from **proprioceptors** are conveyed to the motor cortex.

The parts of SI receiving sensory signals from the feet, hands, and face are much larger than those receiving signals from other parts of the body (Fig. 14.8). Further, the region devoted to the thumb is larger than that devoted to the palm of the hand, which, in turn, is larger than that devoted to the forearm, and so on. This is mainly a reflection of the much higher **density of sensory units** that supply the skin at distal parts of the extremities

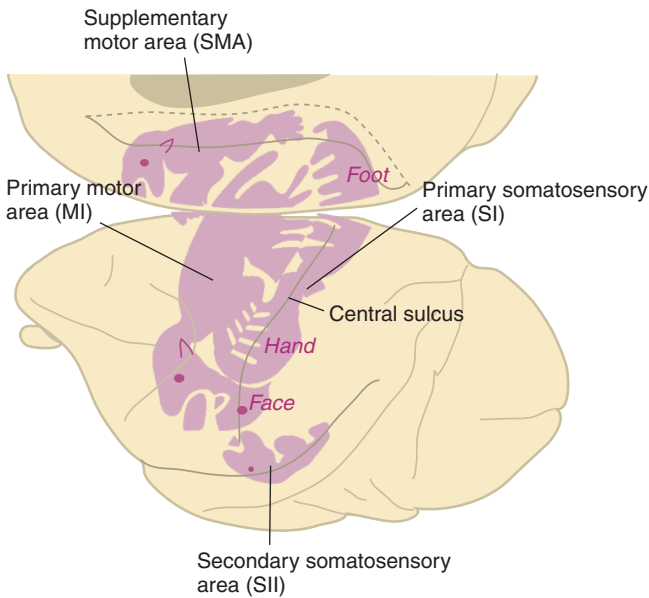


FIGURE 14.7 The somatosensory regions SI and SII and their somatotopic organization (monkey). Motor areas (MI and SMA) are also shown. (After Woolsey 1958.)

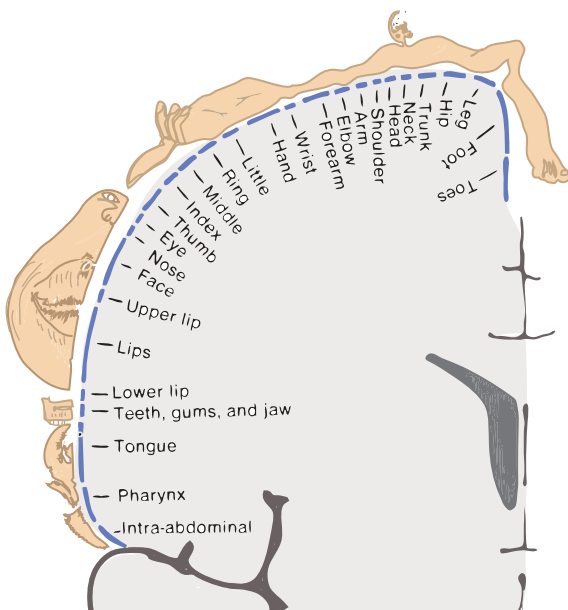


FIGURE 14.8 Relative size of the cortical regions representing various body parts. Schematic section through the postcentral gyrus (SI) of the human brain. Based on electrical stimulation during brain surgery under local anesthesia. (After Penfield and Rasmussen 1950.)

(and parts of the face) than more proximal parts of the trunk. To use this very detailed information from the most densely innervated parts of the body, a large volume of cortical gray matter—that is, many neurons—has to be available for information processing. A **magnification factor** gives a numerical representation

of the cortical representation of certain body parts. Similar overrepresentations exist within the visual and auditory systems.

Various **brain-imaging techniques**, such as fMRI and PET, have brought a wealth of information on the contribution of specific cortical areas in motor, sensory, and cognitive processes. We refer to results from such studies throughout this book. A brief description of these and other methods for study of the living human brain are found in Chapter 11 (see under “Methods to Study Neuronal Activity and Connectivity in the Living Brain”).

The Primary Somatosensory Area (SI)

The primary somatosensory area, in particular, has been the subject of intense anatomic and physiologic investigations. The subdivision of SI into different **cytoarchitectonic areas**—3, 1, and 2 of Brodmann (see Figs. 22.4 and 33.2)—corresponds to functional differences. These areas extend as narrow strips from the midline laterally along the postcentral gyrus: that is, perpendicular to the somatotopic arrangement (Fig. 14.6; see also Fig. 33.3). Animal experiments, particularly by the American neurophysiologist Mountcastle, show that the cytoarchitectonic subdivisions differ with regard to the kinds of receptor from which they receive information. **Area 3a**, on the transition to the MI (see Fig. 22.4), receives sensory signals from **muscle spindles** in particular.⁷ Neurons in **area 3b** are mainly activated by stimulation of **cutaneous receptors** (predominantly by low-threshold mechanoreceptors). Neurons receiving information from rapidly adapting receptors appear to be separated from those receiving from slowly adapting receptors. **Area 2** is influenced by **proprioceptors** to a larger extent than area 3b is; for example, many neurons are most easily activated by bending of a joint. Within each of the cytoarchitectonic subdivisions it appears that the whole body has its representation; thus there are probably three **body maps** within SI. The map in Fig. 14.7 therefore gives a somewhat misleadingly simplified presentation.

Figure 14.8 shows the representation of body parts based on electric stimulation of the cortical surface in conscious patients (during neurosurgery for therapeutic reasons). The patient tells where she feels something when a certain cortical site is stimulated. Although later studies overall agree with the findings of Penfield and Rasmussen (Fig. 14.8), an exception seems to concern the representation of the **genitals**, that most likely are represented on the convexity near the lower part of the

⁷ Some have argued that area 3a should be considered a part of the MI rather than of SI. In certain respects, anatomically and physiologically, area 3a represents at least a transitional zone. Like the other subdivision of SI, area 3a receives thalamic afferents from VPL, whereas MI (area 4) receives them from VL. The afferents from other parts of the cortex are more like those of MI than of the other parts of SI, however.

abdomen (and not on the medial aspect of the hemisphere close to the foot, as originally indicated by Penfield and Rasmussen).⁸

Even though many neurons in SI are activated only or most easily from one receptor type—that is, they are **modality-specific**—there are other neurons in SI with more **complex properties**. For example, many neurons have large receptive fields, indicating that they receive convergent inputs from many primary sensory neurons. Further, movement of just one joint in one direction activates some neurons, while other neurons are activated by several joints. Still other neurons in SI require specific combinations of receptor inputs to be activated. Thus, processing of the “raw” sensory information already begins at the first cortical stage; SI is not merely a simple receiver of sensory information.

Efferent **association connections** from SI pass posteriorly to the **posterior parietal cortex**, which processes the sensory information (see below), and anteriorly to the **motor cortex** (MI). The latter connections appear to be of particular importance while **learning** new movements, whereas they are not crucial for the performance of well-rehearsed movements (as judging from lesion experiments in monkeys). This may be explained by an extra need for fast and precise feedback from the moving parts during learning. The connections from SI to MI, furthermore, are necessary for motor **recovery** after cutting the connections from the cerebellum to MI, as discussed in Chapter 11 (see under “Examples of Substitution from Animal Experiments”). This may also be regarded as a learning situation.

Further Processing of Sensory Information Outside SI

Although processing somatic sensory information starts in SI, clinical and experimental observations show that the cortex posterior to SI is necessary for comprehensive utilization. The posterior parietal cortex comprises **area 5** and **area 7** (Fig. 14.6; see also Fig. 33.7) and belongs to the association areas of the cerebral cortex (these will be further discussed in Chapters 33 and 34). Areas 5 and 7 do not receive direct sensory information from the large somatosensory pathways but via numerous association fibers from SI and SII. They also receive numerous connections from other parts of the cortex. Broadly speaking, in areas 5 and 7 the bits of information reaching SI are put together and compared with other inputs, such as visual information and information about the salience of a stimulus and about intentions. Neurons in area 5 often have large receptive fields and

respond to complex combinations of stimuli, as shown in monkeys. Their activity depends not only on what is occurring in the periphery but also on whether the **attention** of the monkey is directed toward the stimulus. The posterior parietal cortex sends **efferent** connections to motor areas in the frontal lobe, thereby linking sensory information with **goal-directed movements**. Accordingly, some neurons are active in conjunction with the monkey stretching its arm toward something it wants.

In addition to the posterior parietal cortex, **SII** (Fig. 14.6) and adjoining areas in the **insula** (Fig. 14.9; see also Fig. 6.29) also process information from SI. The anterior part of insula integrates somatosensory information with other sensory modalities (taste and smell, and signals from vestibular receptors). Sensory units in these areas typically have large receptive fields and are activated from both sides of the body. Insula is, however, more strongly linked with processing of visceral sensory information and pain (see below, and Chapter 34, under “Insula”).

Symptoms after Lesions of the Somatosensory Areas

Lesions of **SI** in humans entail reduced sensation in the opposite half of the body. A localized destruction of the SI, or of the fibers reaching it from the thalamus, may produce loss of sensation in a restricted area (corresponding to the somatotopic localization within SI). Not all sensory qualities are affected equally, however. **Discriminative** cutaneous sensation and **kinesthesia** are particularly disturbed; much less reduced (if at all) is pain sensation. As is the case with lesions of the dorsal columns, the sensory deficits gradually diminish after the time of the lesion. The least improvement with time is seen in the discriminative aspects of sensation, whereas pain sensation improves considerably. This can perhaps be explained by the fact that the pathways for signals from nociceptors are to a larger extent bilateral than are the pathways from low-threshold mechanoreceptors.

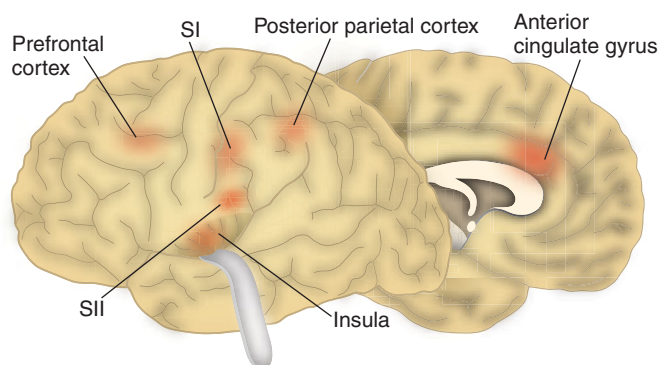


FIGURE 14.9 Regions of the cerebral cortex showing increased activity during pain perception. (Based on data from a meta-analysis published by Peyron et al. 2000.)

⁸ Representation of the genitals on the medial aspect of the hemisphere would be in conflict with the principle of continuous representation of body parts (Fig. 14.7). Indeed, a recent study using natural peripheral stimulation and fMRI (rather than stimulation of the cortical surface) concluded that the penis is represented on the convexity in the transition zone between the lower abdomen and the thighs (Kell et al. 2005).

It might also be explained by signals from nociceptors being distributed to areas outside the SI.

Whereas lesions of the **posterior parietal cortex** produce difficulties with the use of objects (**apraxia**), lesions of **SII** and neighboring regions in the **insula** impair the ability to recognize objects by touch (**tactile agnosia**). Similar problems with the visual recognition of objects occur after lesions of dorsal and ventral divisions of the extrastriate visual areas, respectively.

Which Parts of the Cerebral Cortex Process Nociceptive Information?

Animal experiments show that noxious stimuli activate neuronal groups in many parts of the brain, both subcortically (in the amygdala, hypothalamus, PAG, basal ganglia, and cerebellum) and in the cerebral cortex. SI does not appear to play a central role in pain perception, however, in contrast to its crucial role in other aspects of somatosensory sensation. Although they have been elicited by electrical stimulation of SI in humans, pain sensations are an infrequent effect of such stimulation. Curiously, stimulation of other parts of the cortex did not evoke pain, either. Ablations of SI do not necessarily reduce pain perception, and only occasionally has it been reported to relieve chronic pain. Some even questioned the importance of the cerebral cortex for pain perception. However, recent studies in humans, using PET and fMRI, have demonstrated robust activation of the cerebral cortex on stimulation of nociceptors and, further, that this activation is associated with the subjective experience of pain.⁹ The most consistently activated

regions are the **insula**, **SII**, and the **anterior cingulate gyrus** (Fig. 14.9). In addition, most studies show activation in **SI**, the **premotor area**, and the **thalamus**. Often the **prefrontal cortex** and the **posterior parietal cortex** (area 7b) are activated as well.

Most experimental studies evoked activation of the anterior cingulate gyrus (and other areas) via stimulation of skin nociceptors (injecting irritating substances or radiation heat). In fact, surgical ablation of the anterior cingulate gyrus has been used to alleviate chronic pain. Afterward, some patients report that the pain is still there, but it is less bothersome: it seems as if the affective component of the pain has been reduced (this is similar to pain perception after large lesions of the frontal lobe). This may perhaps be explained by other mental processes that involve the cingulate gyrus. For example, activation of the anterior cingulate gyrus is associated with the person directing his attention toward a stimulus. Further, activation of the anterior cingulate gyrus is associated with monitoring of cognitive and bodily processes: there is a change of activity in relation to errors. Indeed, signals from nociceptors are strong warnings that something is wrong in the body, and they forcefully direct our attention to the painful site.

The cortical pattern of activation upon nociceptor stimulation, as revealed by PET and fMRI, depends less on the mode of stimulation than on the **context** and the subject's **state of mind** (anxiety, expectations, feeling of control, and so forth). Many of the areas activated by painful stimuli may be activated in conjunction with other kinds of sensory processing and mental activities. Therefore, most likely no cortical regions are *specific* to pain processing. It seems, rather, that we experience pain when a distributed **network** of cortical and subcortical regions reaches a certain level and pattern of activity. This "pain network" (pain matrix) overlaps other cortical networks, such as networks for attention, emotional processing, and body image.

⁹ The interpretation of such data is not straightforward, however. For example, it is not always clear whether a change of cortical activity is an expression of altered sensory, affective, or cognitive processing. Furthermore, change of activity may be due to facilitation or inhibition of movements in response to a painful stimulus rather than to the experience of pain as such.

OVERVIEW

While the previous chapter dealt with the basic neurobiological aspects of pain, we here discuss pain in a broader context.

First we emphasize that the ability to feel pain is of vital importance; without it we do not get the necessary warnings to protect ourselves and maintain bodily homeostasis. Nevertheless, in a clinical setting long-lasting pain is a major problem. The pain system has several characteristic properties. One is its pronounced **plasticity**, considering that with ongoing pain neuronal properties are quickly changed in the direction of **hyperexcitability**. This helps explain the phenomenon of **hyperalgesia** (stronger than normal experience of pain upon nociceptor activation). **Pain** is furthermore a strong **learning** signal; we cannot afford to forget potentially harmful stimuli.

Another notable characteristic is that the association between nociceptor activity and pain perception is clear cut only in case of acute exposure to noxious stimuli. In pain of longer duration, it may be experienced more or less independent of nociceptor activity. This is most obvious in cases of **deafferentation** pain where intense pain is felt in a body part in spite of complete (or incomplete) loss of sensory information from that part. We use the term **pathologic pain** about conditions in which pain occurs without any nociceptor activation (or peripheral tissue pathology)—that is, pain that has no obvious biologic function (in contrast to “normal” or “physiological” pain).

Signals from nociceptors are transmitted to the cerebral cortex but also many other parts of the central nervous system (CNS). These include the **PAG** (periaqueductal gray) and other nuclei in the brain stem, as well as the **hypothalamus** and the **amygdala**. These nuclei are important for coordination of autonomic, endocrine, somatic, and emotional responses that optimize our total response to physical and mental challenges.

The transmission of nociceptive signals from the cord to higher levels is subject to **descending control**. The responsible systems are complex, involving networks interconnecting parts of the brain. Among transmitters, **serotonin**, **norepinephrine**, and **endorphins** (opioid peptides) appear to be of special importance. Depending on, among other factors, the emotional state and expectations of the subject, descending systems may inhibit or facilitate transmission at spinal and higher levels. Thus, perception of pain may

be completely abolished or greatly enhanced, depending on the situation.

The **placebo** effect is prominent in relation to pain. That is, pain is often reduced when the person expects that a treatment will help. This is not a matter of belief only, but relates to biologic processes in the brain. For example, the placebo effect is associated with altered activity in the “pain networks” and with release of endorphins in key, pain-related neuronal groups. The opposite phenomenon is called **nocebo**, and means that expectation of a harmful treatment worsens pain.

SOME DISTINCTIVE FEATURES OF PAIN

Due to the great clinical importance of pain, we will treat this subject in some more depth. In several ways—such as cortical processing and its strong affective coloring—pain differs from other kinds of somatic sensation and has proved much more difficult to study scientifically. In the previous chapter, we dealt with the neurobiological aspects of pain. Here, we will discuss some of the characteristics that make pain systems so special, such as the tendency of pain to outlast its initial cause, the complex control mechanisms of the brain, and the strong dependence of pain perception on context and expectation. The latter underscores that the experience of pain—even more than other sensory modalities—depends on an interpretation of available information from various external and internal sources. As phrased by Ramachandran and Blakeslee in their book *Phantoms in the Brain* (1998, p. 54): “. . . pain is an *opinion* on the organism’s state of health rather than a mere reflexive response to an injury.”

Pain Is of Vital Importance

In the clinical situation, where pain is often the main presenting complaint of patients, it is easy to forget the good sides of pain perception. Children born without the ability to feel pain have severely reduced quality of life, and often die young. The following description in the book, *Pain: the gift nobody wants* (Brand and Yancey 1993 p. 3), tells us why: “Tanya, now eleven, was living a pathetic existence in an institution. She had lost both legs to amputation . . . her failure to limp or shift weight when standing (because she felt no discomfort) had eventually put intolerable pressure on her joints. Tanya had also

lost most of her fingers. Her elbows were constantly dislocated. She suffered the effects of chronic sepsis from ulcers . . . Her tongue was lacerated and badly scarred from her nervous habit of chewing it.” The ability to feel pain is indeed necessary for us to be able to prevent injuries from innumerable small (and occasional large) physical threats. Also lowered pain threshold and spontaneous pain in conjunction with tissue damage are biologically meaningful, because they ensure protection and optimal conditions for healing. In a wider context, the pain system constitutes an integral part of the bodily systems for **defense** and **homeostatic control**, as discussed in Chapter 14, under “Homeostatic Surveillance: A Task of Ascending Tracts from Lamina I?”.

The vital importance of the pain system might perhaps give us a clue to why it so often goes awry. In the inevitable balance between **sensitivity** and **specificity**, the system is biased heavily in favor of sensitivity: we cannot afford to miss alarms of potential life-threatening events. Thus, the specificity would be correspondingly low, leaving the system open to false alarms. Indeed, some poorly understood pain conditions, such as **fibromyalgia**, might perhaps be understood in such terms.

“Change Behavior and Remember What Happened!”

The above example of a child born without the ability to feel pain also exemplifies that the feeling of pain is as a strong signal to **change behavior**. The lack of adaptation in the pain system—on the contrary, it easily **sensitizes**—is meaningful: it might be disastrous if we stop paying attention to a steady stream of signals from nociceptors. This is very different from the situation for other sensory systems where adaptation is a characteristic property. Further, pain is a forceful stimulus to **learning**: we quickly learn to avoid everything that previously led to tissue injury or threatened to do so. When the child—mostly by learning by doing—develops mastering and control of its environment, experience of pain is a central guide. In this sense, physical pain is just one aspect of discomfort and suffering and belongs perhaps conceptually more under brain systems handling **punishment** and **reward** (suffering and pleasure) than under sensory systems. Interestingly, functional magnetic resonance imaging fMRI studies show overlap between brain regions that are activated when we feel pleasure and pain. Overlap also exists between the brain activity related to physical and **social pain** (e.g., when social relationships are threatened or lost). Especially the anterior **cingulate gyrus**, the **orbitofrontal cortex**, and the **insula** are sites where pain and emotions meet.

Plasticity of the Pain System

For the development of various maladaptive pain conditions, the pain system’s well-developed plasticity seems

to be crucial. For example, even a relatively short train of signals from nociceptors alters the properties of the receiving dorsal horn neurons. Presumably, this property is necessary for proper functioning of the pain system, for example, to ensure the necessary high level of sensitivity. Nevertheless, plasticity too often seems to go awry causing **pathologic pain**, that is, pain that cannot be explained by adequate nociceptor activation and has no protective or reparative function. In cases of pathologic pain, plastic changes occur at all levels of the pain system, from the dorsal horn to the cerebral cortex.

Nociceptors and the Perception of Pain

It is important to realize that pain perception and nociceptor activity are not synonymous terms. Thus, nociceptor activity and the feeling of pain may occur independently of each other. The usual **definition** of a **nociceptor** is purely physiological: a receptor that is activated by stimuli that produce tissue damage, or would do so if the stimulus continued. In contrast, **pain** is a subjective experience with a psychological definition: an unpleasant sensory and emotional experience, which occurs together with actual or threatening tissue damage, or is described as if it were caused by tissue damage. Usually, of course, nociceptor activation causes the pain, and when we feel pain, we more or less automatically ascribe it to something that harms our body. Indeed, pain is always felt somewhere in our body, even if it is caused solely by abnormal activity of neurons in the brain. Thus, on the one hand, a person may suffer the most intense pain, yet there may be no evidence of nociceptor stimulation; on the other hand, there are many examples of persons exposed to massive nociceptor stimulation who feel no pain. Examples of the latter situation are seen in serious accidents in which the injured person may experience no pain immediately afterward, in spite of considerable tissue damage.¹ This is most likely explained by mechanisms of the brain preventing nociceptive signals from reaching consciousness (this is discussed further later in this chapter). In certain situations, such mechanisms may be necessary to survive.

WHEN THE PAIN SYSTEM GETS OUT OF CONTROL

Acute and Enduring (Chronic) Pain

Many observations suggest that acute, “ordinary” pain differs from pain of longer duration (chronic) regarding

1 Of patients admitted to an emergency ward, 40% reported no pain at the time of injury, 40% reported pain that was judged (by the doctors) as stronger than expected on the basis of the injury, but only 20% reported pain that was judged as “adequate” (Hardcastle 1999).

central mechanisms.² The former is clearly related to nociceptor activation; it ends when (or shortly after) the stimulus ends. The threshold for eliciting acute pain is high (see the preceding definition of a nociceptor). There is good correspondence between the intensity of nociceptor stimulation and the experience of pain. In such instances, pain is a homeostatic factor, serving as a signal to change behavior to avoid tissue damage. Chronic pain, in contrast, is characterized by a weak or absent correlation between stimulus and experience of pain. For example, **hyperalgesia**—the experience of pain on nociceptor stimulation—is more intense than normal—is often present.³ Thus, the somatosensory system is abnormally sensitive. We all experience this altered state of the somatosensory system with, for example, a sprained ankle or a local infection, that is, in situations with inflammation. Even slight movement or touching of the injured part evokes intense pain, or it may be painful also at rest. In this situation the pain can be seen as biologically meaningful, as it ensures that the injured part gets the necessary rest; moreover, the pain subsides in parallel with the inflammation as the tissue heals. Hyperalgesia is due to **sensitization** of both the nociceptors (primary hyperalgesia) and of neurons in the dorsal horn and at higher levels (secondary hyperalgesia). Experimentally, dorsal horn neurons can be made hyperexcitable by inducing inflammation in a joint or in the skin.

Some, but not all, chronic pain patients have a lowered threshold, so that normal innocuous stimuli (such as touching the skin) can elicit intense and long-lasting pain. This phenomenon is called **allodynia** and appears to be caused by (abnormal) activation of nociceptive neurons in the cord by dorsal root fibers (A β) leading from low-threshold mechanoreceptors. Change of pre-synaptic inhibition is probably instrumental in causing such a switch in the signal traffic. (The normal situation is that activity in thick myelinated dorsal root fibers *inhibits* nociceptive neurons, as discussed later under “Analgesia Can Be Produced by Nerve Stimulation and Acupuncture”).

Another characteristic of long-lasting pain is the tendency to **radiate**—the pain spreads out from the original painful site. This is most likely due to sensitization of

(among others) spinothalamic neurons in the segments above and below the dorsal roots leading from the inflamed region (as mentioned, the dorsal root fibers divide in an ascending and a descending branch that may pass for several segments).

Glial cells seem to be implicated in development and maintenance of chronic pain. Thus, in the cord astrocytes and activated microglia release neuroactive substances (interleukins, tumor necrosis factor, nitric oxide [NO], ATP, and others). These substances may increase transmitter release from central terminals of nociceptive dorsal root fibers. Further, activated glial cells can sensitize dorsal horn neurons, among them spinothalamic ones. Activation of glial cells occurs with infections in the CNS, but also in response to release of substance P and amino acid transmitters from nociceptive dorsal root fibers.

Plastic Changes and Altered Brain Networks

Hyperalgesia, allodynia, and radiating pain are at least partly due to altered synaptic transmission in the CNS: chronic pain is associated with **plastic changes** that make neurons in many parts of the somatosensory system hyperexcitable. This is best documented in the dorsal horn but occurs also at higher levels. As with plastic changes in other systems, *N*-methyl-D-aspartate (NMDA) **receptors** have a crucial role. **Substance P** increases the sensitivity of NMDA receptors to glutamate, so that wide-dynamic range spinothalamic neurons might react more vigorously to inputs from low-threshold mechanoreceptors.

Intense pain of some duration may leave “memory traces” in the brain, so that later, minimal provocation may suffice to revive the pain. For example, a man who had a painful spine fracture reexperienced the pain when, many years later, he suffered a myocardial infarction. Electric stimulation during brain surgery indicates that part of the somatosensory **thalamus** becomes hyperexcitable in patients with chronic pain syndromes. For example, patients with panic attacks accompanied by chest pain and patients with deafferentation pain reported their usual pain on thalamic stimulation. In contrast, patients undergoing surgery for movement disorders reported no pain on stimulation at the same thalamic sites.

It appears from morphometric studies that chronic pain conditions may cause **gray and white matter alterations** in pain-related networks. Although findings differ somewhat among studies, most report gray-matter reductions in the anterior **cingulate cortex**, the **insula**, and the **orbitofrontal cortex** (the specific cellular changes that underlie these findings are unknown, however). Further, fMRI studies show altered activation of networks related to attention in chronic pain patients. Finally, in patients with complex regional pain syndrome (CRPS), the

2 The indiscriminate use of the term “chronic pain” has been criticized because it lacks precision and encompasses a variety of conditions with different causes, courses, and prognoses. Indeed, definitions of chronic pain differ widely. Some define it as pain that persists longer than the course of natural healing, others that it lasts longer than three, alternatively six months. Some even use as a criterion that the pain has not responded to available (drug) treatment. The International Association for the Study of Pain defines chronic pain as: “. . . pain which persists past the normal time of healing . . . With non-malignant pain, three months is the most convenient point of division between acute and chronic pain, but for research purposes six months will often be preferred.”

3 The International Association for the Study of Pain has proposed that **hyperalgesia** be used as an umbrella term for all conditions of increased pain sensitivity (Loeser and Treede 2008). Allodynia would then be a special case of hyperalgesia.

somatosensory-cortex representation of the painful part seems to shrink but normalizes during successful therapy. Curiously, in chronic back pain patients, an *enlarged* back representation appears to occur. It seems, therefore, fair to say that while cortical reorganization in chronic pain states is well documented, its functional significance is less than clear.

Pathologic Pain

Chronic pain can occur not only on increased nociceptor activity but also after loss of sensory information, either from nociceptors or from low-threshold mechanoreceptors. We use the term **pathologic pain** here rather loosely to describe conditions in which pain occurs without any nociceptor activation (or peripheral tissue pathology)—that is, pain that has no obvious biologic function (in contrast to “normal” or “physiological” pain). Pathologic pain may have quite different causes, such as partial damage to peripheral nerves or destruction of central somatosensory pathways, for example, in the cord or in the thalamus (usually termed **neuropathic pain**). **Deafferentation pain** is pain that, paradoxically, occurs after loss of sensory information from a body part. A striking example of the latter is patients with **avulsion** of dorsal roots (this occurs sometimes with the roots of the brachial plexus upon a violent pull of the arm). In spite of no sensory nerves entering the cord, such patients often develop excruciating pain in the denervated arm. Pathologic pain also occurs in some patients below a **transverse lesion of the cord**, even though all ascending sensory pathways are interrupted. Pathologic pain may also be felt in an area of the skin with reduced sensibility in patients who have had shingles (**postherpetic neuralgia**). A stroke destroying parts of the thalamus leading to reduced sensibility in a body part is sometimes associated with chronic pain (**thalamic pain**). **Phantom pain**, often occurring for shorter or longer periods after amputations, refers to pain felt in the missing body part. Clinical evidence indicates that this peculiar phenomenon is associated with plastic changes in the brain (especially in the somatosensory cortex). It is not due to abnormal activity in peripheral nerves or ascending sensory pathways. Often, the pain occurs together with a vivid experience of abnormal movements or postures of the missing limb. Probably, the brain misinterprets the lack of sensory information from the missing body, drawing the conclusion that something is seriously “wrong” out there.

It is fair to say that our understanding of pathologic pain syndromes is far from satisfactory. It is particularly enigmatic why apparently identical injuries cause chronic pain in some patients but not in others (the majority). An association between pathologic pain syndromes and certain personality traits has not been convincingly demonstrated.

Complex Regional Pain Syndromes (CRPS): A Special Type of Pathologic Pain

In some patients, pain continues in spite of complete healing of an injury (most often in the extremities). What started as a “normal,” nociceptor-driven pain continues for unknown reasons as a pathologic pain. For example, an apparently trivial fracture of the radius is sometimes followed by pain for years after the fracture has healed. Similar persistence of pain can occur after partial lesions of peripheral nerves (neuropathic pain). Usually, such patients also suffer from hyperalgesia and allodynia, and even light touch may provoke excruciating pain. Often they also show signs of **autonomic dysfunction**, such as abnormal sweating and circulatory disturbances. This condition used to be called reflex sympathetic dystrophy (RSD), reflex dystrophy, or sympathetically mediated pain. It is especially unfortunate, however, that names of poorly understood diseases implicate an etiology, like “reflex” and “sympathetic.” In some cases, especially after nerve injury, the pain has a peculiar burning quality, and the name **causalgia** refers to this condition (Greek: *kausos*, heat; *algos*, pain).

The term **complex regional pain syndrome** (CRPS) was introduced in an attempt to avoid the many confusing terms for these kinds of pathologic pain conditions. CRPS is a purely descriptive term, reflecting that we do not know the pathophysiological mechanisms leading to the various symptoms. It has two subgroups, with reasonably precise definitions, corresponding largely to reflex dystrophy (CRPS type I) and causalgia (CRPS type II), respectively.

CRPS and the Sympathetic System

As mentioned, patients with complex regional pain syndromes often show evidence of autonomic dysfunction in the painful part—mainly hyperactivity of the sympathetic system. The often-used term “reflex sympathetic dystrophy” (RSD) implies that sympathetic dysfunction *causes* the syndrome. The pain relief achieved in some patients by a sympathetic block (e.g., of the stellate ganglion in case of pain in the hand) would seem to support this assumption. Nevertheless, **microneurographic** and other kinds of studies have not confirmed abnormally increased activity of sympathetic postganglionic fibers in such patients, even in those with obvious signs of sympathetic hyperactivity such as profuse sweating and extreme cutaneous vasoconstriction. This seeming paradox may perhaps be explained by **adrenergic receptors** starting to be expressed by **spinal ganglion cells** (Fig. 15.1). Thus, normal levels circulating catecholamines may activate sensory neurons. There is furthermore evidence that in some CRPS patients neuropeptides (substance P and cholecystokinin [CCK] in particular) are released from sensory nerve endings in the skin of the painful parts.

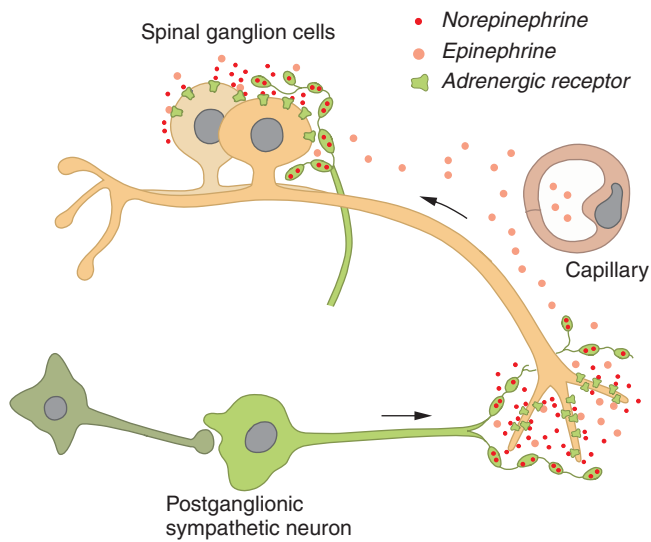


FIGURE 15.1 Sensory neurons (spinal ganglion cells) express adrenergic receptors in certain chronic pain conditions. Receptors are located on the cell body and the peripheral branches.

This causes so-called **neurogenic inflammation**, which may produce some (but not all) of the autonomic signs in CRPS. The relationship between neurogenic inflammation and pain in CRPS is not clear, however.

CENTRAL CONTROL OF TRANSMISSION FROM NOCICEPTORS AND PAIN SENSATION

We now realize that a complex network—stretching from the spinal cord to the cerebral cortex—modulates transmission of signals from nociceptors and pain perception. As mentioned, both the context of the nociceptive signals and how the brain interprets the situation determine how we perceive signals from nociceptors. In the modulatory network, **serotonin**, **norepinephrine**, and **opioid peptides** seem to play crucial roles, although several other neurotransmitters also participate.

Descending Control from the Brain Stem

A dramatic observation by Reynolds (1969) was the starting point for much later research on central control of nociception. He showed that by electrical stimulation of a mesencephalic region, the **periaqueductal gray matter (PAG)** (Fig. 15.2; see also Fig. 6.20), conscious rats could be subjected to major surgery without apparently feeling any pain (as judged from their behavior and other observations). The stimulation produced **analgesia** (no experience of pain on noxious stimuli). Subsequent research indicates that the effect of PAG stimulation can be explained, at least in part, by activation of descending connections to the dorsal horn. Although a sparse direct

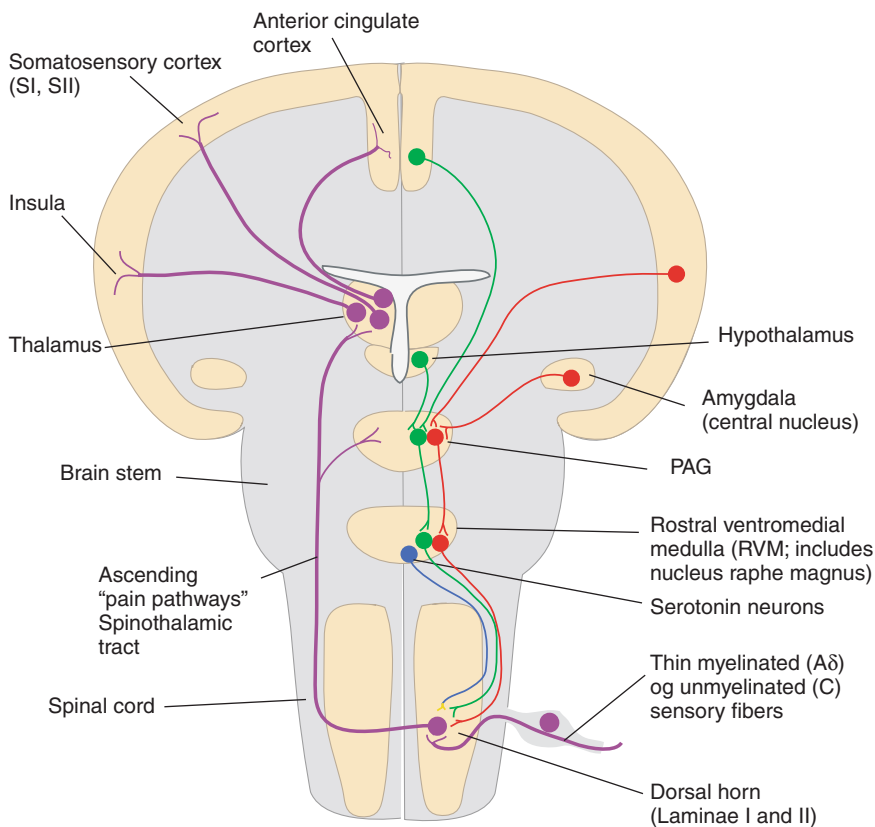


FIGURE 15.2 Descending systems for modulation of nociceptive signal transmission. **Left:** Ascending “pain pathways.” **Right:** Green neurons mediate inhibition of ascending transmission whereas red neurons mediate facilitation. PAG has a pivotal position in the network for pain modulation. The anterior cingulate cortex, the prefrontal cortex, amygdala, and the hypothalamus send instructions to PAG depending on the emotional state and the expectations of the individual. Opioids influence the transmission at all levels of the network. Simplified. (Based on Fields 2004.)

pathway from the PAG to the spinal cord exists, the main pathway appears to be synaptically interrupted in cell groups in the medulla, especially the **raphe magnus nucleus (NRM)** (Fig. 15.2; see also Figs. 26.5 and 26.6) and other nearby cell groups in the **medullary reticular formation**. The descending fibers from these nuclei (raphespinal fibers) lie in the dorsolateral funiculus of the cord, and cutting the latter abolishes the effects of PAG stimulation almost completely. Further, electrical stimulation of NRM inhibits spinothalamic cells.

We do not know in detail the synaptic arrangements in the spinal cord that mediate these effects from the brain stem, but it is of interest that fibers from the NRM establish synapses in the dorsalmost part of the dorsal horn (**laminae I and II**), which receives most dorsal root fibers conveying signals from nociceptors (Fig. 15.3). Many of the NRM neurons contain **serotonin**, and this is apparently a transmitter at synapses established by raphespinal fibers in the cord. The actions of serotonin in the dorsal horn are complex, but experiments with microinjections indicate that it inhibits nociceptive spinothalamic cells. Probably, the inhibitory effects are mediated partly by direct synapses on spinothalamic cells and partly by synapses on dorsal horn interneurons. Transmitters other than serotonin—**norepinephrine** and **opioid peptides** in particular—contribute to the effects on pain transmission obtained by stimulation of NRM and nearby structures. Indeed, the effects of the neurotransmitters used in the descending control systems are not fixed: depending on the cause of the pain and the context in which it occurs, the same transmitter may have opposite effects.

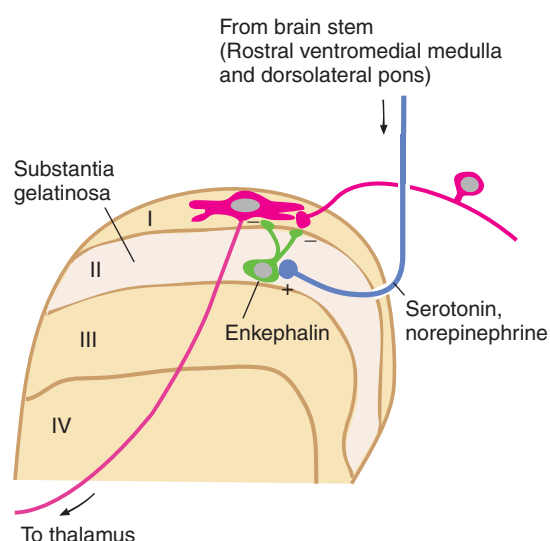


FIGURE 15.3 The substantia gelatinosa. One of several possible wiring patterns in the dorsal horn, which can contribute to inhibition of spinothalamic cells.

Although descending systems that inhibit signal traffic from nociceptors have attracted most attention, there are also connections with **facilitating** effects (Fig. 15.2). Animal experiments suggest that descending connections from the brain stem can contribute to the development of hyperalgesia in cases of peripheral inflammation. Further, descending connections may contribute to the maintenance of pain in some chronic conditions.

PAG Coordinates Stress Responses

Studies of the PAG suggest that this nuclear complex plays a pivotal role in coordinating bodily reactions to painful and potentially harmful somatosensory stimuli—that is, endocrine, autonomic, and somatic responses with the common goal to reestablish or maintain **homeostasis**. Interestingly, different parts of the PAG mediate different kinds of responses to painful and other stressful stimuli. Stimulation of the **ventral part** of the PAG elicits an active, defensive behavior, whereas stimulation of **dorsal part** causes an immobility (**freezing**) reaction. Further, the ventral part mediates opioid-dependent **analgesia**, as judged from abolition of the effect by injections of **naloxone** (which prevents binding of some endorphins to their receptors). Stimulation of the dorsal part, in contrast, produces an analgesia that cannot be reversed by naloxone, suggesting that transmitters other than endorphins are responsible. This dual response pattern elicited from the PAG in experimental animals corresponds to typical human reactions to pain originating from skin and from the viscera, respectively. Pain of cutaneous origin typically produces movements and increased heart rate and respiration, whereas immobility, lowered heart rate, sweating, and nausea often accompany pain arising in deep structures.

Ascending Connections that Modulate Pain Perception

The effects of PAG stimulation on pain sensation are probably mediated not only by descending connections to the cord. There are also ascending connections from the PAG and the **parabrachial nucleus**, which may influence signal transmission at the thalamic level (**intralaminar thalamic nuclei**) and in other cell groups related to pain sensation. Nociceptive signals reaching the **amygdala** are relayed in PAG and some other brain stem nuclei. Together with the cerebral cortex, the amygdala integrates nociceptive signals and other information about the stimulus and its context. In this way, the significance of a stimulus is interpreted in relation to the needs of the whole organism. The amygdala send efferent fibers to PAG that may initiate appropriate responses—among them, suppression (or facilitation) of transmission from nociceptors. The **nucleus accumbens** in the forebrain (see Fig. 23.15)—involved in control of reward-dependent behavior—plays a part in analgesia

evoked by painful stimuli.⁴ In pain-induced analgesia, pain in one part of the body seems to be “rewarded” by analgesia in other parts.

Opiates, Opioids, and Endorphins

Some of the dorsal horn interneurons contain opioid peptides. These neuropeptides, found in many parts of the brain, got their name because they have actions that resemble closely those of the opiate-type drugs, such as morphine. The naturally occurring (endogenous) peptides of this kind are also called **endorphins**, that is, endogenous morphine. Opioids, when injected locally in the dorsal horn, have inhibitory effects on spinothalamic cells, but it is not known whether the inhibition is caused by a direct action or is mediated by interneurons (or both). Three main groups of endorphins have so far been discovered, each with its characteristic distribution in the brain and spinal cord. Best characterized are β **endorphin**, two varieties of **enkephalin**, and **dynorphin**. Whereas β endorphin is restricted to one neuronal group in the hypothalamus (the arcuate nucleus), the enkephalins and dynorphin are present in interneurons in many parts of the CNS (among them, parts that are not concerned with aspects of pain sensation).

That **morphine** and other **opiates** can alleviate severe pain has been known for hundreds of years. Today we know that there are receptors in the brain to which opiates can bind and exert their effects. These receptors are actually receptors for the endorphins, and at least five different receptors, each with affinity for a certain subgroup of endorphins, have so far been identified. **Opiate receptors** (binding morphine) are present in, among other places, the PAG, the NRM, and parts of the spinal dorsal horn (especially laminae I, II, and V). Microinjection of morphine in the PAG can produce analgesia in experimental animals that depend at least in part on connections from the NRM to the spinal cord. The actions of morphine appear thus to be exerted both in the spinal cord and in the PAG. In the cord, binding of morphine to opiate receptors on inhibitory interneurons in the substantia gelatinosa and, presumably, on primary afferent terminals, leads to inhibition of spinothalamic cells. The action in the PAG occurs most likely by activation of the descending pathway from NRM and probably also by activation of ascending connections from the PAG acting on higher levels of the CNS. Binding of morphine in other parts of the brain, such as the amygdala and other limbic structures, is probably also important in explaining the actions of morphine.

⁴ Both dopamine and opioid peptides in the nucleus accumbens appear to be necessary for pain-induced analgesia. The nucleus accumbens probably exerts its effects by indirect connections to NRM or nearby regions. The PAG and the hypothalamus may be intercalated in the pathway.

Analgesia Can Be Produced by Nerve Stimulation and Acupuncture

Analgesia may also be produced therapeutically by stimulation of peripheral nerves. When done with surface electrodes, the procedure is called **transcutaneous nerve stimulation** (TNS). One kind of analgesia occurs immediately and is mediated by activity of **thick myelinated fibers** in the stimulated nerves (i.e., fibers leading from low-threshold mechanoreceptors). Selective stimulation of such fibers is elicited by electrical stimulation with **high frequency** and low intensity or by natural stimuli such as vibration, light touch, or pressure. The analgesia is restricted to parts of the body innervated by the peripheral nerves, and it usually disappears when the stimulation stops. It is mediated by activity in collaterals of the thick dorsal root fibers that inhibit impulse transmission from the thin (A δ and C) fibers in the dorsal horn, via inhibitory interneurons and presynaptic inhibition. Melzack and Wall (1965) proposed such interaction between thick and thin dorsal root fibers in their gate-control theory. It is presumably the basis of the everyday observation that it helps to blow at a finger that hurts, that it may help to move the part that has received an acute injury, that labor pains may be alleviated by rubbing over the lower back, and so forth.

Another kind of stimulation-induced analgesia requires that **thin sensory fibers** be activated. This can be achieved by electrical stimulation of **low frequency** with relative high intensity. Classical **acupuncture** probably obtains the same effect by rotation of thin needles in the tissue. The analgesia produced by this kind of stimulation occurs with latency but may last for hours after termination of the stimulation. Analgesia is not limited to the parts of the body that were stimulated. This kind of stimulation-induced analgesia is most likely due, at least in part, to activation of the descending connections from the brain stem. The activation may happen by way of collaterals from spinothalamic and spinoreticular cells to the PAG and adjacent parts of the reticular formation. The analgesia can be reversed or prevented by intravenous injection of **naloxone**. Further, in animal experiments it has been blocked by sectioning the dorsolateral fascicle. The results of such control experiments are in part contradictory, however, and indicate that not all aspects of stimulation-induced analgesia can be explained by liberation of endorphins.

Functional Role of Pain-Modulating Systems

A dominating view today is that modulation of pain is just one among several bodily adjustments—controlled by the nervous system—to physical and mental challenges. The mental “set” and the individual’s interpretation of the situation determine the choice of responses. In other words, **expectation** of what a stimulus will

cause is more decisive for the response than the stimulus itself. Nevertheless, we do not fully understand the functional role of the pain-modulating systems, nor under which conditions they are activated. It is reasonable to assume, however, that **inhibiting** systems are active in situations with severe injuries and no experience of pain (as, e.g., in war and major civil accidents). In such situations, suppression of pain may enable continuation of intense physical activity, which may be of vital importance. Markedly reduced pain sensitivity may also occur in more peaceful situations, such as sport competitions. Further, pain-suppressing systems (at several levels) appear to be active in expectation-dependent analgesia, although expectation may also increase pain perception (see next section, “Placebo and Nocebo”). Animal experiments indicate that analgesia can occur in stressful situations characterized by the inability to escape (**stress-induced analgesia**). However, the mental state of the animal seems to decide whether analgesia occurs. Thus, in one study analgesia occurred only if the animal was calm at the start of the period of stress, whereas anxious animals became hyperalgesic. This resembles the well-known effect of anxiety on the experience of pain in humans.

PLACEBO AND NOCEBO

The **placebo** effect is probably best understood as one part of a repertoire of responses to challenges that threaten our mental or physical stability (see also Chapter 30, under “Psychosomatic Relations,” and Chapter 31, under “Amygdala and Conditioned Fear”). Although it has attracted most attention in connection with the treatment of pain, we now know that placebo effects concern several physiological processes. The beneficial placebo effect depends on the patient’s positive expectations, but negative expectations may evoke increased pain and harmful physiological alterations. This is called the **nocebo** effect. However, placebo (and nocebo) effects may in addition involve **conditioned responses**, for example elicited by the taste of a drug that previously was experienced as effective (e.g., a painkiller).⁵ In this respect, the placebo response is **learned**, depending on prior experience. Even the observation of another person receiving a beneficial treatment induces a placebo response in the observer.

As discussed, descending connections from the brain stem can inhibit or enhance nociceptive signal

transmission, depending on the situation (Fig. 15.2). Such mechanisms are most likely involved in the placebo and nocebo effects on pain. One likely pathway goes from the amygdala to the PAG. At the cortical level, expectation of pain relief reduces the activity of the pain network (Fig. 15.4A). Further, during the expectation phase cortical regions usually involved in cognitive and evaluative processes show enhanced activity (Figure 15.4B).

A common misconception is that the placebo effect works only with moderate pains and only in comparison with “weak” drugs. However, if the patient believes that the drug she gets is morphine, the placebo effect is much greater than if the patient believes it to be aspirin. Indeed, postoperative, intravenous administration of placebo (saline) in full view of the patient had the same analgesic effect as 6 to 8 mg of morphine given covertly.

The question remains, however, how **expectancy** controls the “pain network” and pain-suppressing systems. Put more broadly: how do thoughts and feelings express themselves through the body? We discuss this theme further in Chapters 31 and 32 although, admittedly, we are far from a full understanding of the **mind–body problem**.

Placebo

The word placebo (literally: “I shall please”) has been used for a long time about mock medicine—that is, drugs and procedures that the doctor knows have no effect. In clinical trials, for example, inert tablets are given as a placebo to decide whether a drug has a specific effect on a disease. Thus, it is well known that a treatment without any specific effect may influence the disease and the experience of pain. We know that the effect is only present if the patient expects the treatment to work. Further, surgery produces a larger placebo effect than drugs, and injection of a drug is more efficient than oral administration.

An operation introduced in the 1950s to cure **angina pectoris** exemplifies the placebo effect of surgery. On a dubious theoretical basis, the internal mammary arteries were ligated to improve the blood supply to the ailing heart. Many patients noted improvement in their condition after the surgery. Later pathological examinations showed, however, that no improvement of the blood supply had occurred. Therefore, a study was initiated to compare two groups of patients with angina pectoris: one group underwent ligation of the artery, the other group underwent the same procedure but without ligation (a sham operation). Many of the patients in both groups noted improvement, and the improvement lasted for the observation period of 6 months. Remarkably, the improvement concerned not only reduction of pain but also objectively measurable variables, such as walking distance, drug consumption, and (in some) even the electrocardiogram. Several later studies have shown

⁵ In one study, patients received the immunosuppressant cyclosporine in association with a flavored drink. After a number of repetitions of the association, the flavored drink alone induced immunosuppression. Several other studies in experimental animals and in humans have shown similar conditioning of immune responses. The endocrine system is also subject to placebo conditioning. The conditioned immune and endocrine responses do not require conscious expectations.

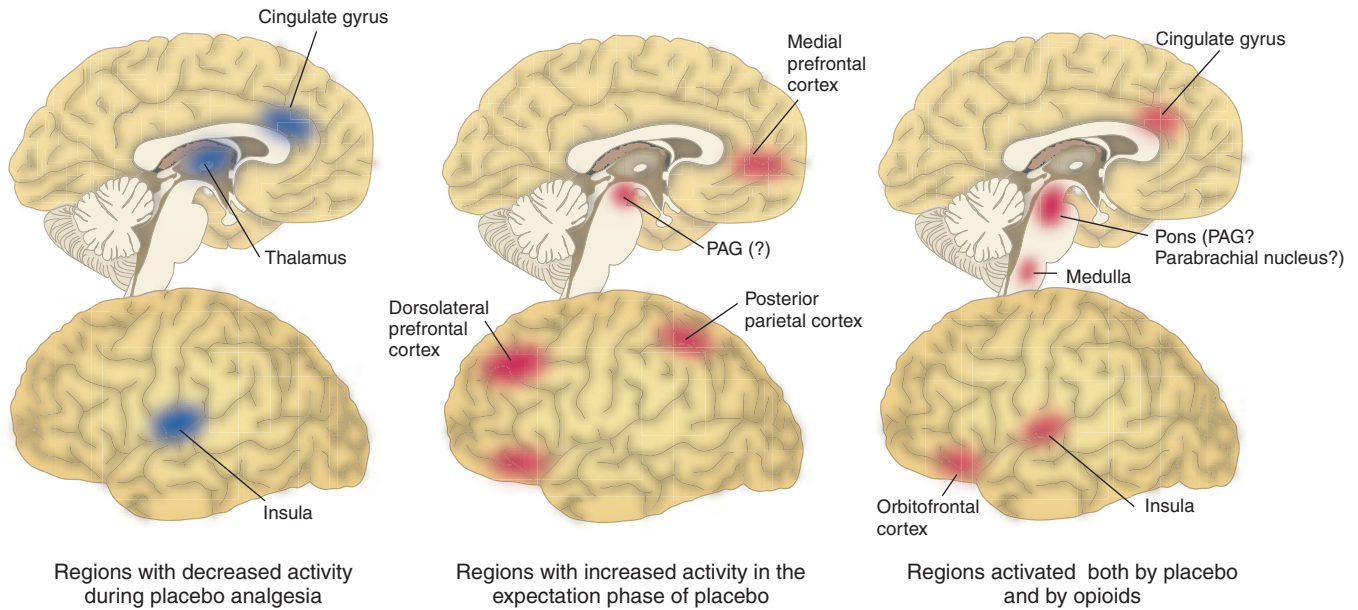


FIGURE 15.4 Brain regions involved in placebo analgesia. The size and position of the relevant regions are only approximations, and other regions than those shown here are involved in placebo analgesia.

The regions shown in **B** are activated by cognitive and evaluative processes, as shown in other fMRI studies. (Based on fMRI data published by Petrovic et al. 2005 and Wager et al. 2004.)

similar results. For example, in one study the effect of **ultrasound** on pain after tooth extractions was compared with placebo. Not only did the patients receiving the probe without any ultrasound (the placebo group) feel less pain, they also experienced less local edema and could open their mouths wider than patients in the placebo group. Thus, the placebo effect is not limited to subjective experiences related to disease but can also affect **physiological processes**. Obviously, the placebo effect is not merely a question of imagination, and its presence in a patient is not related to whether the disease is organic or “functional”. Interestingly, fMRI studies indicate that analgesia obtained by an opioid drug and by placebo activate the same brain sites (Fig. 15.4C).

The Endorphin Hypothesis and Somatotopic Placebo Effects

The “endorphin hypothesis” became popular when it was reported that **naloxone** (an opiate antagonist) could prevent the placebo effect. However, as emphasized by Wall (1993, p. 197), “. . . it is not clear what insight into the overall placebo phenomenon is provided by showing that some link in the machinery involves endorphins.” We now know that the placebo effect, rather than being mediated by a diffuse release of endorphins, is associated with altered opioid transmission in specific brain sites. Further, the placebo effect can apparently be rather specific and restricted to only

certain parts of the body. In a recent study, pain was evoked at different locations by injections of capsaicin. A placebo cream—said to have a strong anesthetic effect—was applied at one of the painful sites. At this site, but not at the others, the persons reported a marked reduction of pain, which was prevented by naloxone. This fits with the fact that the **PAG** is somatotopically organized and that electric stimulation of different parts produce analgesia restricted to certain body parts. Thus, the placebo effect is probably not mediated by diffuse release of opioids but by activation of specific neuronal groups, some of them containing opioid peptides.

Nocebo

As discussed in the preceding text, expectation of a positive effect influences markedly how the brain modulates pain and several physiological processes. Thus, the placebo effect depends on whether the patient believes that the treatment can help. As mentioned, the opposite effect, called **nocebo**, can occur if the patient expects no help or believes that the treatment may be harmful. In such instances, a neutral substance may worsen the pain and produce unwanted biologic effects. The neurobiological basis of nocebo effects is so far largely unknown. However, while placebo responses are associated with increased opioid and dopaminergic activity in several cortical and subcortical sites (e.g., the nucleus accumbens), nocebo responses seems to be associated

with decreased activity in the same sites. Further, **nocebo** seems to be associated with increased **cholecystokinin** (CCK) activity, as a CCK antagonist (proglumide) blocks **nocebo**-induced hyperalgesia. Interestingly, CCK acts as a neuromodulator in the brain sites that show increased opioid activity during placebo analgesia. A close link seems to exist between **anxiety** and **nocebo** hyperalgesia. It may seem that, “. . . verbal suggestions of a positive outcome (pain decrease) activate endogenous μ -opioid neurotransmission, while suggestions of a negative outcome (pain increase) activate CCK-A and/or CCK-B receptors” (Benedetti et al. 2007, p. 265).

In any case, the fact that verbal instructions alone can induce improvement or worsening of a patient’s condition will influence every therapeutic situation. Besides the specific therapeutic effect of the treatment initiated by the doctor, the outcome for the patient depends on whether placebo or **nocebo** effects are evoked.

MODERN VIEWS ON PAIN AND PAIN TREATMENT

Recent years of intensive interdisciplinary research have led to important new insights in pain mechanisms, along with improved pain therapy. Some aspects have been mentioned in this chapter. We now realize that many more parts of the brain are involved in pain processing than was formerly believed. Although we are still far from a complete understanding, we know many of the pieces that form the basis of subjective pain experience and the physical and mental reactions to pain. Some pain conditions, previously rejected as “imagination” or ignored because they were incomprehensible—such as **deafferentation pain**—can now be at least partly explained in a neurobiological context. The pain is equally real to the individual, regardless of whether its cause is a stroke, a nerve injury, a panic attack, or a broken leg, and it is always felt as arising somewhere in the body.

We also know that the experience of pain is associated with activity in a distributed **cortical** and **subcortical network**—not a specific “pain center.” Conceivably, when this network enters a state of synchronized activity we experience the unpleasant emotional and bodily state we call pain. Complex neural networks are typically densely interconnected, so that a specific neuronal group may take part in different tasks, depending on the situation. Thus, many other networks can presumably access the “pain network” and contribute to its activation (Fig. 15.5). This implies that the feeling of pain can be evoked not only by signals from **nociceptors** but also by activity in neuronal groups related to motivation, attention, expectations, emotions, memories, and so forth. Rather than making sharp distinctions between “organic” and “psychological” pain, we realize that all pain encompasses both aspects, albeit in varying proportions.

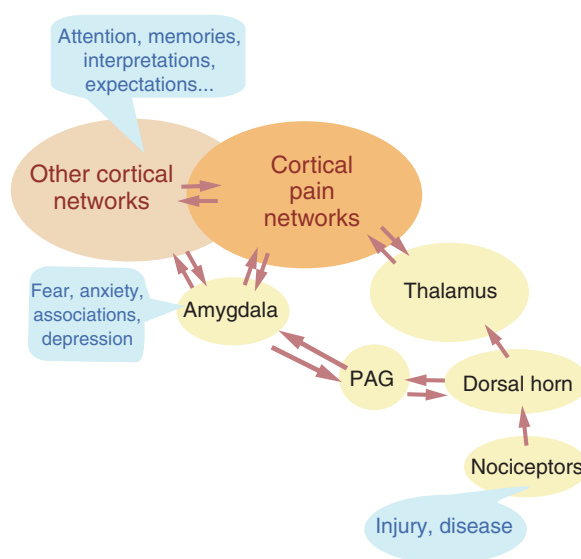


FIGURE 15.5 *The cortical pain network and modes of activation.* Schematic to show how quite different inputs can activate the pain network—from peripheral **nociceptors** triggered by tissue injury to purely mental processes.

Both forms are expressions of dynamic interactions between many brain systems. For example, depression and anxiety strongly influence pain perception, while chronic pain by itself can cause the same symptoms. Psychological **reward** mechanisms can in some instances suppress pain, in others contribute to its maintenance.

For the brain to interpret the multitude of signals arising in the body correctly, a proper balance among the various somatosensory modalities seems necessary. Thus, **loss of sensory information**—regardless of whether it concerns low-threshold mechanoreceptors or **nociceptors**—can cause long-lasting pain. The brain apparently misinterprets what is going on “out there.” Probably, the same occurs when a patient experiences bizarre distortions of the body image after an amputation or even when an extremity is immobilized (Oliver Sacks describes in the book *A Leg to Stand On* [1994] his own experiences of this kind after a knee injury with immobilization). Interestingly, attempts to correct with mirrors the experience of abnormal postures in amputees led to immediate pain relief as soon as the “body image” was normalized. The basic problem may probably be not so much loss of sensory information per se, but rather a **mismatch** between the **motor commands** and the **sensory feedback**. Thus, when the sensory feedback does not match the expectations (based on previous experience), the alarm goes.

We also have learned that the central pain system is **plastic**, so that its behavior can change quickly. This has led to a better understanding of the differences between acute and chronic pain. Pain can obviously leave permanent traces in the brain, even though, mercifully, most

pains are forgotten very quickly. Plastic changes may produce chronic pain in some persons, while in others they may enable reactivation of a long-forgotten pain in a certain body part.

Advances in basic pain research have influenced **clinical practice**; for example, new drugs are designed to attack the central synaptic changes in chronic pain directly. Further, an important aim for modern pain

therapy is to prevent the occurrence of the plastic changes in the dorsal horn and elsewhere. Thus, analgesic drugs are often administered before surgery, and afterward the dosage is individualized according to the need of the patient rather than to a fixed scheme. In addition, the importance of early and efficient pain relief in patients with injuries that are apt to produce persisting pain is being realized.

16 | The Visual System

OVERVIEW

Humans are “visual animals.” Like in other primates, our visual system is highly developed, and accordingly large parts of the cerebral cortex are devoted to processing of visual information. There are approximately 1 million axons in the optic nerve, constituting almost 40% of the total number of axons in the cranial nerves. The receptors for sight, **photoreceptors**, are the rods and cones of the retina. Their adequate stimulus is electromagnetic waves with a wavelength between 400 and 700 nm. The photoreceptors do not react to light with shorter (ultraviolet light) or longer (infrared light) wavelengths. The photoreceptors transform light energy to graded changes of the membrane potential with ensuing release of **glutamate**. From the photoreceptors, the signals pass to **bipolar cells** and from these to **retinal ganglion cells**. Several kinds of **interneurons** enable considerable information processing in the retina. The **rods** are responsible for vision in dim light, whereas the **cones** require daylight and are necessary for perception of visual details and colors.

The **visual pathways** start with the retinal ganglion cells sending their axons to the **lateral geniculate body** of the thalamus. The ganglion cell axons leave the eye in the **optic nerve** and pass through the **optic chiasm**, where axons from nasal half of the retina cross while axons from the temporal half pass through uncrossed. The axons continue in the **optic tract** to the lateral geniculate body. Axons from neurons in the lateral geniculate form the **optic radiation**, and end in the primary visual cortex—the **striate area**—in the occipital lobe of the same side. In this way, stimuli from the left visual field reach the right occipital lobe. The visual pathways show a precise, **retinotopical** organization at all levels. The area striata performs the first analysis of the visual information, while further processing takes place in **extrastriate** visual areas in the occipital, temporal, and parietal lobes. The processing is largely **segregated**, so that, for example, different areas deal with color and motion. Visual information from the extrastriate areas is integrated with other sensory modalities, and finally reaches the frontal lobe where visual information contributes to the control of behavior.

To understand the visual system, it is not sufficient to know the conscious use of visual information; the many reflex effects elicited by visual stimuli must also be taken into account. Among the reflex effects are those

ensuring **fixation of our gaze** on the object we want to examine and follows it if it moves, and those ensuring that the visual images formed at the retina are always in focus. Such **visual reflexes**, however, are only briefly mentioned in this chapter. They are discussed more thoroughly in Chapter 25, under “Central Control of Eye Movements,” and in Chapter 27, under “The Light Reflex and the Accommodation Reflex.”

THE EYEBALL AND THE REFRACTING MEDIA

The Eye and a Camera Have Certain Features in Common

The **eyeball** (*bulbus oculi*) (Figs. 16.1 and 16.2) has a firm outer wall of dense connective tissue covered on the inside by the light-sensitive retina. Between these two layers is a vascular layer, the **choroid**, that is highly pigmented, thus ensuring that light enters the eye only through the pupil and preventing reflection of light (compare with the dull black inside a camera). The diameter of the **pupil** (the shutter) controls the amount of light allowed into the eye. Refraction of the light takes place on its way through the cornea and the lens. The curvature of the lens can be varied by the use of the **ciliary muscles**, so that the retinal image is always sharply focused (in a camera, the focusing is brought about by varying the distance between the lens and the light-sensitive film). **Extraocular muscles** (external eye muscles) attach to the eyeballs and can move them to coordinate their positions, so that the visual images hit corresponding points of the two retinas (Fig. 16.3). This is a prerequisite for our perception of one image, and not two (two slightly different images are formed in the eyes, however). We discuss the extraocular muscles more comprehensively in Chapter 25.

Structural Features of the Eye

The wall of the eye consists of three layers. Outermost is the **sclera**; then follows the **choroid**, and the **retina** is the innermost layer. The eye keeps its spherical shape because the sclera has some stiffness, but mainly because the **pressure** inside the eye is higher than outside (approximately 15 cm H₂O).

Anteriorly, the sclera has a circular opening, in which the transparent **cornea** is positioned like a glass of

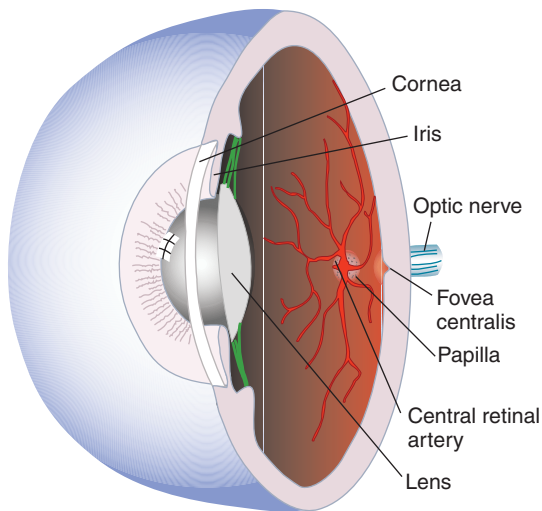


FIGURE 16.1 *The left eye.* The eye is divided in two along the visual axis (cf. Fig. 16.2).

a wristwatch. The cornea consists of a special kind of connective tissue with densely packed collagen fibrils arranged strictly geometrically. This arrangement is a prerequisite for the transparency of the cornea, as is also its lack of blood vessels. The cornea is nourished by the tear fluid and by the fluid inside the eye. The cornea contains numerous unmyelinated sensory nerve fibers, providing information about even the slightest touch. A stratified squamous epithelium that is only a few cells thick covers the outside of the cornea.

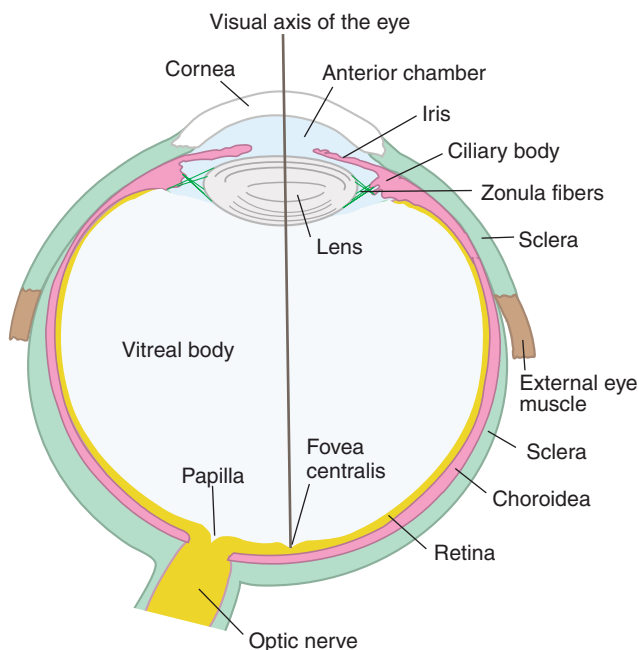


FIGURE 16.2 *Transverse (horizontal) section through the right eye, as viewed from above.*

The **choroid** (the vascular coat) is rich in vessels that nourish the cells of the outer parts of the retina. In addition, the connective tissue of the choroid contains numerous highly **pigmented cells**. Anteriorly, close to the opening in the sclera, the choroid is thickened and forms a ring (Figs. 16.1 and 16.2) called the **ciliary body**. This contains the smooth **ciliary muscle**, which controls the curvature of the lens. From the anterior edge of the ciliary body, approximately at the junction between the sclera and the cornea, the choroid continues as the **iris**. This is a circular disk with a central opening, the **pupil**. The iris contains smooth musculature that serves to regulate the size of the pupil. Smooth muscle fibers arranged circularly form the **pupillary sphincter muscle** (m. sphincter pupillae), which, when it contracts, makes the pupil smaller so that less light enters the interior of the eye. Radially arranged smooth muscle fibers in the iris, forming the **pupillary dilatator muscle** (m. dilatator pupillae) widen the pupil when they contract.

The space inside the eye in front of the lens and the iris is called the **anterior chamber**. It is filled with a clear watery **fluid** that is produced by small processes of the ciliary body. The space behind the lens is filled with a clear jellylike substance called the **vitreous body** (corpus vitreum).

The eyeball is moved inside the orbit by six small striated muscles, the **extraocular muscles** (see Figs. 25.1 and 27.16). The muscles originate in the wall of the orbit, and their tendons insert in the sclera. The extraocular muscles are innervated by the **third, fourth, and sixth cranial nerves**. They cooperate very precisely to produce the movements of the eyes that ensure that the visual axes (Fig. 16.2) of the eyes are always directed toward the point of fixation (Fig. 16.3C).

The Visual Field

The visual field is the part of our surroundings from which the eyes can perceive light (without movement of the eyes or the head). Together, the two eyes cover a large area (Fig. 16.3A and B). Light from a particular point in the visual field falls on a particular point on the retina. Because of the refraction of the light when it passes the optic media of the eye, the image on the retina is upside down. For convenience, we divide the retina and the visual field vertically in two halves: the **temporal** parts—that is, lateral parts, toward the temple—and the **nasal** parts (Fig. 16.3C). The nasal halves of the retina receive light from the temporal halves of the visual field, and the temporal halves of the retina receive light from the nasal visual field. The situation is the same for the upper and lower halves of the retina and the corresponding parts of the visual field. Thus, light from the lower half of the visual field reaches the upper half of the retina. The lateral 30 degrees of the visual field is viewed by one eye only, because the nose prevents light

from reaching the anterior part of the temporal retina (Fig. 16.3C), and is therefore called the **monocular zone**. Both eyes view the rest of the visual field, the **binocular zone**.

Damage to parts of the nasal retina in one eye produces a blind spot in the temporal visual field; damage in the upper half of the retina in one eye produces a blind spot in the lower visual field; and so forth. Damage that affects the binocular zone usually goes unnoticed by the patient. Therefore, the visual field must be examined for each eye separately. This is done with the person fixating the gaze at a target straight ahead while one eye is covered. Often it suffices to test the outer

borders of the visual field by moving an object (e.g., the finger of the examiner) from well outside the visual field toward its center. A systematic examination covering all parts of the visual field for each eye is done by so-called **perimetry**.

The Lens and the Far and Near Points of the Eye: Accommodation

The lens is transparent and built of cells that form long fibers. It is elastic and, when loosened from the ciliary body to which it is attached with thin **zonular fibers** (Figs. 16.1 and 2), it becomes rounder. Contraction of

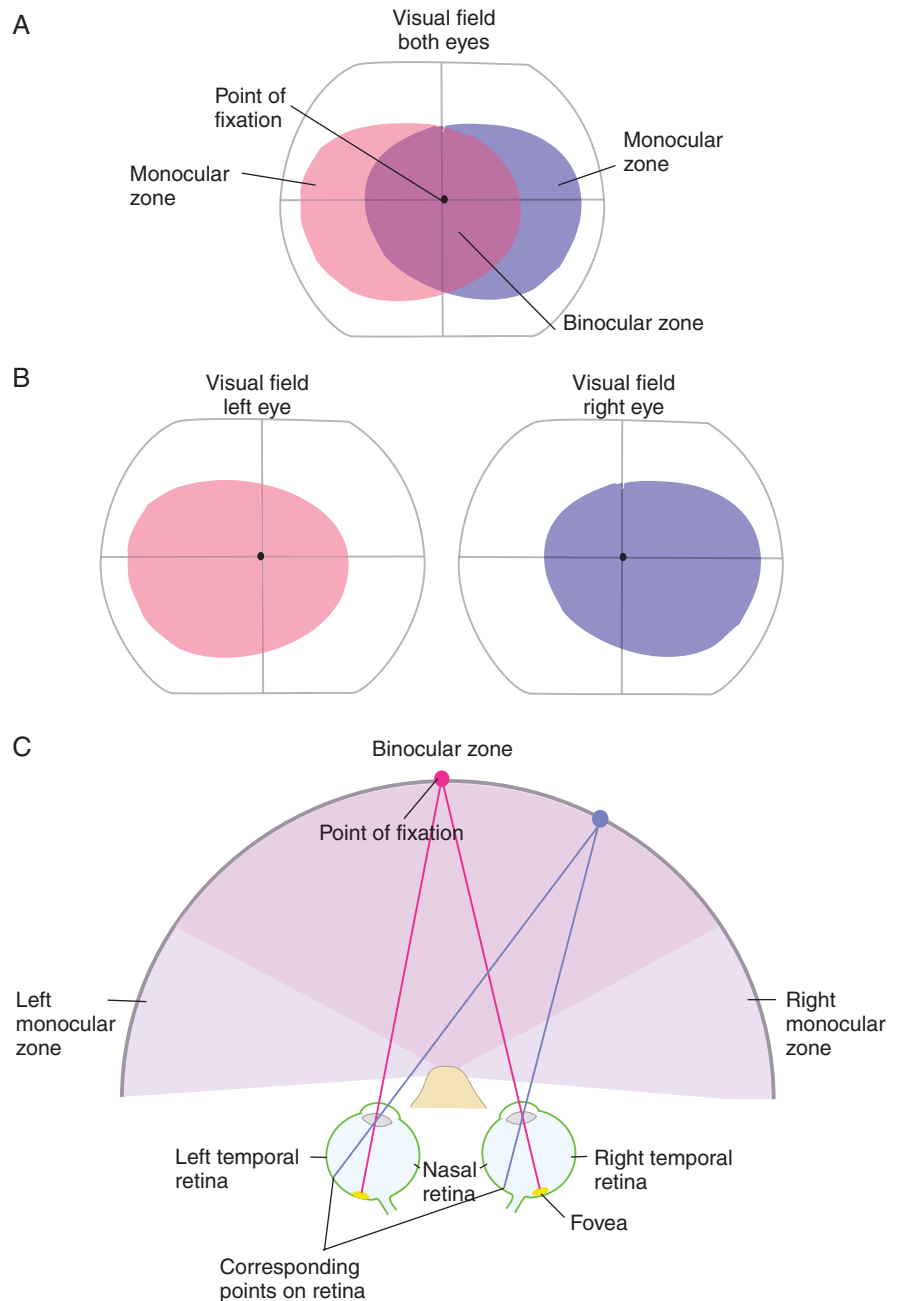


FIGURE 16.3 *The visual field.* **A:** The visual fields of both eyes when the gaze is directed straight ahead. **B:** The visual field as determined for each eye separately. **C:** The positions of the eyes ensure that the images fall on corresponding parts of the two retinas.

the ciliary muscle reduces the diameter of the ring formed by the ciliary body. This slackens the zonular fibers and enables the lens to become rounder; that is, its curvature (convexity) and thus also the refraction of the light increase. Contraction of the ciliary muscle is required sharply to see objects that are closer to the eye than approximately 6 m. This distance is called the **far point of the eye**. In a normal—**emmetropic**—eye, the length of the eyeball is accurately adjusted to the refraction of the cornea and the lens in the relaxed state. When viewing objects at distances greater than about 6 m, the lens maintains the same convexity, and yet the image is always focused on the retina. This is because the light rays entering the eye from points at such distances are all virtually parallel and are therefore collected in the plane of the retina (like a camera focused at infinite distance). If the length of the eyeball differs from the normal (even by only a few hundred microns), the light rays are not collected in the plane of the eye and the sight is blurred. If the eyeball is too long, the light rays are collected in front of the retina. This condition is called **myopia** and is corrected by concave (–) glasses.¹ If the eyeball is too short, the light rays meet behind the retina. Convex (+) glasses correct this **hypermetropia** (in children, because of their elastic lenses, the error is easily corrected by constant accommodation).

The closer an object comes within the far point of the eye, the more the convexity of the lens must be increased by contraction of the ciliary muscle. Such adjustment of the lens for near sight is called **accommodation**. The closest distance from the eye at which we can see an object sharply is called the **near point of the eye**. One's own near point can be easily determined by fixing the eyes on an object (e.g., a finger) that is gradually moved closer to the eye, until it no longer can be viewed sharply.

The far point of the eye depends on the curvature of the lens in its “relaxed” state—that is, with no contraction of the ciliary muscle—and remains stable throughout life. The near point, however, depends on the ability of the lens to increase its curvature and moves gradually away from the eye from birth until about the age of 60. This happens because the lens becomes gradually stiffer and less elastic, so that the ability to increase its convexity declines steadily. At about the age of 45 so much accommodation is lost—or, in other words, the near point is so far away—that it is difficult to read fine print. This condition, called **presbyopia**, is corrected by the use of convex (+) glasses of appropriate strength.

1 A correlation exists between much reading (i.e., accommodating for long periods) and development of myopia in adolescents. Animal experiments confirm that how the eye is used influences the growth of the eyeball. Thus, when 3-month-old monkeys were equipped with +3 glasses, the length of the eye changed to compensate for the refraction error.

THE RETINA

The retina forms the innermost layer of the eye (Fig. 16.2). The outer part of the retina, which adjoins the choroid, is the **pigmented epithelium** consisting of one layer of cuboid cells with large amounts of pigmented granules in their cytoplasm. Internal to the pigmented epithelium follows a layer with **photoreceptors**, and then two further layers with neurons (Fig. 16.4). The processes of the photoreceptors contact the **bipolar cells**, which, in turn, transmit signals to the **retinal ganglion cells**. The axons of the ganglion cells leave the eye in the **optic nerve** to end in nuclei in the diencephalon and the mesencephalon. The pigmented epithelium extends forward to the edge of the pupil (Fig. 16.2), whereas the photoreceptors, bipolars, and ganglion cells are present only in the parts of the retina situated posterior to the ciliary body (pars nervosa retinae).

Unlike many other receptors, the photoreceptors are not of peripheral origin but belong to the central nervous system (CNS). The retina develops in embryonic life as an evagination of the diencephalon (see Fig. 9.3).

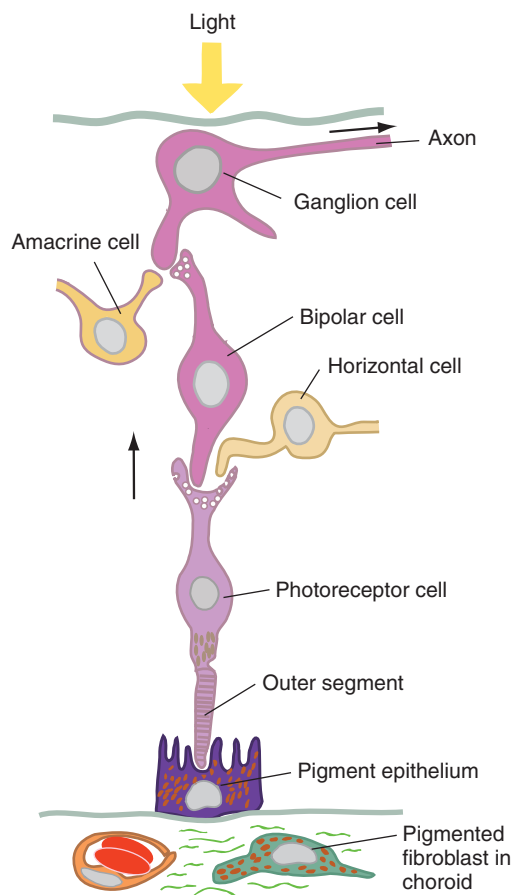


FIGURE 16.4 *The retina.* The main cell types and their interconnections (highly simplified).

Strictly speaking, the term “retinal ganglion cell” is therefore not correct, but it is nevertheless maintained.

Because the photoreceptors are located external to the two other neuronal layers, the light has to pass through the latter to reach the photoreceptors. Because there are no myelinated axons in the retina, however, the layers internal to the photoreceptors are sufficiently translucent.

In addition to the aforementioned neuronal types, the retina also contains many **interneurons**, the **amacrine cells**, and the **horizontal cells** (Fig. 16.4; these neurons are treated in more detail later in this chapter, under “Interneurons in the Retina”). The horizontal cells are responsible for **lateral inhibition** (see Fig. 13.4), among other things. More processing of sensory information takes place in the retina than in any other sense organ. Thus, the visual information transmitted to higher centers of the brain from the retina is already “distorted” by enhancement of the contrast between light and darkness and by preference for signals caused by light from moving objects.

The Retina Has a Layered Structure

Under the microscope, several distinct layers of the retina can be identified in sections cut perpendicular to its surface (Figs. 16.5 and 16.12). Externally, toward the pigmented epithelium, lie the light-sensitive parts of the photoreceptors—their **external segments**. The two types of photoreceptors, the **rods** and the **cones**, can be distinguished because the external segments of the cones are thicker and usually somewhat shorter than those of the rods. Internal to the layer of the external segments, there are three distinct layers with cell nuclei. The **outer nuclear layer** consists of the nuclei of the photoreceptors. The nuclei of the bipolar cells (and many of the interneurons) form the **inner nuclear layer**. The innermost layer of nuclei belongs to the ganglion cells—the **ganglion cell layer**. Between the nuclear layers lie processes of the neurons and their synapses, consequently termed the **outer** and **inner synaptic layers** (or plexiform layers). In the outer synaptic layer, the processes of the bipolar cells end in depressions in the processes of the photoreceptors (Fig. 16.4). The photoreceptor processes contain synaptic vesicles close to the presynaptic membrane.

A special kind of glial cell—the **Müller cells**—extends through the retina from the pigmented epithelium to the vitreous body. They are most likely a form of astrocyte.

Photoreceptors and the Photopigment

Electron microscopy reveals that the outer segments of the rods and cones are packed with folded membrane, forming a large surface containing the light-sensitive photopigment. In the rods, the folds of membrane lie

mostly intracellularly, whereas in the cones they are partly invaginations of the surface membrane. The photoreceptors constantly remove and resynthesize the membrane folds.

The rods and the cones contain different kinds of photopigment. The **rods** contain **rhodopsin**, which has been the most studied. It consists of a protein part, **opsin**, and **retinal**, which is an aldehyde of the **vitamin A** molecule. Retinal is light absorbing and is changed by light (absorption of photons). Simultaneously, the opsin part is changed, and this leads to alteration of the membrane potential of the photoreceptor (hyperpolarization by closure of Na^+ channels). The **transduction** mechanism involves activation of G proteins and intracellular signal molecules, and structurally the photopigments resemble closely other G protein-coupled receptors (e.g., muscarinic receptors, and receptors for smell). The hyperpolarization of the photoreceptors by light stimuli affects the bipolar cells (and retinal interneurons), which then act on the ganglion cells to alter the frequency of action potentials conducted in the optic nerve to the visual centers of the brain.

The photopigment of the **cones** differs slightly from rhodopsin in the structure of the opsin molecules. Further, there are three varieties of **cone opsin** molecules, which explain why we have three kinds of cones absorbing light of different wavelengths (Fig. 16.6). The opsin of the cones is also bound to retinal, but the opsin molecule determines the wavelength sensitivity of retinal.

Dark Adaptation and Light Adaptation

When looking into the eye (e.g., through an ophthalmoscope), the color of the retina is a deep purple because of the content of rhodopsin. The reflection of light from the retina produces the red eyes of flash photography. The color bleaches quickly on illumination of the retina, but it returns slowly in the dark. The light has broken down the rhodopsin, and it takes some time to resynthesize it. We experience the time needed for this process of **dark adaptation** when entering a dark room from strong sunlight. In the beginning, we can hardly see anything, but gradually the ability to see returns. This happens in two stages; first, there is a rapid stage of improvement of about 10 minutes, and thereafter a slower stage of almost 1 hour until full light sensitivity has been restored (if the initial illumination was very intense). Because the rods are responsible for vision in dim light (scotopic vision), the dark adaptation depends on resynthesis of rhodopsin in the rods.

We experience the opposite phenomenon of dark adaptation, **light adaptation**, when moving from darkness into strong light. Also then, after first seeing nothing, vision gradually returns. The strong light bleaches the photopigment massively—that is, there is an intense,

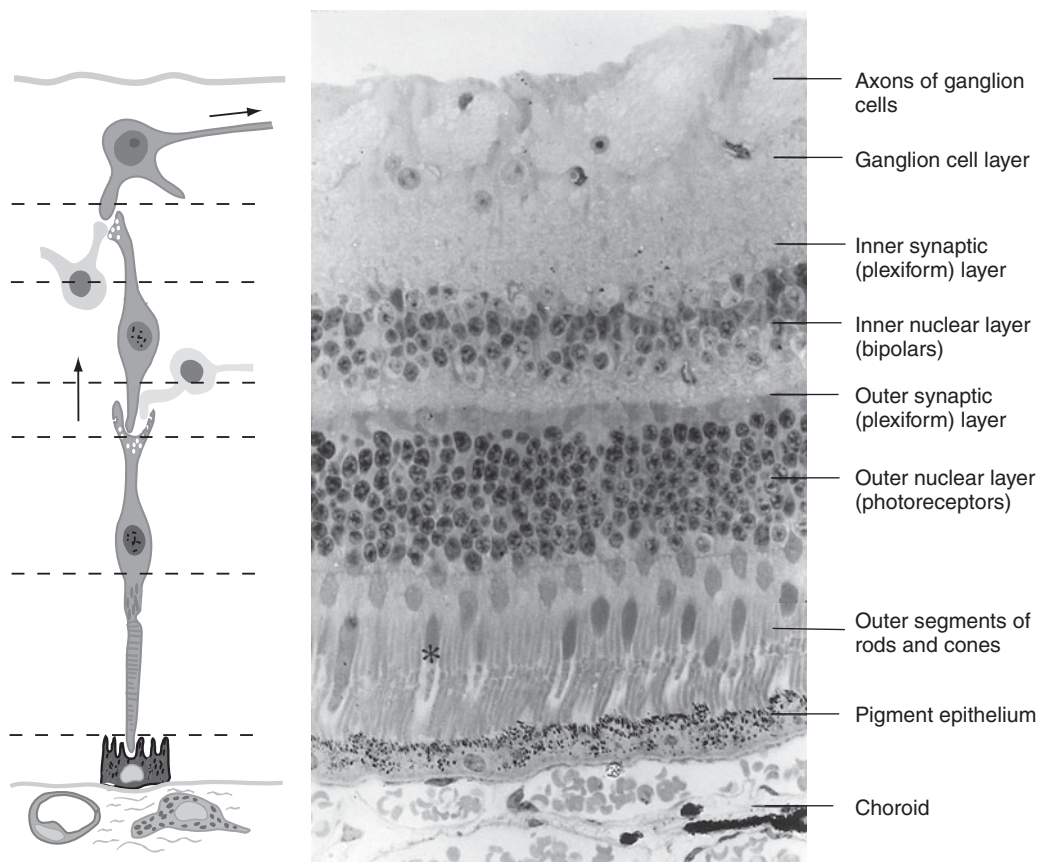


FIGURE 16.5 *The retina.* Photomicrograph of a microscopic section showing the various layers (monkey). The section is from the peripheral part of the retina (this explains the lower density of ganglion cells

here than in Fig. 16.12). The outer segment of a cone is marked with an asterisk. Magnification, $\times 175$.

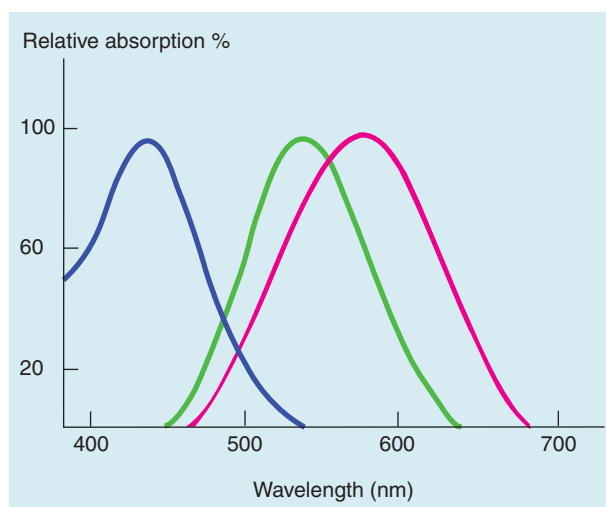


FIGURE 16.6 *The three different kinds of visual pigment.* The diagram shows the efficiency with which the three kinds of cones absorb light of different wavelengths. Note the marked overlapping of the absorption curves.

brief stimulation of many receptors, producing the almost painful sensation of blinding light. The rhodopsin of the rods remains bleached as long as it is exposed to strong light and the rods cannot contribute to vision under these conditions. The cones are first maximally hyperpolarized and are therefore unable to respond. This explains why we cannot discriminate forms and colors at first. However, during a few seconds the outer segments of the cones depolarize to a level where they again can respond to changes of illumination. Change of intracellular Ca^{2+} concentration is of crucial importance for this depolarization. Light adaptation, then, depends on change of the cone-membrane potential. Additional adaptations occur when passing from darkness to light, such as **constriction of the pupils** to reduce light entering the eye (anyone having tried to enter a sunlit street after widening the pupils by atropine has experienced the importance of this adaptation).

Rods and Cones Have Different Properties

The rods are much more sensitive than the cones and react to extremely small amounts of light, whereas the

cones need strong light to react. This is partly because the outer (external) segment of the rod is longer and contains more photopigment than that of the cone. The rods are thus responsible for vision when the light is dim, called **scotopic vision**, whereas the cones are responsible for vision in good light—**photopic vision**.² (That vitamin A is necessary for the synthesis of rhodopsin explains why vitamin A deficiency causes night blindness.) The distribution of light sensitivity for different wavelengths of light is the same for all the rods (maximal sensitivity for wavelengths around 500 nm); hence, they cannot help us **discriminate** between light of different **wavelengths**, which is a prerequisite for color vision. The cones, in contrast, are as mentioned of three kinds, each with a particular variety of photopigment with maximal light sensitivity to different wavelengths. One kind of cone responds best to light with wavelengths in the **blue** part of the spectrum, another in the **red** part, and the other in the **green** part (considerably fewer cones react to blue than to red and to green). One kind of cone alone cannot inform about color, however. This is so because each photopigment is bleached not only by light with wavelengths to which it is maximally sensitive but also by stronger light with shorter and longer wavelengths (Fig. 16.6). Only by **comparing the degree of activation** of the different kinds of cones can the neurons receiving signals from them extract information about the distribution of wavelengths in the light falling on the retina. Together, the three kinds of cones are responsible for **color vision**. As mentioned, however, the cones are not very sensitive to light; from daily experience, we know that we need good light to perceive the color of objects. In poor light, everything appears as a variation of gray.

Other important differences between rods and cones concern their interconnections with other neurons in the retina. Notably, many rods connect to each bipolar cell—that is, there is a high degree of **convergence**. For the cones, on the other hand, there is typically much less convergence, with a few cones connected to one bipolar cell. This also helps us understand why less light is required to convey signals through the optic nerve from the rods than from the cones. The difference in convergence means that the cones provide information with a higher **spatial resolution** than the rods—that is, two points must be farther apart to be perceived as two when the rods are responsible for transmitting the

information than when the cones are responsible. This explains our inability to perceive visual details, such as small letters, in dim light. Therefore, the cones are responsible not only for **color** vision but also for our ability to perceive visual details: they are responsible for precise perception of patterns and **form**.

Color Blindness

In most cases color blindness is an inheritable condition, due to either lack of one kind of cones or an error affecting one of the three cone photopigments. Green or red blindness (or weakness) are the most common forms, affecting about 3% of the male population, whereas blue blindness is very rare. Genes at the X chromosome code the photopigments of the red- and green-sensitive cones, whereas the genes for the blue-sensitive photopigment and for rhodopsin are at autosomal chromosomes. Since the condition is recessively heritable, it is understandable that almost only men are color blind.

Rarely are two kinds of cones lacking, or even all three. Finally, deficient or absent color vision can sometimes be caused by disease of the retina or a lesion of the visual cortex.

Signal Transmission in the Retina

The photoreceptors do not behave like other receptors when exposed to their adequate stimulus: as mentioned, they are hyperpolarized instead of depolarized. Here we briefly discuss how hyperpolarization of the receptors can elicit action potentials in the neurons conducting the signals to the brain.

One important point is that the **photoreceptors** are unusual also in another respect: they are in a **depolarized** state in the dark, with a membrane potential around -30 mV. Na^+ channels that are open in darkness probably cause this. Like other receptors, the photoreceptors (and the bipolar cells) do not produce action potentials but produce only graded changes of the membrane potential. Because the distance is very short from the outer segment—where the membrane potential changes arise—to the synapses between the photoreceptors and the bipolar cells, even small membrane-potential fluctuations cause alterations of the transmitter release from the photoreceptors. (There is thus no need for the production of action potentials, which are necessary only when signals are to be propagated over long distances.) Light falling on the retina causes closure of the photoreceptor- Na^+ channels by degradation of the photopigment and cyclic GMP. Closing of Na^+ channels hyperpolarizes the cell. Transmitter release from the photoreceptors (as from other neurons) is caused by membrane depolarization without any definite threshold that has to be exceeded. Thus, in the dark, the photoreceptors release transmitter continuously, whereas the

² Two kinds of photoreceptors with different sensitivities to light enable the visual system to give meaningful information even with being exposed to extreme variations of light intensities. Release of **dopamine** from one kind of amacrine seems actively to support the change from scotopic (rods) to photopic (cones) vision. Light induces release of dopamine that enhances the signal transmission from the cones compared with the rods. Further, dopamine uncouples electric synapses between amacrine and a certain kind of bipolars, so that signals are less widely distributed horizontally in the retina. This presumably contributes to the enhanced spatial resolution with photopic vision.

release is reduced by light (as if darkness were the adequate stimulus). Recording from **bipolar cells** has shown that they are of two kinds: one is depolarized by light, and the other is hyperpolarized. **Glutamate**—which is the transmitter released from the photoreceptors—has a depolarizing (and therefore excitatory) effect on neurons in other parts of the CNS. With regard to the bipolars, however, one kind is hyperpolarized by glutamate, whereas another is depolarized. This is presumably due to the existence of two different kinds of postsynaptic glutamate receptor.

We can **summarize** the events as follows. When light hyperpolarizes the photoreceptors, the release of glutamate is reduced, as mentioned. This leads to less hyperpolarization, which is the same as depolarization, of one kind of bipolar; thus, some of the bipolars are depolarized and therefore increase their own transmitter release. This is an example of **disinhibition**. The opposite happens with the other kind of bipolar cell, which is hyperpolarized (receives less depolarization) and therefore reduces its transmitter release. Thus, one kind of bipolar reacts with increased transmitter release when light is turned on, the other kind when the light is turned off (Fig. 16.7).

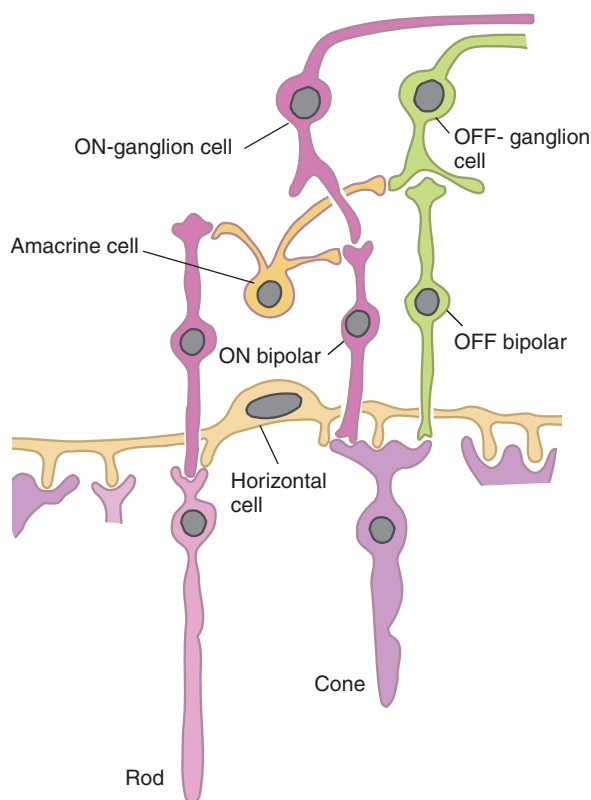


FIGURE 16.7 ON and OFF bipolar and ganglion cells in the retina. Simplified diagram showing the coupling of a cone to two different bipolar cells, and further coupling of the bipolars to ganglion cells that increase or decrease their activity, respectively, when light falls on their receptive fields. Amacrine cells are intercalated in the coupling of rods to ganglion cells.

The bipolar cells have depolarizing (excitatory) effects on the **retinal ganglion cells** (and on amacrine cells), and we can then understand why there are also two kinds of ganglion cells: one that is excited, and one that is inhibited by light hitting the photoreceptors to which they are coupled. We therefore use the terms **ON** and **OFF bipolars** and **ganglion cells**. In contrast to the photoreceptors and the bipolars, the ganglion cells produce action potentials (conducted in the optic nerve to the higher visual centers).

Couplings from Rods and Cones Are Different

Figure 16.7 shows that there are two **parallel signal pathways** from the **cones**. The **ON** ganglion cells increase their firing frequency with increasing intensity of light hitting the cones with which they are connected, whereas the **OFF** ganglion cells increase their firing frequency with increasing darkness. These two channels enable the ganglion cells to inform of a much wider range of light intensities than if there were only one channel. However, the bipolars and the ganglion cells do not inform about the absolute light intensity but the intensity in a small spot on the retina in comparison to the surroundings. This is caused by **lateral inhibition** produced by **horizontal cells** (Fig. 16.7), which are electrically coupled.

The coupling from the **rods** to the ganglion cells is more complicated than from the cones. Thus, bipolars excited by rods do not influence ganglion cells directly but via a special kind of **amacrine cells** (Fig. 16.7). These amacrine cells excite **ON** bipolars and inhibit **OFF** bipolars. It should be noticed that the rods and cones are coupled to the same ganglion cells. Consequently, a ganglion cell transmitting signals from cones in daylight transmits from rods in the dusk.

Interneurons in the Retina

As mentioned, the retina contains interneurons in addition to the photoreceptors, bipolars, and ganglion cells (Fig. 16.4). The **horizontal cells** send their processes in the plane of the retina—that is, perpendicular to the orientation of the photoreceptors and the bipolars (Fig. 16.7). The horizontal cell processes establish contact with the inner segments of the photoreceptors and with the dendrites of the bipolars. They therefore regulate the transmission from the photoreceptors to the bipolars. There is good evidence that the horizontal cells are responsible for the typical receptive fields of the bipolars and ganglion cells with central excitation and peripheral inhibition (or vice versa). This involves complex and unusual synaptic mechanisms that are not fully understood. The horizontal cells are interconnected with **electric synapses**, probably enabling transmission for several millimeters in the plane of the retina.

They release γ -aminobutyric acid (**GABA**) in their efferent synapses onto the inner segments of the photoreceptors. Each horizontal cell is depolarized by **glutamate** released from many photoreceptors. Thus, light hitting the photoreceptors leads to hyperpolarization of the horizontal cells; that is, they release less GABA so that the photoreceptors are disinhibited. Conversely, the horizontal cells hyperpolarize the photoreceptors in darkness.

The other kind of retinal interneuron, the **amacrine cell**, is located with its cell body in the inner nuclear layer and establishes contact with both the axons of the bipolar cells and the dendrites of the ganglion cells (Fig. 16.7). Amacrine cells are thus intercalated between bipolar cells and ganglion cells, and many bipolar cells exert their effect on ganglion cells only or mainly via amacrine cells. As mentioned, this is the rule for **rod bipolars** (bipolars connected with rods). Such amacrine cells (**AII**) form **electric synapses** with **ON** (depolarizing) bipolars and chemical **glycinergic** synapses with **OFF** (hyperpolarizing) bipolars. Some of the processes of the amacrine cells also extend horizontally for considerable distances. The actions of the amacrine cells are varied and complex, and there are numerous morphological varieties. They are also heterogeneous with regard to their transmitter content, and, as mentioned, some establish both chemical and electric synapses. One subgroup of amacrine cells contains **GABA**, for example; others contain **acetylcholine** or **dopamine**. At least seven different **neuropeptides** have been associated with amacrine cells. Twenty different kinds of amacrine cells have so far been identified, considering differences in both synaptic connectivity and transmitter content. The amacrine cells play an important role in influencing the activity of many ganglion cells with properties that cannot be explained by transmission directly from bipolars to ganglion cells. For example, amacrine cells appear to be partly responsible for making certain ganglion cells sensitive to light stimuli (contrasts) with a specific orientation.

Receptive Fields of Retinal Ganglion Cells

Most retinal ganglion cells have in common that they are excited most effectively by shining light on small circular spots on the retina. These are the **receptive fields** of the ganglion cells and can be defined as the **area** of the retina from which a ganglion cell can be influenced. The receptive field can of course be determined not only for ganglion cells but also for neurons at all levels of the visual pathways. The ganglion cells are typically excited from a small central circle and inhibited from a peripheral circular zone, or vice versa (Fig. 16.8A). The American neurophysiologist Stephen Kuffler first demonstrated this in the early 1950s. He introduced the terms **on-center** and **off-center** for gan-

glion cells that are activated and inhibited, respectively, by light hitting the central zone of the receptive field. These correspond to **ON** and **OFF** ganglion cells, described above with reference to the center of their receptive fields (Fig. 16.7). Thus, as mentioned, illumination of a small spot on the retina can lead to increased activity in one chain of neurons—forming, as it were, a channel for signal transmission to the higher visual centers—and reduced activity in another. (The arrangement of a central excitatory field and a peripheral inhibitory zone is also found in the somatosensory system.) For receptive fields of the retinal ganglion cells, the central excitatory or inhibitory part of the receptive field can be explained by direct coupling from photoreceptors to bipolars and further to ganglion cells, whereas the peripheral zone with opposite effects must involve **horizontal cells** producing lateral inhibition (Fig. 16.7). As expected, illumination of the complete receptive field—the central and peripheral zones simultaneously—gives a much weaker response from the ganglion cells than illumination of the central zone only (Fig. 16.8).

In conclusion, each ganglion cell brings information to higher visual centers about a particular small, round area in a definite position on the retina, and thus in the visual field. Together, the receptive fields of all ganglion cells cover the whole visual field with the same type of concentrically arranged receptive fields. Ganglion cells that lie side by side in the retina have overlapping, but not identical, receptive fields.

Retinal Ganglion Cells Exaggerate Contours

Recording of ganglion cell activity under different lighting conditions show that the retinal ganglion cells do not give information about absolute light intensity. Rather, their activity depends on the **contrast** of intensity between the light falling on the central and the peripheral parts of the receptive field. For example, for an on-center cell, a narrow beam of strong light hitting precisely the center of the receptive field while the peripheral zone is in darkness evokes maximal firing frequency.

This preference of the visual system for contrast in light intensity can be demonstrated, for example, by looking at a gray circular spot surrounded by black (Fig. 16.9A). Exchanging the black surrounding with a light gray makes the gray spot appear darker (even though the amount of light the eye receives from the central gray area is unchanged). This property of the visual system makes it particularly suited to detect **contours**, which are especially important for analysis of form. Our ability to judge contrasts of light intensities does not just depend on retinal mechanisms, however. For example, the brightness of an area compared with its neighbors depends on our (subconscious) judgment of how the light falls—that is, whether we assume that

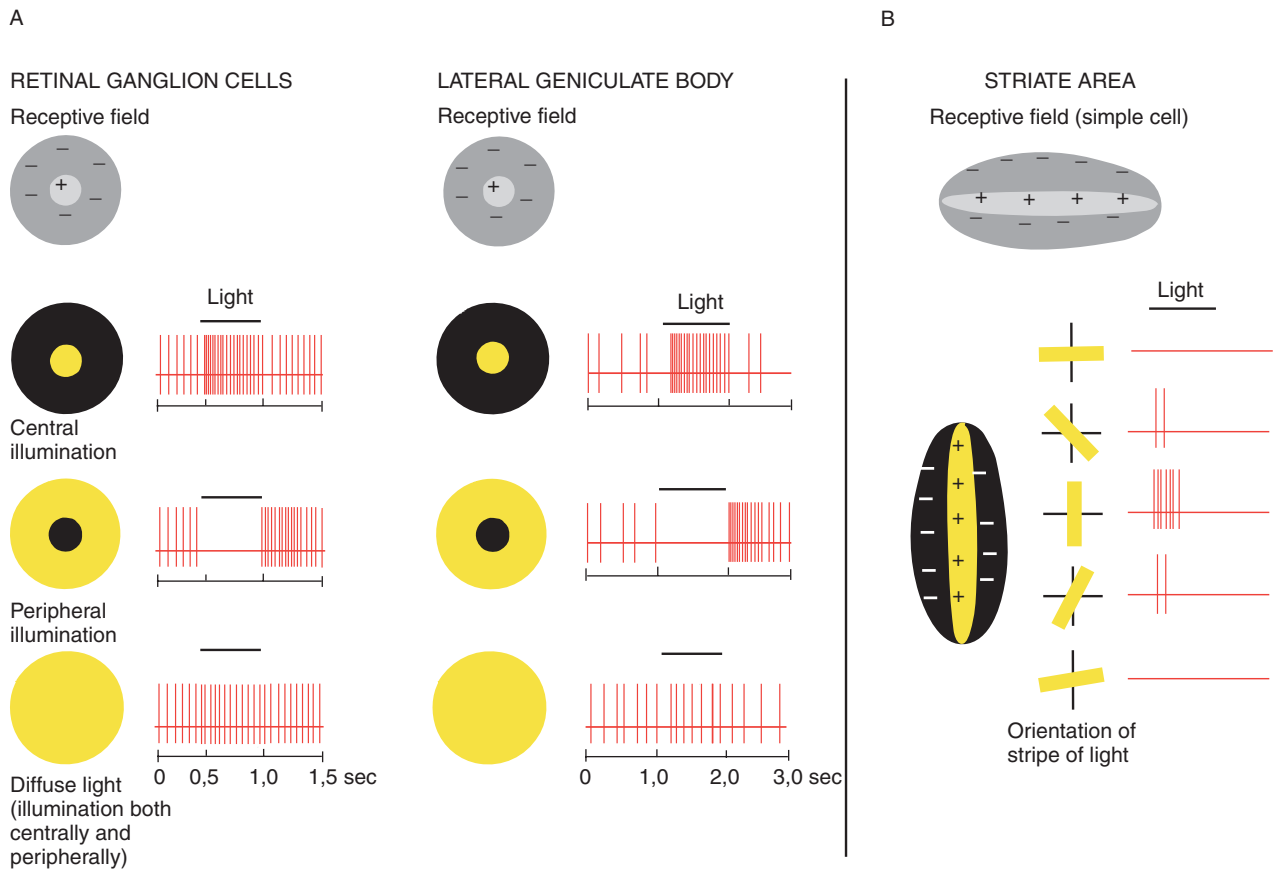


FIGURE 16.8 *Receptive fields of cells at various levels of the visual pathways.* **A:** Retinal ganglion cells and cells of the lateral geniculate body have similar receptive fields. The firing frequencies of the neurons when subjected to different kinds of light stimuli are shown in the graphs. Only cells that are excited by shining light on the central part of the receptive field are shown here (on-center field), but cells with the opposite properties—that is, inhibition from the central

field and excitation from the periphery—also exist (off-center field). **B:** The receptive fields of simple cells of the striate area are typically oblong with an excitatory and an inhibitory zone. The cells are called orientation-specific because they respond preferentially to a stripe of light with a specific orientation. The figure shows only one orientation of the receptive fields, but all orientations are represented among cells of the striate area. (Based on Kuffler et al. 1984.)

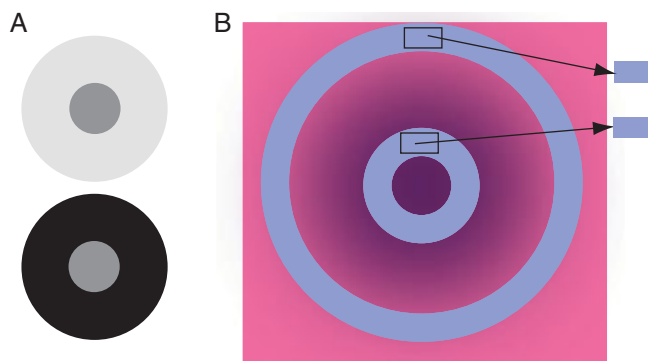


FIGURE 16.9 *Perceived light intensity and color depend on contrast and context.* **A:** The central circle reflects the same amount of light in both examples; yet we perceive the upper one as darker than the lower. **B:** Perceived color intensity differs dramatically with the context. The small colored rectangles are copied from the central and peripheral rings, respectively.

the area is in direct light or in shadow. Processes at the **cortical level** must cause this, since it requires assumptions of the three-dimensional form of the object and from where the light comes. If we misinterpret such conditions, we make false judgments about relative light intensities. Figure 16.7B shows that also our judgment of the color of a field depends on its context.

Visual Acuity and the Size of Receptive Fields

We know from everyday experience that in order to perceive visual details, we must direct the eyes toward an object. The visual axes have to be oriented so that light from the object falls on a small region of the retina in the back of the eye, the **macula lutea** (*macula*, spot; *lutea*, yellow). The macula is about 2 mm in diameter and has a yellowish color when viewed through an ophthalmoscope, distinguishing it from the purple color of the surrounding retina. Here the visual acuity is greatest,

and it decreases steeply when moving peripherally on the retina. The **visual acuity** is expressed as the distance (in degrees) two points in the visual field have to be apart to be perceived as two and not one. In clinical work, visual acuity is usually determined by viewing letters of different sizes at a fixed distance (at the far point of the eye). The center of the macula has a small depression, the **fovea centralis** (Figs. 16.1, 16.2, and 16.10). The depression exists because the bipolars and the ganglion cells are “pushed” aside to enable maximal access for light to the photoreceptors. The small region is also devoid of capillaries. In the central part of the macula, only cones are present, and only in the macular part of the retina is the visual acuity sufficient to enable us to read ordinary print (e.g., a newspaper). The farther we move peripherally from the macula, the lower the visual acuity. One can demonstrate this easily by trying to determine how far out in the visual field one can recognize a face. Closely linked with these differences in visual acuity are differences in the size of the **receptive fields** of ganglion cells in the central and peripheral parts of the retina.

Where the optic nerve leaves the eye, there is a circular area devoid of photoreceptors, appropriately called the **blind spot** (Figs. 16.1, 16.2, and 16.11). We do not notice these blind areas of the two retinas, however, because they are not located at corresponding points (Fig. 16.3).³

Differences between the Central and Peripheral Parts of the Retina

What is the basis for the striking differences in receptive-field size of ganglion cells in central and peripheral parts of the retina? One important factor is that the degree of **convergence** varies dramatically. There are about 100 million photoreceptors in the human retina and only 1 million ganglion cells; on the average, 100 photoreceptors connect to one ganglion cell. In peripheral parts

of the retina, however, the convergence is much greater than 100:1, whereas it is much smaller in central parts (Fig. 16.12). In the central parts of the fovea (foveola; Fig. 16.10), there are even 1:1 connections—that is, one photoreceptor connects to one bipolar, which is connected to only one ganglion cell. As mentioned, the rods show a much greater convergence than the cones, which is in accordance with the distribution of the rods and cones: the macula contains almost only cones, whereas the most peripheral parts contain almost only rods. The relative absence of cones in the peripheral parts of the retina (receiving light from peripheral parts of the visual field) can be demonstrated by how far out in the visual field the color of an object can be perceived. It then appears that the visual field for color is considerably smaller than for moving objects. Another factor that contributes to the higher visual acuity in the central parts of the retina—particularly in the region of the fovea, is that the **density of photoreceptors** is higher there than more peripherally. Figure 16.13 illustrates a third contributing factor: the **dendritic arborizations** of the ganglion cells are more restricted in central than in peripheral parts of the retina, which is also important for the degree of convergence on each ganglion cell. Finally, differences between the retinal **interneurons** in central and peripheral parts of the retina also play an important role.

³ Even when using only one eye we usually do not notice the blind spot, however. This is due to the capacity of the visual system to fill in missing parts of a visual scene (if the size of the missing part is not too big).

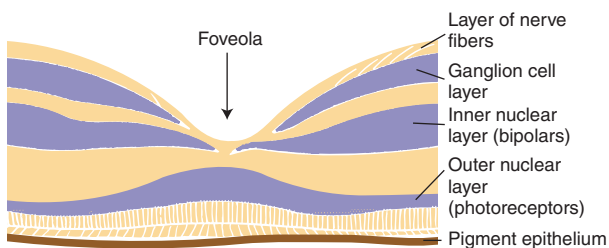


FIGURE 16.10 *The fovea centralis.* Schematized drawing of microscopic section through the posterior pole of the eye. The bipolar and the ganglion cells are “pushed” aside in the most central part of the fovea (the foveola), whereas the density of photoreceptors is at its highest in this same area.

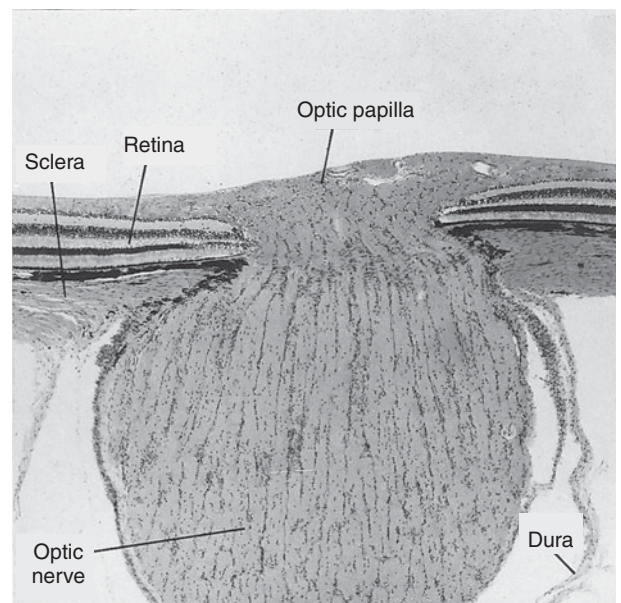


FIGURE 16.11 *The optic nerve and the optic papillae.* Photomicrograph of section through the posterior pole of the eye at the exit of the optic nerve (the blind spot). The optic nerve swells immediately outside the eye because the axons become myelinated. The ganglion cell axons are unmyelinated as long as they course through the innermost layer of the retina.

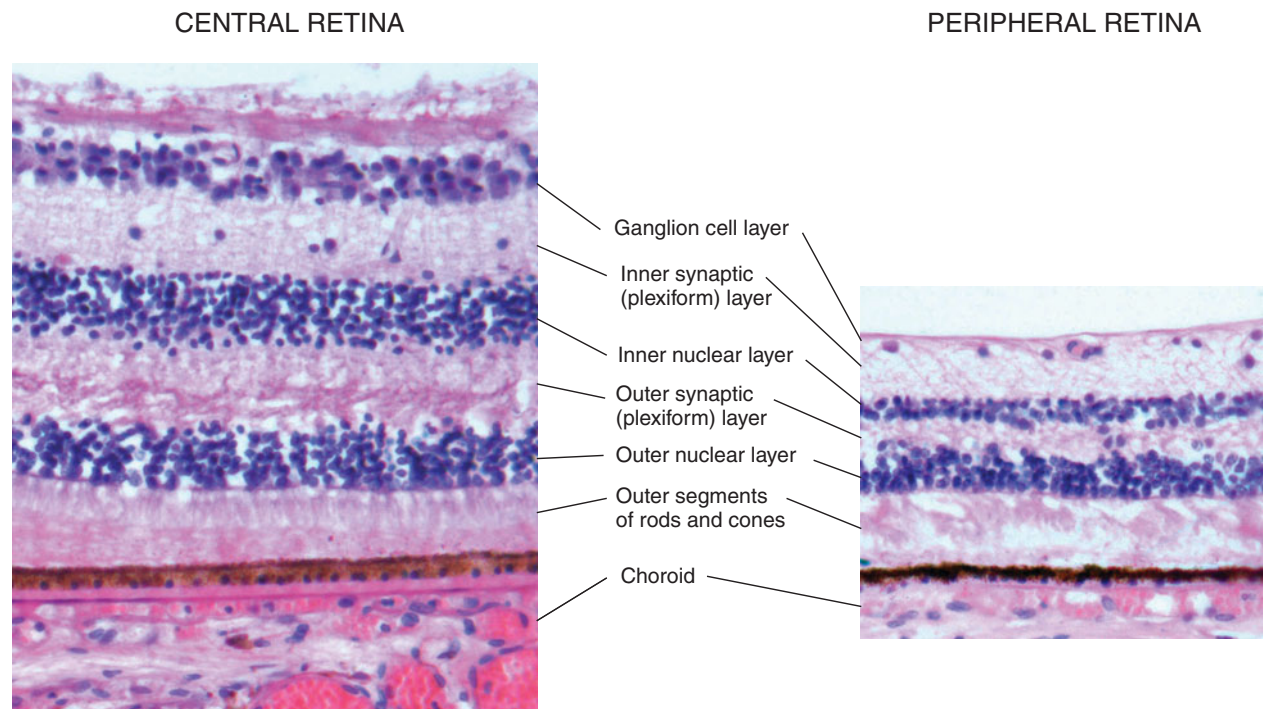


FIGURE 16.12 Central and peripheral parts of the retina. Photomicrographs illustrating how the various retinal layers differ in thickness when moving from the central to the peripheral parts of the retina. The density of ganglion cells is quite different in the two areas.

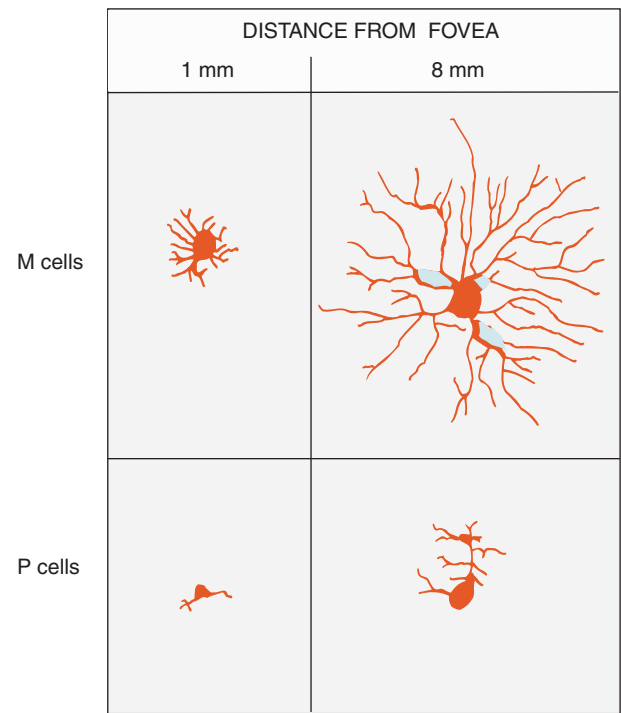


FIGURE 16.13 Two main kinds of retinal ganglion cells (monkey). Both types increase in size in the peripheral direction (distance from the fovea). The extension of the dendritic tree is related to the size of the receptive fields of the ganglion cells. The cells have been visualized by intracellular injection of HRP. (Based on Shapley and Perry 1986.)

There Are Two Main Kinds of Retinal Ganglion Cell

We described two kinds of retinal ganglion cell that differ according to whether they signal light or darkness (ON and OFF ganglion cells). There are, however, further specializations among ganglion cells that we should know to understand the information sent from the retina to higher visual centers.

Anatomic studies showed many years ago that the retinal ganglion cells differ greatly in size. One tendency, mentioned above, is that the **dendrites** of the ganglion cells are longer peripherally than centrally (Fig. 16.13). This relates to differences in the size of their receptive fields. However, ganglion cells with the same placement with regard to eccentricity on the retina also vary in size. It is now customary to recognize two main kinds of retinal ganglion cell, together constituting about 90% of all cells: the **M cells** and the **P cells**. As seen in Fig. 16.13, the P cells are tiny compared with the M cells, but both types are much smaller centrally than peripherally. Physiological studies of their properties indicate that the M cells primarily signal movement and contrasts of illumination, whereas the P cells are responsible for providing information about fine features (high visual acuity) and color. Both kinds can have either ON or OFF properties and can be activated under both scotopic and photopic light conditions. This is what one would expect, because signals from rods and cones

converge on the same ganglion cells (Fig. 16.7). It is nevertheless possible that the M cells play a more important role than P cells in scotopic vision.

Next, we discuss the organization of the pathways followed by the signals from the retina, and we will see that information from the two main types of ganglion cells is kept separate—at least to some extent—up to the cortical level.

More about Retinal Ganglion Cells

The functional properties of the two main morphological types of ganglion cells have been clarified by intracellular staining of cells that first have been characterized by their response to various kinds of light stimuli. Particularly the cat's and the monkey's retinal ganglion cells have been studied in depth. Even though cat and monkey (and presumably human) ganglion cells have several features in common—for example, with regard to the organization of their receptive fields—there are also important differences (notably that cats lack ability to differentiate colors, and their visual acuity is much lower than in monkeys and man). What follows here is based on findings in the monkey. The **M cells** (called A cells by some authors) have a large cell body and a fairly extensive dendritic tree (Fig. 16.13). The axon is relatively thick. The **P cells** (or B cells) have smaller cell bodies, a less extensive dendritic tree, and a thinner axon than the M cells. The P cells are most numerous and probably constitute about 80% of all of the ganglion cells. A major difference is that many of the P cells respond preferentially to light with a particular wavelength—that is, they are **color-specific**—whereas M cells do not have such specificity. The M cells are more sensitive than the P cells to **contrasts** in **intensity** of illumination, however.

A further difference is that the M cells appear to respond better than the P cells to **moving stimuli**. In general, the M cells tend to respond especially when a stimulus starts and stops, whereas the P cells tend to give a signal as long as the stimulus lasts. In spite of anatomic differences between M and P cells, however, each group contains a wide variety of properties. For most properties, the two groups overlap, so that some M cells are more like typical P cells with regard to certain functional properties, and vice versa.

A **third kind** of retinal ganglion cell does not fit into the M and P groups described so far. In the monkey, they constitute about 10% of all ganglion cells. Both anatomically and physiologically, this group is heterogeneous, but the cells are usually smaller than the P cells and have thinner axons. The common feature of this group is that the cells send their axons to the **mesencephalon** rather than to the thalamus, like the M and P cells. A small fraction of the M cells sends an axon (most likely a collateral of the axon going to the thalamus) to the

mesencephalon. It gives some insight into the development of the visual system to compare the proportion of retinal ganglion cells that sends axons to the mesencephalon in various species. Thus, in the cat, about 50% of the axons in the optic nerve pass to the mesencephalon, and the proportion is most likely even higher in lower mammals like the rat. In humans, it is probable that even less than 10% pass to the mesencephalon. With increasing development of the cerebral cortex, more and more of the analysis of visual information takes place at the cortical level rather than in the brain stem visual centers.

ORGANIZATION OF THE VISUAL PATHWAYS

The Visual Pathways

The axons of the retinal ganglion cells constitute the first link in the central visual pathways. All ganglion cell axons run toward the posterior pole of the eye, where they pass through the wall of the eyeball at the **optic papilla** (Figs. 16.1 and 16.11). They then form the **optic nerve**, which passes through the orbit and enters the cranial cavity. Here the two optic nerves unite to form the **optic chiasm** (Fig. 16.14; see also Fig. 6.13). In the optic chiasm some of the axons cross, and crossed and uncrossed fibers continue in the **optic tract**, which curves around the crus cerebri to end in the **lateral geniculate body** (corpus geniculatum laterale) of the thalamus (Fig. 16.15; see also Figs. 6.21 and 6.27). Here the axon terminals of the retinal ganglion cells establish synaptic contact with neurons that send their axons posteriorly into the occipital lobe. These efferent fibers of the lateral geniculate body form the **optic radiation** (radiatio optica) (Fig. 16.16). The optic radiation curves anteriorly and laterally to the posterior horn of the lateral ventricle and ends in the **primary visual cortical area**, which is situated around the **calcarine sulcus** (see Fig. 6.26). Some of the fibers of the optic radiation lie in the posterior part of the internal capsule, where they can be damaged together with the fibers of the pyramidal tract—for example, by bleeding or infarction—thus producing a combination of weakness (paresis) of the muscles of the opposite side of the body and blind areas in the opposite visual hemifield. The primary visual area is **area 17** of Brodmann, which is also called the **striate area**. The latter name (which we will use here) refers to a white stripe in the cortex, running parallel to the cortical surface. The stripe consists of myelinated fibers and is therefore whitish in a cut brain (Fig. 16.17).

Not all fibers of the optic nerve terminate in the lateral geniculate body. Some (about 10% in the monkey) terminate in the **mesencephalon**, especially in the **superior colliculus** and the **pretectal nuclei**. These fibers are of importance primarily for reflex adjustments of the

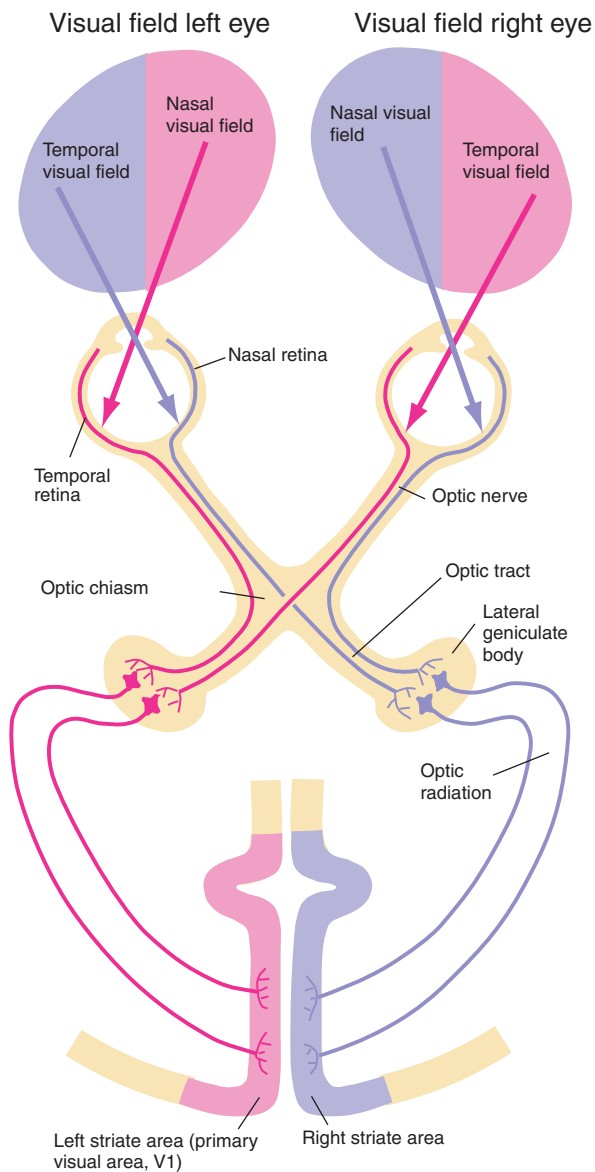


FIGURE 16.14 *The visual pathways.* For didactic reasons, the visual field of each eye is shown separately (cf. Fig. 16.3).

position of the head and the eyes. Some fibers of the optic nerve pass to the **hypothalamus** where they contribute to regulation of circadian rhythms (see Chapter 30, under “Hypothalamus and Circadian Rhythms”).

Axons from the M and P Cells Terminate in Different Layers of the Lateral Geniculate Body

The human lateral geniculate body (and that of other primates, like monkeys) consists of six cell layers (Figs. 16.18 and 16.19). The two ventral-most laminae (1 and 2) are composed of large cells and are therefore called the **magnocellular layers**, whereas the dorsal four are composed of small cells and are called the **parvocellular layers** (Fig. 16.16). Anatomic and physiological studies have shown that the large retinal ganglion cells, the **M cells**, send their axons to the magnocellular layers of the lateral geniculate body, whereas the small ganglion cells, the **P cells**, send their axons (at least preferentially) to the parvocellular layers (Fig. 16.18). There is thus a division of the lateral geniculate body that largely corresponds to the functional division among retinal ganglion cells. It is now usual to speak of the two **parallel pathways**—M and P—from the retina to the lateral geniculate and further to the visual cortex. The significance of this will be discussed in connection with the visual cortex.

Signal Processing in the Lateral Geniculate Body: Corticothalamic Connections

Although the receptive fields of neurons in the lateral geniculate body are closely similar to those of retinal ganglion cells, the lateral geniculate is not merely a simple relay station. Signals from the retina are subject to modification before being forwarded to the striate area. Of special importance are strong, retinotopically organized **corticothalamic projections** from the visual cortex to the lateral geniculate. These connections most likely can control the signal traffic from the retina through the lateral geniculate and have been shown physiologically to influence the properties of the neurons. This modulation probably relates to mental states

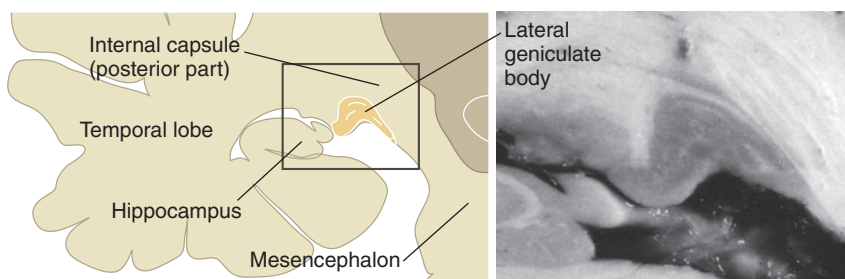


FIGURE 16.15 *The lateral geniculate body.* Drawing and photograph of frontal section through the right hemisphere. The whitish bands separating the lamellae are partly visible.

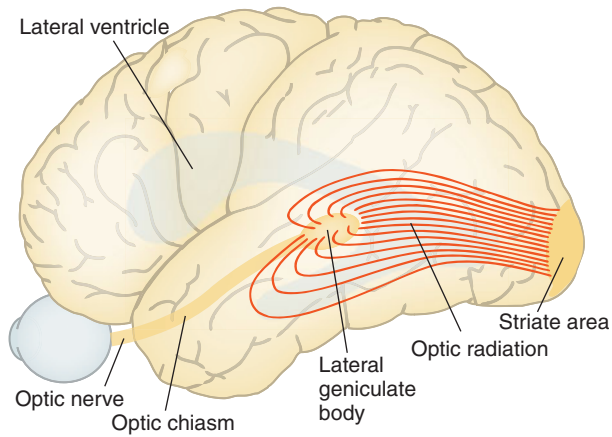


FIGURE 16.16 *The optic radiation.* The course of the fibers from the lateral geniculate body to the striate area shows how the fibers bend around the lateral ventricle and extend partly into the temporal lobe.

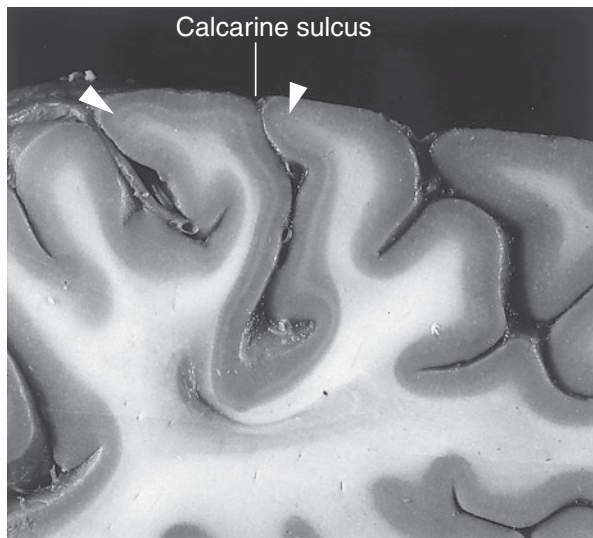


FIGURE 16.17 *The striate area.* Photograph of a section through the human occipital lobe showing the characteristic whitish stripe in layer 4 (the line of Gennari) of the striate area. Arrows mark the border between the striate area and neighboring extrastriate areas. See Fig. 33.5 for photomicrographs of a thionine-stained and myelin-stained sections from the striate area.

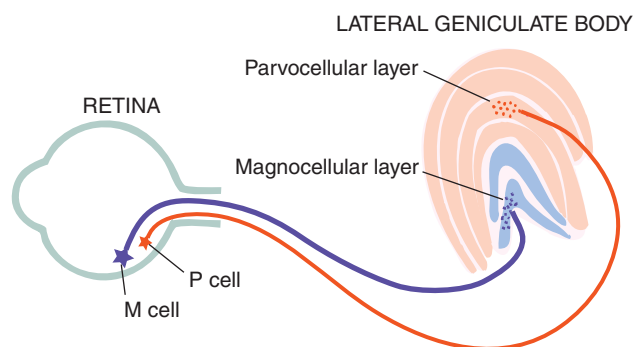


FIGURE 16.18 *The lateral geniculate body.* The two main kinds of retinal ganglion cells end in different layers of the lateral geniculate. (Based on Shapley and Perry 1986.)

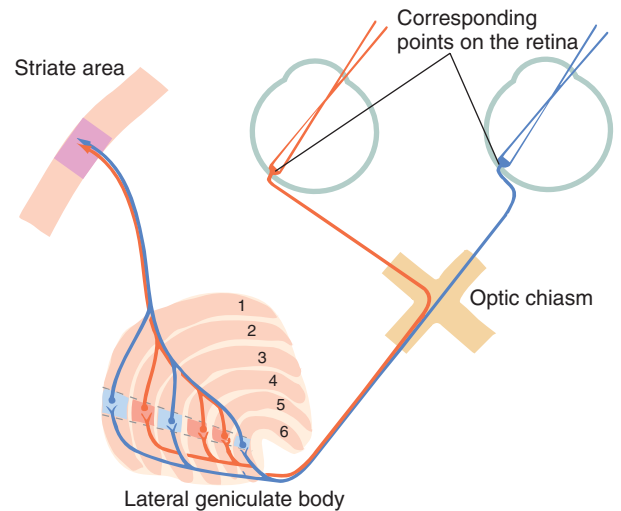


FIGURE 16.19 *Fusion of the visual images.* The signals from corresponding points on the two retinæ end in different layers of the geniculate—that is, signals from the two eyes are kept separate at this level. The convergence of signals takes place in the striate area.

such as **motivation**, **alertness**, **expectation**, and **stimulus context**. Corticothalamic fibers may contribute to the suppression of vision of one eye in patients with **strabismus** (squint, cross-eyed); this has the obvious advantage of avoiding double vision. In addition, the lateral geniculate receives fibers from other sources, notably from **cholinergic** cell groups of the pontine reticular formation; these connections probably regulate the signal transmission through the lateral geniculate to the striate area, in accordance with the level of consciousness and attention.

Finally, a large number of **GABAergic interneurons** and numerous **dendrodendritic synapses** in the lateral geniculate presumably enable neurons with somewhat different receptive fields to influence each other.

Visual Signals from One Side of the Visual Field Reach the Hemisphere of the Other Side

Figure 16.14 shows how the fibers are arranged in the **optic chiasm**. The fibers coming from the nasal halves of the two retinas cross, whereas the fibers from the temporal halves pass through without crossing. In this way, the left lateral geniculate body receives fibers from the temporal retina of the left eye and from the nasal retina of the right eye. The **lateral geniculate body** thus receives light from the contralateral half of the visual field. In functional terms, the crossing of signals corresponds to that taking place in the somatosensory system.

Optic nerve fibers from the two eyes are kept separate at the level of the lateral geniculate body, since three of the six layers receive fibers from the ipsilateral eye, and the three others receive fibers from the contralateral eye (Fig. 16.19). After cutting one optic nerve, almost all cells in three of the layers degenerate

(transneuronal degeneration), whereas the other three layers remain normal. Physiological experiments also show that neurons within each layer of the lateral geniculate body are influenced from one eye only: these cells are **monocular**. We first encounter cells that are influenced from both eyes—**binocular cells**—at the level of the striate area.

Fusion of Visual Images

Normally, we perceive one image of the objects we look at, even though two (slightly different) images are formed on the retina. The two images are perceived as one, and the phenomenon is called **fusion**. Fusion requires that the visual axes of the two eyes be properly aligned, so that the images fall on corresponding points on the retina (Fig. 16.3). The two maculae are obviously corresponding points, and the images fall on them when we fix the gaze on a point to see it as sharply as possible. As mentioned, the signals from the two eyes are kept separate in the lateral geniculate body, but at the cortical level many cells are influenced from both eyes—that is, they are binocular (Fig. 16.19). Convergence in the cortex of signals from corresponding points in the two eyes is a prerequisite for fusion. Fusion is not present from birth but develops gradually from about the age of 3 to 7 months. During this period, the movements of the eyes become coordinated, so that all movements are conjugated and the images fall on corresponding points when the gaze is fixed.

Strabismus (Squint)

Strabismus (squint, cross-eyed) means that the visual axes of the eyes are not properly aligned, and, accordingly, the images do not fall on corresponding points. This may be due to problems with the extraocular muscles or their nervous control, or the “pressure” on the brain to produce fusion may be too weak. The reason for weak pressure may be reduced vision on one or both eyes (e.g., due to retinal disease or a cataract). The lack of fusion in children with a squint leads to underdevelopment or suppression of vision for the eye not used for fixation. In this manner, bothersome double vision is avoided, but even a relatively brief period of strabismus in early childhood may lead to permanently reduced visual acuity. It has been shown in monkeys that strabismus (produced experimentally) leads to a reduced number of cells in the striate area that is influenced by both eyes. In one kind of squint, the child uses the two eyes alternatively for fixation, and in such patients, the visual acuity is usually conserved for both eyes.

The Visual Pathways Are Retinotopically Organized

The arrangement of the visual pathways just described concerns merely retinal halves—that is, a crude retinotopic

localization ensuring that signals from different parts of the visual field are kept separate (compare with somatotopic localization within the somatosensory pathways). But the **retinotopic localization** is much more fine-grained than what appears in Fig. 16.20. Although many fiber systems of the brain are topographically organized, no one is as sharply localized as the visual pathways, which show a true point-to-point localization.

In the **lateral geniculate body**, fibers from differently placed tiny parts of the retina end differently. Each small spot in the retina—and thus in the visual field—is “represented” in its own part of the lateral geniculate (Fig. 16.19). The retinotopic localization in the lateral geniculate is such that neurons influenced from the same part of the visual field (that is, from corresponding points on the retina) lie stacked in a column perpendicular to the layers. This has been demonstrated with various anatomic techniques, and physiologically by inserting microelectrodes perpendicular to the layers and determining the receptive fields of the cells that are encountered. The receptive fields of neurons in the lateral geniculate are quite similar to those of retinal ganglion cells (Fig. 16.8A).

The **thalamocortical** connections from the lateral geniculate body to the **striate area** are also organized with a precise retinotopic arrangement. This has been demonstrated, for example, by injection of a small amount of horseradish peroxidase in the striate area: retrogradely labeled cells are then confined to a narrow column that extends through all six layers of the lateral geniculate.

That all links of the visual pathways are retinotopically organized can be verified by shining light on a small spot on the retina. This evokes increased neuronal activity in a small region of the striate area, and when the light is shone on other parts of the retina, the evoked cortical activity changes position systematically. This kind of experiment has clarified how the **visual field** is represented in the cerebral cortex of animals. Careful examination of patients with circumscribed cortical lesions (often gunshot wounds) provides the basis for maps of the human visual cortex, as shown in Fig. 16.20. Electrical stimulation of the human occipital lobe confirms the retinotopic arrangement within the striate area. Stimulation with a needle electrode usually evokes the sensation of a flash of light in a certain part of the visual field. When the electrode is placed close to the occipital pole of the cerebral hemisphere, the person reports that the flash is located straight ahead, in agreement with the fact that fibers carrying signals from the macula end near the occipital pole. As the electrode is moved forward along the calcarine sulcus, the light flash is perceived as occurring progressively more peripherally in the visual field (at the opposite side of the stimulated hemisphere). If the electrode is placed above (dorsal to) the calcarine sulcus, the light occurs in the lower visual field, whereas stimulation below the calcarine sulcus elicits a sensation of light

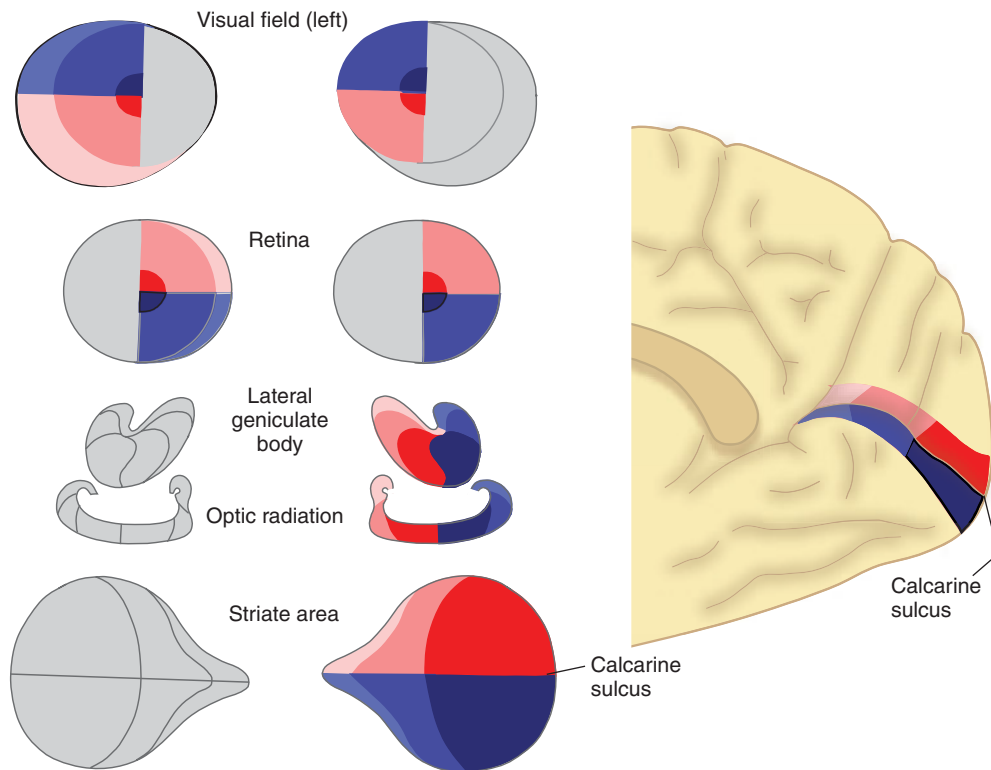


FIGURE 16.20 *Retinotopic localization of the visual pathways.* **Left:** The striate area has been unfolded. Note that information from the upper half of the visual field reaches the part of the striate area below the calcarine sulcus, whereas the lower visual field projects above. Central parts of the visual field are represented most posteriorly and

peripheral parts most anteriorly in the striate area. **Right:** The extension of the striate area on the surface of the occipital lobe; most of it is buried in the calcarine sulcus. The striate area is similarly oriented in the left and the right figures.

in the upper visual field. Studies in healthy humans with positron emission tomography (PET) and functional magnetic resonance imaging (fMRI) have confirmed the main features of the retinotopic organization of the striate area.

Disease processes (e.g., a tumor) involving the visual cortex may also at times elicit sensations of light because the neurons are abnormally irritated. **Epileptic** seizures originating in the visual cortex often start with a **visual aura**—that is, the muscular convulsions are preceded by bizarre patterns of light in the visual field opposite the diseased hemisphere.

Visual Information Can Reach the Cortex via the Superior Colliculus and the Pulvinar

Not all fibers in the optic nerve end in the lateral geniculate body, as mentioned. About 10% (in the monkey) leave the optic tract to terminate in the **pretectal nuclei** and in the **superior colliculus** (see Fig. 27.19). Some fibers end in the **pulvinar** (see Figs. 6.21 and 6.22), and this nucleus (together with another thalamic nucleus, the lateral posterior nucleus, LP; see Fig. 33.7) also receives afferents from the superior colliculus. Like other thalamic

nuclei, these send their efferents to the cortex, notably to the **extrastriate visual areas** (cortical areas processing information from the striate area). Thus visual information may reach the cortex even when the optic radiation or the striate area is damaged. Even though these visual pathways—which circumvent the lateral geniculate body—are retinotopically organized, they are apparently capable only of giving crude information about **movement** in the visual field. Thus, after bilateral damage of the striate area in monkeys, the animals react easily to moving stimuli, even though in other respects they behave as if they were blind. Studies of patients with damage at various levels of the visual pathways and of the visual cortex (localized with the use of MRI) indicate that, as long as parts of the extrastriate visual areas on the convexity are intact, the patients retain some capacity to recognize movements in the visual field. When sitting in front of a large screen with a random pattern of moving dots, patients with damage of the striate area (and surrounding areas on the medial aspect of the hemisphere) reacted with movements of the eyes, apparently following the moving objects. They reported that they felt something moving in front of them, and they had some ability to identify the movement direction. They had no

feeling of seeing anything, however, and when tested with perimetry, they were completely blind. This peculiar condition—termed **blindsight**—is of considerable theoretical interest in the search for the neurobiological basis of consciousness (see later, “Consciousness and Visual Experience”).

The visual connections of the **superior colliculus** are primarily concerned with reflex movements of the eyes and the head, as already mentioned. Thus, most of the efferent connections from the superior colliculus pass to premotor cell groups in the brain stem concerned with control of such movements. The **pretectal nuclei** constitute a link in the pathway for the light reflex (Fig. 27.19).

Central Parts of the Visual Field Are Overrepresented

In addition to the extremely precise retinotopic arrangement of the visual pathways, another important feature must be mentioned. We have described that the density of retinal ganglion cells is considerably higher in central than in peripheral parts of the retina (particularly high in the macula). This corresponds to conditions in the somatosensory system, where the density of receptors is highest at the fingertips (“the somatosensory macula”). Figure 16.20 illustrates that axons from central parts of the retina end in a disproportionately large part of the lateral geniculate body and that this overrepresentation of the central parts of the retina becomes even more marked in the striate area. Again, conditions are similar to those in the somatosensory system (see Fig. 14.8). Thus, the parts of the body and the visual field in which we have the best somatosensory and visual abilities are

provided with a higher density of receptors than other parts, and, further, a much larger number of neurons at higher levels are devoted to the analysis of information from these parts. A disproportionately large part of the striate area treats information from the small macular region (Fig. 16.20). The **magnification factor** quantifies the relation between the cortical area devoted to different parts of the visual field. If information from all parts of the retina were to be treated with similar accuracy, the cerebral cortex would have to be several times larger. Precise control of eye movements, however, ensures that light from the most interesting part of the visual field always falls on the macula.

Interruption of the Visual Pathways

Partial damage to the visual pathways produces symptoms that confirm the arrangement of the cells and fibers at various levels of the optic system (Fig. 16.21). Interruption of the **optic nerve** prevents any visual signals from reaching the brain from that eye. If only the crossing fibers are damaged at the level of the **optic chiasm**, signals from the two nasal halves of the retinas are interrupted, and the patient is blind in the temporal parts of the visual field on both sides. This is called **bitemporal hemianopsia**. The patient may not notice this, however, because the blind part of the visual field is not experienced as darkness but rather as “nothing.” The visual defect may be discovered incidentally by a tendency to bump into objects located a little to the side and perhaps by being hit by a car coming from the side when driving. This kind of visual defect may be caused by a **tumor of the pituitary** (located just below the

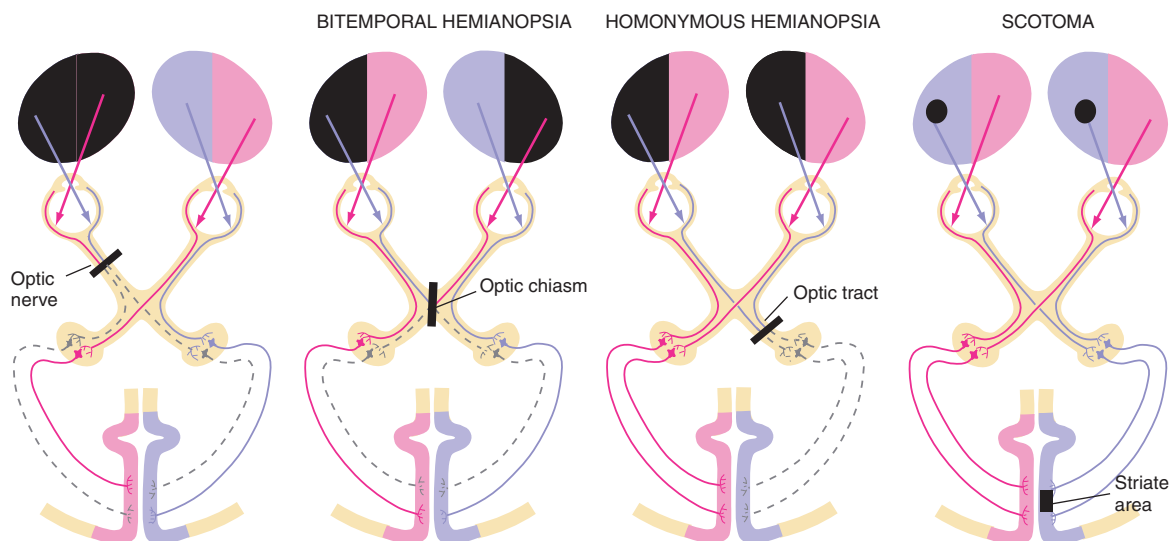


FIGURE 16.21 Visual field defects after lesions of the visual pathways. The black areas indicate the blind parts of the visual field. The visual

fields of the two eyes are shown separately for didactic reasons.

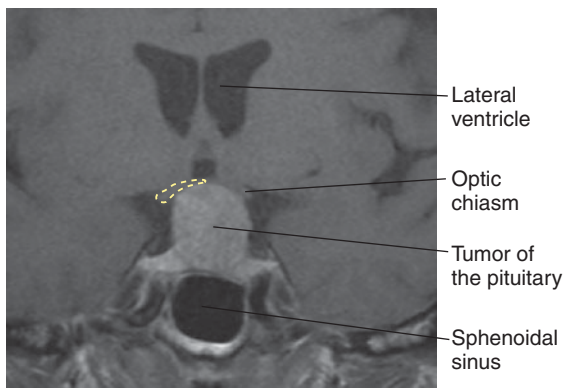


FIGURE 16.22 *Tumor of the pituitary compressing the chiasma.* Frontal MRI. The right part of the chiasma is outlined in yellow. Note how the tumor preferentially hits the middle part of the chiasma, damaging the crossing axons (cf. Fig. 16.21). (Courtesy of Dr. S.J. Bakke, Rikshospitalet University Hospital, Norway.)

optic chiasm). When the tumor grows, it has to expand upward because it is located in a bony excavation (the sella turcica) and thus first compresses the middle part of the chiasm (Fig. 16.22). Damage to the **optic tract** produces a different clinical picture. If the damage is on the right side, visual signals from the temporal half of the right retina and the nasal half of the left retina are prevented from reaching the cortex: that is, the patient is blind in the left half of the visual field. This is called **homonymous hemianopsia**. The same visual defect occurs when the **optic radiation**, or the **striate area**, is totally destroyed. More frequently, however, there are incomplete lesions of the optic radiation (note its position in the posterior part of the internal capsule) or of the striate area, producing blind spots or **scotoma** in the opposite visual field (at corresponding points). Because of the accurate retinotopic arrangement within the visual pathways, mapping of such blind spots enables a precise determination of the site of the lesion.

THE VISUAL CORTEX AND THE FINAL PROCESSING OF VISUAL INFORMATION

The retinotopic localization within the visual pathways may make it appear that a copy of the two retinal images is formed in the striate area. Indeed, the striate area was formerly sometimes called “the cortical retina.” We now know, however, that such a view represents an undue oversimplification; considerable processing and integration of the visual information take place in the striate area. Thus, most neurons have properties that are different from those encountered at lower levels of the visual pathways. Therefore, when we use the term “visual images,” we do not mean images or pictures in the usual sense, either in the retina or in the visual cortex.

Rather, there are temporally and spatially specific patterns of neuronal activity that represent the patterns of light falling on the retina. So far, however, we have only vague ideas about how the activity of the neuronal populations engaged in visual processing at the cortical level is related to our subjective visual experiences.

We will now look at certain fundamental features of the functional organization of the striate area and also mention the regions surrounding it, the so-called **extrastriate visual areas**. These take part in the further processing of visual information. Thus, the visual cortex consists of much more than just the primary visual cortex (V1), the striate area, even though most of the fibers from the lateral geniculate body terminate there. Schematically, the signals from the retina first reach the striate area and are then forwarded to other cortical areas. Different aspects of visual processing such as analysis of color, form, and movement take place in at least partly different subdivisions of the extrastriate areas. Thus, damage to the striate area produces complete blindness in a part of the visual field, whereas damage restricted to extrastriate areas produces defects of restricted aspects of visual analysis. Some of the extrastriate areas receive only indirect projections from the striate area (see Figs. 33.11 and 33.13).

Properties of Neurons in the Striate Area

The retinal ganglion cells and cells of the lateral geniculate body have relatively simple, round receptive fields with a central zone that elicits excitation or inhibition when they are illuminated, as well as a peripheral zone with the opposite effect (Fig. 16.8B). But round spots of light are not an effective stimulus for the neurons of the visual cortex. What constituted the adequate stimulus for the cortical cells remained a mystery for a long time. Diffuse light was not found to be effective, and neither was the kind of stimulus that so effectively affected the cells of the lateral geniculate. In 1962, however, the Nobel laureates (1981) David Hubel and Torsten Wiesel from the United States were able to show that many cells in the striate area respond briskly to elongated fields of light or elongated contrasts between light and darkness. It was furthermore striking that many cells required that the light stimulus be oriented in a specific direction: turning a bar of light some degrees reduced the response markedly. This property was termed **orientation selectivity**. Some cells required a bar of light or a straight light/darkness transition of a specific orientation in a specific part of the visual field to respond, whereas other cells responded to a properly oriented stimulus within a larger area. Such cells thus appear to detect **contours** with a certain orientation regardless of their position within a larger part of the visual field. Hubel and Wiesel called the first type **simple cells** (Fig. 16.8B) and the latter type **complex cells**. Within each group there are several

subtypes according to details of their properties. Why the properties of the complex cells are more complex than those of the simple ones may not be obvious, but presumably the terms were used because Hubel and Wiesel assumed that the properties of the complex cells could be explained by several simple cells acting on one complex cell. The properties of the simple cells, Hubel and Wiesel suggested, could be explained if several neurons in the lateral geniculate with round receptive fields in a row (together forming a stripe) converge on one cortical cell. It is, however, not yet entirely clear which neuronal interconnections underlie the properties of cells in the striate cortex.

Hubel and Wiesel also discovered other fundamental properties of cells in the striate area. Among other things, cells generally respond much better to a **moving** than to a stationary stimulus. Many cells respond preferentially to a line or contour that is moving in a specific direction. Such **direction-selective** cells thus can detect not only the orientation of a contour but also in which direction it is moving. Other kinds of specificities have also been described for cells in the striate area. For example, many **binocular cells** (cells that require input from both eyes) are sensitive to **disparity** of the images: that is, they require that the images from the two eyes are slightly different (disparate). This is always the case for images falling on corresponding points of the retina because the angle of view is different for the two eyes (except in the center of the macula) and also for images of objects that are nearer or farther from the fixation point. The ability to detect **binocular disparity** is important for perception of **depth** and for **stereoscopic** vision.

In addition to single cells being specific or selective with regard to their adequate stimulus, there is also a strong tendency for cells with similar properties in this respect to be located together, more or less clearly separated from cells with other properties. This is called **modular organization** (Figs. 16.23 and 16.24) and concerns properties such as orientation selectivity, wavelength (color) selectivity, and ocular dominance (i.e., which eye has the strongest influence). Such segregation of neurons in the striate area requires that fibers carrying different aspects of visual information from the lateral geniculate end at least to some extent differentially in the cortex. In agreement with this, fibers from the parvocellular layers of the lateral geniculate (Fig. 16.18) terminate deeper in lamina 4 than fibers from the magnocellular layers.

Modular Organization of the Visual Cortex

The first example of modular organization discovered by Hubel and Wiesel was the tendency for cells with similar **orientation selectivity** to be grouped together in **columns** perpendicular to the cortical surface. If we imagine that the striate area is unfolded and we are

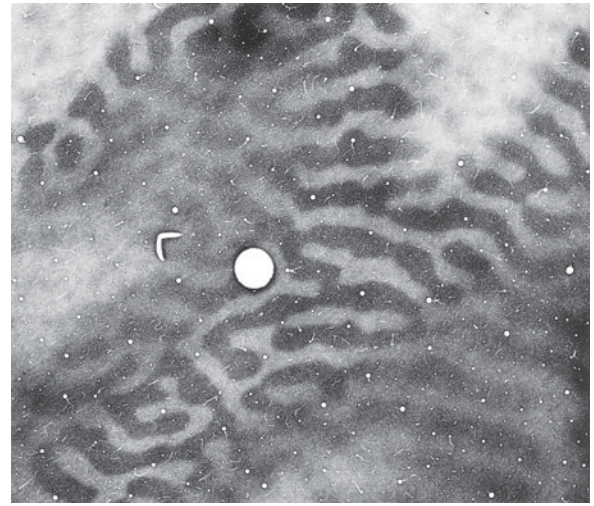


FIGURE 16.23 *Ocular dominance columns.* Photomicrograph of a section cut tangentially to the cortical surface of the striate area and stained to reveal differences in cytochrome oxidase activity. The section is from a monkey that was blind in one eye, causing reduced cytochrome oxidase activity in the regions (light stripes) of the striate area connected mainly with the blind eye. The dark stripes receive their main input from the normal eye. Magnification, $\times 25$. (Courtesy of Dr. J.G. Bjaalie, Department of Anatomy, Institute of Basic Medical Sciences, University of Oslo.)

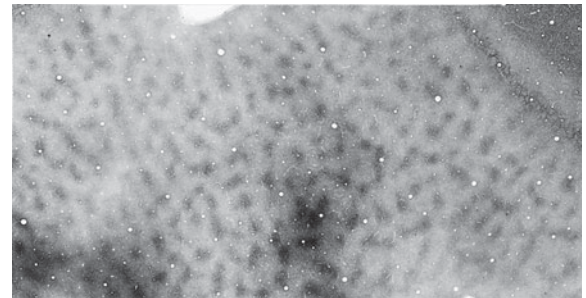


FIGURE 16.24 *Color-specific "blobs" in the striate area.* Photomicrograph of a section cut tangentially to the surface of the striate area of a normal monkey and stained to show differences in cytochrome oxidase activity. The section passes through laminas 2 to 3 and shows numerous small, darkly stained patches or blobs, which correspond to regions with color-specific neurons. Magnification, $\times 25$. (Courtesy of Dr. J. G. Bjaalie, Department of Anatomy, Institute of Basic Medical Sciences, University of Oslo.)

viewing it from above, the groups of cells with similar orientation selectivity are located in an irregular pattern of curving bands. This has been demonstrated with the deoxyglucose method, which demonstrates the neurons that are most active at a certain time. When an experimental animal is exposed for some time to parallel stripes of light, increased glucose uptake takes place in cells distributed in bands in the striate area. Another modular organization concerns **ocular dominance**. As mentioned, many cells in the striate area respond to

light from corresponding points in the two retinas, but for most cells the influence is strongest from one of the eyes. The use of various tracer techniques and the deoxyglucose method has shown that neurons sharing ocular dominance are also distributed in bands within the striate area. Such ocular dominance columns can also be demonstrated in animals with monocular blindness (Fig. 16.23). A final example of modular organization concerns **color-specific** cells (strictly speaking: wavelength specific). They are not aggregated in bands but in clumps or **blobs** in laminae 2 to 3 of the striate area of monkeys. For some unknown reason, these blobs have a higher cytochrome oxidase activity than the surrounding tissue and can thus be easily identified in sections with a simple histochemical procedure (Fig. 16.24).

Extrastriate Visual Areas

As mentioned, several areas around the striate area take part in visual processing. These are collectively called the **extrastriate visual areas** and consist mainly of Brodmann's areas 18 and 19, which in the monkey each consist of several subdivisions. Whereas V1 is often used for area 17 (the striate area), parts of the extrastriate areas are termed V2 to V5. Other parts have specific names. In total, about 30 visual areas have been characterized so far in the monkey, and each is in some way involved in the processing of visual information. Many have a more or less complete representation of the visual field and are retinotopically organized, although with different degrees of precision. Together, the visual areas occupy more than half of all of the cerebral cortex in the monkey. The interconnections between the striate area and the extrastriate areas, and among the extrastriate areas, are numerous and as a rule **reciprocal**. The complete scheme of visual association connections is therefore extremely complex and explains why it is difficult to determine the contribution of each individual area to the processing of visual information.⁴

Further Processing of Visual Information: Segregation and Integration

The properties of single neurons in the striate area suggest that these neurons together are the basis for the cortical analysis of **form, depth, movement, and color**. Their properties, for example, fit predictions made on the basis of psychophysical experiments in humans, such as the preference for contours and for moving

stimuli. Nevertheless, what is taking place in the striate area appears to be mainly a first analysis and sorting of raw data, which must be further processed elsewhere to form the basis for our conscious visual experiences. There is now much evidence of separate, or **segregated**, treatment of the various features of visual images (such as form, color, movement, and location in the visual field) outside the striate area, and we discuss a few examples in the next section. At some stage in the processing, however, different features of the visual image must be brought together and **integrated**. We return to this later in this chapter.

Segregation: Dorsal and Ventral Pathways (Streams) Out of the Striate Area

It has been proposed that a **ventral** stream of information concerning **object identification** ("what") passes downward from the occipital lobe to the temporal lobe, whereas another, **dorsal** stream concerned with **spatial features and movement** ("where") passes upward to the parietal lobe (see Fig. 16.25). This concept is partly based on results from experiments in monkeys with lesions that have been restricted to "visual" parts of either the temporal lobe or the posterior parietal cortex. Monkeys with bilateral lesions of temporal visual areas have reduced ability to identify objects; for example, they can no longer distinguish between a pyramid and a cube. In contrast, lesions of visual parts of the posterior parietal cortex reduce the ability to localize an object in space and in relation to other objects; among other symptoms, such monkeys have difficulties with performing **goal-directed movements**. It suffices with a unilateral lesion of the parietal lobe to produce deficits in the contralateral half of the visual field. This resembles

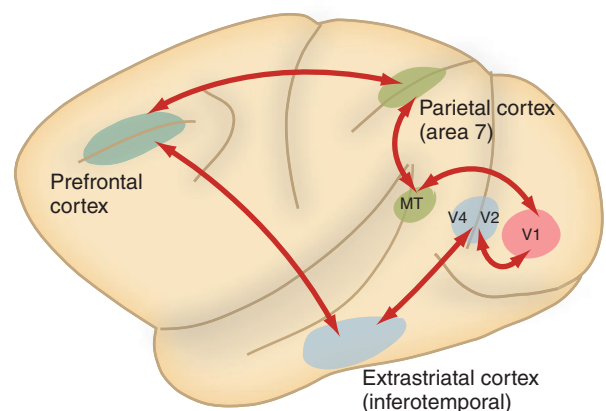


FIGURE 16.25 Dorsal and ventral pathways out of striate area (V1). Monkey. The ventral pathway is especially important for conscious object identification, whereas the dorsal pathway is crucial for perception of movement and space. Note convergence of information from the two pathways in the prefrontal cortex. (Based on data published by Deco and Rolls 2005.)

⁴ It is not immediately clear why the cerebral cortex is organized so that the visual field is represented repeatedly in different parts. It may be a result of the adoption of novel functions by the visual cortex during evolution, and because this probably occurs more easily by adding new areas (or duplicating an old one) than by already existing areas taking up new functions. It is presumably also a simpler solution to have several separate areas than one large area with regard to arrangement of the necessary fiber connections.

the symptoms occurring in humans after damage to the posterior parietal cortex, such as reduced ability to judge movements in the visual field and disturbed eye movements (see Chapter 34, under “More about Symptoms after Lesions of the Posterior Parietal Cortex”). The pathways taken by the signals from the striate area to the temporal and posterior parietal visual regions are not known in detail, but several visual areas are intercalated in the pathways (Fig. 16.25).

It is often stated without reservation that the ventral (temporal) and dorsal (parietal) streams out of the striate area segregate information from **P** and **M** cells, respectively. This is an oversimplification, however. For example, some convergence of information from **M** and **P** cells takes place already in the striate area (besides the more prominent segregation). The sum of evidence suggests that the subcortical pathways from the retina to the striate area are specialized for signaling simple stimulus features, whereas the further pathways do more advanced processing using as, a rule, information from both **M** and **P** cells.⁵

More about Segregated Information Processing in the Extrastriate Visual Areas

While there is no doubt that different aspects of visual information are to some degree segregated in the striate area and in the extrastriate areas, how far the segregation goes is a contentious issue. One striking example of anatomic segregation is the termination in **patches** and **bands** of projections from the striate area to other visual areas. Closer examination of such patterns and correlation with the physiological properties of cells within the patches and bands indicate the existence functionally different information channels out of the striate area, as described in the preceding text with regard to a dorsal and a ventral pathway or stream. More detailed analysis revealed a pathway from cells in the striate area that are predominantly influenced by the **magnocellular layers** of the lateral geniculate. The properties of these striate cells indicate that they signal **movement** and **depth** cues. Another pathway comes from cells that appear to be influenced by the **parvocellular layers** of the lateral geniculate. Accordingly, these striate cells have small receptive fields and are orientation-selective.

⁵ Although behavioral studies suggest that the “parietal” pathway be made nonfunctional by destruction of the magnocellular layers of the lateral geniculate nucleus, the “temporal” pathway is not correspondingly affected by destruction of the parvocellular layers. Thus, whereas selective lesions of the parvocellular layers affect color vision, acuity, and contrast sensitivity, destruction of the temporal visual areas primarily affects form recognition and discrimination. Selective lesions of the magnocellular layers produce reduced perception of contrasts with fast-moving stimuli, while this defect does not occur after lesions of the parietal visual areas. Further, experiments with lesions of the magnocellular layers suggest that depth perception does not depend on information from **M** cells alone.

This pathway presumably signals **forms** and **patterns** and would seem particularly important for our ability to discern visual details. A third “parvocellular” pathway originates in striate neurons that are, at least to a large extent, **wavelength-specific**, signaling information about color.

These three pathways from the striate area appear to be kept separate, at least partly, also at the next station—that is, in **area V2**, which is adjacent to the striate area. From **V2**, information about **movement** is channeled to **area V5** (also called **MT**, the **middle temporal visual area**), whereas information about **color** is channeled primarily to **area V4**. The major outflow from **V5** has been traced to the **posterior parietal cortex**. Information about forms and patterns is channeled primarily from **V2** to **inferotemporal visual areas** (i.e., situated inferiorly in the temporal lobe). How far the specialization goes within each of these areas is not clear, however. The numerous interconnections among the extrastriate areas suggest that they cooperate extensively. Accordingly, single neurons in **area V5**, for example, are sensitive not only to movement but also to certain other visual features. Similarly, neurons in **area V4** are not purely color-specific.

Color Vision and Color Opponency

We discussed parts of the elements responsible for color vision—namely, the three kinds of **cones** with sensitivities for light of different wavelengths (Fig. 16.6). We also emphasized that the brain must compare the degree of stimulation of three kinds of cone to “know” the wavelength composition of the light (and therefore the color of an object). Figure 16.6 shows the large overlap between the sensitivity curves of the three kinds of cone, especially between those with sensitivities in the red and green parts of the spectrum, respectively. How can we then perceive so many nuances of each color? Part of the explanation is found in how the cones are coupled to the next links in the pathway to the cortex. Thus, **retinal ganglion cells** and neurons in the **lateral geniculate body** have narrower sensitivity curves than the cones, making them better at discriminating wavelengths. Many ganglion cells and lateral geniculate cells respond to light of different wavelengths but with opposite signs—one wavelength exciting the cell, the other inhibiting it. This phenomenon is called **color opponency** and must be due to convergence on one ganglion cell (or lateral geniculate cell) of signals from cones with different wavelength sensitivities. Some neurons are excited (**ON** response) by red light in the central zone of the receptive field (Fig. 16.8) and inhibited (**OFF** response) by green light in the peripheral zone; others are inhibited by red light centrally and excited by green peripherally, and so forth. Such combinations improve the ability to **discriminate** wavelengths in the red–green part of the

spectrum. Some neurons exhibit color opponency to blue and yellow light (a combination of red and green), others to white light (stimulation of all three kinds of cones) and darkness.

Even more complex kinds of color opponency are found among neurons in the **visual cortex**. For example, a neuron may respond with an **ON** response to red light and an **OFF** response to green light in the center of its receptive field, whereas the reverse responses are evoked from the peripheral zone (double opponency). Such neurons are found in the cytochrome-rich patches in laminae 2–3 of the striate area (Fig. 16.24). Together, all the color opponency neurons provide accurate information of the wavelength composition of the light hitting a spot on the retina.

Color Constancy

We are used to, and take for granted, that an object has a certain color, regardless of whether we see it in direct sunlight, in the shadow, or in artificial light. For example, we consistently identify a banana as yellow, an apple as red, the grass as green, and so forth (although we perceive differences in nuances). This property of our visual system, called **color constancy**, is by no means self-evident. Thus, the wavelength composition of the light reflected from an object depends not only on the physical properties of the surface but also on the light shining on the object. Different light sources produce light with quite different wavelength composition. The light received by the eye from an object thus differs markedly under different lighting conditions.

Color constancy is obviously of great **biologic importance**. The color of an object gives us essential information as to its nature, but only if the color is an invariant, typical property. For example, we know that a yellow banana is edible, whereas green or brown ones are not. Even under quite different illuminations, we easily make this kind of choice based on color.

We do not fully understand how the brain accomplishes this remarkable task, although it must depend on processing at the cortical level. Color constancy occurs only if the object we look at is part of a **complex, multicolored scene**. Experiments with patterned, multicolored surfaces show that the composition of the light reflected from one part of the visual scene may be changed considerably without changing the color perceived by an observer. If the same square is seen against an evenly dark background, however, the perceived color changes according to the composition of the reflected light—for example, from green to white. This must mean that under natural conditions the brain determines the color of an object by comparing the wavelength composition of the light from its surface with the composition of light from all other surfaces in the visual field.

Afterimages

If we look at a red surface for a few seconds and then move to a white, we see a green surface where we just saw the red. This is called an **afterimage**. The afterimage of yellow is blue, and for white it is black. This phenomenon can be partly explained by selective bleaching of a particular kind of cone. For example, a red light bleaches mainly the cones with wavelength sensitivity in the red part of the spectrum. When the eye then receives white light, the “red” cones are, for a short while, less sensitive than the others. This creates a relative dominance of signals from “green” cones, and the surface is perceived as green. What we have said so far might suggest that the color of the afterimage depends only on the light first hitting the retina. It is not that simple, however. The afterimage depends on the color perceived by the subject, not the absolute wavelength composition—that is, **color constancy** occurs also for after images. The afterimage effect can also occur when looking for a while at a moving object and then at a stationary one. We then perceive an **illusory movement** of the stationary object in the opposite direction.

What Is Color?

Even this fragmentary discussion of color vision may give some thoughts as to the real meaning of the term *color*. Color is, strictly speaking, not a property of an object but a subjective experience of a person seeing the object. The experience is based on how the brain processes information about the wavelength composition of the light reflected from the object (and from all other objects in the same visual scene). That we also can **imagine** colors raises the question of where and how colors (as percepts) are represented in our memory. Now we can only give fragmentary answers to this and other questions about how the brain creates colors or even visual images at all. Nevertheless, the considerations discussed here might serve to emphasize that there is no absolute correlation between a visual stimulus and the perception it evokes. Of course, this is not specific to seeing but pertains to the other senses as well. Fortunately, relationships between certain patterns of stimuli and external events are reasonably constant. Therefore, we usually take the right decisions by relying on our brain’s interpretation (our percept) of the stimuli (the color of a traffic light, the movements of a snake, the contours of a horse, and so forth).

Lesions of the Extrastriate Areas: Visual Agnosia

Humans with lesions of the temporal extrastriate areas show various forms of reduced ability to **recognize objects** (agnosia). A patient may be unable to recognize

faces, another may have no problem with faces but cannot recognize fingers (finger agnosia), and so on (see Chapter 34, under “Lesions of the Association Areas: Agnosia and Apraxia”). High-resolution fMRI in humans and recording from single neurons in monkeys indicate that a small region in the **fusiform gyrus** contains a particularly high proportion of **face-specific** neurons (Fig. 16.26). Nevertheless, recordings with multiple surface electrodes in patients (to localize epileptic foci) suggest that several, separate small areas in the inferior temporal cortex participate in **face recognition**. These areas appear to be parts of a mosaic of areas with different specializations. This agrees with observations after lesions of extensive parts of the inferotemporal extrastriate areas: such patients are deficient in several specific kinds of object recognition and suffer from not only **prosopagnosia** (inability to recognize faces; from Greek, *prosopon* = face). Selective loss of **color vision** (**achromatopsia**) after lesions restricted to a region below the calcarine sulcus (in the **lingual gyrus**) has been convincingly documented (Fig. 16.26). This corresponds most likely to **area V4** in monkeys. Patients with unilateral lesions in this region are reported to see everything in the opposite half of the visual field in black and white, whereas the other half has normal colors.

Damage at the junction between the occipital and the parietal lobes can produce selective impairment of the ability to recognize **movements** (**akinetopsia**). One patient with a bilateral lesion of this region could not see moving objects; when the same objects were stationary,

however, he could easily describe their form and color, and he could judge depth in the visual scene.

Patients with selective loss of **depth perception** after cortical damage have been described. To such patients other people look completely flat, as made from cardboard; patients can recognize their color, contours, and shading, however. The exact site of the lesion in such patients has not been determined, but PET studies indicate that tasks requiring depth perception activate several areas in the cerebral cortex (and in the cerebellar hemispheres), and that these areas overlap with areas involved in other visual tasks.

Integration of Visual Information: One Final Area?

The fact that there is much evidence of separate processing in the visual cortex of different aspects of visual information must not make us forget that at some level a synthesis has to occur. Information about form, position, movement, and color must in some way be linked together. After all, the color “belongs” to a certain object, with a certain form and position and with a certain speed and direction of movement. Our percept is unitary, although the brain first has “dissected” out and analyzed the bits and pieces of information in the light falling on the retina. What does integration of information mean in the context of sensory processing? One might think that there must exist one final, site (area) in which all aspects of visual information are brought together, but **anatomic** data do not favor the existence of such cortical areas. Even in the **prefrontal cortex**, visual information about “what” and “where” is treated separately (in conjunction with coupling of visual features of objects to voluntary movements).

There are also theoretical problems with the notion of a **final integrative area**. Thus, although our conscious experience is unitary, it consists of components to which we have separate, conscious access. For example, a car may be characterized by its color, by its shape, or by its movements. The components are kept separate yet linked in such a way that we perceive them as belonging together. Another theoretical problem with a final integrative area is where to “place” awareness of the visual image. As expressed by the British neurobiologist Semir Zeki (1993): “If all the visual areas report to a single master cortical area, who or what does that single area report to? Put more visually, who is ‘looking’ at the visual image provided by that master area?”

Role of the Striate Area in Visual Awareness and Visual Imagery

We mentioned patients with **blindsight** above (under “Visual Information Can Reach the Cortex via the Superior Colliculus and the Pulvinar”). Such patients have no awareness of having seen anything, yet they

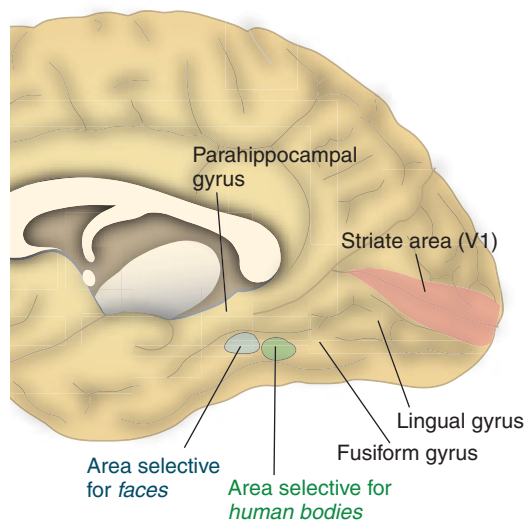


FIGURE 16.26 Subdivisions of the extrastriate visual areas that are selective for recognition of human faces and human bodies, respectively. Surface marking is based on high-resolution fMRI from normal persons. These two areas could not be discerned with standard fMRI with lower resolution. (Based on data published by Schwarzlose et al. 2005.)

can respond to movement in the visual field. In such cases, visual information reaches the extrastriate areas that are specialized for analysis of movements without going through the striate area. We may then ask whether conscious visual experience requires that visual information first go through the **striate area**—in other words, that activation of the striate area is necessary for visual awareness. One approach to this question is to study brain activity with the PET technique while subjects imagine visual scenes, such as a red car or their home street. In such situations, the subject has retrieved the information from memory and “sees” the scene with closed eyes. Which parts of the brain are then specifically activated? It appears that largely the same extrastriate areas are activated during **visual imagery** as when seeing. The PET data are conflicting as to whether the striate area is activated with visual imagery, but the fact that patients with cortical blindness (damage to the striate area) are able to imagine visual scenes strongly suggests that the striate area is not necessary for visual imagery. Also, clinical observations suggest that visual imagery and seeing may not use identical cortical structures. Thus, a few patients have been reported who could recognize objects when seeing them but could not imagine the same objects. The reverse situation has also been described.

Visual Awareness and Synchronized Network Activity

We have treated integration at early stages of the visual processing, such as cells in the striate area responding to contours with a certain orientation regardless where it occurs within a larger part of the visual field. Further integration produces more complex properties, so that cells respond only when several characteristics coincide (such as a contour with a specific orientation moving in a certain direction). Such and more complex integration of data need not occur on single cells, however; it may also occur by coordinated activity in separate neurons that are synaptically linked (neuronal networks). As mentioned, it is noteworthy that the many visual areas are so extensively interconnected. For example, the striate area receives connections from most of the areas to which it projects, and the same holds for many extrastriate areas and their relations to other cortical areas. Presumably, our **conscious visual image** is a product of the total pattern of impulses in many, interconnected cortical areas at any moment. **Synchronized activity** of neuronal groups receiving the same kind of information (e.g., about the direction of movement) may play an important part, and has been shown to occur in cortical areas.

fMRI studies suggest that an extended **network** connecting ventral (temporal) visual areas and parts of parietal and prefrontal areas is especially active in conjunction

with visual **awareness**. A possible common denominator for awareness of sensory information in general, was formulated as follows by Zeman (2004, p. 324): “. . . awareness occurs as the result of physiologically appropriate interactions between neural systems which serve sensation, memory, and action; activity which remains within a single system . . . can influence behavior but will not enter awareness”.

In conclusion, both clinical and brain imaging studies clearly show that extensive parts of the cerebral cortex are involved in conscious visual experience. It should be noted, however, that visual information that is not consciously perceived can nevertheless be used for preparation of voluntary (conscious) movements.

Subconscious Use of Visual Information

A striking example was published by Goodale and Milner in 1992. A woman suffered damage to parts of her occipital lobes (with sparing of the striate cortex) because of carbon monoxide poisoning. Afterward she was unable to recognize objects (**visual agnosia**). For example, she could not decide whether an object was oriented vertically or horizontally, nor could she show the size of the object with her fingers. When asked to grasp it, however, she accomplished this easily and with normal preparatory adjustment of the fingers. Thus, she could adapt her grip to specific visual features, yet she had no awareness of these features. Obviously, the parts of the brain preparing voluntary, **goal-directed movements** have access to information about shape, size, and orientation, no matter whether these features are consciously perceived or not.

Development of Normal Vision Requires Proper Use

There may be other causes than a squint for lack of normal visual development. For example, if the eye does not receive proper stimulation because errors of refraction produce a retinal image that is out of focus or because light for some reason does not reach the retina, vision is not developed normally. In humans, the first 2 to 3 years (in particular the first year) are especially important in this respect; lack of **meaningful use** of the eye even for a short period may then give permanently reduced visual acuity (as discussed in Chapter 9). Animal experiments furthermore show that, contrary to what might be expected, vision is better preserved if both eyes are covered for a period than if only one is covered. This is so because the two eyes “compete” during the early development of the visual system. If only one eye is used, it acquires an advantage and takes over neurons in the visual cortex that normally would have been used by the other eye.

17 | The Auditory System

OVERVIEW

The sense of hearing is of great importance in higher animals—not least in humans, for whom speech is the most important means of communication. The adequate stimulus for the auditory receptors is **sound waves** with frequencies between 20 and about 20,000 Hz. The sensitivity is greatest, however, between 1000 and 4000 Hz and declines steeply toward the highest and the lowest frequencies; that is, a tone of 15,000 Hz must be much stronger than a sound of 1000 Hz to be perceived. The range of frequencies to which the ear is most sensitive corresponds fairly well to the range of frequencies for human speech. The **frequency** of sound waves determines the pitch, whereas the **amplitude** of the waves determines the intensity. Many animals can perceive sound over a much wider range of frequencies than humans can. For example, dogs can hear a whistle hardly noticed by humans, and bats use extremely high-pitched sounds for echolocation. Sound waves pass through the air to the tympanic membrane, which transmits them via a chain of three tiny bones to the **cochlea**. The sensory cells—the **hair cells**—of the cochlea are low-threshold mechanoreceptors sensitive to the bending of stereocilia on their surface. From the cochlea, the signals are conducted to the **cochlear nuclei** in the brain stem through the eighth cranial nerve, the **vestibulocochlear nerve**. This nerve also carries signals from the sense organ for equilibrium—the vestibular apparatus—that anatomically and evolutionarily is closely related to the cochlea. Functionally, however, these two parts have little in common, and we describe the sense of equilibrium together with other aspects of vestibular function in Chapter 18.

From the cochlear nuclei, the **auditory pathways** carry signals to the **inferior colliculus** (and some other brain stem nuclei). Neurons in the inferior colliculus send their axons to the **medial geniculate body** of the **thalamus**. Thalamocortical axons reach the **primary auditory area** (A1) situated on the upper face of the temporal lobe (buried in the lateral sulcus). The auditory pathways from one ear reach both hemispheres, in contrast to the almost complete crossing of somatosensory pathways. Further processing of auditory information takes place in cortical areas surrounding A1 in the temporal lobe. Outward connections from these areas ensure integration of auditory information with other sensory modalities. Nuclei in the brain stem receiving

signals from both ears with a time difference are crucial for our ability to **localize sounds**.

THE COCHLEA

The Cochlea Is Part of the Labyrinth

The **labyrinth** consists of an outer bony part surrounding an irregular canal in the temporal bone, and an inner **membranous part** following and partly filling the canal (Figs. 17.1, 17.2, and 17.3). The membranous canal is filled with a fluid called the **endolymph** (Figs. 17.1 and 17.4). Between the membranous and the bony parts is a space filled with a fluid called the **perilymph**. The composition of the endolymph and of the perilymph differs: the concentrations of sodium and potassium ions in the perilymph are similar to those in the cerebrospinal fluid (i.e., similar to those in the extracellular fluid), whereas the concentrations of these ions in the endolymph are like those found intracellularly. Thus, the cilia of the sensory cells—surrounded by the endolymph (Fig. 17.5)—are embedded in an unusual extracellular fluid with K^+ as the dominating cation. The protein concentration, however, is much higher in the perilymph than in the endolymph and the cerebrospinal fluid. The high K^+ concentration creates a potential of about 90 mV between the endolymph and the perilymph, called the **endocochlear potential** (the endolymph is positive in relation to the perilymph). The special composition of the endolymph and the endocochlear potential are of crucial importance for the transduction mechanism of the hair cells, as discussed later in this chapter.

The labyrinth has two main parts. One is the cochlea and the other is the vestibular apparatus consisting of three semicircular ducts and two round dilatations (Fig. 17.2). Here we consider only the organ of hearing, the cochlea. The **membranous part** of the cochlea—the **cochlear duct**—forms a thin-walled tube with a triangular shape (in cross section), surrounded by the bony part of the cochlea. The duct forms a spiral with two and a half to three turns (Figs. 17.2, 17.3, and 17.4). The lowermost wall of the cochlear duct is formed by the **basilar membrane** (*membrana basilaris*), which is suspended between the two facing sides of the bony canal (Figs. 17.4 and 17.5). At the inner side of the turns, the basilar membrane is attached to a bony

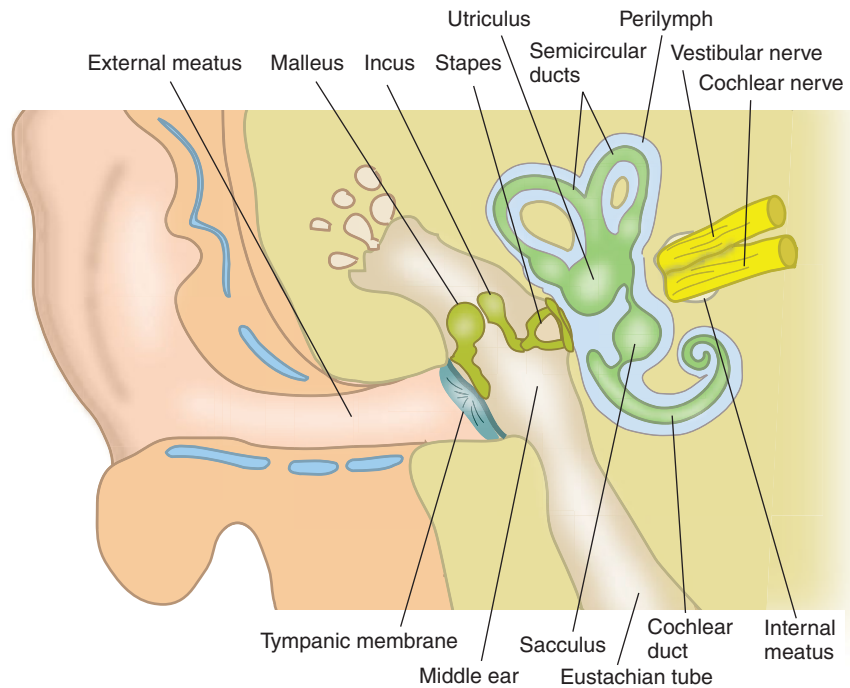


FIGURE 17.1 *The ear.* The middle ear has three ear ossicles, and the inner ear has the membranous labyrinth (located in the temporal bone). The eustachian tube connects the middle ear with the pharynx.

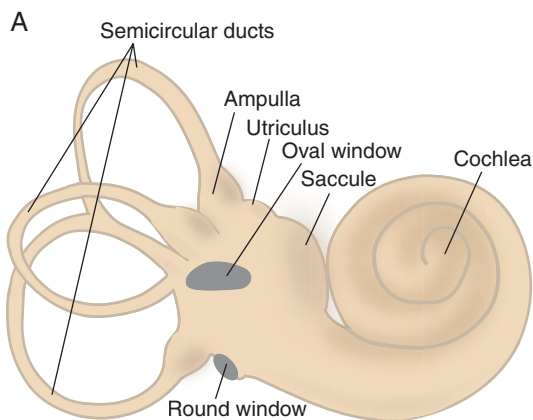


FIGURE 17.2 A: *The membranous labyrinth with its vestibular part (the semicircular ducts, the saccule, and the utricle) and the auditory part (the cochlea).* The stapes is attached in the oval window.



B: *The fluid in the labyrinth visualized via three dimensional reconstruction of MRIs.* (Courtesy of Dr. Einar Hopp, Rikshospitalet University Hospital, Oslo, Norway.)

prominence—the **bony spiral lamina** (lamina spiralis ossea)—which follows the cochlea in its spiraling course (Figs. 17.4 and 17.5). The sensory epithelium, forming the **organ of Corti**, rests on the basilar membrane (Figs. 17.4, 17.5, and 17.6). The length of the cochlear duct, and thus of the basilar membrane, is about 3.5 cm (Fig. 17.7). The thin **vestibular membrane** forms the upper wall of the cochlear duct (Figs. 17.4 and 17.5). The third, lateral, or outer wall of the cochlear duct lies on the bony wall of the canal and is formed by a specialized, stratified epithelium, the **vascular stria** (Fig. 17.6). As the name implies, there are **capillaries** among the

epithelial cells. The vascular stria is responsible for the high K^+ of the endolymph and the positive electric potential between the endolymph and the perilymph.¹

The room outside the cochlear duct consists of two parallel canals. The one situated below the basilar membrane is the **scala tympani**; the one above the

¹ In experimental animals, atrophy of the vascular stria causes hearing loss, and the severity seems to be proportional to reduction of the endocochlear potential. There is evidence that—in addition to loss of hair cells—atrophy of the vascular stria can contribute to presbycusis (age-related hearing loss) in humans.

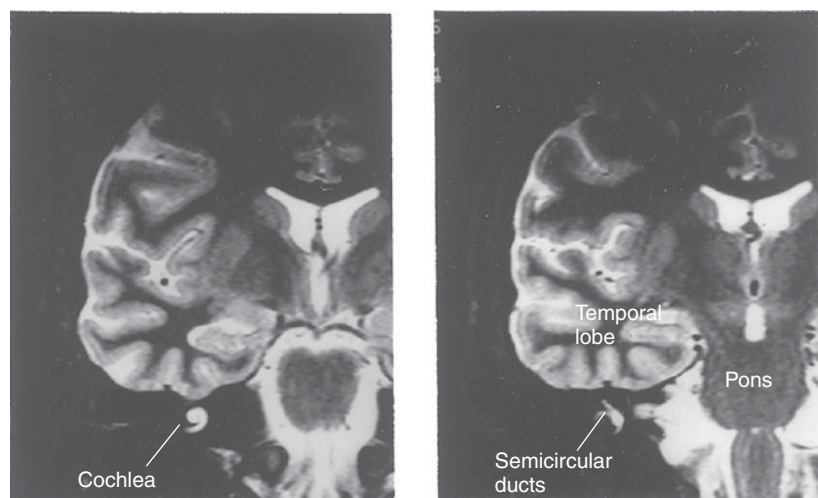


FIGURE 17.3 *The labyrinth.* Two closely spaced, frontal MRIs. The fluid-filled cochlea (left) and semicircular canals (right) are clearly seen just below the temporal lobe (the image is weighted so that water appears white).

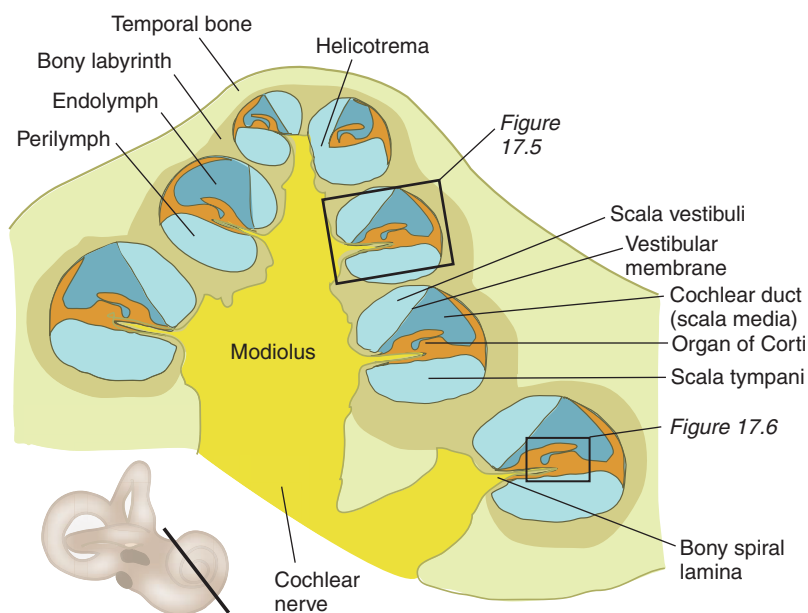


FIGURE 17.4 *Section through the cochlea.*

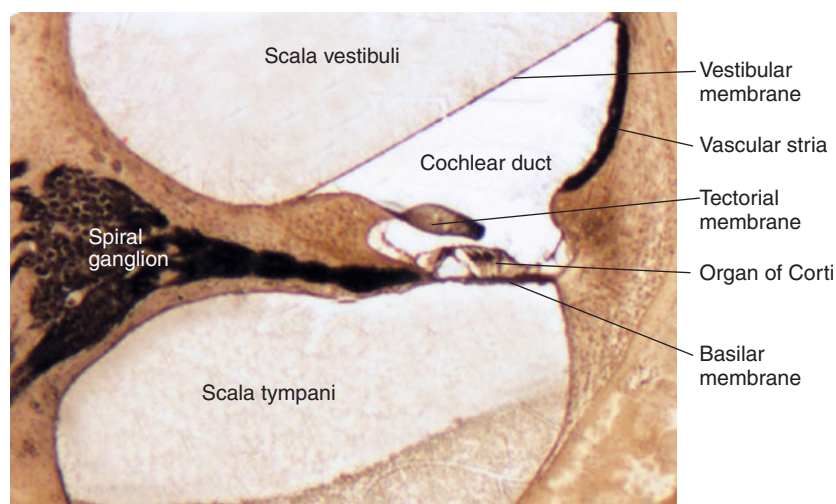


FIGURE 17.5 *Section through the cochlea.* Photomicrograph. Cf. Fig. 17.4.

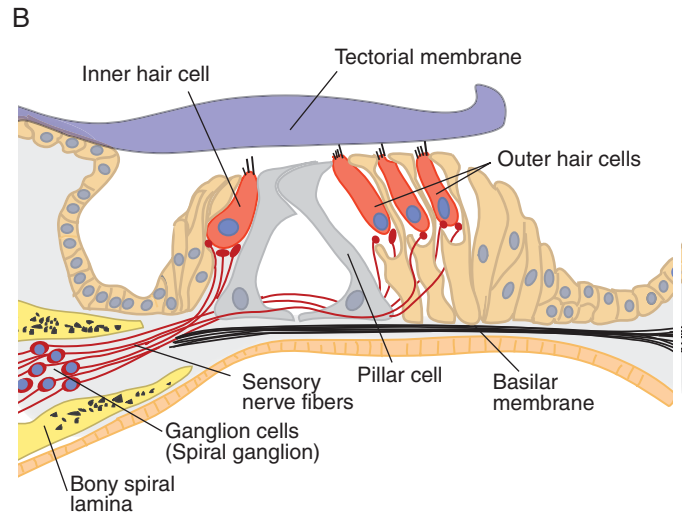
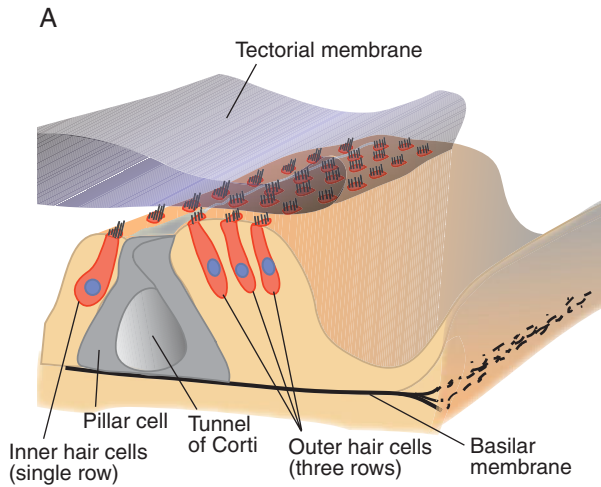


FIGURE 17.6 *The organ of Corti.* **A:** Three-dimensional representation of a short segment of the organ of Corti (the whole organ extends the full length of the cochlear duct). The inner hair cells are in a single row, and the outer hair cells are in three parallel rows. The pillar cells

appear to change their form in relation to loud sounds, possibly to prevent damage of hair cells. **B:** Cf. Fig. 17.4. Note that only the tallest stereocilia of the outer hair cells are in contact with the tectorial membrane.

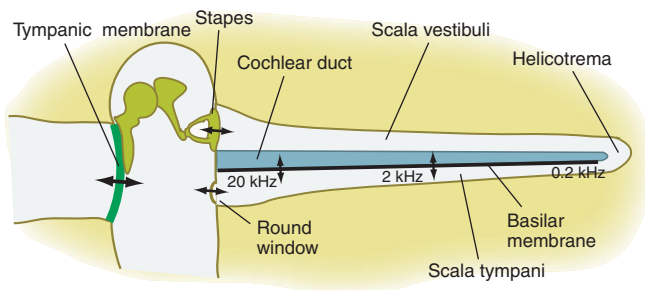


FIGURE 17.7 *The middle ear and the cochlea.* The cochlear duct is pictured as if straightened (length about 3.5 cm). The oscillations of the stapes are transmitted to the fluid in the scala vestibuli and from there to the cochlear duct. Different tone frequencies set different parts of the basilar membrane in motion. Note that the highest frequencies stimulate the hair cells near the base of the cochlea, whereas the lowest frequencies stimulate hair cells near the apex (near helicotrema). (Based on Fettiplace and Hackney 2006.)

vestibular membrane is the **scala vestibuli** (Figs. 17.4 and 17.5). Both have openings or windows in the bone facing the middle ear (Figs. 17.2 and 17.7). The **oval window** (fenestra vestibuli) is situated at the end of the scala vestibuli, whereas the **round window** (fenestra cochleae) is at the end of the scala tympani. The stapes and a thin membrane of connective tissue close the windows, respectively (Fig. 17.7).

How Sound Waves Are Transmitted to the Sensory Cells in the Cochlea

Conduction of sound waves from the air to the receptor cells in the cochlea occurs through the **external ear** (the auricle and the external auditory meatus) and the middle ear or **tympenic cavity** (Fig. 17.1). Sound waves

hitting the skull can also be transmitted through the bone directly to the receptors. This kind of transmission, however, is very inefficient with regard to airborne sound waves and therefore plays no role in normal hearing (bone conduction of sound waves is used for testing the function of the cochlea and also for certain hearing aid devices).

Sound waves hit the eardrum or **tympenic membrane** located at the bottom of the external meatus (Fig. 17.1). The eardrum consists of a thin, tense connective tissue membrane covered by a thin layer of epithelium on both sides; it is richly supplied with **nociceptors**, like the tight skin of the inner part of the external meatus. The three **ossicles** form a chain through the middle ear and connect the eardrum with the oval window (Figs. 17.1, 17.2, and 17.7). The **malleus** (the hammer) has a shaft that is attached to the inner side of the eardrum. The head of the malleus connects to the **incus** (the anvil) by a joint, and the incus is further connected to the **stapes** (the stirrup) by a joint. The basal plate of the stapes inserts in the oval window, thus closing the scala vestibuli (Fig. 17.7). The sound waves make the eardrum and the ossicles vibrate with the frequency of the waves, and thus the movement transmits to the fluid in the cochlea. Because the area of the eardrum is so much larger than that of the basal plate of the stapes, the **pressure** per square unit increases 20 times. This amplification mechanism increases the sensitivity for sound dramatically, compared to a situation without the ossicles. Normally, even the slightest movement of the eardrum is sufficient to cause stimulation of the receptors in the cochlea: sound waves with **amplitude** of only 0.01 nm suffice to produce the weakest perceptible sound with the frequency to which the ear is most sensitive. If the sound waves

were to be transmitted directly from the air to the fluid in the cochlea, a large proportion would be reflected without reaching the sensory cells of the cochlea. The sound would have to be much stronger to be perceived in such a situation. This is the case when diseases of the middle ear destroy the ossicles or stiffen their joints and thus eliminate the amplification mechanism. The ensuing hearing loss is called **conduction deafness**. A prerequisite for the free movement of the eardrum is that the pressure be equal on the two sides. This is ensured by the **eustachian tube** (tuba auditoria), which connects the middle ear cavity with the pharynx (Fig. 17.1).

When the **stapes** is pressed (slightly) into the **oval window**, the pressure of the sound waves is transmitted directly to the fluid (the perilymph) in the scala vestibuli. Because water is incompressible, the sound waves can cause movement of the fluid only because the room can expand at some other point. The thin, compliant membrane that covers the **round window** allows such expansion. The membrane is pressed outward (into the middle ear) each time the stapes is pressed into the oval window. Movement of the perilymph in the scala vestibuli transmits immediately to the endolymph in the cochlear duct through the thin **vestibular membrane**. The movement thus propagates to the basilar membrane, which is pressed downward, and transmits the movement to the perilymph in the scala tympani. In short, movements of the stapes in and out of the oval window—in pace with the sound waves—produce corresponding movements of the basilar membrane. Movements of the basilar membrane stimulate the receptor (hair) cells.

Next, we describe how the receptor cells are arranged in the organ of Corti and the mechanism by which movements of the basilar membrane lead to excitation of the receptor cells.

Sound Pressure Is Measured in Decibels

The **amplitude** of the sound waves determines the **sound pressure**—that is, the pressure of air molecules on the tympanic membrane. The most intense sound that the human ear can perceive is about 10^{12} times stronger than the weakest. A logarithmic scale is used for sound pressure. One decibel (dB) represents the pressure necessary to produce the weakest perceptible sound, whereas just below 130 dB represents the strongest (the pain threshold is at 130–140 dB). Ordinary speech produces a sound pressure between 30 and 70 dB. This scale gives relative measures of intensity because the sensitivity of the ear differs for different frequencies.

The Organ of Corti

The receptor cells in the cochlea are called **hair cells** because they are equipped with sensory hairs, or **stereocilia**, on their apical surface (Figs. 17.6 and 17.8; see also

Chapter 18, under “More about the Vestibular Hair Cells”). Along the basilar membrane, there are two populations of receptor cells: one formed by the **outer hair cells**, the other by the **inner hair cells** (Fig. 17.5). The inner hair cells, closest to the bony spiral lamina, form a single row, while the outer hair cells form three parallel rows. There are approximately 3500 inner and 15,000 outer hair cells in the human cochlea. The two kinds of cells differ in both morphology and innervation. Many more sensory nerve fibers contact the inner than the outer hair cells, whereas the reverse situation exists for the efferent innervation. There is good evidence that the inner hair cells are responsible for signaling sound, whereas the outer hair cells regulate the sensitivity of the sense organ.

Supporting cells surround the hair cells. Two rows of especially large supporting cells, the **pillar cells**,² separate the inner and outer hair cells and form the **tunnel of Corti**. Above the hair cells lies a thick plate, the **tectorial membrane**, which is indirectly attached to the bony wall of the cochlea (Figs. 17.5 and 17.6). Sensory (afferent) nerve fibers of the **eighth cranial nerve** contact the basal parts of the hair cells (Figs. 17.6 and 17.8). The cell bodies of the sensory neurons are located in the **bony spiral lamina** close to the midportion of the cochlea (the **modiolus**, Figs. 17.4, 17.5, and 17.6). **Efferent nerve fibers** also contact the hair cells (Fig. 17.8), enabling the central nervous system (CNS) to control the sensitivity of the auditory receptors.

Deflection of the stereocilia, caused by movement of the basilar membrane, produces **receptor potentials** in both inner and outer hair cells. The further events differ, however: in the inner hair cells, depolarization facilitates transmitter release, whereas depolarization stimulates contractile activity in the outer hair cells.

The Inner Hair Cells and Mechanoelectric Transduction

Unlike the stereocilia of the outer hair cells, those of the inner hair cells are not in direct contact with the tectorial membrane (Fig. 17.6B). How can then the sound waves lead to deflection of the stereocilia of the inner hair cells? Most likely, movement of the fluid surrounding the stereocilia is sufficient, causing the stereocilia to move back and forth, in pace with the vibrations of the basilar membrane. The stereocilia are stiff, due to their content of actin filaments. Thus, each cilium moves like a rod around its point of insertion in the top plate of the hair cell (Fig. 17.8). Further, the stereocilia are coupled so that they all move together. Thereby, the sensitivity

2 The pillar cells and other supporting cells in the organ of Corti appear to contribute to control of cochlear sound sensitivity. In animals exposed to loud sounds, the form and mutual positions of the supporting cells change in pace with changes in cochlear sensitivity. This may be a mechanism to protect the hair cells.

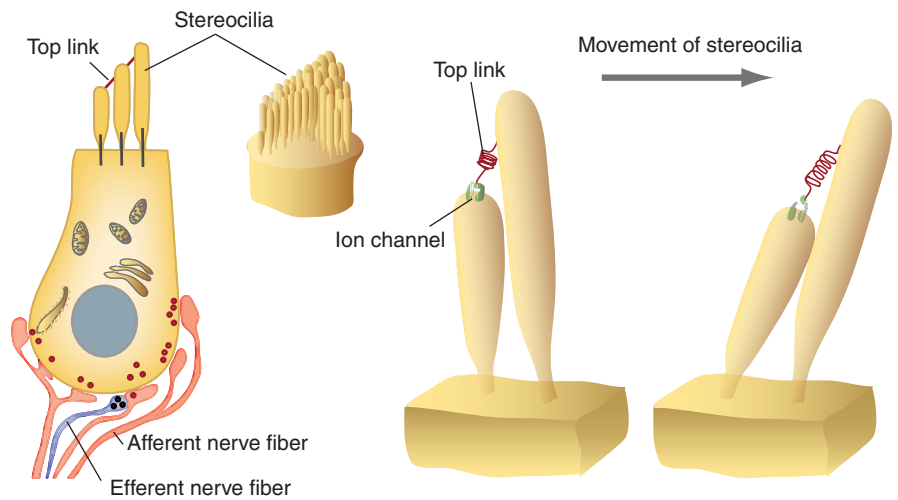


FIGURE 17.8 Inner hair cell. **Left:** Compare with Fig. 17.5. **Right:** The most likely mechanism of sensory transduction in hair cells. The tip link pulls open the ion channel when the stereocilia are deflected. The tip link is thought to act like a spring that gates the opening of the channel. (Based on Pickles and Corey 1992.)

of each hair cell becomes much higher than if the stereocilia moved independently.

The stereocilia are of unequal length and orderly arranged from the shortest to the longest (Fig. 17.8).³ This structural polarization corresponds to a functional one: bending of the stereocilia toward the longest cilium **depolarizes** the hair cell, whereas the opposite movement **hyperpolarizes** it. The depolarization produces a graded **receptor potential**. Thus, bending the stereocilia one way increases the frequency of firing in the afferent nerve fibers. Depolarization of the hair cell is most likely caused by opening of **mechanosensitive ion channels** near the tip of the stereocilia.

The mechanism for channel opening appears to be as follows. The stereocilia are interconnected near their tips by a thin protein thread called a **tip link** (Fig. 17.8). The tip links attach to the channel proteins and opens the channels by a direct pull when the stereocilia are deflected. The depolarization is presumably mediated by K^+ ions entering the hair cell, since the endolymph that surrounds the stereocilia contains a very high concentration of K^+ and low Na^+ . In addition, the endocochlear potential speeds the flow of cations because it adds to the membrane potential of the stereocilia. The required movement of the stereocilia is minimal: a barely perceptible sound moves the tip of each stereocilium about 1 nm (1/100,000,000 mm), whereas the strongest perceptible sound moves the stereocilia about 1 μ m—that is, corresponding to the thickness of the stereocilium.

Depolarization leads to release of **glutamate** from the basal part of the inner hair cells. This transmits the signal to the sensory nerve endings (acting on amino-methylisoxazole propionic acid [AMPA] receptors).

Release of sufficient amounts of the transmitter elicits action potentials in the sensory nerve fibers, and the signals are transmitted to the brain stem (see Fig. 12.1). When the stereocilia move back and forth, the ensuing receptor potential of the hair cell follows a sinus curve; that is, the membrane potential oscillates in pace with the vibrations of the basilar membrane. Because the receptor potential modulates the transmitter release, the action potentials conducted in the sensory fibers follow the frequency of the sound waves.

The Outer Hair Cells Amplify Sound Waves

The motion of the basilar membrane differs from that expected of a passive mechanical structure. There must be mechanisms intrinsic to the cochlea that can **amplify the vibrations** of the basilar membrane almost 100 times. Much evidence points to the outer hair cells as the **motors** of the amplification; that is, they can rapidly transmit mechanical energy to a very narrow strip of the basilar membrane.

Because the stereocilia of the outer hair cells are attached to tectorial membrane, their deflection by basilar membrane vibration seems easy to explain.⁴ When the basilar membrane vibrates, the hair cells are displaced relative to the tectorial membrane, which is relatively immobile because it is anchored to the bony wall. Thus, the stereocilia are moved back and forth, in pace with the frequency of the sound waves. The receptor potential initiates **contractile** activity, changing the form of the cell.⁵ The ensuing fast contractions of the

³ Vestibular hair cells have an extra tall kinocilium (non-motile) marking the highest point in the row of stereocilia (see Fig. 18.6). This disappears from the cochlear hair cells shortly after birth, however.

⁴ Although only the tallest stereocilia attach to the tectorial membrane, the linking of the stereocilia to each other make them move together as a unity.

⁵ The voltage-sensitive contractile protein **prestin** expands or compresses the cell body in response to changes of the membrane potential. The effects of manipulating the *prestin* gene (e.g., studying the effects on hearing of making the gene nonfunctional in knockout mice) suggest that this protein is of critical importance for the amplification mechanism.

outer hair cells move the basilar membrane, producing both an **amplification** of the vibrations and a narrowing of the vibrating part, which sharpens the **tuning** of the **frequency curve**. Together, these changes ensure more precise signal transmission. Whether the amplification is due solely to deformation of the outer hair cell body (elongation and shortening) or whether movements of the cilia also contribute, is not settled.

The **efferent fibers** ending on the outer hair cells release **acetylcholine** (and most likely ATP) that hyperpolarizes the cells by binding to muscarinic receptors. This reduces the active movements of the stereocilia leading to less amplification and broader tuning of the frequency curve.

Otoacoustic Emissions

The ear actually emits faint sounds—**otoacoustic emissions**—that appear to be produced by the contractions of the outer hair cells. A microphone in the external meatus can record such sounds, which can occur spontaneously or be evoked by a click. The hearing system works in reverse, as it were: the outer hair cells move the basilar membrane, which in turn moves the fluid in the cochlea; this moves the ossicles, which then move the tympanic membrane that produces the sound. The phenomenon is used diagnostically—for example, in infants—to decide whether the cochlea functions normally.

Different Frequencies are Registered at Different Sites along the Basilar Membrane: Tonotopic Localization

The ordered arrangement of neurons and nerve fibers signaling different pitches of sound (frequencies) is called **tonotopic localization** (compare with somatotopic localization in the somatosensory system and retinotopic localization in the visual system). As discussed later, the auditory pathways are tonotopically organized all the way from the cochlea to the cerebral cortex. The tonotopic localization in the cochlea has been demonstrated in several ways. After receiving lesions restricted to a small part of the organ of Corti (which extends along the full length of the basilar membrane), experimental animals no longer react to sound in a certain, narrow range of frequencies (pitches), whereas they react normally to sounds of other frequencies. In humans, similar selective deafness may occur after prolonged exposure to noise, for example, in factories. Anatomic examination of the cochlea after death in such persons has shown that the hair cells have disappeared in a restricted region on the basilar membrane, the position of the region differing with the frequency to which the person was deaf. The tonotopic localization has been determined in detail by recording the response of single hair cells to sounds of different frequencies.

Each hair cell is best activated by tones within a very narrow range of frequencies. Together, the hair cells and the sensory fibers leading from them cover the total range of frequencies we can perceive.

The **tonotopic** localization is such that the tones with the **highest pitch** (highest frequencies) are registered by the hair cells closest to the oval window (i.e., on the basal part of the basilar membrane), whereas the lowest frequencies are registered at the top of the cochlea (i.e., at the apical part of the basilar membrane). This can be explained, at least partly, by the physical properties of the basilar membrane, as proposed by the German physicist Hermann Helmholtz in the nineteenth century. The basilar membrane is most narrow basally and becomes progressively wider in the apical direction. The fibers of the basilar membrane are oriented transversely to the long axis of the basilar membrane (Fig. 17.6) and are therefore longer apically than basally. In analogy with the strings of an instrument (e.g., a piano), basal parts would be expected to vibrate with a higher frequency than apical parts. This is the main basis of the **resonance theory** of Helmholtz, which postulates that each position along the basilar membrane corresponds to a certain frequency. Although later research has shown that even a pure tone makes large parts of the basilar membrane vibrate, the region in which the maximal amplitude occurs is very narrow. This appears to be caused by the amplification produced by contractions of the outer hair cells, as discussed in the preceding text. Thus, the hair cells differ in accordance with their position on the basilar membrane, so that their thresholds are lower for certain frequencies than for others.

Cochlear and Brain Stem Implants to Restore Hearing

When deafness is due to loss of hair cells without affection of the afferent nerve fibers, hearing can be restored by a so-called **cochlear implant**. During surgery, a thin, isolated electrode is inserted along the basilar membrane. The isolation is removed at around 20 points, enabling electric stimulation of the cochlear nerve fibers at these sites. Hence, a range of frequencies can be transmitted to the brain; in accordance with the tonotopic localization along the basilar membrane (the frequency resolution is of course inferior to what is possible with a normal number of hair cells). A microphone and microprocessor transform the sound waves to electric signals that are sent to the electrode.

In **children** who are **born deaf**, a cochlear implant can enable development of speech and speech comprehension. Most of the children are able to attend normal school classes. However, the implant must be inserted as early as possible, and not later than 3 to 4 years of age. If the cortical networks necessary for analysis and use of sounds are not developed before this age, they

cannot develop later, probably because the auditory cortex has been taken over by other systems. Indeed, there is evidence from functional magnetic resonance imaging (fMRI) studies that visual stimuli may activate the auditory cortex in congenitally blind persons. This corresponds to the situation in children who are born blind and later regain sight (e.g., by removing a cataract), as discussed in Chapter 9. In **deaf adults**, who have had normal hearing earlier, a cochlear implant can restore hearing so they can comprehend speech. In this situation, the networks responsible for sound processing were presumably established in early childhood and remained essentially intact even after a period without hearing. However, intense training is required after surgery, in order to make sense of the foreign sounds provided by the implant. Further, the longer the time from loss of hearing to implantation, the poorer is the prospect of a satisfactory result.

When deafness is due to destruction of the **cochlear nerve**, a cochlear implant will of course have no effect. In some such patients, restoration of hearing has been attempted by implanting an array of electrodes over the ventral **cochlear nucleus**. Otherwise, the strategy is similar to that used in cochlear implants. The success so far has been much more limited than with cochlear implants, however.

THE AUDITORY PATHWAYS

Distinctive Features of the Central Auditory System

The anatomic organization of the central auditory pathways has some unusual features, different from other sensory systems we have dealt with. More nuclei are intercalated in the auditory pathways, and these nuclei have extensive and complicated interconnections. In addition, some fibers cross the midline at several levels of the auditory pathways. These features have made the auditory pathways more difficult to study than other sensory pathways. The crossing at several levels also renders hearing examinations of limited practical value for determining the site of lesions in the CNS.

Conduction Deafness and Sensorineural Hearing Loss

The most common causes of hearing loss are diseases of the middle ear, either because they compromise the conduction of sounds to the cochlea or because they destroy the neural elements of the cochlea or the eighth nerve. The first kind is called **conductive deafness**, the second **sensorineural (or nerve) deafness**. Transmission of sound to the cochlea can be reduced or abolished by middle ear infections that damage the eardrum or the ossicles. In **otosclerosis**, the basal plate of the stapes becomes fixed in the oval window and therefore cannot transmit sounds. Less frequently, lesions of the central

auditory pathways cause hearing loss; this kind, called **central deafness**, is considered later in this chapter.

To **distinguish** between conductive and sensorineural deafness, one can compare the threshold for sounds conducted through the air with sound conducted through the bones of the skull. In the **Rinne test**, a vibrating tuning fork is first applied to the mastoid process and then a little away from the external ear. Normally, the sound is heard much better when it is conducted through the air than through the bone; however, in conductive deafness, caused by destruction of the eardrum or the ossicles, the bone-conducted sound is heard best. In the **Weber test**, the vibrating tuning fork is applied to the forehead in the midline. Normally, the sound is heard equally in both ears; in conductive deafness, it is heard best in the deaf ear, whereas in sensorineural deafness it is heard best in the normal ear. The reason for the lateralization to the deaf ear in conduction deafness is not clear. It is possible that the airborne sound masks the bone-conducted sound on the normal side.

Destruction of the cochlear nerve or the cochlear hair cells, that is, **peripheral lesions**, produces hearing loss of the ear on the same side (the same happens, of course, with a lesion of the cochlear nuclei). The **hair cells** may be damaged by noise and by certain drugs. There is also a steady loss of hair cells with **aging**, particularly near the base of the cochlear duct (high frequencies). Destruction of the **cochlear nerve** may be caused by a tumor in the internal auditory meatus (Fig. 17.1, called **acoustic neuroma** (arising from the Schwann cells of the eighth cranial nerve)). As the tumor grows, it compresses the cochlear, the vestibular, and the facial nerves (all passing through the internal meatus). Symptoms may therefore be caused by irritation (in the early phase) or destruction of all of these nerves. Thus, the first symptoms may be due to irritation of the cochlear nerve, causing ringing in the ear (tinnitus) and sometimes vertigo due to irritation of the vestibular nerve, but gradually deafness develops. As the tumor grows, it compresses the brain stem with additional symptoms from the trigeminal nerve and ascending and descending long tracts. The cochlear nerve may also be compressed or torn by **skull fractures** passing through the temporal bone. Peripheral lesions of the auditory system, in addition to causing unilateral deafness, also reduce or eliminate the ability to **localize** the source of a sound.

The Cochlear Nerve and the Cochlear Nuclei

The part of the eighth cranial nerve conducting signals from the cochlea is called the **cochlear nerve**. Most of the fibers are afferent and have their cell bodies in the **spiral ganglion**, which is located in the bony spiral lamina (Figs. 17.4–17.6). From the spiral ganglion the fibers pass through the midportion of the cochlea (the modiolus,

Fig. 17.4) and through the **internal acoustic meatus** (internal auditory canal) to the **cerebellopontine angle** (Fig. 17.9). There the nerve enters the two **cochlear nuclei**—the **dorsal** and the **ventral**—located laterally on the medulla (Figs. 17.9 and 17.10; see also Fig. 27.2), external to the inferior cerebellar peduncle (see Fig. 24.1). After entering the nuclei, the cochlear nerve fibers divide and end in precise **tonotopic** order in several parts of the nuclei. The tonotopic localization has been demonstrated, for example, with microelectrode recordings that make it possible to determine the response of single fibers and neurons to tones of different frequencies.

The **efferent fibers** of the cochlear nerve arise in the **superior olivary complex**, located in the rostral part of the medulla (Fig. 17.10). The fibers end, as mentioned, predominantly around the **outer hair cells**. Stimulation of the efferent fibers reduces the signal frequency of the afferent fibers from the inner hair cells. Thus, the CNS can influence the flow of auditory signals at the level of the receptors, not just centrally.

Ascending Pathways from the Cochlear Nuclei

From the cochlear nuclei, the auditory signals are transmitted to the **inferior colliculus** (mainly of the opposite side). The pathway formed by the ascending fibers from the cochlear nuclei is called the **lateral lemniscus** (Figs. 17.9 and 17.10). Many of the fibers end in other nuclei—such as the superior olivary complex—that send fibers to the inferior colliculus. Neurons in the

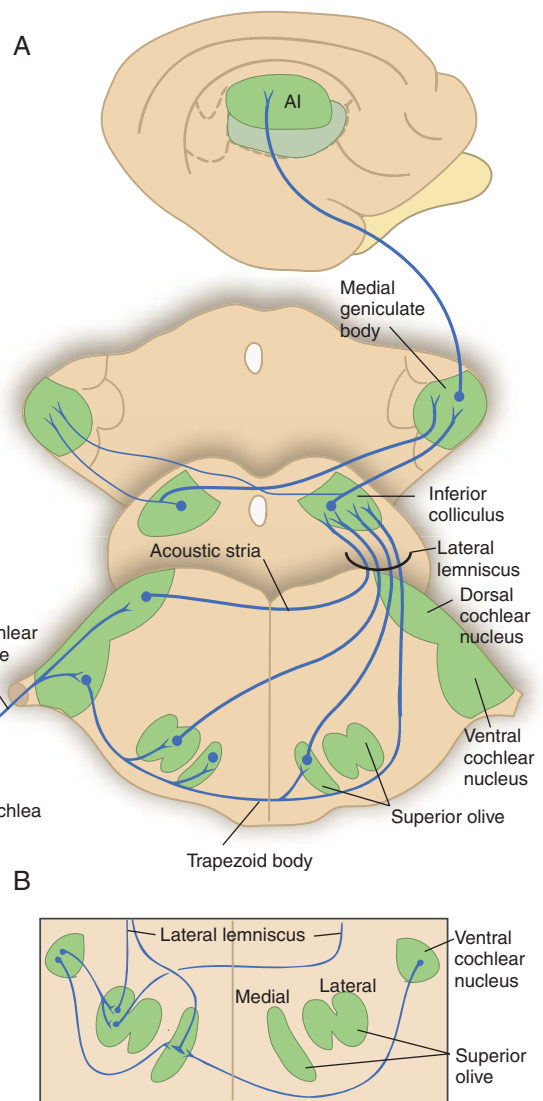


FIGURE 17.10 *Parallel ascending auditory pathways.* Schematic based on experimental studies of the cat. **A:** The connections are bilateral, so that AI on one side receives signals from the cochleae of both sides. **B:** Some main connections of the superior olive. Note that signals from both ears reach the medial part of the superior olive. This arrangement is of importance for localization of sound.

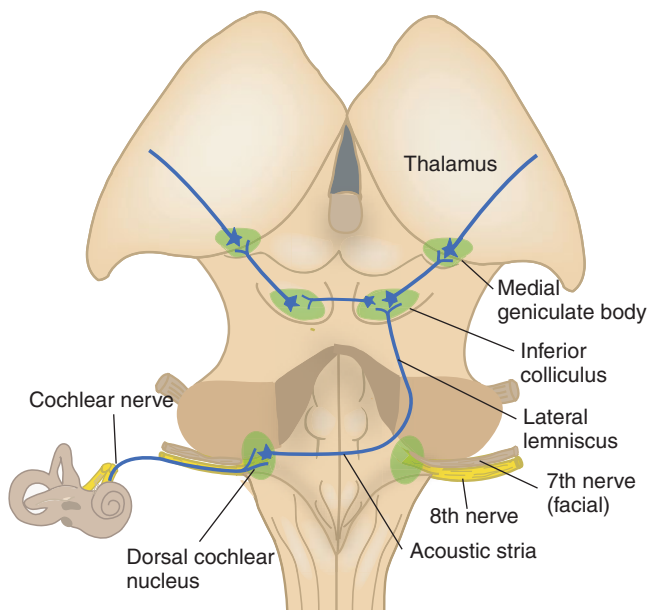
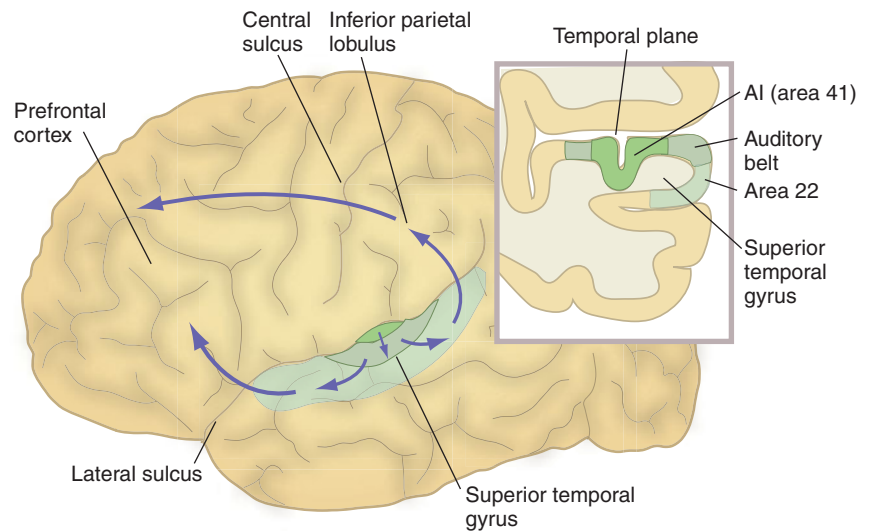


FIGURE 17.9 *Nuclei of the auditory pathways.* Signals from the cochlea of one side reach the medial geniculate bodies of both sides (although not shown in the figure).

inferior colliculus send their axons to the **medial geniculate body** of the thalamus (Figs. 17.9 and 17.10; see also Figs. 6.21 and 6.22). The fibers form an oblong elevation at the dorsal side of the mesencephalon, the **inferior collicular brachium** (brachium quadrigeminum inferius). The efferent fibers from the medial geniculate body end in the **auditory cortex** located in the temporal lobe (in the superior temporal gyrus; Figs. 17.10 and 17.11). The ascending fibers from the medial geniculate body are located in the posterior part of the **internal capsule**. At all levels, the auditory pathway is precisely organized, with cells responding to sounds of different frequencies arranged in parallel lamellae.

FIGURE 17.11 *The human auditory cortex.* The figure shows in a simplified form the concentric arrangement of the auditory areas, and how the connections spread out from AI, finally reaching the prefrontal cortex (there are reciprocal connections at all levels, although not shown here). Dorsal and ventral pathways from auditory areas to the frontal lobe are indicated. The ventral pathway is thought to deal with identification of sounds and their meaning, whereas the dorsal pathway primarily deals with aspects of sound localization and movements. The *inset* shows a frontal section through the temporal lobe, and the distribution of the auditory areas.



Even though the central auditory pathways are predominantly crossed, there is a significant uncrossed component. Therefore, unilateral damage to the pathways does not produce a clear-cut hearing deficit. The ability to **localize** from where a sound comes may be reduced, however.

The Auditory Pathways Consist of Functionally Different Components

Efferent fibers from the cochlear nuclei take different routes (Fig. 17.10A). Fibers from the **dorsal cochlear nucleus** pass in the **acoustic stria** dorsal to the inferior cerebellar peduncle and then cross through the reticular formation and join the **lateral lemniscus** (Figs. 17.9 and 17.10). Most fibers from the **ventral cochlear nucleus** pass ventrally and cross to the other side in the **trapezoid body** in the lowermost part of the pons (Fig. 17.10A). Some of these fibers end in the **superior olivary complex** of both sides, whereas others continue rostrally in the lateral lemniscus to the **inferior colliculus**. The functional significance of these parallel paths out of the cochlear nuclei is not fully understood, but animal experiments (especially in the cat) show that single cells in the dorsal and ventral nuclei have different properties. Schematically, many cells in the **ventral nucleus** respond to sound stimuli much like the primary afferent fibers of the cochlear nerve, whereas cells in the **dorsal nucleus** have more complex response properties. For example, cells in the dorsal nucleus are often excited by sound with one particular frequency and inhibited by another frequency. It has been suggested that the dorsal nucleus forwards signals that are important for **directing attention** toward a sound, whereas information from the ventral nucleus is important for, among other things, **localizing** a sound.

Experimentally, two components of the ascending auditory pathways have been identified. One is called the **core projection** and is a pathway for auditory signals only. It is precisely tonotopically localized at all levels and terminates in the **primary auditory cortex, AI**. The core projection is synaptically interrupted in the central parts of the inferior colliculus and specific parts of the medial geniculate body. The other component is called the **belt projection**. It is synaptically interrupted in the peripheral parts of the inferior colliculus and terminates in the **cortex surrounding the AI**. The cells of this pathway are influenced by visual and somatosensory stimuli in addition to auditory ones. The belt projection is thought to be important for **integrating** of auditory information with other kinds of sensory information.

Generalizations to conditions in humans on the basis of studies done in other species must be made with particular caution, however. Thus, species differences appear to be greater for the auditory system than for other sensory systems. This may presumably be related to the great differences that exist among species with regard to how sound is used as a source of information. For example, certain nuclei that are large in the cat are very small in humans, and vice versa.

Sound Localization

It is not enough that the CNS analyzes the sound frequencies and on this basis interprets the meaning of the sound (e.g., who or what produced the sound). Locating the source of a sound is another important but difficult task of the auditory system. To be able to judge the distance and direction to a sound source (potential prey or a danger) is obviously of vital importance to most animal species. The precision of our ability to localize sound is formidable: humans can distinguish sounds

separated by no more than 2 degrees. To do this, central parts of the auditory system must be able to detect temporal differences of 11 μ sec (microseconds) between sounds reaching the two ears.

Unilateral damage to the **auditory cortex** reduces the ability of experimental animals to locate sounds coming from the opposite side. Thus, a cat with such a lesion does not move toward its prey sending out a brief sound (e.g., a mouse). The head and eyes, however, move toward the prey—even after bilateral damage of the auditory cortex—showing that nuclei at lower levels of the auditory pathways can locate the sound and elicit appropriate reflex movements.

Concerning location of sounds in the **horizontal plane**—that is, to the right or left of the midsagittal plane—basic computations occur in the first synaptic relay after the cochlear nuclei, the **superior olive** (Fig. 17.10). We are less certain regarding mechanisms for location of sounds in the vertical plane, although the **dorsal cochlear nucleus** is probably involved. Neurons in the **inferior colliculus** respond specifically to sound from a certain direction, and the inferior colliculus probably contains a **map** of our **auditory space**.

The Superior Olive and Sound Localization

The **superior olivary complex** (superior olive) is located in the lower part of the pons, in the trapezoid body (Fig. 17.10). A striking feature is that most neurons are influenced from both ears, which led to the assumption that the superior olive is particularly important in localizing the origin of a sound. When a sound hits the head from the right side, it will reach the right ear slightly before it reaches the left, because the head is in the way. The sound will also be slightly weakened before reaching the left ear. Psychophysical experiments in humans indicate that side differences in both time and intensity are used by the auditory system to localize sounds. The time difference is most important for localizing sounds of low frequencies, whereas intensity differences are most important for sounds of higher frequencies (above 4000 Hz).

The superior olivary complex consists of several, tonotopically organized subdivisions. The lateral part receives afferents from the cochlear nuclei of both sides and projects bilaterally to the inferior colliculi. Most cells in the lateral part are excited by signals from the ear of the same side and inhibited by signals from the contralateral ear (via interneurons). The cells respond best when the sounds hitting the two ears are of different intensities. Consequently, the **lateral part** of the superior olive is assumed to use intensity differences for the analysis of sound localization.

The **medial part** of the superior olive appears to be particularly important in localizing **low-frequency sounds**. It receives afferents from a particular subdivision of the ventral cochlear nucleus of both sides (Fig. 17.10B). Each

cell has two long dendrites oriented transversely. One dendrite receives signals from the right ear, the other from the left. These cells are very sensitive to small time differences in synaptic inputs to the two dendrites and are most sensitive to sounds with low frequencies. The efferent fibers of the medial part pass to the central nucleus of the inferior colliculus on the same side.

Descending Control of the Auditory Pathways

There are descending fibers at all levels of the auditory pathways. Numerous fibers pass from the auditory cortex to the medial geniculate body (like other thalamic nuclei) and to the inferior colliculus. Other fibers descend from the inferior colliculus to the nuclei at lower levels. As mentioned, efferent fibers in the cochlear nerve end in contact primarily with the outer hair cells of the cochlea; such fibers come from the **superior olivary complex** and form the **olivocochlear bundle**. The descending connections are, at least in part, precisely organized and can therefore be expected to selectively control subgroups of neurons in the auditory pathways (e.g., neurons transmitting information about a certain frequency).

There are many inhibitory **interneurons** in the nuclei of the auditory pathways, and both γ -aminobutyric acid (GABA) and **glycine** are used as transmitters for such interneurons. Physiological experiments show that the central transmission of auditory signals can be inhibited, probably at all levels from the cochlea to the cerebral cortex. The censoring of the sensory information that is allowed to reach consciousness is perhaps even more pronounced in the auditory system than in other sensory systems. Selective suppression of auditory information is necessary if we are to **select** the relevant sounds among numerous irrelevant ones. Such mechanisms are most likely at work when, for example, at a cocktail party with numerous voices we nevertheless are able to select and pay attention to only one of them.

The brain must be able to distinguish between sounds that are generated externally and sounds we produce ourselves. Indeed, auditory neurons are inhibited during **vocalization**. Although the exact mechanisms are unknown, this most likely involves interpretation of **corollary discharge**—that is, copies of the motor commands producing the sounds are sent to (most likely) the auditory cortex. Descending connections may ensure specific inhibition at several levels of the auditory pathways.

Auditory Reflexes

The ascending auditory pathways convey signals that enable the conscious perception of sounds. Auditory information is also used at a subconscious level to elicit reflex responses. The **reticular formation** receives collaterals from the ascending auditory pathways, and such

connections mediate the sudden muscle activity provoked by a strong, unexpected sound—that is, a **startle response**. Other auditory signals pass to the nuclei of the facial and trigeminal nerves, which innervate two small muscles in the middle ear: the **stapedius** and **tensor tympani muscles**. Contraction of these muscles dampens the movements of the middle ear ossicles and thereby protects the cochlea against sounds of high intensity. Paresis of the **facial nerve** often is accompanied by hypersensitivity to sounds, or **hyperacusis**.

Other, more complex, reflex arcs mediate automatic **movements of the head and eyes**, and even the body, in the direction of an unexpected sound. The centers for such reflexes are probably located in the **inferior** and **superior colliculi**. The inferior colliculus sends fibers to the superior colliculus, which has connections with the relevant motor nuclei in the brain stem and spinal cord. Further, in the superior colliculus **integration** of auditory, visual, and somatosensory information takes place, so that the final motor response is appropriate for the organism as a whole.

THE AUDITORY CORTEX

The cortical auditory areas are less studied than the visual areas, and important aspects of human hearing are still poorly understood. While the auditory cortical areas are organized according to the same general principles as the other sensory areas, notable differences exist. Thus, the auditory system shows a more pronounced parallel organization than, for example, the visual system. This is particularly evident in the auditory pathways and intercalated nuclei but concerns also the cortical areas. Further, more information processing takes place at lower levels in the auditory than in other sensory systems. While the auditory cortex is not crucial for the ability to identify single sounds, it is required when different sounds are to be put together to a meaningful whole.

Core and Belt Areas

The **primary auditory area (AI, core region)** appears to consist of several tonotopically organized subdivisions (in contrast to the striate area that consists of one retinotopically organized representation of visual field).⁶

⁶ In monkeys, three areas in the superior temporal gyrus receive tonotopically organized projections from the ventral part of the medial geniculate body. One is the 'classical' AI area, the two others adjoin AI rostrally (termed R and RT). The reason to include them in the term primary auditory cortex is that all three show structural features typical of primary sensory areas, such as a well-developed lamina 4 consisting of densely packed, small neurons (see Fig. 33.4), and dense afferent projections from a specific thalamic nucleus. The tonotopic arrangement in AI is such that the highest frequencies are represented caudally and the lowest rostrally, whereas the opposite arrangement appears to exist in area R.

AI is situated on the upper face of the temporal lobe in a region called the **temporal plane** (Fig. 17.11). In humans, Brodmann's **area 41** is thought to correspond to area AI of monkeys and other animals. Several other auditory areas are arranged concentrically around AI (in monkey about 15 such areas have been identified; cf. organization of extrastriate visual areas). The areas closest to AI form a belt-like region, and have therefore been termed the **auditory belt**. The belt region receives thalamocortical afferents from other parts of the medial geniculate body than AI, while its main afferents appear to come from AI. Outside the auditory belt, we find additional auditory areas that receive processed auditory information from the auditory belt, but no afferents from the medial geniculate body. Here starts integration of auditory and other sensory modalities. These areas are probably located within area 22 of Brodmann (Fig. 17.11; see also Fig. 33.3). There appears to be several **parallel channels** from the core region to the belt region, and further on to area 22 (similar to the organization of parallel pathways out of the striate area)—each conveying different aspects of auditory information. There are, however, numerous interconnections among areas suggesting that the functional segregation cannot be absolute.

Properties of Neurons in Primary Auditory Cortex

Fibers from the **medial geniculate body** end with precise tonotopic localization in the AI (Figs. 17.10 and 17.11). Accordingly, single neurons in AI respond to sounds with a narrower frequency range than neurons in other auditory areas (sharper tuning). Although many neurons in AI depict **simple**, physical features of sounds (such as pitch and amplitude), others have surprisingly **complex** properties. Some respond, for example, best to a sound when the frequency is increasing or decreasing. Other cells are influenced from both ears, but often such that they are excited by signals from one ear and inhibited from the other. Further, the response of many neurons (as in other primary visual areas) depends on the **context** of a stimulus. It even seems that the **object** with which a sound is associated modulates the activity of many AI neurons.

Asymmetrical Organization of the Auditory Cortex in the Temporal Plane

The auditory cortical areas appear to be asymmetrical in most humans. Thus, according to several MRI and postmortem studies the so-called **Heschl's gyrus**—containing the AI—is larger on the left than on the right side, and the same holds for regions (**area 22**) adjacent to AI in the temporal plane (some studies did not find such asymmetries, however). Intrinsic (horizontal) connections in area 22 show a higher number of separate

clusters of terminal fibers on the left than on the right side, which presumably enables a more fine-grained analysis. It has furthermore been reported that fibers in the white matter are more heavily myelinated in the left than in the right temporal plane, enabling faster processing. Such anatomic differences are compatible with evidence that the left auditory cortex has a higher degree of temporal sensitivity needed for optimal speech discrimination. The right auditory cortex appears to be better than the left in the discrimination of pitch, melody, and sound intensity.

Further Treatment Outside the Primary Auditory Cortex

As mentioned, the **auditory belt** receives its main afferents from AI. The tonotopic localization is less sharp than in AI. Many neurons appear to respond best to **species-specific sounds** (sounds of speech in humans). Other neurons seem to code the **localization** of a sound source, while others combine information about speech sounds and their source (wherefrom and from what). With regard to perception of voices and music, the analysis of **temporal patterns** is of particular importance.

The areas outside AI show **functional specializations**. Thus, at least partly different subregions of the auditory belt deal with “**what**”—that is, the frequency composition of the sounds—and “**where**”—that is, the localization in space of the sound source. fMRI studies show, for example, that especially caudal parts of the auditory cortex are activated by moving sounds, while speech-relevant sounds primarily activate rostral parts.⁷ Further,

⁷ Clinical observations in humans also support specializations among the auditory areas outside AI. For example, in some patients after bilateral, partial lesions of the superior temporal gyrus, speech comprehension may be preserved while the perception of speech melody (**prosody**) and music may be impaired.

there is evidence from both experimental animals and humans that the further projections to the **frontal lobe** are divided into a **ventral** pathway for “what” and a **dorsal** pathway for “where” (Fig. 17.11). This seems to correspond to the organization of visual corticocortical connections into dorsal and ventral pathways or streams (see Fig. 16.24). Projections to the prefrontal cortex enable transformation of auditory information into actions. How far the subdivision of tasks between a ventral and a dorsal pathway goes is nevertheless not clear. Thus, many neurons in posterior parts of the auditory cortex, supposed to be specialized for spatial tasks, respond to various aspects of speech.

Damage to Central Parts of the Auditory System and Acoustic Agnosia

Restricted lesions of the auditory pathways or the auditory cortex—**central lesions**—usually produce no clear-cut symptoms. As mentioned, this is because the connections from the cochlear nuclei to the cerebral cortex are bilateral (although with a contralateral preponderance). Patients with bilateral damage of the auditory cortex are reported to be able to perceive sounds and even discriminate tones with different pitches and intensities (even though not necessarily with normal speed and precision). This corresponds to findings made in monkeys. The ability to recognize and interpret tones in particular patterns, however, is reduced or abolished. Such patients are unable to recognize familiar sounds such as laughter, a bell that tolls, sounds of various animals, and so forth. They are furthermore unable to understand the speech of other people, even though they can speak and read themselves. This is called **acoustic agnosia**.

18 | The Sense of Equilibrium

OVERVIEW

The sense of equilibrium, narrowly defined, depends on signals from sense organs that record the position and movements of the head. Such receptors are found in the **vestibular apparatus** in the inner ear. Together with the cochlea they form the **labyrinth**. As the cochlear duct, the vestibular labyrinth is filled with **endolymph**. The vestibular labyrinth consists of three **semicircular ducts** (canals) and two small expansions, the **sacculus** and the **utricle**. **Cristae** equipped with hair cells are found in expansions of the ducts, and respond to movements of the endolymph. Movements of the endolymph arise when the head rotates, and the hair cells thus signal **angular acceleration**. The semicircular ducts are oriented in three, mutually perpendicular planes. Hair cells in the sacculus and utricle are found on flat surfaces called saccular and utricular **maculae**. The stereocilia of the hair cells are embedded in a gelatinous substance containing **otoliths**. These tiny stones ensure that the hair cells respond to gravitational forces and thus record the position of the head in space.

Afferents from the vestibular apparatus end in the **vestibular nuclei** in the upper medulla and lower pons. From there, signals flow in three main directions: to the **spinal cord**, to neuronal groups controlling **eye movements**, and to the **cerebellum**. These connections mainly control automatic responses aimed at maintaining the upright position (**postural reflexes**) and fixation of visual targets (**vestibulo-ocular reflexes**). In addition, vestibular signals reach several small areas in the **cerebral cortex**. Thereby, signals from vestibular receptors contribute to our conscious **awareness** of the position of the body in space.

The vestibular apparatus is not the only source of sensory information for maintaining equilibrium, however, and neither does it provide all necessary information. Visual information and signals from somatosensory receptors (proprioceptors and cutaneous receptors) also contribute. To control the upright, bipedal position of the human body, the brain needs information not only of the position of the head in space (provided by the vestibular receptors) but also the head's position in relation to the body and the mutual positions of the major body parts. Indeed, the experience of **dizziness** and disequilibrium can have many causes other than dysfunction of the vestibular apparatus. To analyze such problems one must take into account all factors that contribute to maintenance of our upright position.

Postural control and the role of vestibular signals in the control of eye movements are also discussed in Chapter 22 and Chapter 25, respectively.

STRUCTURE AND FUNCTION OF THE VESTIBULAR APPARATUS

The Vestibular Part of the Labyrinth

Unlike the cochlea, the vestibular apparatus does not depend on the external ear and the tympanic cavity to function. Like the cochlear duct with the organ of Corti (see Figs. 17.4 and 17.6), the **membranous labyrinth** containing the vestibular receptors is embedded in the temporal bone and surrounded by a space containing **perilymph**. The membranous labyrinth is filled with **endolymph** (Figs. 18.2 and 18.3).

The vestibular part of the labyrinth consists of two small vesicles, the **utricle** and the **sacculus**, and three circular tubes connected to the utricle, the **semicircular ducts** (see Fig. 17.2). Each of the semicircular ducts has a swelling, the **ampulla**, in the end close to the utriculus. The ducts are oriented in three planes perpendicular to each other. In the erect position and with the head in a neutral position, the lateral semicircular duct lies approximately in the horizontal plane, whereas the posterior and the anterior ducts are oriented vertically (Fig. 18.1).

Sensory epithelium with **hair cells** (Figs. 18.2 and 18.3) is found at five locations in the vestibular labyrinth. The hair cells are similar to those in the cochlea, except that they are equipped with a long kinocilium not present on cochlear hair cells (compare Figs. 18.6 and 17.8). In each of the ampullae of the semicircular ducts, there is a transversely oriented elevation, the **ampullar crista** (Fig. 18.3). Hair cells with long cilia (Fig. 18.4) are present between the epithelial cells that form the crista. The cilia project into a jellylike mass, the **cupula**. In the utricle and the sacculus there are small patches of hair cells like those in the ampullae, the utricular and saccular **maculae** (Figs. 18.2 and 9.5). The macular hair cells are also covered by a jellylike mass, which is special because of its content of small "stones," the **otoliths** (Figs. 18.2, right, and 18.5). The otoliths are crystals of calcium salts. The utricular macula lies in approximately the same plane as the lateral semicircular duct (i.e., horizontally in the neutral position), whereas the saccular macula is oriented almost vertically (Fig. 18.1).

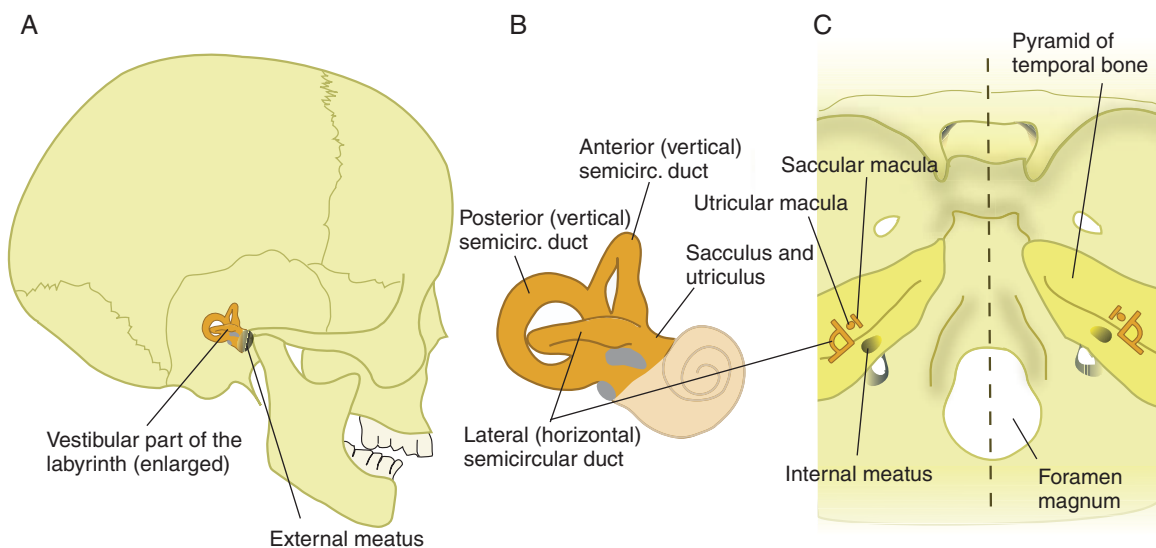


FIGURE 18.1 *The vestibular apparatus.* **A:** The position of the labyrinth in the temporal bone. **B:** The vestibular part of the labyrinth is colored more darkly than the auditory part. Note the orientation of the semicircular ducts in relation to the conventional planes of the

body. **C:** The base of the skull, as viewed from above, with the vestibular apparatus projected to the surface of the pyramid of the temporal bone. Note the orientation of the saccular and utricular maculae.

More about Vestibular Hair Cells

The vestibular and cochlear hair cells have the same basic properties (see Chapter 17, under “The Inner Hair Cells and Mechanoelectric Transduction”), although there are some structural differences. On the apical end of the cells there are 50 to 110 **stereocilia** and one longer and thicker **kinocilium** (Fig. 18.6). The stereocilia are unusually long microvilli and contain actin filaments like other microvilli. The kinocilium contains microtubuli (like cilia of the respiratory epithelial cells). The stereocilia are arranged regularly in accordance with their height (Figs. 18.2, 18.4, and 18.6; see also Fig. 17.8). This structural **polarization** of the receptor cells corresponds with the functional polarization mentioned above. Thus, bending of the stereocilia toward the kinocilium increases the firing frequency of the sensory fibers in contact with the cell, whereas bending in the opposite direction reduces the firing frequency. This is because the receptor cell is depolarized or hyperpolarized by bending of the cilia. The receptor potentials of the hair cells induce release of **glutamate**, which depolarizes the afferent nerve fibers. Deflection of the cilia perpendicular to the direction of the polarization produces no response, whereas oblique displacements give a reduced response compared with a stimulus that is properly aligned with the polarization. This means that a given **firing frequency** of an afferent fiber is **ambiguous**; it can be caused by a weak stimulus in the direction of the polarization or a stronger obliquely oriented one. Further, a given firing frequency may be caused by moving the head forward (e.g., walking) or tilting the head backwards. This may be why the hair

cells in the maculae are arranged differently with regard to the orientation of the polarization axes, so that all the cells together cover 360 degrees (Fig. 18.6B). This ensures that the information received by the brain about head position in space is unambiguous. This requires, of course, that the brain is able to compare the magnitude of the signals from various parts of the maculae to reach a conclusion.

Electron microscopic studies showed more than 50 years ago that there are two kinds of vestibular receptor cells, which are found both in the cristae of the semicircular ducts and in the maculae of the saccule and utricle (Fig. 18.6A). One kind, the **type 1** cell, is bottle-shaped, whereas the **type 2** cell is slender. The sensory fibers (the peripheral process of the vestibular ganglion cells) end differently on the two cell types (Fig. 18.6A). The functional significance of the two types of hair cells is still unknown.

The Adequate Stimulus for the Semicircular Ducts Is Rotation of the Head

Flow of the fluid (the endolymph) inside the semicircular ducts displaces the cupula, thereby bending the cilia (Fig. 18.3). This activates or inhibits the sensory cells, depending on the direction of bending. Rotational movements of the head produce flow of the endolymph in the semicircular ducts. This is explained by the inertia of the fluid: when the head starts to rotate, the fluid “lags behind,” and when the rotation stops, the fluid continues to flow for a moment (like the water in a bowl that is rotated rapidly for a couple of turns). If the head rotation continues at even velocity, the fluid and

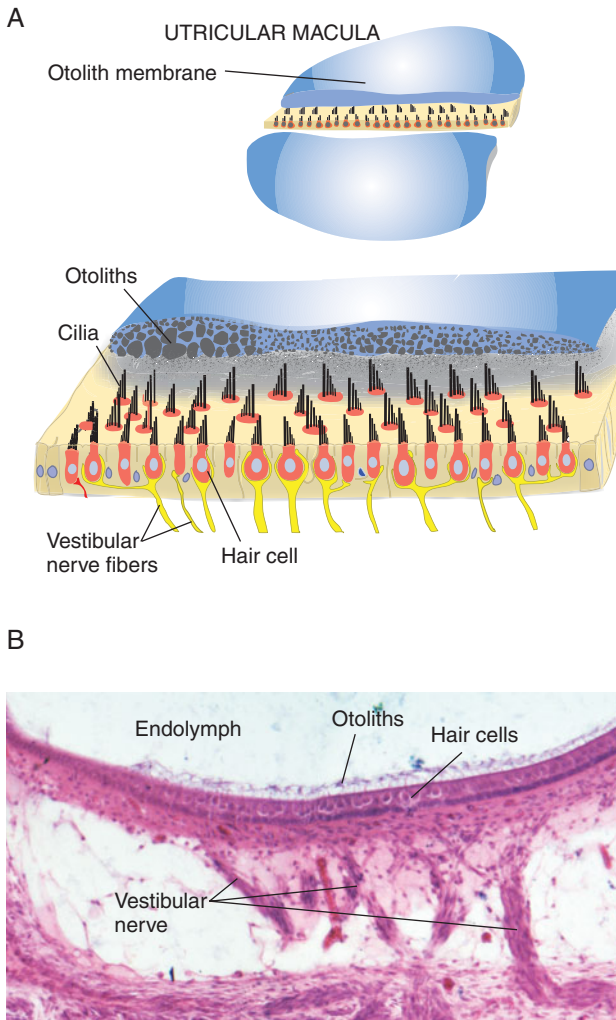


FIGURE 18.2 *The utricular and saccular maculae.* **A:** The utricular macula. Note the polarization of the hair cells. The otoliths are embedded in a gelatinous substance that covers the hair cells (otolith membrane). (Based on Lindeman 1973.) **B:** Photomicrograph of a section through the saccule with the macula.

the head will after a short time move with the same speed and in the same direction, which means that the cilia are not bent. Thus, it is clear that not every rotational movement is recorded by the receptors of the semicircular ducts: alteration of the velocity of the rotational movement—that is, positive or negative **angular acceleration**—is the adequate stimulus for the semicircular duct receptors. Thus, these receptors have **dynamic sensitivity**. Linear acceleration does not affect (or affects only slightly) the semicircular ducts. By linear displacement of the head (a translatory movement) without concomitant rotation, there is no stimulation of the semicircular duct receptors. An example of linear acceleration is a car that starts or brakes on a flat, straight road. If the road goes up and down and is curved, rotational accelerations of the car (and the heads of its passengers) are superimposed on the linear ones.

The orientation of the semicircular ducts ensures that rotation of the head in any conceivable plane produces change of activity of the receptors in one or more of the ducts. The brain monitors the rotational movements of the head by reading the pattern of activity produced by all the receptors. The semicircular ducts of both sides must function normally to supply the brain with the necessary information. A pair of ducts (e.g., the right and left lateral ones) gives **complementary signals** to a given rotation—that is, increased signal frequency from the one and reduced from the other.

Sacculus and Utriculus Signal the Position of the Head and Linear Acceleration

The small **otoliths** have higher specific weight than the endolymph and the substance in which they are embedded and are therefore more influenced by gravitational forces. Taking as an example the utricle, which in the neutral head position is oriented horizontally (Fig. 18.1), a change of head position tilts the macula, and the otoliths pull the cilia in that direction. Different angles of tilt produce different patterns of activity of the macular hair cells. Owing to their different orientations in space, the utricular and saccular hair cells together provide information about all possible head positions. The **utricle** records especially **lateral tilt** (i.e., head positions that vary around a sagittal axis), whereas the **sacculus** probably records mainly flexion–extension of the head (i.e., movement in the cervical joints around a transverse axis). Its ability to provide information about the position of the head at any one time (in the absence of movement) shows that the vestibular apparatus has **static sensitivity**. This property depends on the presence of receptors that adapt slowly or not at all, so that they give a constant signal as long as a certain position is maintained. The static sensitivity of the vestibular apparatus depends, as we have seen, on the force of gravity and disappears in a state of weightlessness (such as in space travel). Because the static sensitivity is a property of the utricle and the sacculus, these parts of the vestibular apparatus are called the **static labyrinth**. We will see, however, that this part of the labyrinth also has dynamic properties.

The **dynamic sensitivity** of the utricle and the sacculus is seen when their activity is recorded during **linear acceleration**. The response increases (i.e., the firing frequency of the afferent fibers) with increasing acceleration. This is again explained by the inertia of the otoliths. On linear displacement of the head with changing velocity, as in a car that is accelerating, the otoliths lag behind and thereby bend the cilia backward (in relation to the direction of the movement). The opposite happens when the car slows down; the otoliths continue to move for a moment and bend the cilia forward (compare with the forces acting on a loose object in a car when the car speeds up or slows down).

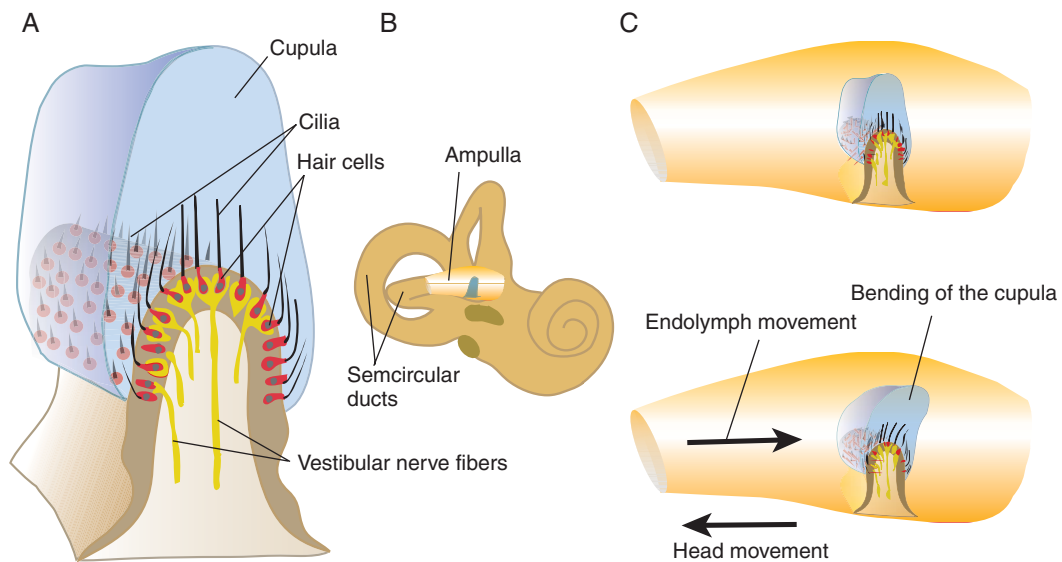


FIGURE 18.3 *The ampullar crista.* A: Three-dimensional drawing showing the relationship between the crista, the hair cells, and the cupula. (Based on Wersäll 1956.) B: The labyrinth with the position

of the crista in the horizontal semicircular duct. C: Movement of the endolymph deflects the cupula and the stereocilia of the hair cells. See text for further details.

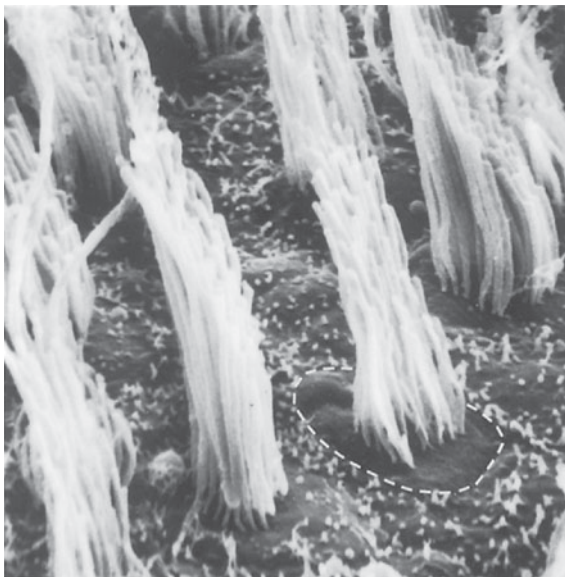


FIGURE 18.4 *Sensory (hair) cells of the utricle macula.* Scanning electron micrograph. The apical surface of one sensory cell is indicated. Magnification, $\times 10,000$. Compare with Fig. 9.2B. (Courtesy of Dr. H. Lindeman, Rikshospitalet National Hospital, Norway.)

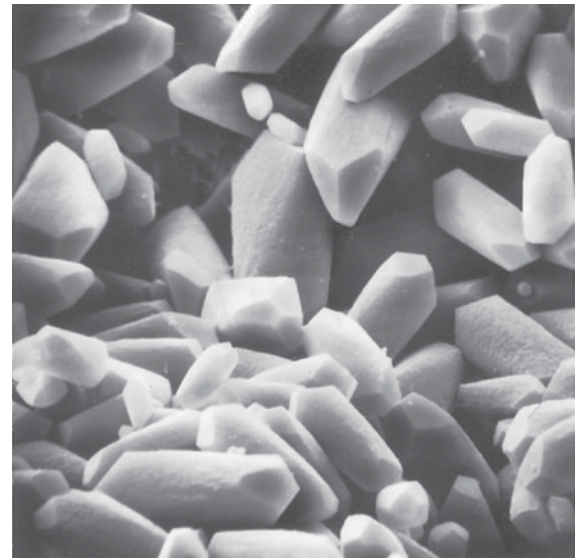


FIGURE 18.5 *Otoliths from the utricle macula.* Scanning electron micrograph. Magnification, $\times 2100$. (Courtesy of Dr. H. Lindeman, Rikshospitalet National Hospital, Norway.)

CONNECTIONS OF THE VESTIBULAR NUCLEI

The vestibular system differs from other systems in its high degree of **multisensory integration**. Thus, signals from the semicircular ducts and from the saccular and utricular maculae, that is, information about movements and head positions, converge in the vestibular nuclei and in the cerebellum. Further, integration of

visual, proprioceptive, and vestibular signals occurs at all levels—from the first synapses of the primary afferent fibers in the brain stem to the cerebral cortex. This is obviously necessary because the vestibular signals—less than other sensory signals—are behaviorally useful on their own. The position and movements of the head must continuously be related to the positions and movements of the eyes and of the body. Accordingly, vestibular

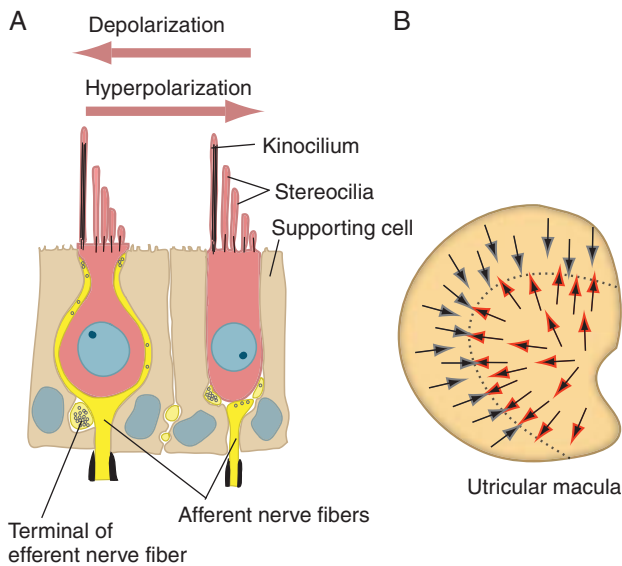


FIGURE 18.6 *Sensory cells of the vestibular apparatus.* **A:** Two types of hair cells with different shapes and relations to the afferent nerves. Note the polarization of the cilia. **B:** The utricle macula, as viewed from above. Arrows indicate the direction of polarization of the sensory cells in the various parts, which altogether cover all directions of deflection of the sensory hairs.

stimulation does not evoke a conscious “vestibular” sensation that can be easily distinguished from other sensory modalities.

The Vestibular Nuclei and Primary Afferent Fibers

The primary afferent fibers of the eight cranial nerve end in various parts of the **vestibular nuclei**.¹ The collection of large and small vestibular nuclei is collectively called the **vestibular complex**. It covers a large area in the floor of the fourth ventricle and consists of four large nuclei and several small cell groups. (Not all cell groups within the vestibular complex receive primary vestibular fibers, however, and are therefore not vestibular, strictly speaking.) Figure 18.7 shows the location of the

¹ There are also **efferent cholinergic fibers** in the vestibular nerve, ending in contact with the vestibular receptors (Fig. 18.6A). The efferent fibers come from a small cell group in the lower pons just lateral to the abducens nucleus. The projection is bilateral and ends diffusely. Rotational head movements evoke efferent activity, with mainly excitatory effects on the vestibular primary afferents. Although its functional role is still unknown, one possibility is that the efferent innervation increases regularity of afferent firing.

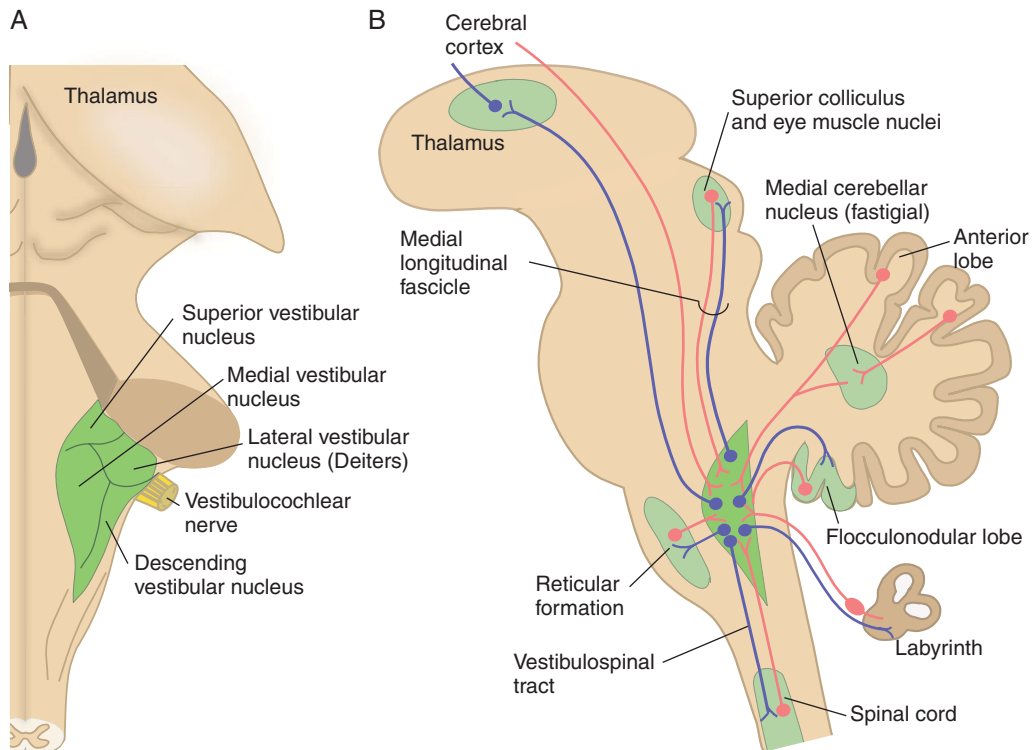


FIGURE 18.7 *The vestibular nuclei.* **A:** Location of the nuclei in a dorsal view of the brain stem. **B:** The main afferent and efferent connections. Note reciprocal connections with the spinal cord, the cerebellum, the reticular formation, and the nuclei of the extraocular

muscles and other visually related nuclei of the mesencephalon. Note also ascending connections to the thalamus (and from there to the cerebral cortex), and descending connections from the cerebral cortex to the vestibular nuclei.

four major nuclei: the **superior**, the **lateral** (or nucleus of Deiters), the **medial**, and the **descending** (or inferior).

Primary afferent vestibular fibers divide into an ascending and a descending branch when entering the brain stem. Together, they end in large parts of the vestibular complex and in parts of the cerebellum (see Fig. 24.4). Although terminations of fibers from the ampullar cristae and the maculae **overlap** in the vestibular nuclei, they also show notable differences in distribution. Thus, afferents from the **cristae** (that is, from the semicircular ducts) end in the **superior nucleus** and the rostral part of the **medial nucleus** but not in the lateral nucleus, whereas the fibers from the **utricle and saccular maculae** end in the **lateral nucleus** but not (or only sparsely) in the superior nucleus. In agreement with the distribution of primary afferents, neurons in the superior nucleus respond best to rotational head movements (angular acceleration), whereas the cells in the lateral nucleus are particularly sensitive to static head position. Nevertheless, physiological studies show that a substantial proportion of neurons in the vestibular nuclei integrate information about angular acceleration and static position/linear acceleration.²

The Vestibular Nuclei Receive Afferents from Regions Other than the Labyrinth

The physiological properties of neurons in the vestibular nuclei are not copies of those of the primary afferent fibers. This is due to convergence on the cells of various kinds of afferents (such as fibers from the semicircular ducts and the utricle) and by interconnections between the nuclei (e.g., commissural fibers linking the two sides). Further, the vestibular complex receives afferents from other parts of the central nervous system (CNS), especially the **spinal cord**, the **reticular formation**, certain mesencephalic nuclei, and the **cerebellum**. Afferents from the mesencephalon arise, for example, in the **superior colliculus**, and the cerebellar fibers come from both the flocculonodular lobe and the anterior lobe (Fig. 18.7; see also Fig. 24.4), and contribute to adaptation of vestibular reflexes to changed conditions for example during growth of the head, wearing of glasses, and so forth.

The vestibular nuclei receive signals from the **cerebral cortex** (mainly indirectly via the reticular formation but also some direct fibers). The corticovestibular fibers arise in parts of the cortex, such as the **SI** (areas 2 and 3a) and the **insula**, which receive converging information from the labyrinth and proprioceptors. The vestibular nuclei are also influenced from the **posterior parietal cortex**, which

is concerned with spatial orientation and goal-directed movements. Presumably, these cortical regions are important for building internal representations of the position and movements of the body, necessary for the control of movements. Accordingly, physiological experiments show that **vestibular reflexes**—the vestibulospinal ones in particular—are modulated in conjunction with voluntary movements. In this way, the reflex responses are subordinated to the overall plan for the movements, presumably by way of corticovestibular connections (among others).

Efferent Connections of the Vestibular Nuclei

Schematically, the vestibular nuclei (and therefore also the vestibular receptors) act on three main regions (Figs. 18.7 and 18.8), namely **motoneurons in the spinal cord**, **motoneurons in the nuclei of the extraocular muscles**, and the **cerebellum**. Accordingly, information from the vestibular apparatus is used primarily to influence muscles that maintain our **upright position** (equilibrium) and muscles that produce **eye movements**. The latter movements ensure that the retinal image is kept stationary when the head moves.

Most of the fibers to the **spinal cord** come from the **lateral vestibular nucleus** and form the **lateral vestibulospinal tract**. The fibers descend in the ventral funiculus on the same side as the nuclei from which they come. In the ventral horn they end—in part monosynaptically—on α and γ **motoneurons** (Fig. 18.8A). The tract is somatotopically organized, so that various body parts can be selectively controlled. The vestibulospinal tract has strong effects on the muscles that contribute to equilibrium and posture (see also Chapter 22, under “Vestibulospinal Tracts”). As mentioned, the lateral nucleus receives afferents from the utricle; these provide information about the static position of the head in space and thereby indirectly about the position of the body. Change of body position also changes its center of gravity, with a resulting need to adjust muscle tone to maintain equilibrium.

A smaller, **medial vestibulospinal tract** arises in the **medial vestibular nucleus**. It also descends in the ventral funiculus and acts on motoneurons. The fibers do not reach below the upper thoracic segments, however, and are thought to be of importance primarily for **head movements** elicited by the vestibular receptors. The so-called **vestibulocollic reflex** serves to stabilize the position of the head in space.

Fibers to the **nuclei of the extraocular muscles** arise mainly in the **superior and medial nuclei**, which receive many primary afferent fibers from the semicircular ducts (Fig. 18.8B). The fibers leave the nuclei medially and join to form a distinct fiber bundle, the **medial longitudinal fasciculus**, which is located close to the midline below the floor of the fourth ventricle (Figs. 18.7 and 18.8; see also Fig. 6.18). Some of the fibers cross to the

2 For example, a study in the cat with electric stimulation of separate divisions of the vestibular nerve found that about one-third of all neurons received convergent inputs from the vertical semicircular canal and sacculus/utricle. Another one-third received convergent input from sacculus and utricle, and one fifth from the horizontal canal and sacculus/utricle.

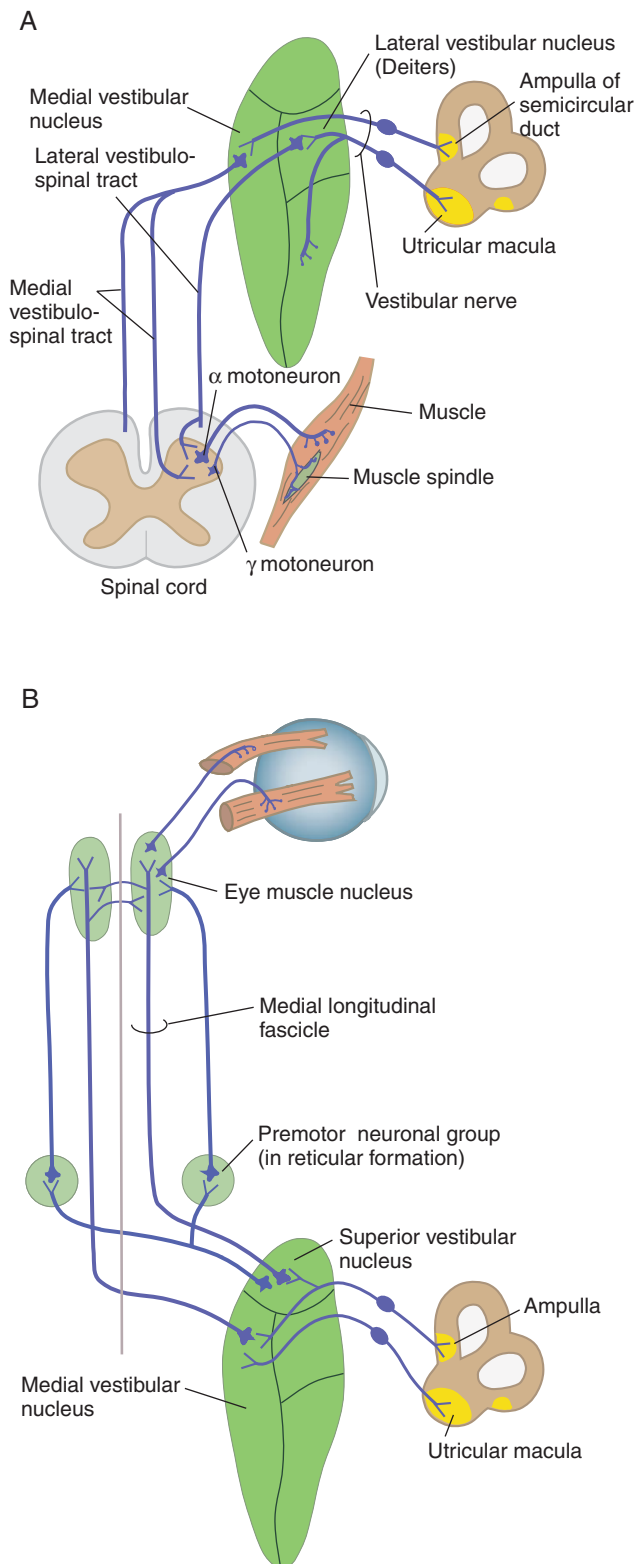


FIGURE 18.8 Main features of vestibular connections. **A:** Descending connections acting on α and γ motoneurons in the spinal cord. The medial vestibulospinal tract reach only cervical levels. **B:** Ascending connections controlling eye movements.

other side as they ascend. They end in the abducent, the trochlear, and the oculomotor nuclei and are precisely organized. In addition to the direct connections in the medial longitudinal fasciculus, there are **indirect pathways** from the vestibular nuclei to the eye muscle nuclei via the **reticular formation** (Fig. 18.8B). We return to this below when discussing vestibular reflexes.

Vestibulocerebellar fibers (arising in the medial and descending vestibular nuclei) end primarily in the **vestibulocerebellum** (Fig. 18.7; see also Fig. 24.4). Fibers from the vestibulocerebellum to the vestibular nuclei can adjust the gain (sensitivity) of the **vestibulo-ocular reflex**, as will be discussed next (see also Fig. 25.4).

Connections to the **thalamus** from the vestibular nuclei have been demonstrated anatomically and end in the VPL and nearby nuclei. Physiologically, scattered neurons in a fairly large regions respond to signals from the vestibular apparatus. This probably explains why vestibular signals reach several, discrete regions of the cerebral cortex.

VESTIBULAR REFLEXES: CONTROL OF EYE MOVEMENTS AND BODILY POSTURE

The special problems encountered in space journeys aroused great interest in the mechanisms that underlie vestibular reflexes. As with other parts of the brain being comprehensively investigated, conditions turn out to be more complex than initially thought. We will discuss only a few salient points. There are two kinds of reflexes that are elicited from the vestibular apparatus: **vestibulo-ocular reflexes** that control eye movements and so-called **labyrinthine reflexes** mediated by the vestibulospinal tracts controlling bodily postures (especially the upright position). In general, signals from the utricle and the saccule elicit **tonic** reflex effects, whereas **phasic** reflex responses are caused by signals from the semicircular ducts when the stimulus is rotation of the head (angular acceleration) and from the utricle or saccule when the stimulus is linear acceleration. Because the vestibular receptors inform only about the head, other receptors must inform about the movements and positions of other bodily parts. To control our upright position, the brain must integrate these various sources of information and issue commands that are appropriate for the whole body. Therefore, the labyrinthine reflexes must operate in concert with other postural reflexes, and we will treat these together below.

Vestibulo-Ocular Reflexes

There are several vestibulo-ocular reflexes, mediated by reflex arcs of various complexities. This topic is also discussed in Chapter 25 dealing with the control of eye movements. In general, the vestibulo-ocular reflexes

ensure that the **image is kept stationary** on the retina when the head moves (rotates).

The simplest vestibulo-ocular reflex is mediated by a chain of three neurons (Fig. 18.8; see also Fig. 25.4):

1. Primary afferent fibers from the cristae of the semicircular ducts
2. Neurons in the vestibular nuclei that send their axons to the nuclei of the extraocular muscles (passing in the medial longitudinal fasciculus)
3. Motoneurons in these nuclei, which send their axons to the extraocular muscles

In addition, there are other pathways from the vestibular nuclei to the nuclei of the extraocular muscles that are synaptically interrupted in the reticular formation and some other brain stem nuclei (Fig. 18.8B).

A movement of the head in any direction is accompanied by a **compensatory movement** of the eyes in the opposite direction and with the same velocity as the head movement. Rotation of the head produces movement of the endolymph inside the semicircular ducts. Taking a rotation in the horizontal plane (turning the head from one side to the other) as an example, mainly the lateral semicircular duct records the movement and elicits a compensatory eye movement in the horizontal plane.

When the head movement is relatively small, the eyes move with exactly the same velocity as and in the opposite direction of the head, and the image is kept in the same position on the retina all the time. When the head movement becomes larger, so that it becomes impossible to keep the image stationary even with maximal excursion of the eyes, a fast, or **saccadic**, movement occurs in the same direction as the head movement. Then the gaze is fixed again on the object, and another slow movement follows (as long as the head continues to move in the same direction). Such an alternation between slow and fast, saccadic eye movements is called **nystagmus**. In this case, the nystagmus was produced by stimulation of the semicircular ducts (rotation of the head) and is therefore called **vestibular nystagmus**. Movement of the surroundings can also elicit nystagmus when the head is stationary. This **optokinetic nystagmus** occurs, for example, in a train-passenger watching the landscape pass by.

Vestibular Signals Must Be Integrated with Other Sensory Modalities for Control of Eye Movements

As mentioned, the vestibular nuclei receive afferents from sources other than the labyrinth, such as nuclei in the mesencephalon, the reticular formation, and the cerebellum. Some of these sources mediate visual information that can modify the vestibular reflex responses. This convergence of various inputs seems logical. Thus, to achieve optimal control of the eye movements, the responsible neural cell groups must receive and integrate

vestibular information about movements of the head, visual signals about movements of the image on the retina, and proprioceptive signals about movements of the eyes relative to the head.

Vestibular Stimulation Produces Nystagmus and Falling Tendency

When an upright person rotates fairly rapidly a few times around his axis and then stops, the eyes can be seen to move rapidly one way (saccade) and slowly the other for some seconds afterward. Obviously, the rotation has induced **nystagmus**. By using special instruments, it can be seen that there is nystagmus also at the start of the rotation, but in the opposite direction of that occurring after the rotation has stopped. When the person rotates to the right, the saccade movement is to the right and the slow movement is to the left, as if the person fixes her gaze on a stationary point and then moves the eyes rapidly when this point is slipping out of the visual field. The eyes then move to a new fixation point, and the same sequence of events is repeated. This **postrotatory nystagmus** is caused by stimulation of the receptors in the semicircular ducts. As mentioned, at the start of the movement the inertia of the endolymph makes it lag behind, thereby bending the cilia of the receptor cells, whereas the endolymph continues to flow for a moment after the rotation has stopped. The person who had just stopped rotating feels as if he were still rotating, but now in the opposite direction. The direction of the nystagmus corresponds to the **illusion** of such a rotation—that is, with the saccade phase to the left after a rotation to the right. If the rotation of the body continues for some time, the nystagmus disappears and the person gets dizzy (compare with ballet dancers who deliberately ensure that the head does not move with even velocity during pirouettes; this way they have sufficient time for fixation so that the brain gets information to determine the orientation of the body in space).

After stopping the rotation, the person is also unsteady and tends to fall to one side, especially if he is asked to keep his eyes closed. Further, the arm deviates to the right if the person is asked to point straight ahead (with his eyes closed). This is called **postrotational past pointing**. After the rotation stops, the illusion of the opposite movement (i.e., the person feels he is turning to the left) causes the past pointing to the right: the person feels that the room is moving to the right.

The postrotational effects on postural muscles are mediated via the **vestibulospinal tracts** (Fig. 18.8A) and show that the receptors of the semicircular ducts also influence the spinal cord and the postural muscles, not just the cranial nerve nuclei and the extraocular muscles.

Nystagmus, falling tendency, and past pointing can also be produced by irrigation of the external auditory meatus with hot or cold water. The change of temperature

makes the endolymph flow in the semicircular ducts and thus produces stimulation of the receptors. Such a **caloric test** is used clinically to examine the function of the vestibular labyrinth and the conduction of signals to the brain stem.

Various **diseases** affecting the vestibular receptors or the signal pathways to the motoneurons of the extraocular muscles (the vestibular nerve, the vestibular nuclei, the medial longitudinal fasciculus, and the cerebellum) can produce nystagmus in the absence of vestibular or visual stimulation. This is called **spontaneous nystagmus**. In certain cases, the nystagmus may be present only in certain positions of the head (positional nystagmus).

Neck and Labyrinthine Reflexes

The vestibular receptors inform about the position and movements of the head in space, whereas neck proprioceptors can inform about the position and movements of the body in relation to the head. Based on information from both kinds of receptors, the brain can decide whether the head is moving in isolation or whether it moves together with the rest of the body. Obviously, different kinds of postural responses are needed in these two situations. The **labyrinthine reflexes** are elicited by stimulation of the sensory receptors of the **semicircular ducts** and the **utricle** of the labyrinth. In the **neck reflexes**, the response is a change of muscle tension, especially in the extremities supporting upright stance, induced by a change in the position of the head relative to the body (such movements take place primarily in the upper cervical joints). The labyrinthine reflexes when operating alone produce muscle contractions in the trunk and extremities that serve to keep the position of the head constant. The neck reflexes, as mentioned, serve to keep the position of the body constant in relation to the head. The latter is a prerequisite for the labyrinthine reflexes to function properly; the vestibular apparatus can provide information only about the position of the head in space and not about its position in relation to the body. Thus, the labyrinthine reflexes work on the assumption that the head has a constant position relative to the body, and the neck reflexes ensure that this position is constant.

The reflexes may be either **tonic** or **phasic**. A phasic neck or labyrinthine reflex consists of a rapid, transient change of muscle tension in postural muscles as a response to a change of posture (usually a disturbance of the equilibrium). We experience phasic labyrinthine reflexes when we trip over and a coordinated set of compensatory movements occur before we consciously perceive what is going on. In a tonic reflex, the change of muscle tension lasts as long as the new position is maintained.

Vestibulospinal tracts are the most likely candidates for mediation of the labyrinthine reflexes. The **reflex**

center of the neck reflexes is located in the medulla, and the effects on the motoneurons are most likely mediated by both the **reticulospinal** and vestibulospinal tracts.

More about the Neck and Labyrinthine Reflexes

Studies of **decerebrate**, four-legged animals have elucidated the neck and labyrinthine reflexes (the brain stem is transected just below the red nucleus in the mesencephalon). To demonstrate clearly the **neck reflexes** in decerebrate animals (mostly cats have been studied), the vestibular receptors must have been eliminated. When then the head is bent backward (extension of the neck), the muscle tension is increased in the extensor muscles of the forelimbs and decreased in the extensors of the hind limbs. Forward bending of the head (flexion) induces the opposite pattern of changes (extension of the hind limbs and flexion of the forelimbs). Tilting the head sideways increases the extensor tone on the same side and reduces it on the other side, as does turning the head sideways. These changes of the muscle tone aim at reestablishing the position of the body relative to the head. The receptors for these reflexes are located near the upper cervical joints, because they disappear after transection of the upper three cervical dorsal roots. **Muscle spindles** are the most likely candidates, but joint receptors may also contribute. As mentioned, the functional role of the neck reflexes cannot be understood when observed in isolation, however. Only in conjunction with the labyrinthine reflexes are their effects appropriate for the whole body.

To demonstrate clearly the **labyrinthine reflexes**, the neck reflexes must have been eliminated by cutting the upper dorsal roots (in a decerebrate animal). It then appears that the effects produced by the labyrinthine reflexes are the opposite of those of the neck reflexes when the latter act alone. Thus, bending the head backward elicits flexion of the forelimbs and extension of the hind limbs, and vice versa when the head is bent forward. The purpose of these changes of muscle tone is to bring the head back to the position held before the movement—that is, to keep the position of the head in space constant. Provided the neck reflexes ensure that the body stays in a constant position relative to the head, the labyrinthine reflexes will serve to maintain both the position of the head in space and the upright position of the whole body. The labyrinthine reflexes are shown clearly when the experimental animal stands on a platform that can be tilted in various directions. Tilting the platform forward increases the extensor tone in the forelimbs and decreases the tone in the hind limbs. Tilting the platform sideways increases the muscle tone in the extensors on the side to which the tilt is directed. Both responses are obviously appropriate for the maintenance of body balance. When the platform is moved quickly in one direction and then back, the

reflex response is transient (**phasic reflex**). When the platform is maintained in the new position, the altered muscle tone is upheld (**tonic reflex**).

When **both reflexes work together** in an intact organism, backward bending of the head, for example (with movement taking place only in the upper cervical joints and no change of body position), produces no change in muscle tension of the extremities. The tendency of the labyrinthine reflexes to produce forelimb flexion and hind limb extension is canceled by the opposite tendency of the neck reflexes. In contrast, when the same movement of the head is produced by a backward movement of the whole body (with no movement of the head relative to the body, like a horse that is rearing), the labyrinthine reflexes act alone to produce extension of the hind limbs and flexion of the forelimbs. Another example is an animal standing on a platform tilting forward with no movement of the head relative to the body. In that case, the labyrinthine reflexes produce forelimb extension and hind limb flexion. This is an appropriate response to maintain balance when standing on a downhill slope. However, if the position of the head in space is kept constant and the body is moved in relation to the head, the neck reflexes act alone. An example is a cat jumping down from a table: the neck is extended (keeping the head position constant), producing extension of the forelimbs, which is appropriate for landing.

Postural Reflexes: Various Receptors Contribute

In order to control posture and balance, the CNS must receive sufficient information about positions and movements in various body parts as well as the nature of the support surface. This information enables the brain to compute the position of the center of gravity and the movements needed to keep it in the right position relative to our supporting base. Many sense organs can provide relevant information (visual, proprioceptive, cutaneous, and vestibular). The signals are analyzed in the CNS, which in response initiates **postural reflexes**, that is, automatic, coordinated contractions of muscles that maintain our upright position. Postural reflexes can be elicited from the labyrinth, from proprioceptors and by vision. The contribution of various receptors to control of the upright position is not static, however. This is because the contribution of each receptor type varies with the nature, magnitude, and context of a postural perturbation.³ For example, somatosensory information from the sole of the foot is less reliable when the support surface is slippery than when it is firm and

even. Further, visual information may be unreliable if we are in an environment with moving objects, because it may be difficult to distinguish own movements from those of the surrounds. Persons with loss of vestibular function manage well as long as they can see but they have serious problems in maintaining the body's equilibrium in the dark. If one source is unreliable, its contribution tends to be ignored, and information from other sources becomes more important. We also treat postural reflexes in Chapter 22 in conjunction with the control of automatic movements.

More about Receptor Types and Their Contribution to Postural Control

Persons standing on a platform that can be tilted or moved forward or backward have been much used in studies of postural control. In this situation, the experimenter can control the direction, speed, and amplitude of perturbations, and specific kinds of sensory information can be removed temporarily. In addition, patients with loss of one or the other kind of receptor have been much studied.

Signals from **muscle spindles**—providing rapid information about joint position and movements—are important for postural control. This is, for example, evident from persons who lose the proprioceptive sense (cf. Chapter 13, under “Clinical Examples of Loss of Somatosensory Information”). Further evidence that muscle spindles play a role for posture comes from experiments with **vibration** (activating Ia afferents) of leg muscles and neck muscles. Such vibration elicits postural adjustments, which are appropriate, considering that the brain “believes” that the muscles are being stretched. The signals are obviously interpreted as if the center of gravity were moving. In this case, the muscle spindle information is not primarily used at the segmental level for quick postural responses but is integrated with other inputs at higher levels. Platform experiments further support that the contribution of muscle spindles to postural control does not depend mainly on simple, **spinal reflexes** (i.e., reflex contractions in response to muscle stretch that do not depend on higher levels). When the platform is suddenly displaced backward (without tilting), the body first sways forward with movement primarily at the ankle joints. The balance is regained mainly because the calf muscles at the back of the leg contract (muscles of the hip, the back, and in the neck also contribute, especially if the displacement is large). The first part of the contraction of the leg muscles occurs so early after the perturbation that a reflex must mediate it. (Somewhat later there is also a voluntary contraction, which contributes to the final outcome of the postural adjustment.) In this case, a reflex elicited by stretch of muscle spindles in the calf muscles would seem appropriate. In another situation, however, such a reflex would worsen

3 There furthermore seems to be individual variation among normal subjects with regard to which kind of information they rely on for the reflex adjustments in quiet standing. In a study of healthy volunteers, only half of the subjects increased their body sway when the eyes were closed. This suggests that persons differ with regard to how much they rely on visual information to stabilize the quiet standing position.

the balance: if the platform is suddenly tilted backward, the calf muscles are stretched (as in the former example) but the center of gravity is now displaced backward instead of forward. Consequently, a contraction of the calf muscles would worsen the imbalance. To regain balance, the muscles at the front of the leg have to contract as rapidly as possible, in spite of being shortened by the tilting of the platform, and this is what happens. Thus, whether a segmental, spinal reflex is elicited depends on the situation, and inappropriate reflex responses to stretch are generally suppressed. Indeed, for the functionally important reflex response in the leg muscles (starting about 90 msec after the balance perturbation), signals from proprioceptors around the **vertebral column** may be more important than signals from leg muscle spindles. Thus, the contraction of the leg muscles correlates less with their own length change than with the displacement of the trunk.

Signals from **cutaneous** low-threshold mechanoreceptors in the **sole of the foot** seem to contribute to the reflex contractions of the leg muscles in platform experiments. The dorsal column–medial lemniscus pathway transmits such signals very rapidly. Presumably, the brain determines the center of pressure by calculating the difference between the pressure applied to the heel and the forefoot. The center of pressure informs about the position of the body center of mass. Patients with **neuropathies** who have reduced sensibility in the sole of the foot witness the importance of this input for postural control and for normal gait.

Vision also contributes to adjustment of muscle tone with the purpose of maintaining body equilibrium. Platform experiments indicate that the muscle contractions in response to a movement of the platform depend on whether the subject can see that the body moves in relation to the surroundings. If the experimental setup is such that it appears as though the surroundings do not move (i.e., they are made to move in the same direction as the head), the earliest reflex contractions of the legs are weaker than when the sense of vision also informs about the movement. We do not fully know the pathways involved, but the final commands are probably sent from the vestibular nuclei in the vestibulospinal tract.

In platform experiments with moderate postural perturbations, signals from **vestibular receptors** seem not to be essential for the corrective contractions of leg muscles (ankle strategy). With larger perturbations, however, vestibular information contributes to contractions of hip muscles (hip strategy). When visual information is eliminated, signals from the utricle and the saccule are indeed necessary for the proper orientation of the body in space, as can be shown in animals whose otoliths have been removed. **Unexpected fall** from some height elicits a postural reflex that depends on information from the vestibular apparatus. The fall initiates a contraction of the muscles of the leg as an appropriate preparation to

the landing. The contraction begins about 75 msec after start of the fall and can therefore occur before landing (and thus before a stretch reflex can be elicited). The latency is also too short for the contraction to be voluntary. Animal experiments indicate that the receptors of this reflex are located in the labyrinth (probably in the saccular macula).

Postural Reflexes Are under Central Control

Reflex responses occur too early after a balance perturbation to be caused by conscious, high-level decisions. Indeed, the main advantage of a reflex response is that it occurs so rapidly. Nevertheless, even automatic movements are subject to central control. The response to a perturbation challenging our upright position is strongly modulated by expectation and the context in which the perturbation occurs. For example, a contraction of the triceps surae muscle in response to dorsiflexion in the ankle joint may restore balance in one situation while worsen it in another. In general, the individual reflex responses are subordinate an overall, coordinated motor plan designed to attain a certain goal.

In **infants**, comprehensive, high-level motor programs have not yet developed and accordingly, some postural reflexes operate “on their own.” This concerns some vestibular reflexes, the grasp reflex, and others. In comatose patients with lesions of the upper brain stem, exaggerated postural reflexes may appear spontaneously—usually with a strong dominance of extensor tone. This condition is called **decerebrate rigidity**.

CORTICAL PROCESSING OF VESTIBULAR SIGNALS

Several Areas Receive Vestibular Signals

Vestibular signals reach several small areas in the cortex, as shown electrophysiologically in monkeys and with imaging methods in humans (Fig. 18.9). The **parietal insular vestibular cortex—PIVC**—is of particular interest (as the name implies, it lies at the junction between the parietal lobe and the insula).⁴ The PIVC contains neurons that are activated by signals from both the semicircular canals (head rotation) and from proprioceptors around the upper cervical joints. Because, in addition, the neurons are influenced by visual signals it seems likely that they integrate all kinds

⁴ Imaging studies in humans place the PIVC and other presumptive vestibular-activated areas somewhat differently. For example, some find that PIVC includes posterior parts of the insula, while others limit it to a region just posterior to the insula. Such discrepancies are most likely due to differences in experimental conditions. It is very difficult to rule out that cortical activation evoked by caloric, galvanic, or optokinetic stimuli—applied to stimulate vestibular receptors—are due to concomitant stimulation of somatosensory or visual systems.

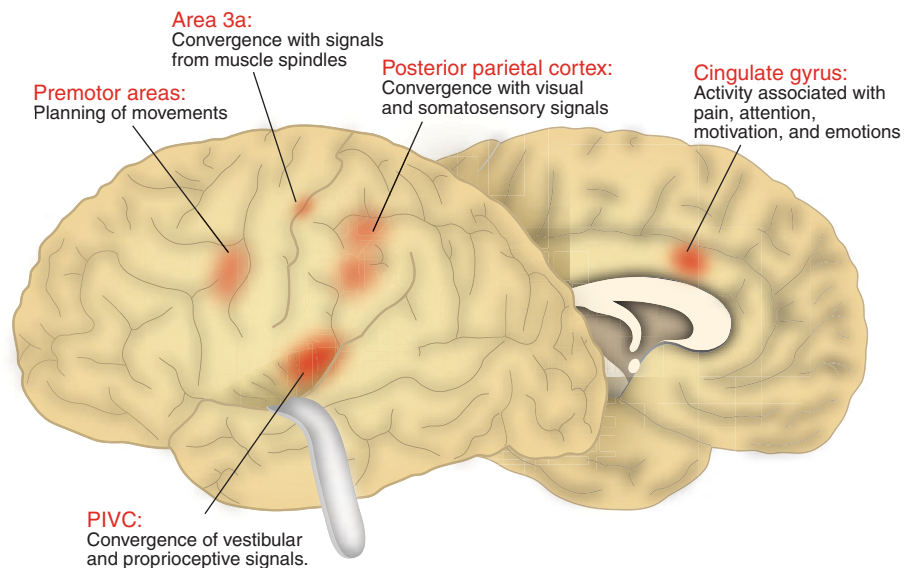


FIGURE 18.9 *Cortical areas receiving signals from the vestibular apparatus.* The locations of regions are only approximations, based on extrapolation from fMRI data published by Bottini et al. 2001.

of **movement-related information**. Signals from the labyrinth would thus contribute to the awareness of bodily posture and movements in space. Interestingly, clinical observations suggest that lesions in this region result in denial of ownership of the contralateral hand. Neurons in other parts of the cortex that are activated from the labyrinth have properties similar to those in PIVC. This concerns neurons in **area 3a** (muscle spindle input), in **area 2** (joint receptor inputs), and in the **cingulate gyrus**. An area close to the PIVC in the **posterior parietal cortex** integrates vestibular and optokinetic information. Vestibular activation furthermore occurs in the **premotor cortex** (and in some other areas). The various “vestibular” cortical areas are interconnected and send efferent fibers to the vestibular nuclei, as mentioned earlier.

Distributed Networks, Body Image, and Body Scheme

Vestibular information seems to be processed in a distributed cortical network, not in a single center. This network integrates several kinds of sensory information related to extrapersonal space and the body. It is connected with and overlaps other networks—notably those that control movements but also networks related to attention, emotions, and pain. By feeding into this network, vestibular receptors contribute to, but are not solely responsible for our awareness of body orientation and movements in space. Other specialized regions of the cortex process other aspects of body-related information. For example, two extrastriatal visual areas appear to be specialized for recognition and analysis of human bodies and body parts. We mentioned the **fusiform body area** (FBA) in Chapter 16 (see Fig. 16.26), which is particularly activated by recognition of whole human bodies (rather than body parts). Another similar area

in the occipital lobe—**extrastriatal body area** (EBA)—responds especially to body parts (e.g., when the person looks at an arm or a hand). Active movement of the body part modulates the EBA activity, and this may enable this region to help decide whether the hand belongs to me or someone else (body ownership).

Together, these multiple, interconnected regions enable us to be aware of our bodies; they provide us with a feeling of **ownership**, and a sense of **agency** (i.e., the experience that *I* am moving my hand, not someone else). They also, presumably, form the basis of **internal models** representing stored information of the neuronal processes needed for specific actions (grasping a glass of water, descending a staircase, and so forth). **Body image** and **body scheme** are terms often used about two different aspects of internal bodily representations. The body image concerns our conscious perception of the appearance of our body, with regard to its form, size, and other characteristics. The body scheme concerns spatial representations of our body parts that do not enter awareness. The body scheme provides a basis for actions, and is continuously **updated** during movements. Putting it simply, the body image concerns “what” whereas the body scheme concerns “how.” At least superficially, the distinction between body image and body scheme resembles the distinction between the ventral and dorsal pathways of the visual system: the ventral pathway processes explicit, conscious knowledge about objects, whereas the dorsal pathway deals with implicit knowledge about the objects’ spatial characteristics and its relation to action.

Normally, **ownership** and **agency** are self-evident: I do not doubt the hand is mine, and that *I* am moving it. The transformation of intention into action is effortless and requires no knowledge of the internal operations underlying it. Further, I always experience my body

as a **unity**: I have a “natural” sense of **coherence**. Nevertheless, these self-evident “facts of life” depend on a complex interplay among numerous specialized regions of the brain. There is no single area responsible for our sense of unity, ownership, or agency. Indeed, there are numerous clinical examples that the body image and body scheme are more fragile than we usually assume. Thus, after a stroke, the patient may deny ownership of an arm, or not recognize that the arm is paralyzed or even experience that someone else is moving the arm. In addition, strong emotions and mental illnesses (e.g., schizophrenia) may disturb the body image or body scheme. Immobilization of joints—presumably producing a mismatch between motor commands and sensory feedback—may cause bizarre experiences of positions and movements of the extremity.

Disequilibrium, Dizziness, and Vertigo

Berthoz and Viaud-Delmond (1999, p. 709) give a concise yet comprehensive description of dizziness; “Dizziness is characterised by a marked distortion of self-world relations and reflects a discrepancy between internal sensation and external reality. Spatial disorientation, as well as dizziness, can be due to a peripheral problem in any of the sensory modalities; or, it may be due to a central problem, involving not one particular sensory modality but rather the integration and weighting of the different modalities and their relation with memory.” Certainly, **disequilibrium** and **dizziness** can arise because one of the usual channels of sensory information is lost or malfunctioning—that is, dizziness of peripheral origin. **Vertigo**—with the very distinctive illusion of movements (usually rotation), either of the body in relation to the surroundings or vice versa—should be distinguished from other kinds of dizziness. True vertigo is usually caused by labyrinthine disease, whereas dizziness without vertigo may have various causes. For example, considerable loss of proprioceptors and vestibular receptors is common in elderly people and may contribute to dizziness. Abnormal stimulation of certain receptors may also provoke dizziness and imbalance. Perhaps the problems arise not so much because of lack of information as from a **conflict** between the various sensory inputs: the internal model expects a certain relationship between them so that if the composition of inputs is altered, it takes some time to update the internal model. The updating presumably depends on use, as do plastic changes in general—that is, improvement depends on specific training of the impaired functions. Elderly people suffering from dizziness may easily enter a vicious circle: they need increased amount of balance training because of reduced sensory inputs or central processing capacity, yet they move less than before because they are afraid of falling. Dizziness is also a common symptom in many **psychiatric**

disorders, and it is a transient reaction in certain situations, as with strong **emotions**. In such cases, malfunctioning receptors are hardly responsible. Rather, the cortical networks for spatial orientation may have a reduced ability to integrate the various sensory inputs—perhaps because inputs from networks related to emotions and memories disturb them.

Unilateral **destruction** of the **vestibular apparatus** in humans usually gives very disturbing symptoms, dominated by dizziness, spontaneous nystagmus, nausea, and disequilibrium. The symptoms generally vanish with time, mainly due to central excitability changes that level out the unequal inputs from the two sides. The normalization of eye movements seems to be due, at least partly, to the patient learning to use proprioceptive signals instead of vestibular ones. Thus, in monkeys with unilateral damage of the labyrinth, signals from neck muscle spindles initiate an almost normal “vestibulo-ocular reflex” when the head rotates.

Motion Sickness

Many people suffer from this condition characterized by nausea, vomiting, dizziness, and autonomic disturbances like cold sweat and low blood pressure. Presumably, the symptoms arise because of **conflict** between motor commands and sensory information (from the semicircular ducts, the maculae, the proprioceptors, and vision); that is, there is a **mismatch** between the sensory information and what was expected based on prior experience. Usually, our own movements produce sensory signals, whereas when sitting in a car signals from vestibular receptors are passively produced and are not accompanied by the expected pattern of proprioceptive signals. Such theories do not explain all aspects of motion sickness, however.

The **symptoms** of motion sickness resemble those evoked by ingestion of certain poisons (see Chapter 27, under “The Vomiting Reflex”) and involve some of the same central structures (indeed, labyrinthectomized dogs show decreased susceptibility to many emetic drugs). This concerns the **solitary nucleus** in the medulla, as well as the pathways that control the preganglionic autonomic neurons and motoneurons responsible for emptying the stomach. Animal experiments confirm that stimulation of vestibular receptors can produce several **autonomic effects**, by connections from the vestibular nuclei to the reticular formation and then to preganglionic autonomic neurons. Signals from the sacculus and utricle are probably particularly important for the occurrence of motion sickness, judging from studies of persons exposed to weightlessness during space travels. The **cerebellum** (especially the nodulus and uvula in the posterior vermis) also seems to play a role. Thus, after removal of the posterior vermis in animals, provocations must be much stronger to produce motion sickness.

19 | Olfaction and Taste

OVERVIEW

The chemical senses of smell and taste enable us to recognize a vast number of different molecules in our surroundings. Obviously, this gives us information of great importance that triggers a range of responses—from avoiding poisonous food to enjoying the fragrance of a rose. Both senses depend on binding of molecules to specific **chemoreceptors**. These receptors are very sensitive, thus permitting recognition and discrimination of molecules in extremely low concentrations (in air or in fluids). Stereospecificity—that is, the shape of the ligand determines which receptor it binds to—is another common property of the chemoreceptors for taste and smell. The transduction mechanisms—that is, how the stimuli translate to graded receptor potentials—are similar for receptors for taste and smell because they (with some exceptions) are coupled to **G proteins** (as are photoreceptors and many neurotransmitter receptors).

The **olfactory system** differs from other sensory systems in certain respects: the **sensory cells**—located in the upper nasal cavity—are neurons with an axon going directly to the **olfactory bulb** (part of the central nervous system (CNS)). Further, the signals in the **olfactory tract** destined for the cerebral cortex are not synaptically interrupted in the thalamus but goes directly to the **primary olfactory cortex** the medial temporal lobe near the uncus.

The taste (gustatory) receptor cells are found in taste **buds** scattered over most of the tongue. The sensory cells are equipped with receptors identifying four basic **taste qualities** (sweet, salty, sour, and bitter). In addition a fifth quality called umami (Japanese: savory, meaty) is evoked by glutamate binding to specific receptors. Signals from the taste buds are transmitted in the **facial** (intermediate) and **glossopharyngeal** nerves to the **solitary nucleus** in the upper medulla. From there signals are distributed to other brain stem nuclei important for food intake and digestion, and to higher levels such as the **hypothalamus**, the **amygdala**, and the primary taste area in the anterior part of the **insula**. Integration of olfactory and gustatory information takes place in the **orbitofrontal cortex** (and some other places). Here, information from the chemical senses is processed in a broader context with the aim to facilitate appropriate behavior.

THE OLFACTORY SYSTEM

The sense of smell does not play the same important role for adult humans as the senses of hearing and vision. This does not mean that the sense of smell is insignificant in daily life. The perception of odors is special by its close association with memories and with emotions and moods. Presumably, this explains why we usually remember odors so well. An enormous industry devoted to producing perfumes and other fragrances shows that for humans the sense of smell has an important role in interpersonal communication.

Going back in the evolution of the species, we find that smell is the most primitive of the senses. It is also the most important one at the early stages of evolution. To understand the evolution of the brain, knowledge of the “olfactory brain” or **rhinencephalon** has been important because in the early, primitive vertebrates almost the whole cerebrum is devoted to the processing of olfactory signals. In higher vertebrates, new parts of the cerebrum emerge that gradually and completely overshadow the phylogenetically old parts. The organization of the central pathways and nuclei that process olfactory information reflects the fact that this system developed earlier than the parts of the cortex (the neocortex) that treat other sensory modalities. The term olfactory brain for these old parts is unfortunate, however. In higher vertebrates, large parts of the regions corresponding to the rhinencephalon in lower animals have nothing to do with the sense of smell but have taken on other important functions, the hippocampus (Chapter 32) being the most striking example. This is a common occurrence during evolution: structures that are no longer used for one function may form the basis for the development of new capacities. Thus, the parts treating olfactory signals have not developed in pace with the rest of the brain during evolution and are therefore relatively much smaller in humans than in, for example, cats and dogs. In absolute terms, however, the differences are not so marked.

Receptor Cells for Smell

The special receptor cells for smell are located in the mucous membrane of the upper part of the nasal cavity,

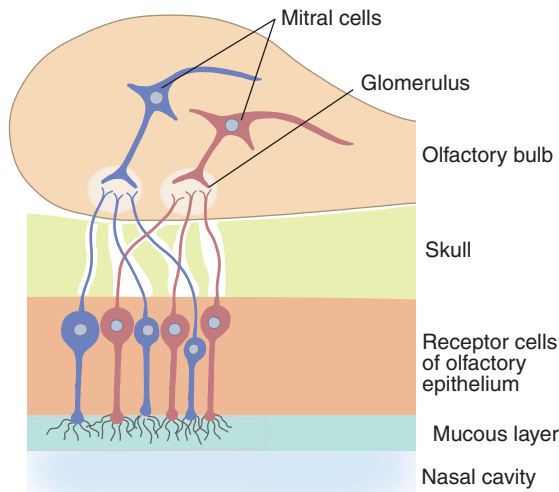


FIGURE 19.1 *The olfactory epithelium and connections to the olfactory bulb.* Neurons (receptor cells) in the olfactory epithelium have cilia embedded in mucus. The central process (axon) of the receptor cells end on mitral cells in the glomeruli of the olfactory bulb. The axons from neurons sharing odorant specificity tend to converge on one or a few glomeruli in the olfactory bulb. Thus, the glomeruli show some odorant specificity. (Based on Mombaerts 1996.)

the **olfactory epithelium**.¹ The total number of receptor cells has been estimated to about 10 million in humans, and the cells are constantly renewed. Because they are in fact primitive neurons, they are exceptions to the rule that neurons that die are not replaced. The olfactory epithelium is pseudostratified and consists of supporting cells and so-called basal cells besides the receptor cells (Fig. 19.1). The **supporting cells** probably insulate the receptor cells electrically, so that signals are not propagated from one cell to another. The **basal cells** divide mitotically and probably give rise to the receptor cells. The **receptor cells** are bipolar and send a dendrite-like branch toward the epithelial surface, and an axon through the base of the skull to the olfactory bulb. The dendrite ends with an expansion densely covered with **cilia** (Fig. 19.1). Because the cilia are embedded in the **mucus** that covers the epithelium, only substances dissolved in the mucus can act on the receptor cells. Many

¹ The area of the **olfactory epithelium** is obviously not easy to determine in humans. Thus, figures in the literature vary from 1 to 5 cm². This variation may at least partly be due to a patchy distribution of the epithelium and age-related loss. Thus, while in the fetus the epithelium is continuous and covering a large proportion of the nasal cavity, it is gradually replaced by respiratory epithelium in a patchy fashion. Further, the epithelium may be more anteriorly located than formerly believed. A combined histological (biopsies) and electrophysiological study located the anterior border of the olfactory epithelium at the level of the anterior end of the middle turbinate—1 to 2 cm anterior to what was formerly believed (Leopold et al. 2000).

odorous substances are hydrophobic, however; that is, they do not dissolve easily in water. Therefore, we assume that there exist mechanisms—such as transport proteins—to bring hydrophobic odorants through the mucus. Several families of **odorant-binding proteins** (OBPs) have in fact been identified in the mucus. It is probable that each protein binds specifically to certain odorants and may serve to concentrate the odorant in the proper part of the olfactory epithelium (i.e., the part containing the receptors specific for the particular odorant). The odorant-binding proteins may also help remove the odorant so that the receptors quickly regain their sensitivity. Specialized glands produce the mucus, which consists of several layers.

Transduction Mechanism

Experiments with a large number of odorants suggest that the shape of the molecule, rather than its chemical composition, determines how it smells (stereospecificity). This **stereochemical theory** of smell proposes that the receptor sites on the receptor cells have different shapes and that only molecules with a complementary shape fit into the receptor site. Binding of the molecules (odorants) to specific **odorant receptor proteins** (ORs) in the membrane of the cilia evokes a **receptor potential**. This involves activation of G proteins and cyclic AMP. Increased intracellular cyclic AMP opens Na⁺-selective cation channels and thus depolarizes the cell. The ORs share structural features with photoreceptors and β adrenergic receptors. In contrast to photoreceptors, however, the olfactory receptors produce action potentials. The action potential arises in the initial segment of the axon and is transmitted to the olfactory bulb.

The Olfactory Receptor Cells Express an Enormous Repertoire of Receptor Molecules

What is the basis of our ability to discriminate several thousand different odors? The olfactory system is special by the expression of a large number of specific **odorant receptor proteins** (ORs), coded by the largest vertebrate gene family comprising around 1000 genes (only the immune system can recognize more different molecules). Human ORs express “only” about 350 different ORs on the surface of the cilia, while rodents and lower primates express many more, probably because a large fraction (60%–70%) of the human odorant-receptor genes appears to be **pseudogenes** (genes that are not expressed). In rodents, the proportion of pseudogenes is only about 5%. Presumably, this evolutionary increase in the number of pseudogenes may reflect diminished importance of olfaction in higher primates.

Although each olfactory receptor cell appears to express only one kind of OR, electrophysiological recordings show that the individual olfactory cell responds to several odorants. This is because each OR can bind several odorants, and each odorant can bind to multiple ORs. Thus, each odorant will evoke a complex pattern of activity among the odorant receptor cells.

Central Pathways for Olfactory Signals

Like other sensory cells—for example, those in the inner ear and in the taste buds—the olfactory receptor cells are surrounded by supporting cells. In other respects, however, the olfactory receptor cells are different from other sensory receptors, showing a more primitive arrangement. Thus, the olfactory cells themselves send a process (an axon) centrally, whereas in the inner ear, for example, a peripheral process of a ganglion cell contacts the receptor cell. The unmyelinated axons of the olfactory cells form many small bundles, together constituting the **olfactory nerve** (the first cranial nerve; Fig. 19.2). The bundles pass through the base of the

skull close to the midline in the anterior cranial fossa—through the **lamina cribrosa** of the ethmoid bone. The olfactory nerve enters the **olfactory bulb** (Figs. 19.1 and 19.2; see also Fig. 6.13), located just above the nasal cavity under the frontal lobe. In the olfactory bulb, the unmyelinated olfactory nerve fibers establish synaptic contacts with the **mitral cells** (Fig. 19.1), which, in turn, send axons to the brain through the **olfactory tract** (Fig. 19.2). The mitral cells are collected in small, round aggregates, called **glomeruli** (Latin: *glomerulus*, small ball of thread). Each glomerulus forms a functional unit.

In view of the enormous repertoire of specific ORs, it is natural to ask to what degree the specificity is maintained in the further projection from the olfactory epithelium. There seems to be some topography within the sheet of olfactory epithelium. Thus, different receptor genes are expressed in somewhat different parts of the olfactory epithelium, as shown with the *in situ* hybridization technique. **Four zones** differing with regard to receptor–gene expressions exist in rodents. Further, anatomic studies with axonal transport methods show that axons from these four zones end differentially in the olfactory bulb (Fig. 19.2). A more pronounced segregation occurs in the glomeruli, however, as each glomerulus receives afferents from receptor cells with closely similar specificity. This implies that axons from receptor cells with quite different positions in the olfactory epithelium converge in one glomerulus. This glomerular specificity appears not to be inborn, however, but results from **use-dependent plasticity**. Thus, in neonatal mice each glomerulus receives axons with different specificities. This condition persists if use of the olfactory system is prevented by closure of the nostrils shortly after birth.

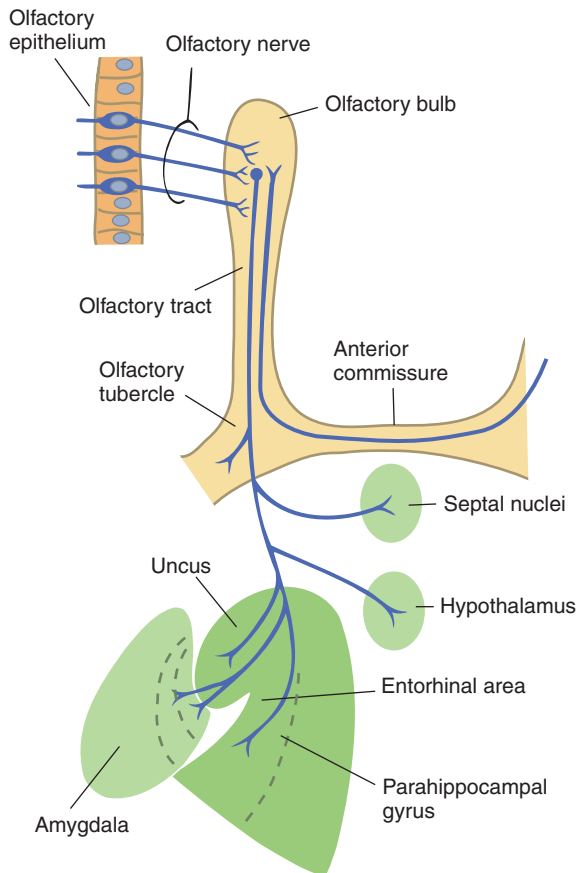


FIGURE 19.2 *The olfactory pathways.* Schematic illustration of some main connections. In the upper left are the olfactory receptors in the nasal mucosa, sending their central processes to the olfactory bulb. The neurons of the olfactory bulb send their axons to the cortex and various nuclei in the vicinity of the tip of the temporal lobe.

The Olfactory Bulb

The structure of the olfactory bulb is complex. It is not a simple relay station but, rather, a small “brain” in itself, carrying out substantial processing of the sensory information reaching it. In this sense, there are similarities with the retina, and both are parts of the CNS that have been “moved” outside the brain. There is some evidence that the olfactory bulb is of decisive importance for the **discriminative** aspect of olfaction, or the ability to distinguish different odors. Thus, lesions of nuclei in which the fibers from the olfactory bulb terminate do not appear to impair simple olfactory discrimination.

As mentioned, the axons from the receptor cells synapse with the dendrites of the **mitral cells**. The **glomeruli** form complex arrangements involving several dendrites and processes of local neurons. The number of glomeruli in the olfactory bulb declines with age, and only a few are said to remain in very old persons.

Besides mitral cells, the olfactory bulb contains numerous small **granule cells**, forming a separate layer. They are local neurons that interconnect mitral cells by way of **dendrodendritic synapses** (the granule cells lack axons). The granule cells are believed to mediate **lateral inhibition** in the olfactory bulb, and this is analogous to the horizontal cells of the retina. Like the horizontal cells, the granule cells are **electrically coupled**.

Many **neurotransmitters** are found in the olfactory bulb. The mitral cells most likely use glutamate, whereas the granule cells release γ -aminobutyric acid (GABA). Other interneurons are believed to be dopaminergic. In addition, norepinephrine and glycine are present in some neuronal processes.

There are also **efferent fibers** in the olfactory tract, as shown anatomically. Accordingly, electrical stimulation of the olfactory cortex can influence (primarily inhibit) the signal transmission through the olfactory bulb. Some of the efferent fibers release norepinephrine and are involved in synaptic changes in the olfactory bulb related to a certain kind of **learning** (see later).

The Terminal Areas of the Olfactory Tract

The fibers in the olfactory tract eventually take various directions, ending in different nuclei, most of which are located close to the tip of the temporal lobe (Figs. 19.2 and 19.3). In contrast to pathways for other sensory modalities, which are synaptically interrupted in the thalamus, fibers from the olfactory bulb pass directly to the cortex. Another difference concerns the cortical lamina in which the fibers terminate: olfactory fibers terminate in the outermost layers, whereas thalamocortical fibers

terminate mainly in deeper layers. Most of the efferent fibers from the olfactory bulb end at the medial aspect of the **temporal lobe**—partly in the **cortex** and partly in the **amygdala** (Figs. 19.2 and 10.3). The amygdala is located just below the cortex in the tip of the temporal lobe (Fig. 19.1; see also Figs. 31.1 and 31.3). In the cortex, fibers terminate both in the so-called **piriform cortex** in the **uncus** and in the adjoining parts of the **entorhinal area** (Figs. 19.2 and 19.3). The fibers to the amygdala end only in the corticomedial nucleus, which sends efferent fibers to the **hypothalamus**. (The amygdala are discussed further in Chapter 32.) The cortical regions in the temporal lobe that receive direct fibers from the olfactory bulb are called the **primary olfactory cortex** (Fig. 19.3). It is believed that olfactory signals come to consciousness in these and nearby cortical areas.

Uncinate Fits and Déjà Vu

Lesions affecting the uncus and the immediately surrounding cortex can be accompanied by subjective olfactory experiences (often unpleasant). Such sensations often occur as so-called **uncinate fits**, which frequently also include a peculiar feeling of experiencing the events in a dream—**dreamy state**. Often the patients feel that they have experienced the event before (**déjà vu**). Such uncinate fits may develop into an epileptic seizure, and the condition is regarded as a form of **epilepsy**.

Further Processing Outside the Primary Olfactory Cortex

The olfactory cortex and the amygdala forward olfactory signals to other parts of the brain. Connections to other parts of the cortex serve to **integrate** olfactory with other kinds of sensory information, leading to the analysis of its meaning. From the primary olfactory cortex, direct fibers reach nearby areas on the underside of the frontal lobe, the **orbitofrontal cortex**. In addition, this part of the cortex might receive olfactory information indirectly by way of connections from the **mediodorsal thalamic nucleus**, which receives fibers from the amygdala. The orbitofrontal cortex receives converging projections from many cortical areas, such as visual and **somatosensory** association areas, parts of the insula in receipt of **taste** information, and areas related to **emotions**, **motivation**, and **memory**. Such connections are generally reciprocal; that is, the sending areas receive information from the orbitofrontal cortex, which presumably concerns the broader meaning of the olfactory information.

One important pathway for olfactory signals reaches parts of the **hypothalamus**. These hypothalamic parts are involved in control of appetite, digestion, and feeding behavior (among other functions). Since also the orbitofrontal cortex sends efferents to the hypothalamus (among other areas), this part of the brain receives both relatively “pure” olfactory information from the

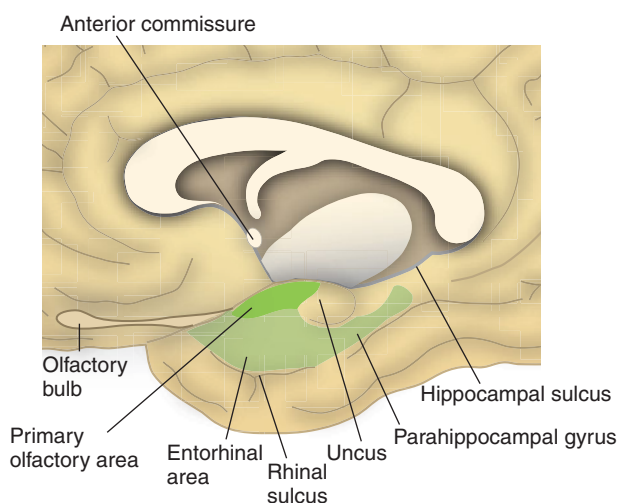


FIGURE 19.3 The olfactory cortex. The medial aspect of the cerebral hemisphere. The primary olfactory cortex is located in the uncus (dark green). The olfactory cortex in humans most likely also encompasses parts of the entorhinal area (light green).

amygdala and highly processed information that has been analyzed regarding its meaning for the organism.

The olfactory nuclei (not the olfactory bulb) on the two sides are interconnected by fibers running through the **anterior commissure** (Fig. 19.2). Thus, olfactory information from both sides of the nasal cavity is treated in each hemisphere.

Olfactory Signals and Behavior

Olfactory connections to the hypothalamus—both direct and indirect ones via the amygdala—are important for eating and for behavior directed at **acquiring food**. Sexual reflexes and sexually related behavior are also influenced by olfactory signals, although more so in lower mammals than in humans. The structural basis for such reflexes and behavior is very complex and not known in detail. Olfactory connections to the amygdala are most likely of importance, not only because the amygdala acts on the hypothalamus but also because it acts on parts of the prefrontal cortex involved in control of emotions and emotional behavior.

Olfaction and Learning

Olfactory sensations in certain **sensitive** (critical) periods of development can induce lasting changes of behavior; that is, learning. This phenomenon is called **olfactory imprinting**. One example concerns **sheep** that establish a strong bond to the lamb shortly after the birth. This depends on odors of the lamb that are present in the amniotic fluid. The mother must be exposed to the odor(s) during the first 4 to 12 hours for the bond to develop. The migration of **salmons** for thousands of miles is an almost incredible example of how memory of odors can control behavior. At 2 years of age, the salmon is imprinted by odors at its birthplace, and these odors guide it when returning “home” several years later. Imprinting by smell or taste can occur also before birth. If **pregnant rabbits** are fed a diet with certain aromatic substances, their offspring prefer foods containing these substances. This happens even if they grow up with a substitute mother on a different diet.

Olfactory imprinting is related to synaptic changes in the **olfactory bulb** (and most likely other places as well). Simultaneous arrival of olfactory signals and signals in **norepinephrine**-containing afferents seems to be necessary for lasting synaptic changes to occur. In the example with sheep mentioned above, more mitral cells responded to odors from the lamb after imprinting.

Pheromones

The term pheromone was introduced by Karlson and Lüscher in 1959 (p. 55), and defined as “. . . substances which are secreted to the outside by an individual and

received by a second individual of the same species, in which they release a specific reaction, for example, a definite behaviour or a developmental process.” Now the term is often used more broadly about all kinds of inborn, chemical communication between individuals of the same species. In animals—rodents have been most studied—several substances in body fluids (such as urine and saliva) have pheromone activity and influence a wide range of social interactions, such as sexual behavior, aggression, recognition of other individuals, and so forth. After destruction of the sense organ for pheromone recognition, rodents show, for example, reduced sexual activity and territorial defense. Pheromones may be volatile substances (small molecules) that move easily with the air, or they may be heavier molecules (peptides) that are exchanged among individuals only by close contact. In general, it seems that the volatile pheromones act as signals for sexual attraction or warning impending danger, whereas the peptide pheromones are of special importance for recognition among individuals.

The **vomer nasal organ** is a tubular structure in the nasal septum with an anterior opening. It is well developed in rodents, whereas it seems to disappear before birth in humans. It contains sensory cells like those in the olfactory epithelium, expressing a distinct class of receptor molecules (In higher primates, the genes coding for these receptors have become nonfunctional). In rodents, the sensory cells send their axons to the **accessory olfactory bulb** and from there connections to cortical structures that partly overlap those treating signals from the olfactory epithelium.

Even though the vomer nasal organ is lacking, there is evidence of pheromone-like actions in humans, where pheromones presumably act via receptors in the olfactory epithelium (in rodents pheromones act via both the vomer nasal organ and the olfactory epithelium). Synchronization of the menstrual cycle in women living close together is believed to be mediated by pheromones. Especially volatile substances present in armpit sweat seem to act as pheromones in humans. For example, some studies suggest that male armpit sweat may influence female menstrual cycle and mood. Mothers (but not fathers) seem to be able to recognize their babies by smell.

The significance of pheromones for human social behavior and development is controversial. It is safe to say, however, that they would play a minor role compared with their effects in rodents and other animals. In a broader context, pheromones are just one means of social communication. As concluded by Swaney and Keverne (2009, p. 239): “. . . the evolution of trichromacy [color vision] as well as huge increases in social complexity have minimised the role of pheromones in the lives of primates, leading to the total inactivation of the vomer nasal organ . . . while the brain increased

in size and the behavior became emancipated from hormonal regulation.”

GUSTATORY SYSTEM (THE SENSE OF TASTE)

The sense of taste is not among the most important of the special senses in humans, and much of what we usually call taste experience is in reality brought about by stimulation of **olfactory receptors**. This happens primarily by expiratory airflow through the nose while eating (compare the reduced sense of taste during a common cold). We nevertheless interpret this olfactory stimulation as taste (a further example of the importance of central interpretations of sensory signals for our conscious perception). In addition, signals from oral **thermoreceptors** and **mechanoreceptors** contribute to what we experience as a unitary sensory phenomenon. Finally, **nociceptors** activated by spicy food (such as chili peppers) contribute to taste perception.

The Taste Receptors and Taste Qualities

The true taste signals come from **chemoreceptors** in the taste buds that are located primarily in the epithelium of the tongue (Fig. 19.4). The taste buds are concentrated along the lateral margins of the tongue and the root of the tongue and are found in small elevations of the mucous membrane called **papillae**. The largest (vallate)

papilla lies along a transverse line posteriorly on the tongue (Fig. 19.5).² The **taste buds** are composed of approximately 100 elongated sensory cells and supporting cells (Fig. 19.4). The **sensory cells** of the taste buds are constantly renewed; each cell lives probably only about 10 days. They have long **microvilli** at their apical surface, protruding into a small opening in the epithelium, the **taste pore**. Here, substances that are dissolved in the saliva contact the membrane of the sensory cell. Terminal endings of sensory (afferent) axons contact the basal ends of the sensory cells. Binding of the tasty substances to receptors in the membrane of the sensory cells depolarizes the cell and thus produces a **receptor potential** (as in other sensory cells).³ **ATP** released from the basal aspect of the sensory cell binds to ionotropic **purinoceptors** (P_{2x}) in the sensory nerve endings and produces action potentials.

2 There are also some taste buds on the soft palate, which are innervated by the intermediate nerve. The few taste buds present on the most posterior part of the tongue and on the upper side of the epiglottis are innervated by the vagus nerve. Extralingual taste receptors are believed to protect the airways from aspiration of fluids.

3 There are two kinds of receptor cell in the taste buds. Unexpectedly, it seems that the kind that expresses taste receptors in high concentrations do not contact sensory fibers, whereas the other kind, expressing fewer taste receptors, makes direct contact. Therefore, several taste cells probably work together as a unit with some cells being responsible for transduction while others transmit signals to the sensory fibers. Indeed, the taste cells seem capable of mutual communication by way of gap junctions and release of neurotransmitters (ATP, serotonin, and others).

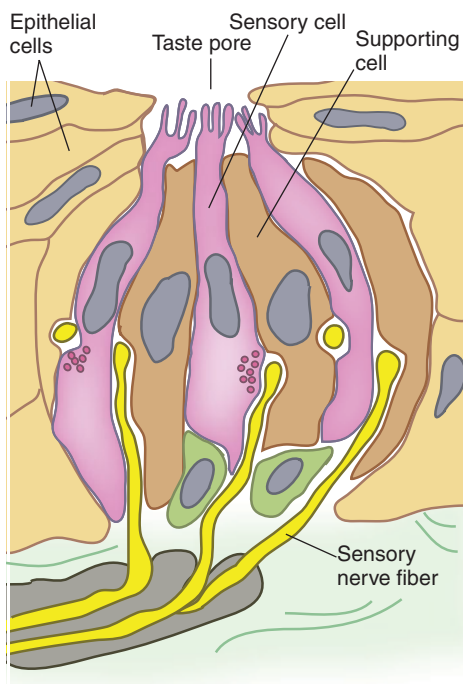


FIGURE 19.4 *Taste bud.* Semischematic drawing based on electron micrographs. The receptor molecules for taste substances sit in the membrane of the receptor cell cilia. In the taste pore, the receptors are exposed to substances dissolved in fluids.

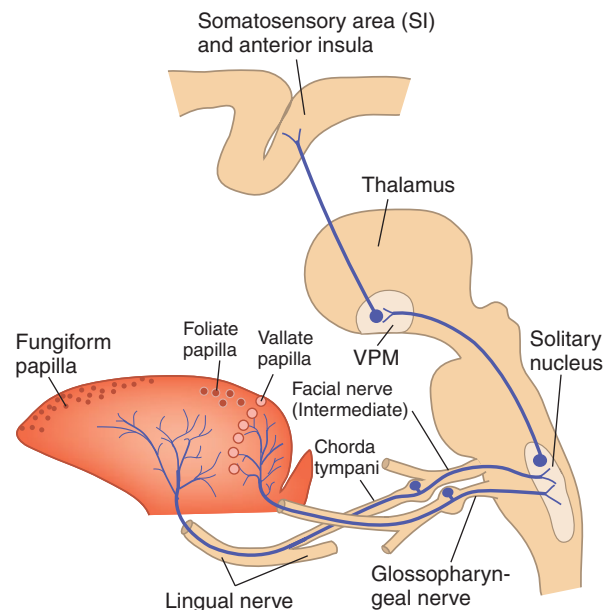


FIGURE 19.5 *Pathways for taste signals.* The various kinds of papilla contain taste buds. In the solitary nucleus, taste information is integrated with somatosensory signals from the oral cavity. The solitary nucleus receives addition input from the viscera via the vagus nerve (not shown in the figure).

We usually distinguish four elementary **taste qualities**: **salty**, **sour**, **sweet**, and **bitter**. In addition, **umami** taste is now often included as a fifth quality. The receptors evoking the umami quality detects **amino acids** and **L-glutamate** in particular. Indeed, the sodium salt of glutamate is widely used as a flavor enhancer. There are probably many more specific receptors than basic taste qualities. Especially bitter tasting substances appear to be detected by several specific receptors. This seems reasonable, as bitter taste often is associated with poisonous foods and their recognition would give special survival value.

Each of the basic taste qualities is most easily perceived (have the lowest threshold for identification) in a particular **region of the tongue**: sweet at the tip of the tongue, then salty, sour, and bitter more posteriorly, in that order. Such differences are only relative, however, and humans are able to perceive all qualities from either the anterior and posterior part of the tongue. Accordingly, receptors for all taste qualities have been identified in all parts of the tongue equipped with taste buds.

Fat is very important for the tastiness of foods. It therefore seemed a paradox that no specific receptors were known, even though the texture of fat—detected by low-threshold mechanoreceptors in the oral cavity—appears to play a role for taste perception. However, now binding of long-chain **fatty acids** to the apical membrane of taste cells have been demonstrated, and furthermore, we know that fatty acids act on taste-cell ion channels.

Flavors Act on Ion Channels and on G Protein–Coupled Receptors in the Apical Membrane

Salty and sour substances depolarize taste cells by direct binding to **ion channels**, whereas sweet and bitter substances and glutamate act mainly or only on **G protein–coupled receptors**. A number of genes code for sweet, bitter, and umami receptors.

No specific receptors are required for **salty** and **bitter** substances, because they act on ubiquitous H^+ and Na^+ ion channels. The specificity of the sensory cell depends on the fact that the cell is exposed to the stimulus only at the apical membrane. The cells of the taste bud are interconnected with **tight junctions** (zonula occludens) that seal the lateral membranes off from the taste pore. Thus, fluid in the mouth cannot penetrate between the cells. **Sour substances** appear to act primarily by virtue of their concentration of hydrogen ions, which act by closing K^+ channels in the apical membrane. This depolarizes the cell. Hydrogen ions, however, act on several kinds of channels, such as channels for Na^+ and Ca^{2+} ; one might think that closure of such channels would counteract the effect on K^+ channels. Specificity is achieved because the concentration of K^+ channels is

particularly high at the apical membrane, whereas the other kinds of channels are evenly distributed. Sour substances will therefore close many K^+ channels but only a few other channels. **Salty substances** act primarily on passive Na^+ channels in the apical membrane. When the concentration of Na^+ increases in the fluid in contact with the apical membrane, Na^+ ions enter the cell and depolarize it.

Sweet substances and **umami** act on members of the same family of G protein–coupled receptors (**T1R**), which also includes metabotropic glutamate receptors and $GABA_B$ receptors. One kind of the T1R receptor binds several different sweet substances, while another binds amino acids. In humans, this receptor is selective for glutamate (in rodents it binds several kinds of amino acids).

Bitter substances another G protein–coupled receptor family than sweet and umami (**T2R**). Whereas only three genes code for T1R, about 30 code for T2R. Each variety of T2R recognizes only a few bitter substances but each sensory cell expresses several kinds of the T2R. Thus, each cell can recognize a wide variety of bitter substances.

Why Cats Don't Like Sweets

Cats do not express the gene coding for one of the sweet receptors (T1R2). This may probably explain why cats show little interest in sweet foods: they probably get no taste perception from it. In an evolutionary perspective, sweet receptors are much more important for primates that feed on fruits and berries than for carnivores feeding primarily on meat. **Aspartame**, used as an artificial sweetener, binds in humans to the T1R2 receptor and therefore gives a sweet sensation. In mice, this receptor is slightly different and unable to bind aspartame. Consequently, mice are unable to detect aspartame in their food. Similar genetic **polymorphisms** may perhaps explain why humans differ concerning their propensity for sweets.

Modulation of Taste-Cell Sensitivity

Various factors, such as hormones and neurotransmitters, and neuropeptides modulate the sensitivity of the sensory cells. These substances can be released from basal cells in the taste buds or enter from the blood. The basal cells of the taste buds contain **serotonin** (among other substances). Although its function is not clear, serotonin increases the sensitivity of the sensory cells to taste stimuli. Depression and anxiety are commonly associated with reduced taste sensation. One study found reduced sensitivity for all taste qualities (but especially sweet) in deeply depressed patients. The sense of taste returns when the depression resolves. Low levels of monoamines in the taste buds may explain

these findings, as suggested also by the finding that **selective serotonin-reuptake inhibitors** (SSRIs) reduces the threshold for detecting flavors by 20% to 30% in normal persons.

Vasopressin (the antidiuretic hormone) and **aldosterone**—both protecting the blood volume by acting on ionic transport across the renal tubular epithelium—**increase the response of the taste cells to sodium chloride** (by acting on Na^+ channels).

Specificity of Taste Cells and Primary Afferent Units

Each sensory cell in the taste buds appears to respond rather specifically to one kind of taste substance (sweet, sour, bitter, or umami). The further transmission of signals in the primary afferent fibers is less specific but retains some segregation. Thus, primary taste afferents (sensory units) fall into two broad categories, “**specialists**” or “**generalists**,” depending on degree of specificity. The specialist sensory units respond best to stimulation with one category of substances, whereas the generalists respond to several with about the same threshold. This conclusion is based on recordings from single afferent fibers in several animal species. In agreement with the existence of specific primary afferents, stimulation of one kind of sensory unit (e.g., sweet specific) evokes behavior in agreement with the nature of the stimulus. The specific response to electric stimulation of the sensory units remains even though their properties are changed (by genetic manipulation) so that they no more express the appropriate receptor. Thus, the central connections of the sensory unit determine the behavioral response evoked upon stimulation of a sensory unit, even if its receptor expression has

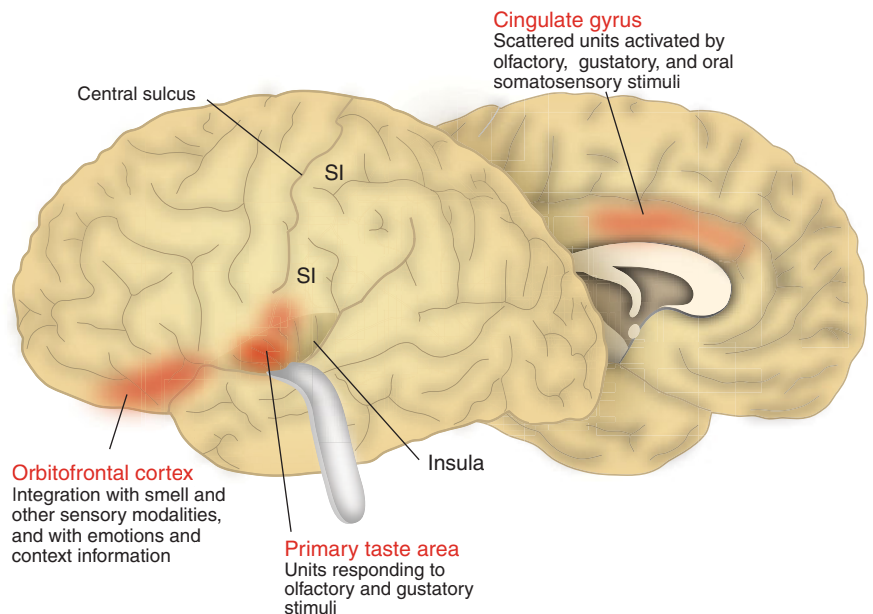
been changed from, for example, sweet to bitter. This implies the existence of mechanisms ensuring that, when the sensory cells are renewed, they retain their “line” to the brain. On the other hand, each sensory fiber must be able to recognize “its” kind of sensory cell, because each sensory fiber branches and supply several taste buds and several sensory cells in each taste bud.

Pathways for Taste Signals and the Primary Taste Area

The **intermediate nerve** (accompanying the **facial nerve**) contains the taste fibers from the anterior two-thirds of the tongue, whereas the fibers from the posterior third follow the **glossopharyngeal nerve** (Fig. 19.5). All taste fibers end in the rostral part of the **solitary nucleus**. The neurons in the solitary nucleus send their axons to several areas, thus mediating reflex effects, coordination of the activity of visceral organs, and perception of taste. Fibers pass to the visceral efferent nuclei of the **salivary glands** and to the dorsal motor nucleus of the **vagus**. Such connections mediate reflex secretion of saliva and gastric juice (and other digestive fluids). Some of the cells in the solitary nucleus send their axons to the **hypothalamus**. This enables taste signals to influence the higher autonomic centers. Taste information reaches the **amygdala**. Finally, direct fibers from the solitary nucleus end in the **thalamus**, enabling taste signals to reach the cerebral cortex. The synaptic interruption in the thalamus is in the **VPM nucleus** (Fig. 19.5; see also Fig. 14.6), which serves as a relay for signals from the face in general.

Taste signals are transmitted from the thalamus to the **face region** of the **SI**, close to the representation of

FIGURE 19.6 Cortical regions processing information from taste receptors. These regions may constitute a “flavor network” responsible for our experience and discrimination of flavors. The density of taste-responsive units is low in all regions. In contrast to primary visual and auditory cortices that contain only modality-specific units, the primary taste area contains neurons integrating several sensory modalities, notably olfactory and oral somatosensory stimuli.



the tongue (see Fig. 14.8) and the anterior part of the **insula**. Together, these areas comprise the **primary taste area** (Fig. 19.6). Sensory units responding to the elementary taste qualities are, as mentioned, to some degree segregated in the first link of the taste pathways. However, a considerable convergence occurs in the **solitary nucleus**. Most neurons respond to several taste qualities, although a few respond specifically to sweet or bitter substances. In the primary taste area, even fewer neurons appear to be specific. Further, many neurons receive convergent inputs of taste and other sensory modalities, so that integration of taste and, for example, olfaction starts at the first cortical station. For example, **chemotopic localization** in the primary taste area (topographically organized representation of different flavors) has not been documented in neurophysiologic studies of primates. Nevertheless, we have no problems discriminating the basic taste qualities and innumerable other nuances of taste. How can a high level of discrimination be achieved if there are no “**labeled lines**” keeping apart signals from different kinds of receptor (as in the somatosensory system)? One view is that discrimination occurs by simultaneous analysis of the activity pattern in many sensory units—so-called **population coding**. The temporal pattern of signals from different receptors is crucial, not their spatial segregation. Nevertheless, the “labeled line” view received support recently from evidence of chemotopic localization in the primary taste area of rodents. This was shown by transsynaptic transport to the cortex from bitter and sweet receptors in the taste buds.⁴ Possibly, there may be elements of both mechanisms—that is, labeled lines and population coding—in the central processing of gustatory signals. It is fair to say, however, that the mechanisms responsible for taste discrimination are not fully understood.

Signal Transmission from Taste Buds Is Modulated in the Brain Stem

The response of neurons in the solitary nucleus to signals from taste buds depends on several other inputs to the solitary nucleus. For example, the response is inhibited by distension of the **stomach**. This is mediated by signals transmitted from the stomach by the **vagus nerve**. Further, solitary neurons respond to alterations in blood levels of **insulin** and **glucose** (presumably by

way of descending fibers from the hypothalamus). The solitary nucleus receives descending fibers from the **amygdala**, probably mediating conditioned **aversion** to certain flavors. Finally, the cortical taste area sends fibers to the rostral part of the solitary nucleus, enabling that the signal traffic is modulated by **context** and **expectation** already at the brain-stem level.

Further Cortical Processing

Our **conscious experience** of taste is due not just to stimulation of taste receptors, as mentioned. Furthermore, numerous taste cells with different specificities are presumably always stimulated while eating or drinking. The synthesis in the cerebral cortex of all these varied signals forms the basis of our subjective experience of taste. Convergence of olfactory and taste signals takes place in the anterior part of the insula. A higher level of integration occurs in the **orbitofrontal cortex** where taste, smell, and other sensory modalities meet (Fig. 19.6). Thus, responses of neurons in the orbitofrontal cortex to flavors depend on whether the animal is hungry or not. This may be caused by connections from cortical areas related to **motivation**—that is, areas that may inform about the significance of a sensory stimulus. That our subjective sensory experience is determined not only by the stimulus holds for all sensory systems. Nevertheless, the emotional influence on the sensory experience is probably more marked for taste and smell than for other modalities. We know from everyday experience, for example, how the same smell or taste may be experienced as pleasant in one situation and nauseating in another.

Conditioned Taste Aversion

Taste is an important learning signal; it may be of vital importance to learn rapidly the association between a taste and its significance—that is, whether it signals something edible or poisonous. Survival of wild animals depends upon their ability to learn such association at the first trial, establishing a stable **conditioned taste aversion** to dangerous foods. Nausea is a very efficient learning signal when evoked by food intake. By just trying a small portion of the food, the animal usually can survive and learn to never again try the same potential food. The close coupling in the solitary nucleus of gustatory information with signals from the stomach most likely is involved in establishing the associations between flavors and their significance. Other neuronal groups believed to participate in conditioned taste aversion are found in the **parabrachial area** (pons), the **amygdala**, and the **orbitofrontal cortex**.

⁴ An functional magnetic resonance imaging (fMRI) study of six persons reported slightly different locations of cortical responses to different flavors. The interpretation of such data is uncertain, however, because the distribution of response may have been influenced by other factors than taste that varied during the experimental sessions.

IV

MOTOR SYSTEMS

THE cell groups and tracts in the central nervous system that control the activity of the skeletal muscles compose the **motor systems**. We may also use the term **somatic** motor systems to distinguish them from the systems that control smooth muscles and glands.

Although the motor systems consists of several interconnected parts, we first discuss some general aspects (Chapter 20). Then we treat the peripheral (**lower**) **motor neurons** (Chapter 21) and the **central (upper) motor neurons** (Chapter 22). These parts are directly

involved in mediating the commands from the motor centers to the muscles, and are necessary for the initiation of voluntary movements; paralysis ensues when they are damaged. Then we treat the **basal ganglia** (Chapter 23) and the **cerebellum** (Chapter 24), which have their main connections with the central motor nuclei and are necessary for the proper execution of movements rather than for their initiation. Finally, in Chapter 25 the **control of eye movements** is treated separately due to the distinctive features of this system.

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Motor Systems and Movements in General

OVERVIEW

This chapter discusses the close interdependence of what we, somewhat arbitrarily, term the motor system and other “systems” of the brain. For example, normal motor control and especially motor learning depend heavily on sensory information. Large parts of the cerebral cortex in the parietal and frontal lobes are engaged in transformation of sensory information into action. Further, the purpose of movements is often not movement per se but movements performed to collect sensory information. Without precise control of our fingers, lips, tongue, and eyes, sensory information would be severely degraded. This also concerns the chemical senses (olfaction and taste). For example, sniffing is a prerequisite for optimal detection of smells. We furthermore need to be able to scan our environments with our eyes, and our fingers move swiftly and accurately back and forth over objects we need to identify. We further discuss movements differing with regard to velocity and to whether they aim at stabilizing or moving parts of the body. Finally, we discuss how we can classify movements according to degree of voluntary control, from fully automatic reflex movements to the most consciously controlled skilled movements of the fingers.

MOTOR AND OTHER SYSTEMS ARE MUTUALLY DEPENDENT

The Term *Motor System* Lacks Precision

Although we include only restricted parts of the nervous system in the term motor system, nevertheless signals from parts of the brain not usually considered motor are necessary for proper movement. Thus, many parts of the central nervous system (CNS) may be considered motor in the sense that they contribute to the activity of the more narrowly defined motor systems. Some of these regions are usually considered part of the sensory systems, as they are crucial for the processing of sensory information (e.g., the posterior parietal association areas, discussed in Chapter 14, which process visual and somatosensory information but also issue commands to motor areas). Other parts of the brain,

which are usually classified as “cognitive,” are nevertheless indispensable for performance of purposeful movements (e.g., parts of the prefrontal cortex involved in the early stages of movement planning and in the mediation of motivated behavior). On the other hand, parts that we here include in the motor systems, notably the basal ganglia and the cerebellum, participate in nonmotor tasks. The basal ganglia, for example, play prominent roles in emotional and cognitive functions. Likewise, recent research, for example, functional magnetic resonance imaging (fMRI) studies in humans, strongly suggest that the cerebellum participates in sensory and cognitive functions.

We may **conclude** that, at the higher levels of the CNS, it becomes arbitrary whether we classify regions as sensory, motor, or cognitive: these concepts are too narrow to encompass the complex tasks undertaken by large parts of the cerebral cortex and many related subcortical nuclei.

Sensory Feedback

Although for didactic reasons we treat the motor and sensory systems as independent entities, this is an oversimplification. For the motor systems to function, they must cooperate closely with the sensory systems. For example, during most movements, the motor centers need constant information from receptors in muscles, around joints, and in the skin about whether the movement is progressing in accordance with the plan. Further, visual information is often crucial for the proper execution of movements. Such **sensory feedback** information enables the CNS to adjust and correct the command signals issued to the muscles either during the movement or during the next time the movement is performed. **Learning** of new skills especially depends critically on reliable sensory feedback. Indeed, we perform unfamiliar movements slowly to allow sufficient time for sensory feedback.

Movements in the Service of Sensation

Many movements—notably of the hands, mouth, and tongue—are executed to collect sensory information.

Indeed, the hand is a sensory organ in its own right. For example, we use delicate, exploratory finger movements to judge the form, surface, consistency, and so forth of objects. Losses of muscle coordination or of hand sensation have equally devastating consequences for hand function. In addition, eye movements are entirely devoted to assist the brain in the acquisition of visual information, and head movements aid the auditory system. As aptly formulated by the Israeli neuroscientist Ehud Ahissar (2008, p. 1370) “Without eye movement, the world becomes uniformly gray; without sniffing, only the initial changes in the odor environment are sensed; and without finger or whisker motion, objects cannot be identified.”

Motor Systems and Self-Recognition

Movements play an important role in the experience and ownership of our bodies. For example, experiments in human volunteers show that self-recognition (e.g., the hand belongs to me) is significantly better during a self-generated movement than when depending on proprioceptive and visual information alone. Our body scheme and body image (cf. Chapter 18) depend on regular updating by sensory information provided by our own, purposeful movements. If there is a mismatch between movement commands and sensory feedback, misconceptions of the body regularly occur (e.g., after amputation, deafferentation, immobilization of joints, and so forth).

CLASSIFICATION OF MOVEMENTS

Stabilizing and Moving

Before we describe the motor systems, some comments on movements in general are pertinent. First, muscle contraction may not necessarily elicit movement (i.e., alter the position of one or more joints); just as often, muscle activity is used to prevent movement—for example, muscles maintain our **posture** by counteracting the force of gravity. In preventing movement, the muscles may be said to **stabilize** a joint (against external forces) and to have a postural function. Further, movement in one part of the body—for example, in an arm—requires that muscles in other parts contract to prevent the body balance from being upset. Therefore, a muscle in one situation may be used as a mover and in another situation as a stabilizer.

Contractions: Concentric, Isometric, and Eccentric

Whether or not a movement is to occur depends on the magnitude of the force produced by the muscle contraction and the external forces acting on the joint. When the

external force is smaller than the muscle force, the muscle shortens and a movement occurs; this is called **concentric** or **isotonic contraction**. When the external forces equal the force of the muscle contraction, no movement occurs; this is called **isometric contraction**. When the external force is greater than the opposing force produced by the muscle contraction, the muscle lengthens; this is called **eccentric contraction**. Eccentric contraction occurs, for example, with the thigh muscles when we walk down a staircase, as the muscles brake the movement produced by the weight of the body.

Ramp and Ballistic Movements

Movements may be classified by the **speed** with which they are performed. **Ramp movements** are performed relatively slowly. The crucial point is that the movement is slow enough to enable sensory feedback information to influence the movement during its execution. **Ballistic movements** are very rapid, and their characteristic feature is that they are too fast to enable feedback control: the name derives from analogy with a bullet shot out of a gun.

Automatic and Voluntary Movements

Movements may also be classified according to whether they are **voluntary** or **automatic**; automatic movements take place without our conscious participation. In reality, this is much too crude a distinction: there is a gradual transition from what Hughlings Jackson in the past century termed the most automatic to the least automatic movements. The **most automatic** movements are basic, simple reflexes, such as the retraction of the arm from a noxious stimulus. Locomotion is an example of a semiautomatic movement—that is, the basic pattern is automatic, but starting and stopping and necessary adjustments may require conscious (voluntary) control. The **least automatic** movements are precision grips with the fingers and delicate manipulatory or exploratory movements such as writing, drawing, playing a musical instrument, and so forth. Equally precise voluntary control exists for the muscles of the larynx, the tongue, and some of the facial muscles. We know that the degree to which a movement is automatic changes with **learning**: in the beginning a new movement requires full voluntary control, and in the process of learning the movement becomes more automatic. When playing a well-rehearsed musical piece on the piano, for example, we do not need to pay attention to the fingers and their movements.

As a rule, the most automatic movements require only the use of relatively simple **reflex arcs** at the spinal level; participation of higher motor centers is not necessary. Somewhat less automatic and more complex movements such as ventilation, locomotion, and postural control

depend, in addition, on the participation of neuronal groups in the brain stem. Such movements do not require our attention directed toward them but can be subjected to voluntary control. The least automatic movements depend on the participation of the highest level—the **cerebral cortex**—to coordinate and control the activity of motor centers in the brain stem and spinal cord. The vast number of neurons and the plasticity of the human brain enable learning of an almost infinite

repertoire of voluntary movements. Also, these features ensure great **flexibility** in how motor tasks are solved. The same task may be solved in different ways, and we can continuously adapt to novel challenges. The great adaptability and flexibility of movements distinguish humans from most animals. Most animals are highly specialized for a limited number of motor tasks, controlled by stereotyped motor programs that develop according to a fixed pattern.

21 | The Peripheral Motor Neurons and Reflexes

OVERVIEW

The peripheral or lower motor neurons (motoneurons) constitute the final and only connection between the central nervous system (CNS) and the muscles. If they are destroyed, paralysis of the muscles ensue. There are two types of lower motor neurons— **α motoneurons** innervating the extrafusal muscle fibers and **γ motoneurons** supplying the muscle spindles. The motoneurons are arranged in **columns** in the spinal cord; each column supplying one or a few muscles with synergistic actions. Each column extends through two or more segments, so that each muscle receives fibers from at least two spinal segments. An α motoneuron and all the muscle fibers it innervates is called a **motor unit**. Large muscles consist of a few hundred to more than a thousand motor units. Muscle fibers are classified according to ATPase activity into **type 1** and **type 2** fibers. In general, type 1 fibers have the highest endurance whereas the type 2 fibers contract with the highest velocity and force. The muscle fibers of one motor unit are all of the same fiber type.

A **reflex** is an involuntary response to a stimulus, which is mediated by the nervous system. The motoneurons are parts of **reflex arcs**, consisting of receptors that capture the stimulus, sensory neurons conducting signals to the CNS, a reflex centrum (e.g., in the spinal cord), and an effector (muscle or glandular cells). In this chapter, we discuss the **flexion reflex** and **stretch reflexes** in particular. Stretch reflexes are of two main kinds: the **monosynaptic** stretch reflex (routinely tested by a tendon tap) and the polysynaptic, **long-latency** stretch reflex. Both consist of a muscle contraction in response to muscle stretch (the muscle spindle is receptor). We also discuss the **resting tone of muscles** (muscle tone), and how it may vary in health and disease. Finally, we treat the ability of peripheral nerves to **regenerate** after severance.

MOTONEURONS AND MUSCLES

The **peripheral motor neurons** are nerve cells that send their axons to skeletal muscles. Another term is **lower motor neurons**. These are the motoneurons in the ventral horn of the spinal cord and in the somatic motor

cranial nerve nuclei. There are two kinds: **α motoneurons** innervate the extrafusal muscle fibers, whereas the **γ motoneurons** innervate the intrafusal muscle fibers of the muscle spindle (see Figs. 13.6 and 13.7).

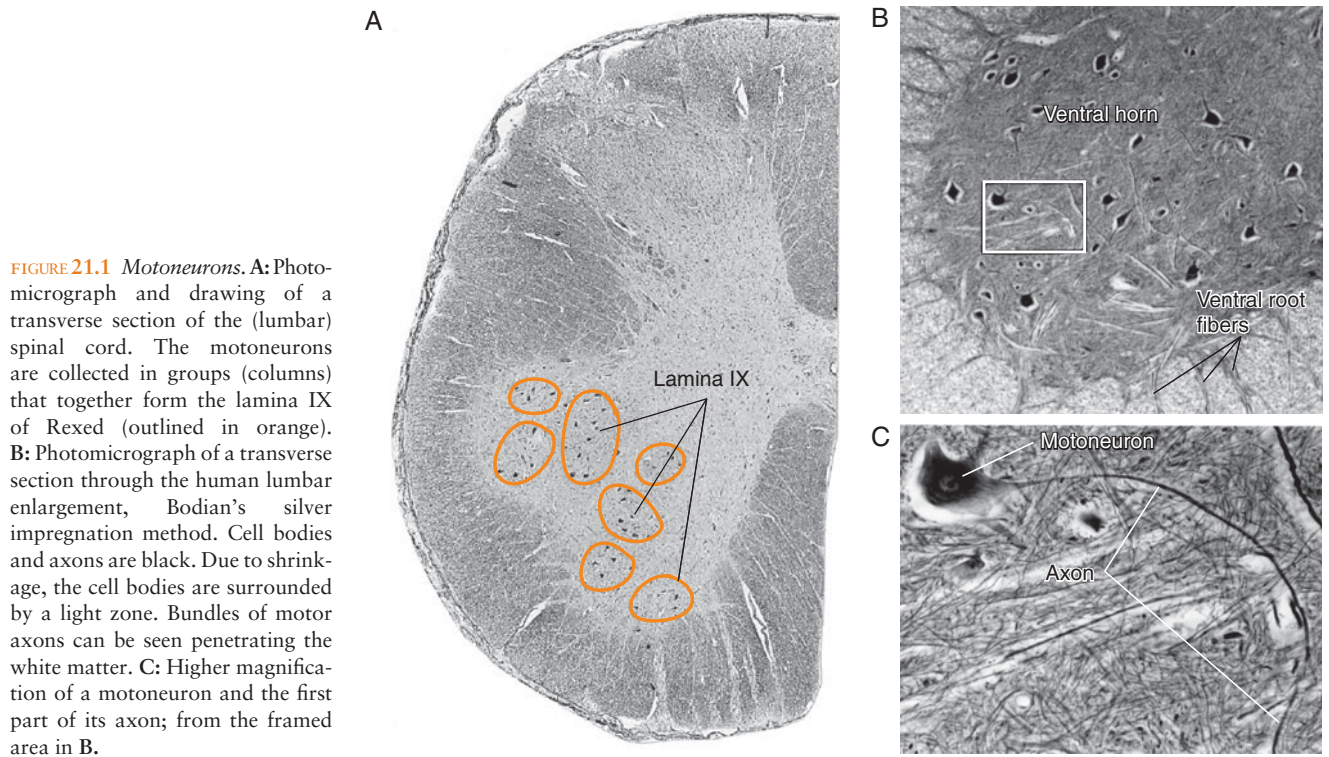
The motoneurons of the ventral horn (Fig. 21.1) and in the cranial nerve nuclei are easily recognized in microscopic sections because of their large size and the big clumps of rough endoplasmic reticulum (rER) in the cytoplasm of their cell bodies (see Fig. 1.2). The rich content of rER indicates that the neurons have a high protein synthesis. These proteins are, for example, enzymes for transmitter synthesis and metabolism and various kinds of membrane proteins. The vast surface of the motoneurons with their large dendritic tree and long axons presumably explains why motoneurons contain more rER than most other neuronal types.

Ventral Roots and Plexuses

The axons of the motoneurons leave the spinal cord through the **ventral roots** and continue into the **ventral and dorsal branches** (rami) of the spinal nerves to innervate skeletal muscles of the trunk and the extremities (see Figs. 6.5 and 6.7). Correspondingly, the axons from the **cranial nerve motor nuclei** supply the muscles of the tongue, pharynx, palate, larynx, and face, as well as the extraocular muscles. The axons of all the motoneurons located in one spinal segment leave the cord through one ventral root and continue into one spinal nerve. The ventral branches of the spinal nerves form **plexuses** so that the motor axons from one spinal segment are distributed to several peripheral nerves (Fig. 21.2).

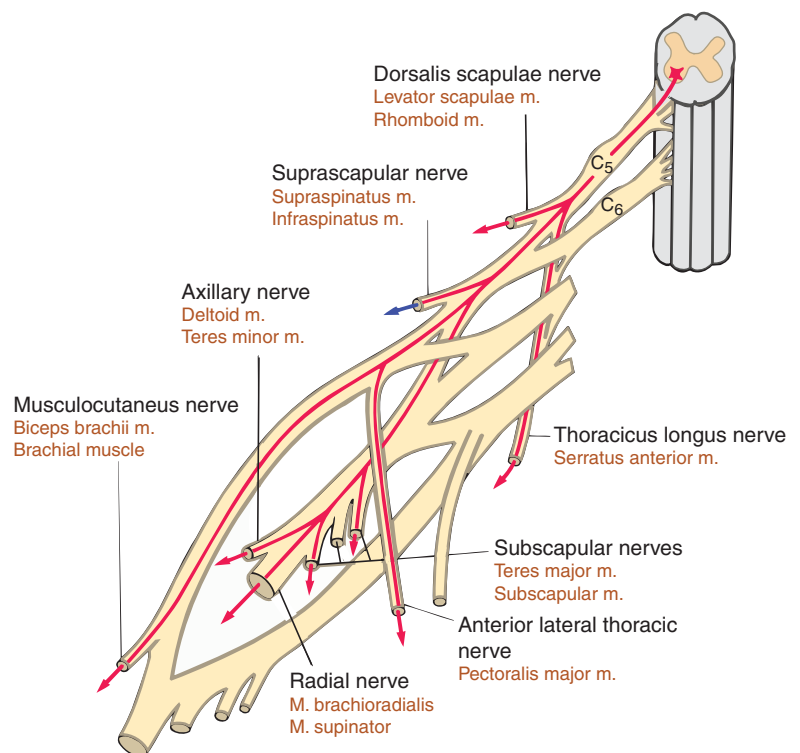
The Final Common Path and Synaptic Contacts on Motoneurons

Contraction of skeletal muscles can be elicited only by signals conducted in the axons of motoneurons. If these axons are interrupted, the muscles become **paralyzed**. The peripheral motor neurons thus constitute the **final common path** for all signals from the CNS to skeletal muscles (the term “final common path” was introduced by the British neurophysiologist and Nobel laureate



Sir Charles Sherrington). The motoneurons may be compared with the keys of a piano on which higher levels of the CNS can play. As we describe later in this chapter, many parts of the CNS cooperate in determining the activity of the motoneurons and thus the contraction of our muscles.

Each motoneuron probably receives about 30,000 nerve terminals—some forming excitatory synapses, others inhibitory; some with fast synaptic actions, others with slow modulatory ones. The sum of these influences determines whether and with what frequency the motoneurons will send action potentials to the muscles. That we can



perform such a wide variety of movements is due to the ability of the CNS to select precisely, by way of the motoneurons, the combinations of muscles to be used and to determine the speed and force with which they are to contract.

Neurotransmitters

The motoneurons use **acetylcholine** as transmitter, and the synthesizing enzyme **choline acetyltransferase** (ChAT) can be demonstrated immunohistochemically in the motoneurons and in their terminals. Motoneurons also contain the neuropeptide calcitonin gene-related peptide (**CGRP**). The level of CGRP in the motoneurons is under the influence of descending connections from higher levels of the CNS; when such connections are transected, the level of CGRP drops. There is experimental evidence that CGRP influences the synthesis of acetylcholine receptors of the muscle cells. If so, this may be one (of several) means by which the CNS can influence the properties of muscle cells.

Motoneurons Are Collected in Columns

The motoneurons are collected in groups, which form **Rexed's lamina IX** in the spinal cord (Fig. 21.1; see also Fig. 6.10). The dendrites of the motoneurons do not respect the boundaries of lamina IX, however, and extend far in the transverse and in the rostrocaudal directions—for example, into lamina VII, where many interneurons are located (see Fig. 6.12). The rostrocaudal (longitudinal) extension of the dendrites enables dorsal root fibers from several segments to act on each motoneuron. The dendritic tree increases the surface of the motoneurons enormously, and, not surprisingly, the vast majority of the nerve terminals contacting motoneurons are axodendritic.

Three-dimensionally, the motoneurons are collected in longitudinally oriented **columns** (Fig. 21.3). Each column contains the α and γ motoneurons to one muscle or a few functionally very similar (synergistic) muscles. Within a column supplying more than one muscle, the motoneurons to each muscle are at least partly segregated. As a rule, each column extends through more than one segment of the cord. Consequently, each muscle receives motor fibers through more than one ventral root and spinal nerve.¹ **Destruction of one root or spinal**

nerve only—for example, by disk protrusion in sciatica or by a tumor growing in the spinal canal—will not produce paralysis of a muscle but only a more or less pronounced paresis (weakness).

The anatomic organization of motoneurons has been studied via transection of muscle nerves in experimental animals. A retrograde reaction, which is easily seen through a microscope, occurs in the cell bodies of the motoneurons. Studies using retrograde transport of tracer substances have detailed the picture considerably. Study of patients with **poliomyelitis** has provided information about conditions in the human cord (the poliovirus infects and kills motoneurons). Because the distribution of paralyzed muscles usually has been determined before death, it can be compared with the distribution of cell loss among the motoneuron groups in the cord and brain stem.

Motoneuron Columns Are Somatotopically Distributed in the Ventral Horn

Groups of motoneurons that supply **axial muscles**—that is, muscles of the back, neck, abdomen, and pelvis—are located most medially within the ventral horn, whereas motoneurons supplying muscles of the **extremities** lie

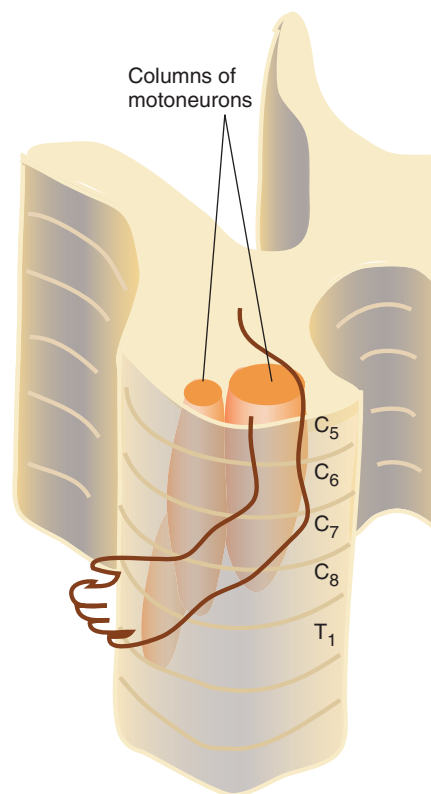


FIGURE 21.3 Columnar arrangement of motoneurons. Schematic of the somatotopic localization of the motoneuronal columns innervating the arm, located in the spinal segments C₅–T₁. Motoneurons supplying the intrinsic muscles of the hand are located most caudally (C₈–T₁) and most dorsally in the ventral horn.

1 About 20 *Hox* genes control the specification of motoneurons with regard to their peripheral target (muscle). In addition, expression of recognition molecules and guidance receptors in the peripheral tissues is required for axons to find their target. When looking at the intricate trajectory of branches from motoneurons in one segment (Fig. 21.2) the task of creating a precise topographic relationship between spinal cord motoneurons and muscles seems daunting. It should be recalled, however, that at the time of outgrowth in early embryonic life, the distances are small and trajectories of axonal growth are usually fairly straight. The complicated structure of the plexuses (Fig. 21.2) arises later in development.

more laterally. This explains why the ventral horn is broader (extends more laterally) in the segments of the cord that send fibers to the extremities (i.e., C_5 – T_1 and L_1 – S_2 ; compare Figs. 6.8, 6.10, and 6.11). There is also a further somatotopic organization: motoneurons supplying **proximal muscles** of the extremities (the shoulder and hip) are located more ventrally than those supplying the **distal muscles** (the hand and foot). This is shown in Figure 21.3, which also shows that the proximal muscles are supplied from motoneurons located more rostrally than those supplying the distal muscles. For example, the shoulder muscles are mainly innervated by the upper parts of the brachial plexus (C_5 – C_6), whereas the lowermost segments (C_8 – T_1) innervate the intrinsic muscles of the hand.

Motoneurons Are of Functionally Different Kinds

As mentioned, the α and γ motoneurons supplying one muscle lie together within one column in the ventral horn. In a microscopic section of the cord it can be seen that the motoneuron cell bodies vary in size (Fig. 21.1; see also Fig. 1.2). As in other parts of the nervous system, the neurons with the largest cell bodies have the thickest (and thus the fastest conducting) axons. The γ motoneurons are the smallest within a group, while the α motoneurons—although larger than the γ motoneurons—vary considerably in size. Such size differences are related to differences among the muscle cells supplied by the α motoneurons. Briefly stated, the smallest α motoneurons control delicate movements with little force, whereas the largest motoneurons come into play only when a movement requires great force. The large α motoneurons also have a much higher maximal **firing frequency** than the small ones, and the large ones tend to fire in brief bursts with a high frequency, whereas the small α motoneurons tend to go on firing for a long time with a low frequency. These differences in **firing pattern** reflect that the large motoneurons are used for forceful, rapid movements of short duration, whereas the small motoneurons can uphold a moderate muscular tension for a long time. For these reasons, we apply the term **phasic α motoneurons** to the large ones, and **tonic α motoneurons** to the small ones. The properties of the α motoneurons are discussed further when we deal with the motor units.

The Motor End Plate and Neuromuscular Transmission

After entering the muscle, the α -motoneuron axon divides into many thin branches or collaterals. Each of these terminal branches contacts one muscle cell only. Each muscle cell is thus contacted by only one branch from one α motoneuron. Such a branch ends on the muscle cell approximately midway between its ends, forming the **motor end plate**, where the signal transfer

from nerve to muscle takes place (Figs. 21.4 and 21.5). Within the end-plate region, the axonal branch divides further and forms up to about 50 nerve terminals (boutons), each establishing synaptic contact with the muscle cell. The postsynaptic side at this **neuromuscular junction** is somewhat special compared with synapses in the CNS, as **junctional folds** and a thin **basal lamina** are intercalated between the presynaptic and postsynaptic membranes (Fig. 21.5B). The boutons contain **acetylcholine**, and the postsynaptic membrane contains **acetylcholine receptors** of the **nicotinic** type. The density of acetylcholine receptors is much higher in the end-plate region than elsewhere on the muscle cell surface, which is appropriate because only at the end plate is the muscle cell normally exposed to acetylcholine. During embryonic **development**, however—before the nerve fibers growing out from the cord have reached the muscle cells—the acetylcholine receptors are evenly distributed all over the muscle membrane. Only after establishment of properly functioning synaptic contacts are the receptors redistributed to attain the mature pattern. During development, spinal motoneurons transiently express the neuropeptide **galanin**, which presumably influences the synapse formation at the motor end plate.

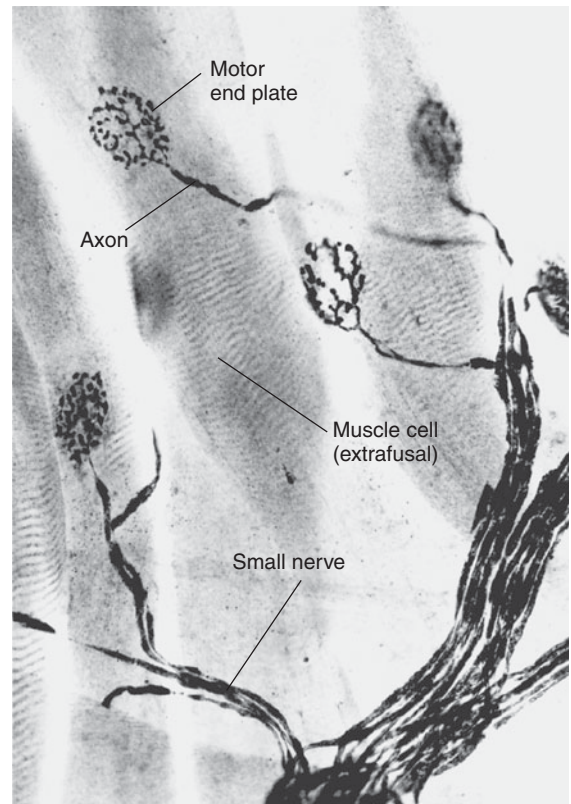


FIGURE 21.4 Motor end plates. Photomicrograph of skeletal muscle cells and a small bundle of nerve fibers innervating the cells. The tissue is stained with gold chloride to darken the nerve fibers and their terminal boutons. Each motor end plate consists of numerous small boutons. Four end plates are seen here. Magnification, $\times 500$.

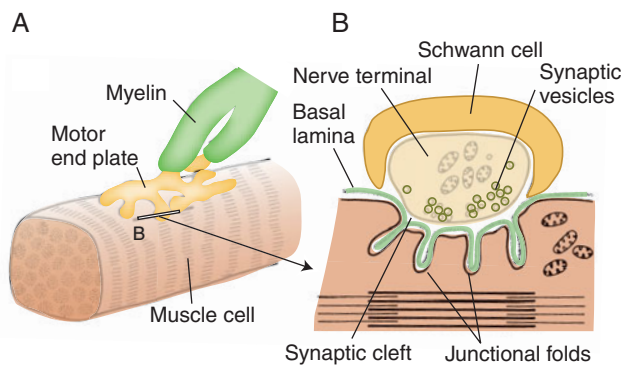


FIGURE 21.5 *The motor end plate.* **A:** Schematic showing how the myelinated nerve fiber loses the myelin sheath before it ramifies in the end-plate area, each terminal branch ending in a nerve terminal. Each muscle cell has only one end plate. **B:** Section through one of the nerve terminals in **A**, based on electron microscopic observations. The synaptic cleft contains a thin basal lamina, and the postsynaptic membrane is thrown into deep folds (junctional folds).

An action potential propagated along the axon of the motoneuron depolarizes all the boutons and elicits release of acetylcholine. The transmitter binds to the acetylcholine receptors, and, as at other excitatory synapses, this depolarizes the postsynaptic membrane. This change in the membrane potential is called the **end-plate potential**. Because of the large number of nerve terminals formed by one terminal fiber, enough transmitter is released by a single nerve impulse to depolarize the muscle cell membrane to the threshold for an action potential. The action potential is propagated over the whole surface of the muscle cell and elicits a brief contraction. The enzyme **acetylcholinesterase** rapidly terminates the action of acetylcholine by degrading it. The enzyme is present in the synaptic cleft and the junctional folds.

Neuromuscular Transmission Can Be Disturbed by Poison and Disease

Various drugs and naturally occurring poisonous substances can influence the signal transmission at the neuromuscular junction and produce involuntary muscle contractions or muscle paralysis. The South American Indian poison **curare** and similar synthetic substances paralyze the muscle cells by blocking the acetylcholine receptors. Such drugs are often used during abdominal surgery to obtain sufficient muscle relaxation. Several kinds of **snake poisons** act by blocking acetylcholine receptors and thereby paralyze the victim. The **botulinum toxin** (produced by a microorganism growing in certain kinds of spoiled food) paralyzes the muscles by preventing the release of acetylcholine from the nerve terminals at the motor end plate. A similar mechanism sometimes produces muscle weakness in patients with **cancer**; apparently, substances preventing release of acetylcholine are produced in the body.

The disease **myasthenia gravis** is characterized by excessive fatigability of striated muscles, which is caused by **autoantibodies** binding to the acetylcholine receptors. Thus, there are fewer than normal acetylcholine receptors available at the neuromuscular junction and release of acetylcholine opens fewer ion channels, leading to less than normal depolarization. The probability of evoking an action potential in the muscle cell membrane is consequently reduced. Drugs that inhibit the acetylcholine esterase (**neostigmine**, **physostigmine**) may lessen the symptoms. With this inhibition, the transmitter gets a longer time to act, and the probability of evoking an action potential is increased. Typically, most severely affected are the muscles of the head, producing symptoms such as double vision (due to paresis of extraocular muscles) and involuntary lowering of the upper eyelids (ptosis). The voice becomes weaker while speaking and swallowing may become difficult. In addition, the respiratory muscles are usually affected.

The Force of Muscle Contraction Is Controlled by the Motoneurons

A single presynaptic action potential at the motor end plate elicits only a brief contraction, a **twitch**, of the muscle cell (Fig. 21.6). The twitch lasts only for about one-tenth of a second. However, if another action potential follows shortly after the first one—that is, before the tension produced by the first twitch is over—the tension produced by the muscle is upheld and, furthermore, may increase considerably. This is called **summation**. Up to a limit, the tension produced by the muscle cell increases with increasing frequency of action potentials; that is, the **force** produced by the muscle cell is determined by the **firing frequency** of the motoneuron. Whereas the twitch is the response of the muscle cell to a single nerve signal, **tetanic** contraction is the term used of the muscle response to a train of signals with the highest frequency to which the muscle cell can respond (Fig. 21.6). The tetanic tension is thus the maximal force the muscle cell can produce. (One may, as shown in Fig. 21.6, differentiate between unfused or incomplete tetanus at submaximal firing frequencies and fused or complete tetanus at the maximal firing frequency. The term “tetanus,” as used here, refers to the complete tetanus.)

There Are Functionally Different Kinds of Striated Muscle Fiber Types

In both animals and humans, the skeletal muscles are composed of different kinds of muscle cells or muscle fibers (these terms are used interchangeably). The most clear-cut evidence is provided by the fact that in some species certain muscles have a dark color (“**red**” muscles), whereas other muscles are light (“**white**” muscles)—for

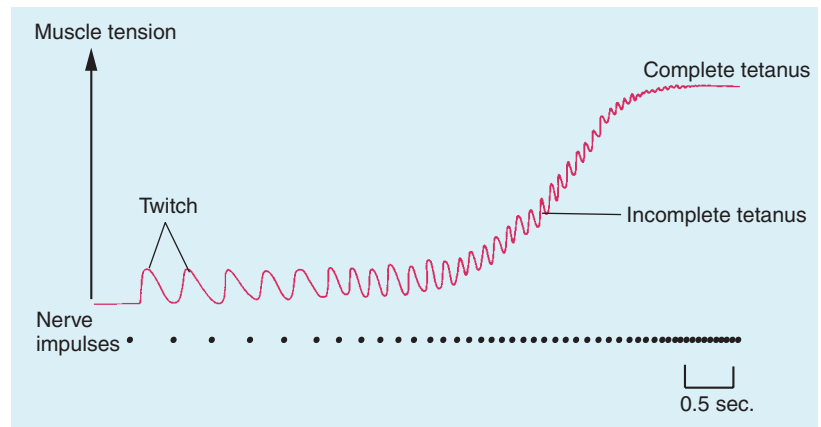


FIGURE 21.6 *Muscle contraction.* The muscle tension increases with increasing firing frequency of the motoneurons innervating the muscle cell. (Redrawn from Kandel and Schwartz 1985.)

example, the almost white breast muscles and the dark leg muscles of the chicken. Such muscles are composed of muscle fibers of only one (or predominantly one) kind, and we classify the muscle cells as either white or red. The color difference is due to differences in the content of **myoglobin**, which is red (closely related to hemoglobin) and transports oxygen within the muscle cell. Further study showed that white and red muscle cells differ with regard to **endurance**—that is, how long they can maintain tension. This is mainly due to differences in the capacity to take up oxygen and to **aerobic ATP production** (oxidative phosphorylation). As one would expect, the red muscles have the highest endurance. In addition to containing more myoglobin than the white muscle fibers (cells), the red ones also contain more mitochondria, which are responsible for the aerobic ATP production. There are also other important differences: the white muscle fibers contract more rapidly and develop greater force than the red fibers. With physiological methods, muscle cells can be classified as **fast twitch** (FT), corresponding largely to white fibers, and **slow twitch** (ST), corresponding to the red fibers.

The differences with regard to **contraction velocity** and maximal force development are related to differences in the amount and type of **myosin ATPase** (enzymes cleaving ATP, thus providing the energy for the muscle contraction). With histochemical staining methods, muscle fibers are classified in accordance with their ATPase activity (Fig. 21.7). On the basis of ATPase staining, human muscle fibers are classified as **type 1**, corresponding largely to the red and the ST fibers mentioned above, and **type 2** fibers, corresponding largely to the white and FT fibers. The type 2 group is heterogeneous, however, and consists of **type 2A** fibers, which resemble the type 1 fibers in having a relatively high oxidative capacity, and the **type 2B** fibers, which are the most typical white fibers with low oxidative capacity.

The CNS thus can **select** muscle fibers in accordance with the requirement of the **task**: one fiber type is best suited for contractions of moderate force that last for a

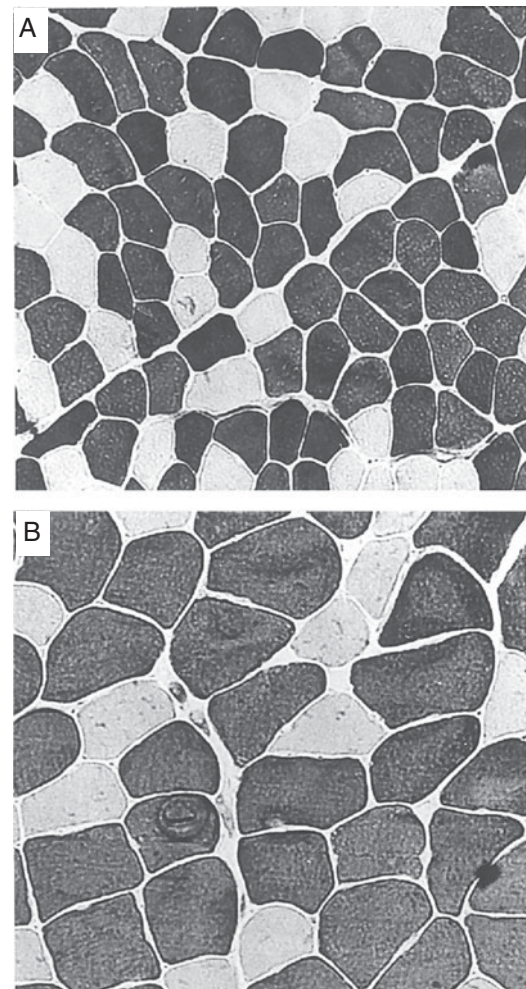


FIGURE 21.7 *Muscle fiber types.* Photomicrographs of cross sections of skeletal muscle (human). The sections are treated so that the color intensity of the muscle fibers depends on their myosin-ATPase activity. With this particular treatment, the type 1 fibers are light and the type 2 fibers are dark. **A:** From the quadriceps muscle of a “normal” person. **B:** From the same muscle of a weight lifter. Note the difference in muscle fiber thickness. Magnification, $\times 200$. (Courtesy of Dr. H.A. Dahl, Norwegian University of Physical Education and Sport, Oslo.)

long time, whereas the other type is used particularly for contractions with high force but short duration.

Muscle Fiber Types in Humans

With few exceptions, human skeletal muscles are composed of a mixture of type 1 and type 2 muscle fibers. However, the thickness difference between type 1 and type 2 fibers is much less marked than in experimental animals. Type 2 fibers are as a rule only slightly thicker than type 2 in males, while in females the reverse situation appears to be the rule.

Histochemical examination of many different muscles in several individuals has indicated that the **composition** of fiber types is fairly constant when comparing muscles from one individual, whereas the **individual differences** are great. Thus, some persons have a high percentage of type 1 in most of their muscles, whereas others have a strong preponderance of type 2 fibers. Studies of fiber composition in successful **athletes** have shown that those engaged in endurance sports (such as marathons and cross-country skiing) usually have a high percentage of type 1 fibers, whereas a high percentage of type 2 fibers is found in those engaged in sports requiring explosive force (such as ice hockey and weight lifting).

Recent studies suggest that the relationship between histochemical classification (ATPase) and functional properties is less clear-cut than formerly believed. Thus, muscle fibers belonging to the same fiber type histochemically can nevertheless have quite different contractile properties in different muscles. For example, the human adductor pollicis and soleus muscles both consist of about 80% type 1 fibers; yet the contraction velocity of is about the double in adductor pollicis.

Muscles Also Contain Connective Tissue: Stretching of Muscles

A muscle consists of a large number of muscle cells, stretching from tendon to tendon (usually at an angle to the longitudinal axis of the muscle). Bundles of muscle fibers are surrounded by connective tissue that contains the vessels and nerves. The whole muscle is wrapped in a connective tissue sheath, the muscle **fascia**. In some muscles the fascia is thick and fairly tight; in others it is thin and loose. The connective tissue of the fascia and within the muscle is continuous with the tendons. Therefore, the passive properties of a muscle—that is, its consistency and resistance to being stretched—depend on both properties of the muscle cells and on the amount and arrangement of the connective tissue.

Passive **stretching** of muscles to maintain or increase joint mobility is used therapeutically and in sports. The reasons for increased mobility after regular stretching are not entirely clear. One study, with 3 weeks of intensive

stretching, found no change of the viscoelastic properties of muscle and tendon in spite of increased range of motion. The muscle tension was higher in extreme joint positions, however, suggesting that the increased range is due to increased tolerance to high tensile forces rather than to changes in the muscle.

Motor Units

As mentioned, the axon of the motoneuron divides into numerous branches when entering the muscle, and each terminal branch makes synaptic contact with one muscle cell. Further, each muscle cell is innervated by one motoneuron only. When an α motoneuron sends action potentials to a muscle, all muscle cells innervated by that motoneuron contract simultaneously. Therefore, in a sense, an α motoneuron and all the muscle cells that it innervates constitute the smallest functional unit of the motor system and were called a **motor unit** by Sherrington (Fig. 21.8). (Compare with the term “sensory unit,” introduced in Chapter 12.) The **size** of a motor unit—that is, the number of muscle cells supplied by one motoneuron—varies greatly. This has been studied by counting the axons in the muscle nerve (after destruction of the dorsal roots, to let the sensory axons degenerate) and counting the muscle cells. An average seems to be around 150 muscle cells per motoneuron, with a range of from less than 10 to more than 1000. As might be expected, the **smallest motor units** are found in muscles that are used for delicate movements, which we must be able to control very precisely. Examples are the intrinsic muscles of the hand, the muscles of the larynx, the facial muscles, and the extraocular muscles. The **largest motor units** occur in large muscles used for movements of considerable force and with less precise control, such as the muscles of the back, the abdomen, and the thigh. Within one muscle, however, the motor units also vary in size.

There is also a relationship between the **size of a motor unit** (in terms of number of muscle fibers) and the size of its **motoneuron**. Thus, the motoneurons with the largest cell bodies and the thickest axons as a rule belong to the largest motor units. This fits with the use of large motor units for fast and forceful movements: the large motoneurons have the highest maximal firing frequencies, and their axons conduct the signals to the muscles with a minimal delay.²

Muscle cells belonging to different motor units lie **intermingled** in the muscle, as is obvious in sections stained to identify fiber types (Fig. 21.7). Correspondingly, muscle

2 The relationship between motoneuron size and motor unit properties is not absolute, however. Motoneurons of the same size may have different maximal firing frequencies. Further, motoneurons of the same size may innervate muscle cells with different contractile properties. The latter is probably because the motoneuron firing frequency (and firing pattern) influences the contractile properties of the muscle cells.

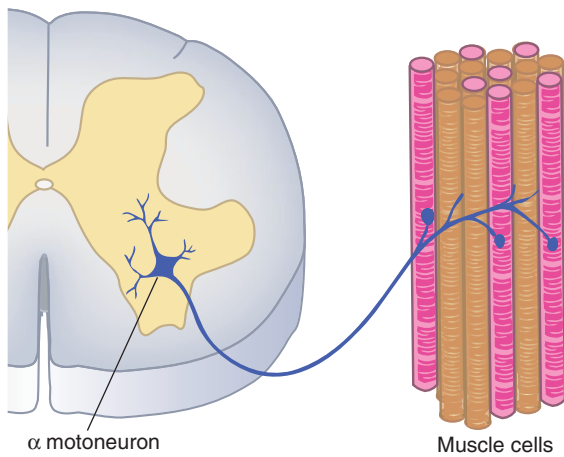


FIGURE 21.8 A motor unit. Schematic of a motoneuron in the ventral horn and its axon, which branches to supply many muscle fibers. The muscle cells belonging to one unit (red) lie intermingled with muscle cells belonging to other motor units (brown).

cells belonging to one motor unit spread over a considerable part of the total cross-sectional area of the muscle.

Motor Units and Fiber Types

Muscle cells belonging to the same motor unit are all of the same **fiber type**. Thus, they not only contract simultaneously but also share properties with regard to **contraction velocity**, **maximal force**, and **endurance**. In general, the smallest motor units consist of **type 1** fibers, whereas the largest ones consist of **type 2B** fibers. This further increases the difference between large and small motor units with regard to their maximal force; not only are there more muscle cells in the large units, but each cell also develops more force. Recruitment of one extra motor unit with type 1 fibers adds just a little extra to the total tension of the muscle, which thus can be graded finely (like an electrical switch with many small steps to vary the heat of a stove). As one might expect, such motor units are used for precisely controlled movements of small force. Recruitment of a large motor unit consisting of type 2 fibers, in contrast, gives a comparatively large increase in the total tension of the muscle (the switch has large steps). Such motor units are recruited only when large force production is needed. Often this concerns fast movements because high acceleration requires a large force.³

³ The fiber type used for a particular kind of movement has been studied by the so-called **glycogen-depletion technique**. By prolonged use, all of the muscle fibers of a motor unit deplete their stores of glycogen, as evident in histochemically treated frozen sections of a small piece of muscle tissue (obtained by biopsy). The type 1 muscle fibers are depleted first when the force exerted is low—that is, no type 2 fibers are recruited. When the force is very high, however, the type 2 fibers are depleted first; although type 1 fibers are also recruited, because of their high endurance, they are not depleted during the short period in which a maximal force can be maintained.

Electromyography

As mentioned, muscle contraction is elicited by an action potential that is propagated along the muscle cell membrane. Like action potentials of nerve cells, the muscle cell action potential can be recorded with an electrode. This is called **electromyography (EMG)** and is performed either with an electrode placed on the skin overlying the muscle—**surface EMG**—or with a thin needle electrode inserted into the muscle. The surface method gives an impression of the total electrical activity of the muscle, whereas needle electrodes sample the activity of a small part of the muscle. Because the EMG is a measure of the electrical activity and not of the mechanical activity of the muscle, it is not well suited to measure muscle-force production. In some instances of prolonged activity, the muscle force may be declining in spite of constant or even increasing EMG activity—the declining force being caused by changed conditions in the muscle itself—whereas muscle action potentials are evoked normally by the nerve to the muscle.

When recording from a normal muscle at the start of a very weak, voluntary contraction, the EMG shows regular, single potentials due, presumably, to the activity of only one motor unit. As the force increases, more potentials occur in the EMG, reflecting the recruitment of more motor units. At a certain level of force, the potentials are so frequent that the picture becomes unclear—that is, it is impossible to recognize a further increase of the EMG activity. This occurs not only because of recruitment but also because of the increase in the frequency of action potentials for each motor unit.

EMG is a valuable tool for studying the participation of various muscles in normal movements: for example, to determine the timing of contraction in muscles of the leg during walking. EMG also aids in the diagnosis of diseases of the peripheral motor neurons and of the muscles themselves, as in determining whether a disease affects the motoneurons or the muscle. If the motoneurons are put out of action, the EMG activity will be reduced or absent, whereas in a disease that affects the contractile apparatus of the muscle, the EMG may be normal. In addition, in cases of injury to peripheral nerves, it can be ascertained whether the lesion is complete (no EMG activity) or incomplete (even though there may be no visible movements).

Gradation of Muscle Force: Recruitment and Frequency

As we all know, the force of muscular contraction can be finely graded within very wide limits, from a barely perceptible contraction to a tension that is high enough to tear the muscle loose from its insertion. This depends on the ability of the CNS to control the activity (the level of excitation) of the motoneurons. There are two means by which the force of muscle contraction can be

gradually increased. One means is to bring more and more α motoneurons to send action potentials to the muscle. This is called **recruitment**, because more and more motor units—and thus muscle fibers—are called into action. The other means is by increasing the **firing frequency** of the motoneurons already recruited and thereby increasing the force developed by each motor unit. Thus, the force exerted by each motor unit (as by each muscle cell) can be graded from the small and brief tension produced by a single twitch to full **tetanic tension** (Fig. 21.6).

When a muscle contraction is initiated, the small motoneurons are always recruited first, and with increasing force the larger ones are recruited successively. This is called the **size principle of recruitment** and was demonstrated by the American neurophysiologist E. Henneman in the 1960s. The excitability of the motoneurons is inversely related to their size; thus, the small motoneurons are more easily excited to the threshold for initiation of action potentials than are the large ones. This is presumably related to differences both in membrane properties of the motoneurons and in the density of excitatory synapses. The size principle ensures selection of the motor units that are best suited for a particular kind of movement.⁴

Recruitment during Movements with Different Force Requirements

The number of motor units that are recruited at the beginning of a voluntary contraction depends on the force requirement. If **maximal force** is needed (as with a movement of maximal speed), most or all motor units are recruited almost simultaneously. Nevertheless, the small ones are activated slightly before the large ones because the small motoneurons reach the threshold for eliciting action potentials first. Thus, the order of recruitment is maintained (according to the size principle) when maximal force is required. To produce the maximal force, besides recruiting as many motor units as possible, the central motor centers must increase the firing frequencies of the motoneurons to the maximal: a muscle develops maximal force when all motor units are recruited and contract tetanically (complete tetanus). Such a contraction, however, can be maintained for only a short period before the muscle fatigues. The situation is different when a movement requires **low force**. Consider, for example, an isometric contraction that should last as long as possible (i.e., the force must be kept constant). Here, more than half the motor units are

recruited at the beginning of the contraction (judging from animal experiments).⁵ Because a relatively large fraction of the motor units is recruited, the required force is achieved with a low firing frequency, so that the motor units can work for a long time without fatigue. If the contraction continues, however, some motor units fatigue and “fresh” ones are recruited to maintain constant force. Finally, so many small or medium large motor units are fatigued that the largest motor units must be recruited to prevent a drop of force. This recruitment pattern is only valid, however, when the contraction force is so low that the blood flow is not impeded (even at 25% of maximal voluntary contraction force the blood flow is significantly reduced).

REFLEXES

When a response to a stimulus is automatic (involuntary), and the response is mediated by the nervous system, we call it a “reflex.” Many of the tasks of the nervous system are carried out automatically or reflexly—that is, independent of conscious interaction. This, of course, frees the higher levels of the brain from handling numerous trivial everyday tasks. The distinction between reflex movements and those initiated voluntarily are not always as clear-cut as it might seem from this definition, however. We discussed in Chapter 20 that movements are best classified on a continuous scale from the most automatic to the least automatic. In the text that follows we discuss mainly some spinal reflexes that are examples of the most automatic movements. Chapters 18 and 22 discuss somewhat less automatic movements, such as postural reflexes and locomotion. Reflexes involving cranial nerve nuclei are discussed in Chapter 27, and some visceral reflexes are treated in Chapter 29.

Even though reflexes differ in many respects, they nevertheless share some **fundamental properties**. Reflexes are stereotyped and constant because the same stimulus always gives the same kind of response. With increasing strength of the stimulus, however, the response usually increases in strength or magnitude. The reflexes are **inborn**: we do not need to learn them from experience. (One example that illustrates the need for inborn motor behavior is the sucking reflex in the newborn child.) Many reflexes occur in large groups of animal species; all mammals, for example, have several reflexes in common. In general, the reflexes are appropriate and useful and ensure that the individual adapts to the environment. Reflexes are also fundamental for reproduction.

⁴ In spite of the general validity of the size principle for recruitment of motoneurons, experiments with the use of biofeedback (EMG) during voluntary movements indicate that humans can to some degree select among the low-threshold motor units, probably depending on the ability of higher motor centers to focus excitation among motoneurons with presumably the same size.

⁵ There are, however, large differences among human muscles with regard to the proportion of motor units that are recruited early. In intrinsic hand muscles, for example, most of the units are recruited even at low forces, whereas in the biceps brachii new units are recruited until maximal force.

Some reflexes are simple with regard to both the stimulus and the response, like the blink reflex (closing of the eye when something touches the cornea). Others are much more complex and require cooperation of many structures, like the swallowing reflex. Some reflexes involve lower parts of the CNS only (spinal cord and brain stem), whereas others involve the higher parts (even the cerebral cortex). Some reflexes are mediated by a chain of only two or three neurons, others by complicated and extensive neuronal networks.

Even though reflexes are independent of our will, some of them can be **suppressed voluntarily** (e.g., the reflexes for emptying the bladder and the rectum). Other reflexes take place without our being aware of them, and with no possibility of influencing them voluntarily (like most of the reflexes related to the control of visceral functions).

Conditioned Responses (Reflexes)

True reflexes should not be confused with another kind of automatic behavior, **conditioned** reflexes or, better, conditioned responses. In conditioned responses the stimulus evoking a true reflex response (the unconditioned stimulus) has been replaced—by learning—by another stimulus. This learning occurs when the unconditioned stimulus (e.g., a puff of air on the cornea which elicits a blink reflex) is regularly preceded by another kind of stimulus (e.g., a tone; the conditioning stimulus). After some time, the conditioning stimulus will elicit the reflex response even when it is not followed by the unconditioned stimulus. The classical examples are the experiments of the Russian physiologist Pavlov, in which gastric secretion was produced by ringing a bell (conditioned stimulus). During the learning phase, the bell was always followed by food being presented (unconditioned stimulus). There are many examples from daily life of such conditioned behavioral responses, and conditioning is an important kind of learning. The **cerebellum** appears to be important for conditioning of movements, such as the blink reflex but also movements that involve skeletal muscles.

Some Reflexes “Disappear” during Development

Some reflexes are present only during certain phases of development, such as the **sucking reflex** and the **grasping reflex** in infants. When the reflexes disappear, they are no longer needed and would only disturb voluntary, goal-directed movements. The reflex arcs do not disappear, but the reflex response is suppressed by higher levels of the CNS. That the nervous apparatus persists is witnessed by the persistence or reappearance of primitive reflexes in brain-damaged children and adults.

The Reflex Arc and Basic Features of Reflexes

The structural basis of a reflex is a **reflex arc** (Fig. 21.9), which consists of the following links:

1. A **receptor**, which records the stimulus and “translates” it to action potentials
2. An **afferent link** (a primary sensory neuron), which conducts the action potentials to the CNS
3. A **reflex center**, in which the signals from the receptor may be modified (increased or decreased) by signals from other receptors and other parts of the CNS, where aftersignals are issued to effectors
4. An **efferent link** (neurons with axons passing out of the CNS), which conducts action potentials to the organ producing the response
5. An **effector**, which may be skeletal (striated) muscle, cardiac muscle, smooth muscle (in the wall of vessels and visceral organs), or glands

Reflexes with their reflex center in the spinal cord are called **spinal reflexes**. **Brain stem reflexes**, as the name implies, have their center in the medulla, pons, or mesencephalon; **cortical reflexes** have a reflex center that involves parts of the cerebral cortex. Some reflex arcs are simple, while others that involve coactivation of several muscle groups may be highly complex—with reflex centers that include neurons in the brain stem and in the cord. Most reflex centers include links of several synaptically coupled neurons; such reflexes are called **polysynaptic**. If only one synapse is intercalated between the afferent and the efferent link, the reflex is called **monosynaptic**.

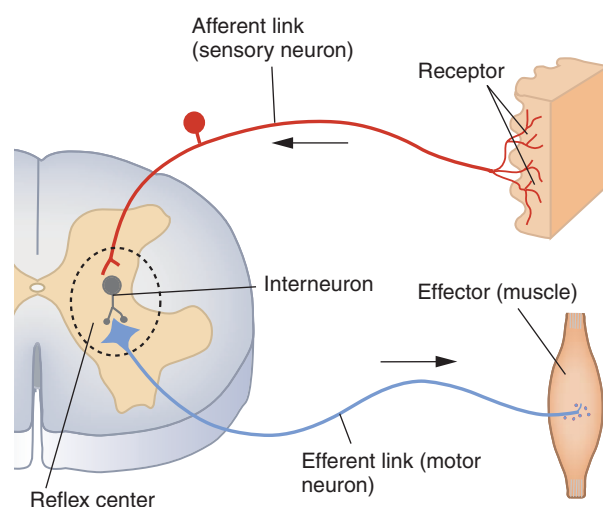


FIGURE 21.9 *Reflex arc.* Schematic. All reflex arcs contain the same elements. The figure shows a spinal reflex arc, with cutaneous nerve endings as receptors and a skeletal muscle as effector. The reflex center has in this example two synapses (a disynaptic reflex).

Although a reflex response is mediated each time by the same set of neurons, the **excitability** of these neurons can be modified from higher levels of the CNS. This is necessary to adapt individual reflexes to the overall plan for bodily movements. Reflex movements that operate on their own would disturb normal voluntary movements. **Modulation** of the excitability of the reflex center can be exerted by presynaptic inhibition of primary afferent fibers, by postsynaptic excitatory or inhibitory actions on interneurons and motoneurons, and by efferent control of the sensitivity of some kinds of receptor (cf. γ motoneurons).

We first discuss two different spinal reflexes with skeletal muscles as effectors. The first—the **flexion or withdrawal reflex**—serves to protect the body; the second, so-called **stretch reflex** (or rather stretch reflexes since there are several varieties), automatically adjusts muscle tension during postural tasks and voluntary movements.

The Flexion Reflex

This reflex is evoked by activation of **nociceptors** in the skin and underlying tissues. For example, when a foot hits a sharp object while walking, the whole leg is immediately withdrawn away from the object (Fig. 21.10). In this case, one or more interneurons are intercalated between the terminals of the afferent sensory fiber and the motoneurons producing the response—that is, the reflex is **polysynaptic**. That this is a reflex is clear from, among other things, the fact that it may be elicited even when the spinal cord is transected above the reflex center. The flexion reflex disappears in deep unconsciousness. For surgery, anesthesia has to be sufficiently deep to abolish flexion reflexes.

As mentioned, the **receptors** for the flexion reflex are nociceptors. The **stimulus** usually hits a small spot on

the skin, whereas the **response** involves a complex array of muscles (such as extensors of the ankle, flexors of the knee and the hip; Fig. 21.10). Contraction of the muscles in the leg that is withdrawn, however, is not sufficient. Muscles of the other leg must also contract (primarily extensor muscles) to prevent loss of balance. Thus, the stimulus must be distributed to motoneurons in many segments of the cord. This happens by way of ascending and descending collaterals of the primary sensory fibers and by way of spinal interneurons (Fig. 21.10). In response to a simple stimulus, a purposeful, harmonious movement occurs, requiring that all the muscles contract at the right time and with the right force. The synaptic couplings in the cord underlying this response must be both complex and precise.

A Flexion Reflex Can Be Evoked from Low-Threshold Receptors

Under certain circumstances, several kinds of receptor in the skin, around the joints, or in muscles can elicit a flexion reflex. Common to these receptors is that their signals converge on **interneurons** that excite **flexor motoneurons**. The sensory fibers from such receptors were therefore termed **flexor reflex afferents (FRAs)**. Their effects have been most studied in so-called spinal animals (the cord is transected and isolated from the rest of the CNS). Many FRAs lead from low-threshold mechanoreceptors (e.g., group II muscle afferents), while others lead from nociceptors. Normally, the convergence of FRAs on interneurons is believed to ensure necessary **feedback** during ongoing movements—perhaps rhythmic movements in particular. The term “flexor reflex afferents” is therefore not quite appropriate because many of the FRAs do not normally evoke the protective flexion reflex. In patients with **transverse lesions of the cord** the situation is different, however.

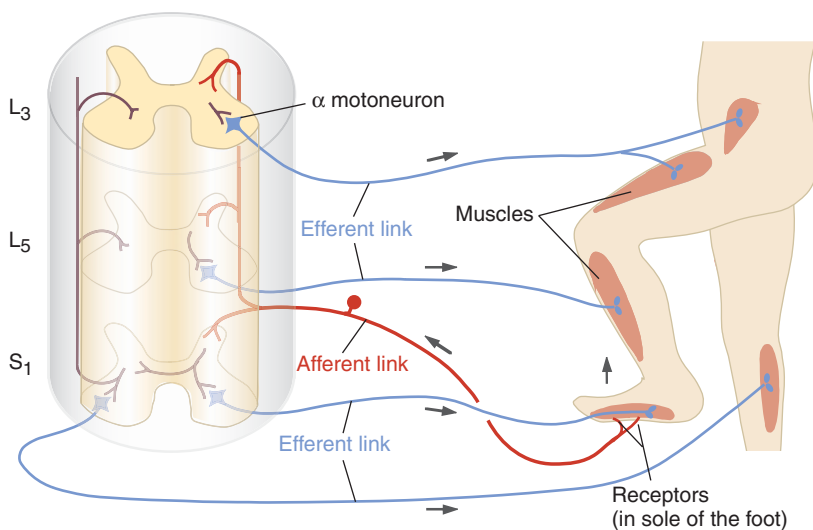


FIGURE 21.10 The flexion (withdrawal) reflex. The diagram is highly simplified. In this case, several synapses are intercalated between the afferent and the efferent links of the reflex arc. Here nociceptors in the sole of the foot are activated (by walking on a sharp object). The signals are conducted centrally in a sensory fiber (red), which sends out several collaterals and, via interneurons and propriospinal fibers, activates α motoneurons at several segmental levels of the cord. In turn, the motoneurons make many muscles contract to lift the foot off the ground (away from the painful stimulus). At the same time, extensor muscles are activated in the other leg to maintain balance.

Reflex changes occur in body parts that have been innervated from the cord below the lesion, including forceful and long-lasting contractions of flexor muscles. Such contractions are elicited by innocuous stimuli, including stimulation of low-threshold mechanoreceptors and thermoreceptors. Most likely, this is due to abnormal excitability of interneurons in receipt of FRA inputs produced by the lack of supraspinal control (see also Chapter 22, under “Mechanisms Responsible for Development of Spasticity”).

Stretch Reflexes

Under certain circumstances a muscle responds with a contraction when it is being stretched. (It is easy to find out by oneself that stretching a muscle does not always produce a contraction.) When the latency of the contraction is too short to be voluntary, we call this a **stretch reflex**. The most obvious purpose of such a reflex might be to keep the muscle length constant. **Receptors** for stretch reflexes are the **muscle spindles**. Stretch reflexes have been much studied due to their relation to movement control.

We speak of different stretch reflexes because muscle contraction occurring as a result of muscle stretching usually consists of several phases of contractions, each with a different latency as recorded by EMG. The reflex **latency** is the time between the stimulus and the response (between stretch and contraction). It results from the conduction time from the muscle spindles to the spinal cord; the delay at the synapse(s) intercalated between the muscle spindle afferent fibers and the α motoneurons; the conduction time from the motoneurons to the muscle; and, finally, the synaptic delay at the motor end plate. Measurement of the latency between the stimulus and the response makes it possible to decide whether a muscle contraction is the result of a **monosynaptic** stretch reflex, or whether other—**polysynaptic**—pathways are involved. The monosynaptic stretch reflex, with only one synapse intercalated between the afferent and the efferent link, is the simplest reflex with the shortest latency (Fig. 21.11). Each intercalated synapse increases the latency with a few milliseconds. Thus, muscle contractions occurring with longer latencies after stretching the muscle indicate that there are more neurons intercalated between the afferent link (the sensory fibers) and the motoneurons. That some stretch reflex responses have longer latency than others may not only be due to more synapses intercalated in the reflex center, however. Thus, while the fast monosynaptic reflex depends on signals in the thick group Ia fibers (see Figs. 13.6 and 13.7), slower conducting group II muscle spindle afferents may contribute to reflex responses with longer latencies.

We discuss the monosynaptic and the polysynaptic stretch reflexes separately next. Much remains to be

clarified, however, concerning the role of the stretch reflexes in movement control.

The Monosynaptic Stretch Reflex

The **patellar reflex** (Fig. 21.11)—tested routinely as part of a clinical examination—is probably the best-known example of a stretch reflex. The **stimulus** is a tap on the patellar ligament (the tendon of the quadriceps muscle). The **response** is a contraction of the quadriceps muscle, thus producing a brief extension movement at the knee joint. The reflex arc is shown schematically in Fig. 21.11, and it can be seen that only one synapse is intercalated in the reflex center—that is, the reflex is **monosynaptic**. The **receptors** are the muscle spindles in the quadriceps muscle that is stretched; the **afferent link** is constituted by the Ia fibers, which end in the spinal cord with some collaterals ending directly (monosynaptically) on the α motoneurons supplying the quadriceps muscle. The axons of the α motoneurons constitute the **efferent link**, which at the motor end plate activates the extrafusal fibers of the quadriceps muscle (the effector). For the patellar reflex, the **latency** between the stimulus and the start of the contraction (as recorded with EMG) is about 30 msec, and a slightly shorter latency pertains

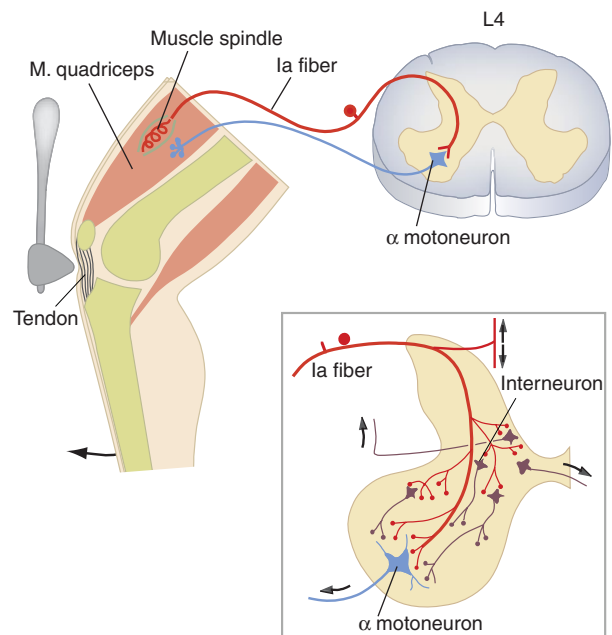


FIGURE 21.11 *Stretch reflex.* The patellar reflex is used here as an example of a monosynaptic reflex—that is, with only one synapse intercalated between the afferent and the efferent link. The stimulus is a tap on the patellar ligament below the patella, which stretches the muscle. The response is a brief muscle contraction. **Bottom right:** The Ia fiber from the muscle spindle contacts the motoneurons monosynaptically, but it also contacts many interneurons. Some of these are excitatory, mediating a polysynaptic activation of some motoneurons; other interneurons are inhibitory, mediating inhibition of other motoneurons.

to reflexes in the upper arm muscles because of a shorter peripheral pathway—for example, the **biceps reflex** has a latency of about 25 msec.

To evoke a muscle contraction through this simple reflex arc, the muscle must be stretched rapidly—that is, by **phasic** stretching. The actual lengthening produced by the stimulus may be very little; the salient point is the **velocity of stretching**. As discussed in Chapter 13, the group Ia afferents of the muscle spindles are especially sensitive to the velocity of stretching, so that even a brief stimulus, stretching the muscle only a fraction of a millimeter, produces a brisk response. A tap on a tendon with a reflex hammer produces this kind of stimulus, although it is far from being a natural stimulus. Because both the stimulus and the response last only for a very short time, we also use the term **phasic stretch reflex**. The monosynaptic stretch reflex can be elicited in most skeletal muscles but much more easily in some muscles than in others. There are also individual differences: in some healthy persons, phasic stretch reflexes cannot be elicited.

The Jendrassik Maneuver and Strengthening of the Monosynaptic Stretch Reflex

As mentioned, in some healthy persons the monosynaptic reflex cannot be elicited by ordinary tendon taps with a reflex hammer. Absent stretch reflexes may also be due to lesions of the reflex arc, however. To help decide whether the reflex is merely weak or absent, there are means to strengthen the reflex response in healthy people. The person forcefully contracts a muscle of the arm (especially the forearm) or the jaws while the examiner tries to evoke a reflex in the lower extremities. It takes about 300 msec from start of the maneuver to maximal facilitation of the reflex. The facilitation of the monosynaptic stretch reflex by such maneuvers must be due to descending signals (pyramidal tract?) that increase the excitability of the reflex center. One possibility with some experimental support is shown in Fig. 21.12. The descending fibers act on the interneurons that inhibit the interneurons that are producing presynaptic inhibition of the Ia terminals. Other possible explanations have also some experimental support, for example, that increased γ activity may play a role. Whatever the mechanism, the effect is obviously not restricted to increased reflex gain for the working muscles but also for muscles in other parts of the body.

The Monosynaptic Stretch Reflex Is Not as Simple as It Appears

Even a monosynaptic reflex is not quite as simple as Fig. 21.11 indicates, however. Stretching of the muscle activates not one but presumably, most of the muscle spindles of the muscle so that many motor units

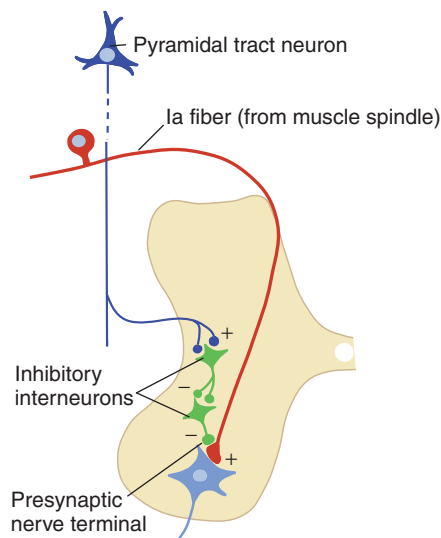


FIGURE 21.12 Strengthening of the stretch reflex by the Jendrassik maneuver. Possible mechanism, in which descending commands reduce presynaptic inhibition of Ia afferents from the muscle spindles. (Based on electrophysiological studies in humans by Zehr and Stein 1999.) See text for further discussion.

contribute to the response. The reflex center is not restricted to one spinal segment. Thus, α motoneurons supplying the quadriceps muscle are present in several lumbar segments (L_2 – L_4). Further, the **strength** of the response—even when the stimulus (the tendon tap) is kept constant—may vary considerably from person to person and even from time to time for one person depending on, for example, the level of **anxiety**. This can be understood when considering that the motoneurons receive influences not only from the muscle spindle afferents but also from other kinds of receptors and higher levels of the CNS (compare the effect of γ motoneurons on the excitability of the muscle spindles). Further, the terminals of the Ia afferents are subject to **presynaptic inhibition**, which reduces the excitation that impinges on the α motoneurons. Altogether, this means that the **gain** of the reflex is modulated in accordance with the overall need of the organism. It also explains why—during **learning** of a motor task—the strength of the monosynaptic reflex response can be strengthened or weakened according to what is functionally appropriate. We return to modulation of reflex responses.

Reciprocal Inhibition

Stretching a muscle does not only lead to excitation of the α motoneurons innervating the muscle (**homonymous excitation**); at the same time, the motoneurons of the antagonists are inhibited. This phenomenon is called **reciprocal inhibition** and is obviously functionally appropriate: it prevents a stretch reflex from being elicited in the antagonists when they are stretched by the

reflex contraction of the agonists (which, in turn, would elicit a new stretch reflex of the agonists, and so on). The reciprocal inhibition thus prevents unwanted **oscillatory** movements. Indeed, after damage to descending motor pathways, repetitive alternate contractions of agonists and antagonists can occur. This phenomenon, called **clonus**, is most likely due to disturbed functioning of spinal inhibitory interneurons. Reciprocal inhibition also reduces the tension of the antagonist muscles, which otherwise would brake the desired movement. In the case of the patellar reflex, the motoneurons of the knee flexors are inhibited (Fig. 21.13). The inhibition of the α motoneurons of the antagonist is mediated by so-called **Ia-inhibitory interneurons**, which are excited by Ia afferents from the agonist muscles.⁶

Motoneurons Can Inhibit Their Own Activity: Renshaw Cells

Together, the many inputs to the motoneurons determine their level of excitability—that is, these inputs also determine the activity of the skeletal muscles. Besides the inhibition of motoneurons by primary afferent fibers (via interneurons) described in the preceding text, the motoneurons can also inhibit their own activity. Thus, the α -motoneuron axons send off collaterals before they leave the ventral horn (Fig. 21.14). Because they turn back, they are called **recurrent collaterals**. The recurrent collaterals end primarily on a group of inhibitory interneurons, the **Renshaw cells**, located ventromedially in the ventral horn. The axons of the Renshaw cells ramify extensively and contact primarily the α motoneurons (but also certain kinds of inhibitory interneurons). In this manner, the activity of the α motoneurons excites the Renshaw cells, which, in turn, inhibit the same α motoneurons and other α motoneurons that supply muscles with similar action (agonists). Further, the Renshaw cells inhibit the inhibitory interneurons mediating the **reciprocal inhibition** (Fig. 21.10)—that is, the inhibition of the α motoneurons supplying the antagonists is reduced (**disinhibition**). In case of a stretch reflex, the activity of the Renshaw cells shortens the reflex contraction of the agonists and, at the same time, shortens the reciprocal inhibition of the antagonists. In general, recurrent inhibition appears to be more important in proximal than in distal muscles.

The Renshaw system seems important to prevent the α motoneurons from sending long trains of action potentials as a response to a brief stimulus. In situations (such as during voluntary movements, when prolonged

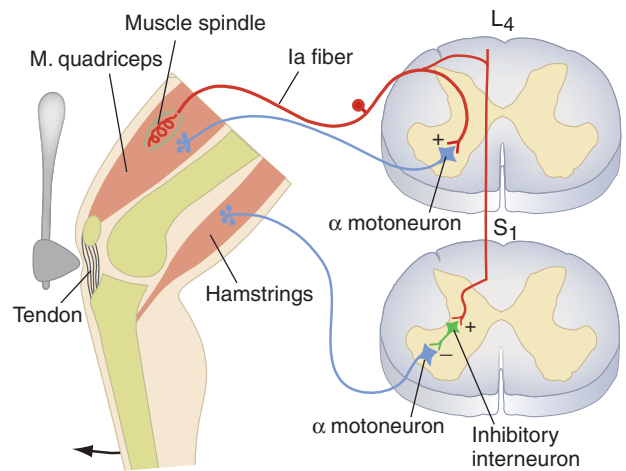


FIGURE 21.13 *Reciprocal inhibition.* Schematic of how the stretching of one muscle—producing a reflex contraction—elicits inhibition of the antagonistic muscles via inhibitory interneurons.

motoneuronal activity is required), **descending connections** from the brain stem and the cerebral cortex may inhibit the Renshaw cells that are supplying the working muscles. Thus, the activity of the Renshaw cells, as that of many other kinds of spinal interneurons, is modulated in accordance with the overall plan for the movements.

The Long-Latency (Polysynaptic) Stretch Reflex

As mentioned, stretching of a muscle may elicit reflex contraction via routes other than the monosynaptic reflex arc. The muscle spindle afferent fibers (Ia and II) contact not only α motoneurons but also many interneurons

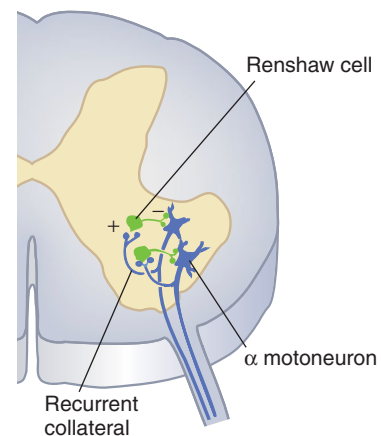


FIGURE 21.14 *Recurrent inhibition.* The α motoneurons inhibit their own activity by sending recurrent collaterals ending on the inhibitory Renshaw cells. The Renshaw cells are also influenced by other spinal neurons and from higher levels, which determine how strongly they will respond to a certain input from the α motoneurons.

⁶ The termination of Ia afferents in the spinal cord are differentiated, so that some collaterals end in contact with interneurons that *inhibit* the muscle in which the muscle spindles lie. Although this inhibitory influence is not noticeable when testing the monosynaptic stretch reflex, it may have a role in performance of natural movements when the eliciting of stretch reflexes may be unwanted.

(Fig. 21.11), establishing **polysynaptic** routes from the muscle spindle afferents to the motoneurons. Rapid stretching of the biceps muscle may elicit two or three reflex responses, as determined with EMG in human subjects. In addition to an early EMG activity with a latency of about 25 msec (the monosynaptic stretch reflex, or **M1** response), another reflex contraction (**M2**) starts at about 50 msec and sometimes a new phase of contraction (**M3**) at about 70 to 80 msec. These are not voluntary responses, since the earliest voluntary muscle contractions occur at about 100 msec after the stretch (in the upper arm). These reflex responses constitute the **long-latency stretch reflex** (other names are the functional stretch reflex, the polysynaptic stretch reflex, and the long-loop stretch reflex).

A striking property of the long-latency stretch reflex is that the strength of the response depends to such a high degree on whether the muscle is relaxed or **active** at the time of stretching. If the muscle is relaxed or only slightly active when stretched, there is usually no long-latency reflex response at all. Further, the strength of the response depends on **prior instruction** to the subject with regard to whether to resist the imposed stretch or to let go. When the person is asked to let go when an imposed movement (at an unpredictable time) stretches the muscle (e.g., an imposed extension at the elbow that stretches the biceps muscle), the reflex response is much smaller than when the person is asked to resist the imposed movement. Thus, the magnitude of the reflex response can be adapted to what is functionally appropriate in a particular situation.

A further characteristic of the long-latency stretch reflex is that the strength of the response may change during **learning** of a motor task. Thus, by repeated trials, the reflex response becomes weaker in muscles in which a contraction in response to stretching is functionally inappropriate and stronger in muscles in which a contraction is appropriate. This learning effect, or adaptation of the stretch reflex, occurs only in connection with the particular learned movement; in connection with other movements, the reflex response of the muscle is unaltered. As mentioned, the monosynaptic stretch reflex is also subject to similar learned, task-related modulation, but the changes appear to be smaller than those obtained with the long-latency reflex.

Is the Long-Latency Stretch Reflex Mediated by the Cerebral Cortex?

The exact central pathway followed by the impulses mediating the long-latency stretch reflex has been much debated. Indirect data indicate that the reflex pathway may involve the motor cortex of the cerebral cortex (therefore, the term “long-loop stretch reflex” is often used). In support of a **transcortical** route are the observations that the reflex is weakened or abolished by

lesions of the descending motor pathways or by lesions of the dorsal columns (presumably carrying the signals from the muscle spindles to the cerebral cortex). Further, the reflex is often weakened after lesions of the **cerebellum**. But such findings may also be explained by a purely spinal reflex that is under strong supraspinal control.

More decisive evidence of a transcortical route for the long-latency stretch reflex—at least regarding certain muscle groups—comes from observations in a few patients with a peculiar inborn abnormality of the pyramidal tract. These persons always perform **mirror movements** of the hands; asked to flex the index finger of the left hand, they always flex the right index finger as well (more proximal movements, e.g., of the shoulders, are performed normally). This behavior appears from electrophysiological studies to be caused by branching of individual pyramidal tract axons to supply motoneurons of both sides of the spinal cord. Thus, stimulation of the hand region of the motor cortex of one hemisphere causes symmetrical movements of both hands (unlike the normal situation, in which such stimulation always cause movements of the opposite hand only). When eliciting stretch reflexes in such subjects, the monosynaptic reflex occurs only on the same side as the stretch is applied (as normal), whereas the long-latency stretch reflex occurs in both hands after a unilateral stimulus. The latter observation is hard to explain unless the reflex arc of the long-latency reflex involves the pyramidal tract.

These findings may not pertain to long-latency stretch reflexes in all muscle groups. For example, the long-latency stretch reflexes appear to be transcortical for distal arm muscles but not (or to a lesser degree) for proximal arm muscles and muscles in the foot. For these latter muscles, the long-latency reflex may be elicited by **group II muscle afferents**, which conduct with about half the velocity of group Ia fibers.

The Function of Stretch Reflexes

One might think that the stretch reflexes—which, after all, are relatively simple—are well understood with regard to their functional roles. For example, the muscle spindles and the motoneurons are among the best-characterized receptors and central neurons, respectively. Nevertheless, we still do not fully understand the role of the stretch reflexes in the control of voluntary movements and in the control of posture and muscle tone. We discuss here only some possible functions.

As mentioned, one likely task of the stretch reflex is to ensure that the **length** of a muscle is kept constant. In many situations, this is of obvious importance—for example, in the upright position when some external perturbation threatens the **body balance**. The sudden displacement of the center of gravity forward stretches

the extensor muscles of the back and thus might elicit a stretch reflex tending to resume the former position. It is furthermore an obvious advantage that such a corrective contraction occurs as quickly as possible. Making such an adjustment depend only on voluntary contraction would lengthen the latency fourfold, with the danger of the corrections occurring too late. Also while **walking**—when external perturbations may disturb the programmed pattern of muscular activity—stretch reflexes may contribute to rapid adjustments. Nevertheless, it is not clear to what extent stretch reflexes really participate in such adjustments. (See also Chapter 18, under “More about Receptor Types and Their Contribution to Postural Control.”)

Another situation in which stretch reflexes may be of importance is during slow, **precise voluntary movements** when the external opposing forces change unpredictably. Again, the advantage would be that the adjustment of muscle tension occurs much earlier than can be achieved by voluntary action alone. (See also Chapter 13, under “Proprioceptors and Voluntary Movements.”)

Stretch Reflexes May Correct for Change in External Resistance during Precision Movements

Studies of slow movements of the thumb by the British neurologist David Marsden have shed light on the contribution of stretch reflexes during precision movements. The subject is asked to flex the thumb with a constant speed against an external opposing force of constant magnitude. The EMG of the flexor pollicis longus muscle is recorded continuously. The external force is then changed suddenly at unpredictable times during the movement—either increased or reduced. When the external force is increased, the movement is immediately slowed down. Because of the **α - γ coactivation** (see Chapter 13, under “Muscle Spindles in Humans and α - γ Coactivation”), the frequency of signals from the muscle spindle increases. The α - γ coactivation ensures that, as the muscle shortens owing to the activation of the α motoneurons, the spindle midportion is stretched. This upholds the firing of muscle spindle afferents in spite of the shortening, which otherwise would have led to reduced firing. To keep up with the steadily shortening muscle, the firing of the γ motoneurons must also increase steadily. When the movement is suddenly halted or slowed down, the firing frequency of the γ motoneurons continues increasing, in anticipation of further shortening of the muscle. Thus, for a moment, the firing of the muscle spindle afferents increases more than what is appropriate with regard to the actual length of the muscle. This increases the excitation of the α motoneurons, and their firing increases, thus increasing the force of the muscle contraction. The result is that the increased external force is rapidly compensated for, and the original speed of the movement is resumed.

When the opposing external force is suddenly reduced, the opposite events take place. The speed of the flexion movement of the thumb increases, and the firing frequency of the spindle afferents decreases for a moment, thus reducing the firing of the α motoneurons and the force of contraction. The speed of movement is adjusted.

It is probable that the stretch reflex functions in this manner especially during **slow precision movements** when we cannot accurately predict the external force at all times. The sensitivity of the muscle spindles is kept at a high level, so that they may record even the slightest perturbations and ensure that the activity of the α motoneurons is adjusted appropriately.

Cutaneous Receptors and the Precision Grip

In some situations, stimulation of low-threshold skin **mechanoreceptors** causes reflex muscular contraction. For example, when (during a precision grip with the fingers) the object slips, a reflex increase of the grip force occurs. The latency from the start of the slip to the muscular response is only 60 to 80 msec—that is, too short to be mediated by a voluntary command. Examination of patients with reduced cutaneous sensation but normal motor apparatus suggests that loss of such rapid, reflex adjustment of the grip force is partly responsible for their difficulties with precision movements (see Chapter 13, under “Clinical Examples of Loss of Somatosensory Information”).

Central Modulation of Reflexes

It should be clear from the preceding discussing that stretching of a muscle does not necessarily elicit a reflex contraction. Many factors influence whether there will be a response, such as the velocity of stretching and whether the muscle is active when being stretched. Further, the response depends heavily on whether a reflex contraction is functionally appropriate. Such **gain modulation** of the stretch reflex is mediated by descending connections from higher levels of the CNS (e.g., from the motor cortex). It appears to be exerted mostly by a precise control of the excitability of specific sets of **spinal interneurons**, which are intercalated in particular reflex arcs. In addition, **presynaptic inhibition** may very selectively “switch off” the input from specific sets of receptors. Thus, reflex arcs may be “opened” or “closed” in accordance with the need of the overall plan for movements (Fig. 21.15). Obviously, the motor programs in the cerebral cortex and the brain stem specify not only the muscular activity but also the gain of spinal reflex arcs at any moment.

Reflex modulation has been studied most during **locomotion** in humans and is found in several muscle groups. Thus, the strength of reflex contractions in many leg muscles depends on whether the leg is in the

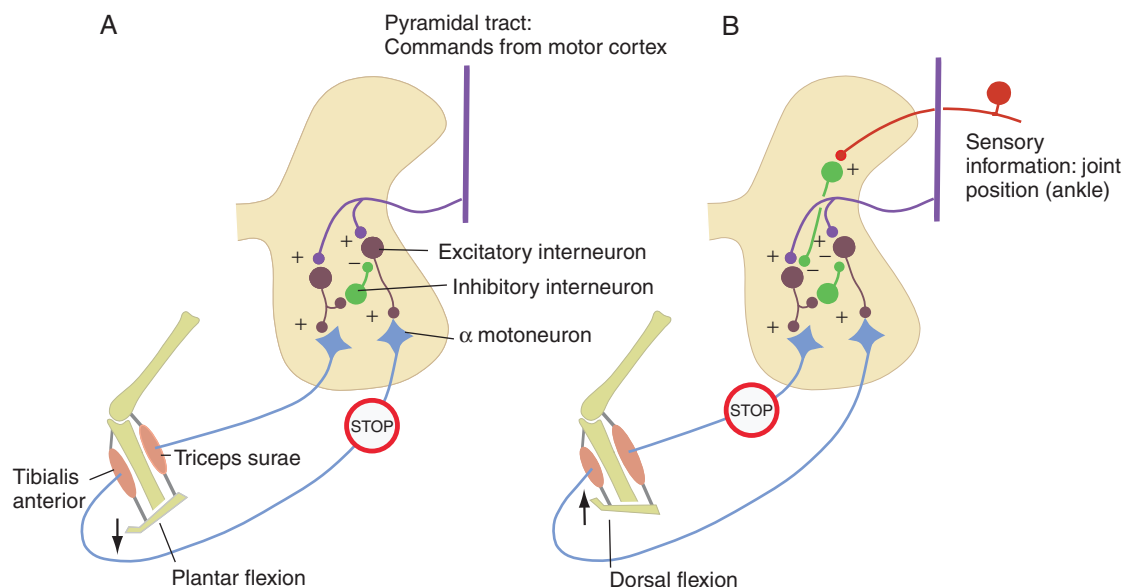


FIGURE 21.15 *Spinal interneurons and switching of signal pathways.* Synaptic couplings (highly schematic) may explain how the position in the ankle joint can determine whether a central command facilitates a flexion or an extension movement. **A:** The central command produces contraction of ankle flexors (triceps surae) because the signal pathway to the extensors is “shut off” by inhibitory interneurons.

B: The position in the ankle joint has changed, giving a different sensory input (from muscle spindles) than in **A**. The sensory signals excite the interneurons that inhibit triceps motoneurons. Thus, the same command from the motor cortex as in **A** produces in **B** contraction of the tibialis anterior muscle rather than the triceps. (Based on McCrea 1992.)

swing or in the stance phase. In the ankle flexors (m. triceps surae), for example, stretch reflexes are weak or absent in the swing phase, whereas they are brisk in the stance phase. This seems appropriate, because in the stance phase the reflex contraction would strengthen the desired plantar flexion, whereas in the swing phase the appropriate movement is dorsal flexion of the foot to enable free swing of the leg.

Arm muscles also show marked reflex modulation in persons catching a ball. In a certain phase of the movement, even the reciprocal inhibition is absent—that is, a stretch reflex can be elicited in a muscle without inhibition of its antagonists. This is functionally appropriate when agonists and antagonists are required to co-contract to stabilize the joint.

In addition, the gain of the **flexion reflex** is under some degree of supraspinal control. This is evident during walking, when a flexion reflex may cause a fall. Thus, noxious stimulation of the sole of the foot elicits a brisk flexion reflex if the leg is in the swing phase, whereas the response is weak or absent when the foot is used for support. Interestingly, this kind of modulation appears not to be present in four-legged animals, presumably because their balance is less vulnerable to changing the support of one leg only.

MUSCLE TONE

The term **muscle tone** refers to the tension in muscles. Usually, it means the slight tension that can be felt in a

relaxed muscle; a more precise term is therefore **resting tone**. In pathological conditions, the resting tone may be either increased or decreased. The term “muscle tone” is ambiguous, however, as witnessed by its use by various authors. (Some authors have suggested that the term should be abandoned because its lack of precision.)

Muscle tone is usually determined by **palpation** (judging the consistency and stiffness of the tissue by pressing the muscle belly between the fingers) and by **stretching** (judging the resistance offered to passive stretch). Some authors use the term only about the tone as judged by stretching. Palpation and stretching do not test identical properties of the muscle, however. For example, after a stroke giving **central pareses** (as in capsular hemiplegia), the muscle tone is commonly reduced when judged by palpation (the consistency is reduced), whereas the resistance offered to passive stretching is increased (spasticity). Because examination of muscle tone is important for the diagnosis of diseases that affect the motor system, it is regrettable that we do not have a clear understanding of the basis of normal muscle tone and of the mechanisms behind changes of muscle tone in disease states.

Muscle Tone Is an Expression of Tissue Stiffness

The confusion about the exact meaning of the term “muscle tone” stems partly from the fact that words like “firmness,” “tension,” “stiffness,” and “elasticity” are often used without definition. Different persons may therefore use them with different meanings.

Unfortunately, this is often the case even when observations made by different examiners are compared. Stiffness and tension can be defined in mathematical terms, however, and can also be objectively recorded and measured. The definition of stiffness (K) is $K = \Delta T / \Delta L$; that is, K expresses how much the muscles are stretched (ΔL) by a certain increase in tension (ΔT). The greater the stiffness, the higher the tension necessary to stretch the muscle. Note that this definition applies only to the component of muscle tension determined by passive stretching and not to the consistency (firmness, elasticity) as judged by palpation. Consistency can hardly be measured objectively and depends on the subjective assessment of the examiner.

What Determines Muscle Tone?

From everyday experience, we know that **contraction** of a muscle (caused by actin–myosin interactions) is the one factor that most markedly alters the muscle tone, as judged by both palpation and stretching. The purpose of the contraction, of course, is to increase the tension of the muscle. The tone felt in a fully **relaxed** muscle (showing no EMG activity), however, must depend on passive properties of the muscle. Small contributions to the tension of the muscle stem from the **viscoelastic properties** of the muscle cells and the connective tissue of the muscle and its tendons. There is evidence that even among persons who are able to relax their muscles completely (as judged by EMG), the muscle tone (as judged by palpation) may vary. What are the bases of such differences? The elastic properties of the muscle explain why, when the muscle is stretched passively beyond a certain length, the tension increases steeply (the slack is taken out of the elastic, as it were). This is experienced when a relaxed muscle becomes firmer on palpation and offers increasing resistance to stretching toward extreme joint positions (therefore, such positions should be avoided when examining muscle tone). Even in a relaxed muscle of medium length, however, properties of the muscle cells themselves can influence the muscle stiffness. Elastic **titin (connectin)** molecules of the cytoskeleton connect the Z discs with the myosin filaments. They extend from the Z disc to the middle of the A band of the sarcomere and contribute to the resistance to stretch offered by a relaxed muscle cell. The titins exist in several varieties (isoforms). For example, there are different isoforms in the soleus and the psoas muscles, and the resting tone differs greatly in these muscles (highest in the psoas). Further, even in relaxed muscle a few **crossbridges** between actin and myosin filaments can exist, judging from experiments with isolated muscle cells. If the number of such crossbridges varies among muscles, it might contribute to differences in resting tone. Further, differences in the amount and composition of **connective tissue** and the **extracellular fluid** may contribute to individual differences in

muscle tone. So far, however, these factors are only hypothetical.

Nevertheless, many healthy persons are unable to relax their muscles completely, at least during an examination. Thus, EMG would show a modest electric activity when the examiner handles the muscles. The contraction is probably not of a reflex nature but is produced by voluntary commands (i.e., by signals issued from the cerebral cortex). Thus, in such persons there is usually muscle activity also in muscles that are shortened passively as the examiner produces a joint movement (the person “helps” the examiner).

In conclusion, individual differences in resting muscle tone among healthy persons may in some cases be caused by differences in the degree of relaxation. In addition, there are most likely also individual differences with regard to the passive viscoelastic properties of the muscles.

Is Muscle Tone Due to Reflex Contraction?

It was formerly assumed that even relaxed muscle was in a state of slight contractile activity, and that this activity was maintained by a steady flow of signals from the muscle spindles driving motoneuron firing. This was based primarily on observations in animals with transection of the brain stem (decerebration), in which the activity of the γ **motoneurons** is enhanced. This increases the signal traffic from the muscle spindles, and thereby the muscles are in a state of tonic contraction. In accordance with these observations, muscle tone in normal human subjects was assumed to be maintained by the long-latency stretch reflex (“tonic stretch reflex”). This view cannot be upheld in light of more recent data, however. In the first place, several **EMG** studies show that in persons who are able to relax properly, there is no electrical activity in their muscles (and, therefore, no contractile activity). Further, **microneurographic** studies have not confirmed that there is signal traffic in the muscle spindle afferents that conduct from relaxed muscles in human subjects. In relaxed subjects, the resistance against even rapid passive stretching of a muscle is very low (except when the monosynaptic stretch reflex is elicited by a tendon tap). There is thus no evidence of a “reflex tone” in relaxed muscles.

Muscle Cramps and Plateau Potentials

Cramps are sudden, involuntary, and painful muscle contractions. The person notices that a part of the muscle becomes tight. Cramps may occur in healthy people during sleep, in pregnancy, and after fatiguing muscular exercise. Cramps may also occur in diseases such as **neuropathies**, **amyotrophic lateral sclerosis (ALS)**, and **vascular diseases**. Stretching the muscle is often the most efficient way to stop the contraction. It was shown

many years ago that the kinds of cramp mentioned here are caused by high-frequency **firing of α motoneurons** (other kinds of cramp may be caused by altered conditions in the muscle itself). An interesting possibility is that cramps arise because of the motoneurons' ability to form **plateau potentials**, as characterized by a stable depolarized state (see also Chapter 22, under "Monoaminergic Pathways from the Brain Stem to the Spinal Cord," and "Mechanism Responsible for the Development of Spasticity"). In this state, a brief excitatory input can trigger a train of action potential lasting for many seconds or even minutes. Experimentally, a brief train of signals in Ia fibers from the muscle spindles can cause sustained motoneuron firing and cramps, possibly because of induction of plateau potentials, but any excitatory input would presumably have the same effect. The plateau potential can be terminated by a brief hyperpolarizing synaptic input. Whether plateau potentials occur in muscle cramps in humans is unknown, but if they do, why they are induced at rest in some healthy persons needs nevertheless to be explained. Conceivably, increased extracellular K^+ due to intense motoneuronal firing during endurance exercise may elicit plateau potentials. Another possibility is that plateau potentials are triggered by altered inputs to motoneurons during muscular fatigue—for example, by increased muscle spindle afferent activity and reduced presynaptic inhibition. That stretching and sometimes massage terminate cramps might be due to stimulation of afferent inputs that inhibit motoneurons (via interneurons). Ia afferents are unlikely to be activated in this situation, as they respond poorly to slow stretching of the muscle. Electrical stimulation of the tendon inhibits experimentally evoked muscle cramp, suggesting that stretching of the cramped muscle works by activation of 1b afferents from the tendon organ.

So-called **writer's cramp** is a task-specific **focal dystonia** of the hand. The cramp usually occurs when trying to do a task that requires fine-motor movements. In this case, it is believed that basal-ganglia dysfunction lies behind the abnormal motoneuronal firing.

Changes of Muscle Tone in Disease

Pathologically changed muscle tone is called hypotonia when it is lower than normal and hypertonia when higher than normal. Of course, to recognize abnormal muscle tone one must first be able to decide what is normal, but from the preceding discussion, it should be clear that "normal muscle tone" is not a precise, well-defined concept. The decision as to whether a muscle has a normal tone is based largely on the subjective judgment of the examiner. This judgment, of course, depends on experience. Whereas hypertonia may be identified with reasonable certainty and even measured in semiquantitative terms by stretching the muscles, the

decision as to whether a muscle has abnormally low tone is more difficult. As mentioned, a normal, fully relaxed muscle will have a very low tension when tested by stretching. Some authors believe that when paretic or paralytic muscles feel softer and more flaccid than normal muscle, it is because of lack of voluntary contraction, which probably occurs to some extent during passive movements and palpation. Indeed, experiments measuring the resistance to passive stretching of normal and alleged hypotonic muscles did not show consistent differences. The experiments were performed by measuring the falling time of the leg (passive flexion of the knee) in healthy persons with the ability to relax fully (determined with EMG) and in patients with pareses of the quadriceps muscle (clinically judged as hypotonic). On the other hand, the individual differences in falling time—that is, muscle tone—among the normal subjects were fairly large, presumably because of differences in the passive viscoelastic properties. Nevertheless, with peripheral pareses, rapid changes of the passive viscoelastic properties of the muscles may occur, which may help explain why the paretic muscles feel softer on palpation. After damage to peripheral motoneurons, the muscles waste rapidly, reducing the muscle volume to sometimes only 20% to 30% of normal in about 3 months. The metabolism of muscle cells is obviously dramatically altered by loss of contact between nerve and muscle.

Abnormally increased muscle tone, **hypertonia**, would imply that the muscles continuously have an increased tone, in spite of attempts to relax. As mentioned, many healthy persons are not able to relax completely, at least not in an examination situation, and the border between normal and pathologically increased muscle tone may not be easy to draw. Fairly characteristic disturbances of muscle tone do occur in certain diseases of the CNS, however.

Spasticity and Rigidity

The term **spasticity** is used in clinical neurology of a condition in which there is increased resistance against rapid stretching of muscles. By palpation, the muscles may feel normal or hypotonic, and there may not be increased resistance against slow, passive movements. Spasticity occurs after damage to the descending motor pathways from the cerebral cortex to the motoneurons and is probably due primarily to changed spinal-interneuron excitability. The increased resistance to rapid stretch is most likely caused by abnormally brisk monosynaptic stretch reflexes, whereas the long-latency stretch reflexes are weaker than normal.

Rigidity is the term used to characterize the increased muscle tone occurring in **Parkinson's disease** (Chapter 23). Even with very slow, passive movements, an increased, "cogwheel"-like resistance is felt by the examiner. This may be caused by increased **long-latency stretch reflexes** that are elicited by abnormally slow movements.

Also with rigidity and spasticity, there is evidence of changed passive **viscoelastic properties** (in addition to the changed stretch reflexes). Thus, in patients with moderately severe Parkinson's disease, increased resistance to slow elbow extension was found, even though there was no EMG activity of the biceps muscle (which was being stretched). In some spastic patients, increased resistance even to slow stretching of relaxed leg muscles was present, but only when the spasticity had lasted for more than a year. Thus, it seems as though an altered pattern of signals from the motoneurons—like those occurring in diseases with rigidity or spasticity—may change the passive, viscoelastic properties of the muscles.

INJURY OF PERIPHERAL MOTOR NEURONS AND REGENERATION

When all motoneurons (or their axons) supplying a muscle are destroyed, the muscle cannot be made to contract: it is **paralyzed**. Both voluntary and reflex movements are abolished. If not all of the motoneurons (or their axons) supplying a muscle are destroyed (Fig. 21.16), the muscle can still contract, although with less speed and force than normal. This is called a partial paralysis, or **paresis**. Typically, the paretic muscle feels soft and flaccid, and there is a marked reduction in the muscle mass. This is called muscle **atrophy** or wasting and is more marked with more complete destruction of the motoneurons. The muscle cells that no longer receive signals from the motoneurons become thinner and eventually disappear if no reinnervation takes place (see later).

Peripheral and Central Pareses

The muscle weakness caused by the loss of the α motoneurons or their axons (**lower motor neurons**) is called a **peripheral paralysis** (paresis), to distinguish it from a **central paralysis** caused by interruption of the central motor pathways (the upper motor neurons). Central pareses are discussed in Chapter 22. Suffice it here to mention that in central pareses the spinal reflex arcs are intact (Fig. 21.16). Characteristic of peripheral pareses—apart from the weakened or abolished voluntary contractions—is that the muscles are flaccid, reflex movements are weakened or abolished, and muscle wasting progresses rapidly and becomes marked.

In cases of peripheral pareses, the **distribution** of affected muscles may tell us where the disease process is located. For example, the distribution will differ, depending on whether the lesion is located in the spinal cord, in the plexuses formed by the spinal nerves, or in the peripheral nerves more peripherally (Fig. 21.2). As a rule, one muscle is supplied with motor fibers from two or more spinal segments, as discussed above (Fig. 21.3). Therefore, damage restricted to one spinal segment or

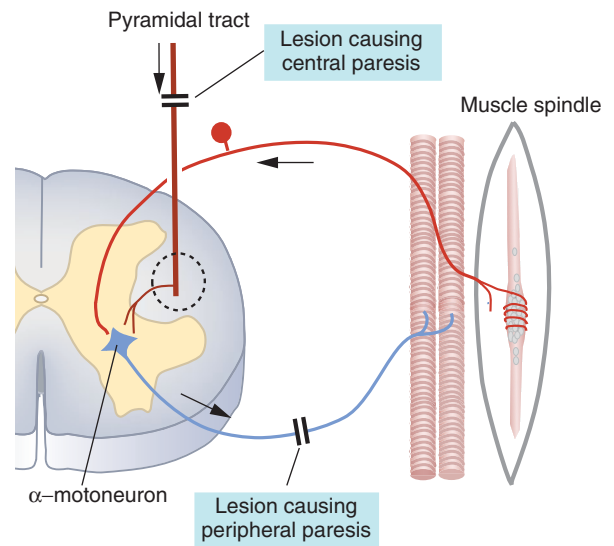


FIGURE 21.16 *Peripheral and central pareses.* Lesions of the motoneurons produce peripheral pareses, characterized by loss of both voluntary and reflex contractions. Central pareses—characterized by loss of voluntary movements but retained reflex contractions—ensue when descending corticospinal pathways are interrupted.

its ventral root (e.g., the C₅ root as in Fig. 21.2) produces only a paresis, not a complete paralysis of a muscle. Alternatively, several muscles become paretic because each segment contains motoneurons supplying several muscles, but if the peripheral nerve (e.g., the thoracicus longus nerve in Fig. 21.2) supplying a muscle (the serratus anterior) is severed, the muscle becomes paralytic.

Peripheral Axons Can Regenerate

When a **peripheral nerve is injured** so that the axons are interrupted, the distal parts of the axons degenerate and are gradually removed (by macrophages). The cell bodies (of the motoneurons and the spinal ganglion cells) show **retrograde** changes and, even though some of the cells die, many survive. The proximal parts of the axons of the surviving neurons start to grow (in contrast to what happens in the CNS; see Chapter 11, under “Can We Help Restitution?”). When possible, the growing axons follow the canals in the nerve left by the degenerated axons. The **Schwann cells** do not die and form a lining of the canals for the growing axons. The axons grow 1 to 2 mm per day, although the growth gets slower the farther peripherally the axon grows.

Factors that Influence Regeneration

The outcome of this regeneration of the axons depends on several factors, among them the conditions at the site of injury and the age of the person. If the continuity

of the nerve is not lost—as when the nerve is **crushed** or damaged by **compression**—the growing axons have a fair chance of finding the right track and reaching the target they innervated formerly. If the nerve is completely **severed** (e.g., by tearing or a cut), many axons will follow a wrong path even though the cut ends are meticulously stitched together. Thus, motor axons may innervate different muscles than previously, and, likewise, sensory axons may innervate different regions of the skin than before the injury. For example, motoneurons formerly supplying an extensor muscle may after the regeneration supply a flexor; sensory nerve fibers formerly supplying the thumb may innervate the index finger; and so forth. The longer the distance an axon has to grow after the damage, the poorer the chances that it will reach its target. Thus, one cannot expect full functional recovery after suture of a cut peripheral nerve. The results are better in **children** than in adults, however. This may be because children have a greater regenerative capacity, but presumably also because their brains adapt more easily to a novel pattern of innervation in the periphery.

In cases of **incomplete severance** of a nerve, the remaining motor axons usually send out new branches within the muscle (**collateral sprouting**). The sprouts grow into and make synaptic contacts at the “empty” motor end plates left by the degenerated axons. In this case, many of the muscle cells, which shortly after the injury were paralyzed, become **reinnervated** and some of the muscle power is regained. The remaining **motor units** of the muscle become larger, and therefore the

control of the muscle may become less precise than before the injury. Animal experiments suggest that the remaining motor units may increase their size four to six times.

After suturing a cut muscle nerve, **reflex** contractions often show less recovery than **voluntary** contractions. This is probably because the sensory fibers (innervating muscle spindles and tendon organs) have greater difficulties in reaching their correct target than the motor axons. Thus, after crushing a nerve—when the regenerating axons more easily find the right path—the reflex contractions show better recovery.

Regeneration of Motor Axons after Avulsion of Spinal Roots

Severed axons of α motoneurons can, in fact, grow for short distances in the spinal cord. This has been shown in animal experiments after avulsion of the nerve roots of the brachial plexus (Fig. 21.1). In this kind of lesion—occurring in humans exposed to forceful traction and depression of the shoulder and arm—the roots are literally torn loose from the cord. If the roots are surgically attached to their former sites of attachment, the motor axons first grow through the ventral horn—that is, the central nervous territory devoid of Schwann cells—and then into the roots and, finally, the muscles. In animals and in humans muscle function has been regained after root avulsions by this technique, even though the innervation is less precise than before. The reason for this unusual central regeneration is so far unknown.

The Motor Cortical Areas and Descending Pathways

OVERVIEW

This chapter deals with the control of complex, purposeful movements—from such that are largely automatic to those that require our full attention. The most direct control is exerted by **upper motor neurons** in the cerebral cortex and the brain stem that send fibers to end in direct synaptic contact with the **lower motor neurons** (motoneurons) or with interneurons that, in turn, contact the motoneurons. (The basal ganglia and the cerebellum—both necessary for proper motor control without being responsible for movement initiation—will be treated in the next two chapters.) The **pyramidal tract** is the only **direct** pathway from the cerebral cortex to lower motor neurons in the brain stem and the spinal cord. The pyramidal tract is indispensable for the ability to perform precise, voluntary movements of the hands (requiring fractionate finger movements), whereas more proximal movements can be performed although with less speed and precision in the absence of the pyramidal tract. For the latter kind of movement, several **indirect** pathways—synaptically interrupted in the brain stem—are especially important. The **corticoreticulospinal** pathways are important for maintaining the upright position (posture), for orienting movements of the body toward external events, and for fairly crude, stereotyped voluntary movements of the extremities. Nevertheless, both the direct and the indirect pathways participate, to a varying degree, in virtually all voluntary movements. The sharing of tasks among them is such that cell groups in the brain stem can largely on their own control many automatic movements, whereas the participation of the cortex increases with increasing degree of voluntary control.

Reticulospinal and vestibulospinal pathways are of special importance for the more **automatic movements**, such as postural adjustments and locomotion. Rhythmic movements, such as locomotion and respiration, depend on the activity of **rhythm generators** in the spinal cord and the brain stem. With regard to locomotion, there is most likely one spinal rhythm generator for each extremity. Supraspinal control of locomotion depends on a mesencephalic locomotor region (MLR). A contribution from the motor cortex is necessary if the ground is uneven and unpredictable.

We also discuss **motor cortical areas**—that is, areas that participate in planning and organizing movements. Many parts of the cortex are motor in this respect, but we usually restrict the term to the **primary motor area** (MI) in the precentral gyrus, the **premotor area** (PMA) just in front of MI, and the **supplementary motor area** (SMA) on the medial aspect of the hemisphere. The MI gives origin to a large part of the pyramidal tract fibers, and has monosynaptic connections with the motoneurons in the brain stem and spinal cord. PMA and SMA act largely by “instructing” the MI what to do. Whereas lesions of MI produce pareses, lesions of PMA and SMA give difficulties with initiation of movements and with purposeful movements that require coordinated sequences of muscle contractions. Posterior parietal areas and regions in the prefrontal cortex are necessary for transforming sensory information into action, and for selection of actions that are behaviorally appropriate.

The final part of the chapter deals with symptoms resulting from **damage** to the central motor pathways. The main characteristics of central pareses—such as **hyperreflexia** (spasticity) and **loss of dexterity**—are described and some tentative explanations are offered.

Customarily, a division has been made between the pyramidal (corticospinal) tract and other nuclei and tracts involved in motor control. The latter have often been collectively called the “**extrapyramidal system**.” So many parts of the brain contribute in different ways to motor control, however, that lumping them together in a “system” confuses more than it clarifies. Therefore, we will not use the term “extrapyramidal” here but rather describe what we regard as the most important components in control of voluntary movements.

THE PYRAMIDAL TRACT (THE CORTICOSPINAL TRACT)

The pyramidal tract is of crucial importance for our ability to perform precise, voluntary movements. The tract consists of axons of neurons with their cell bodies in the cerebral cortex, as indicated by its other name, the **corticospinal tract**. The axons descend through the internal capsule, the crus cerebri (cerebral peduncle), the pons, and the medulla (Figs. 22.2, 22.3, and 22.7). Most of the fibers cross to the other side in the lowermost part

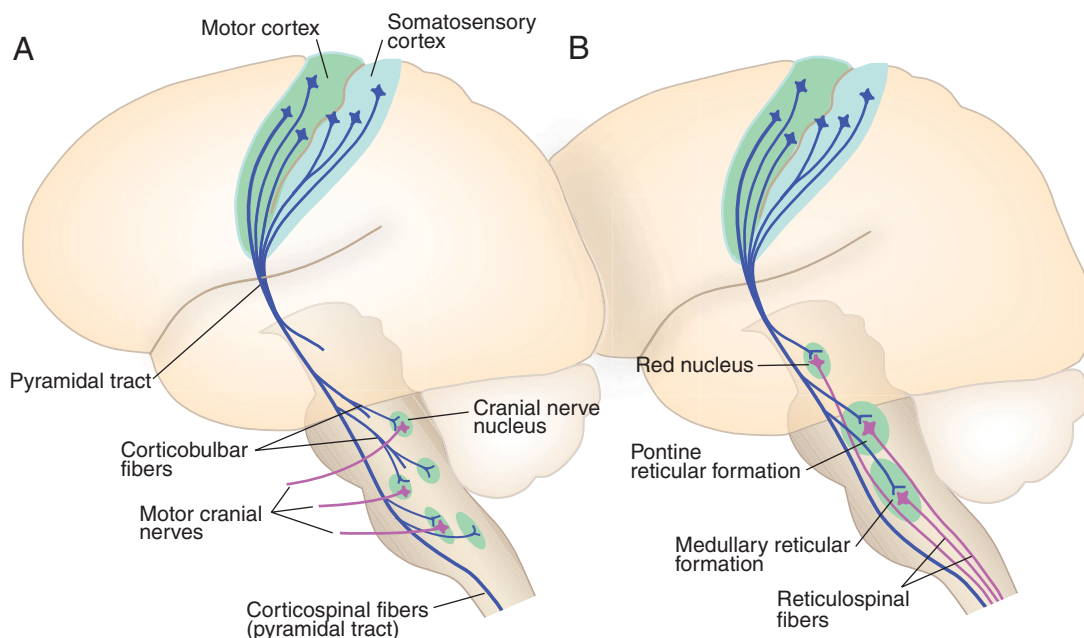


FIGURE 22.1 Direct and indirect motor pathways to the spinal cord. **A:** The pyramidal tract passes directly from the cerebral cortex to the motoneurons in the brain stem (corticobulbar fibers) and in the spinal

cord (corticospinal fibers). **B:** Nuclei in the brain stem with efferent connections acting on motoneurons. Indirect corticospinal pathways are established by corticofugal connections to the brain stem nuclei.

of the medulla and continue downward in the lateral funicle of the cord, to finally establish synaptic contacts in the spinal gray matter.

Fibers also leave the pyramidal tract on their way through the brain stem, to reach cranial nerve motor nuclei (Figs. 22.1 and 22.3). Such fibers form part of the **corticobulbar tract** (other corticobulbar fibers reach the red nucleus, the pontine nuclei, the reticular

formation, the colliculi, the dorsal column nuclei, and other nuclei).

The pyramidal tract derives its name from the **pyramid** of the medulla, which is formed by the fibers of the corticospinal tract (see Figs. 6.15 and 6.16). Strictly speaking, therefore, the term “pyramidal tract” encompasses only the fibers destined for the spinal cord and not those destined for the cranial nerve motor nuclei.

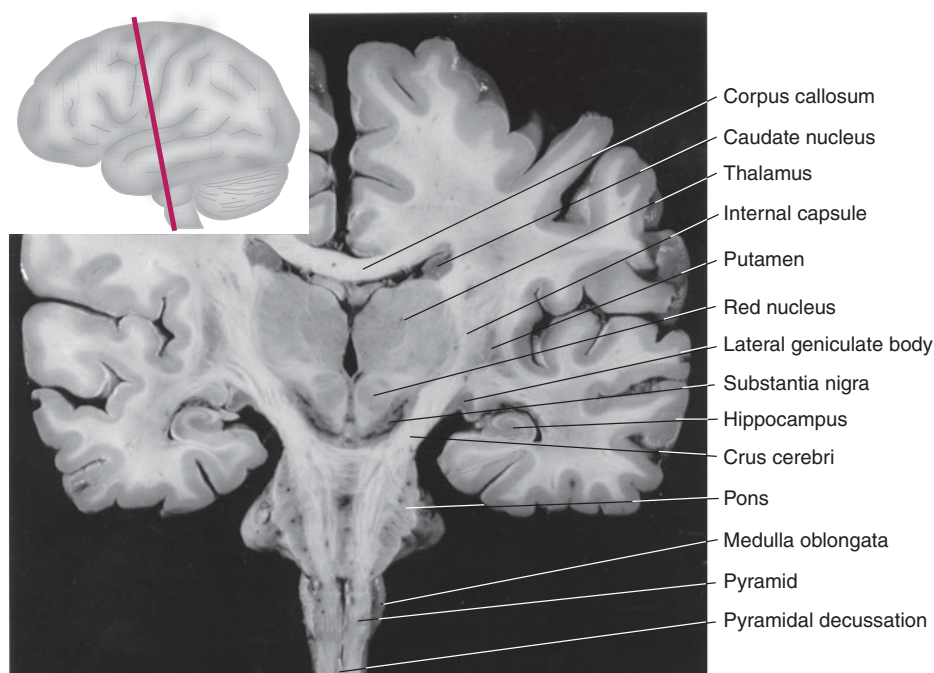


FIGURE 22.2 Course of the pyramidal tract through the internal capsule and the brain stem. Longitudinal bundles of myelinated fibers are evident in the crus, the pons, and the medullary pyramid. Compare with Fig. 22.13.

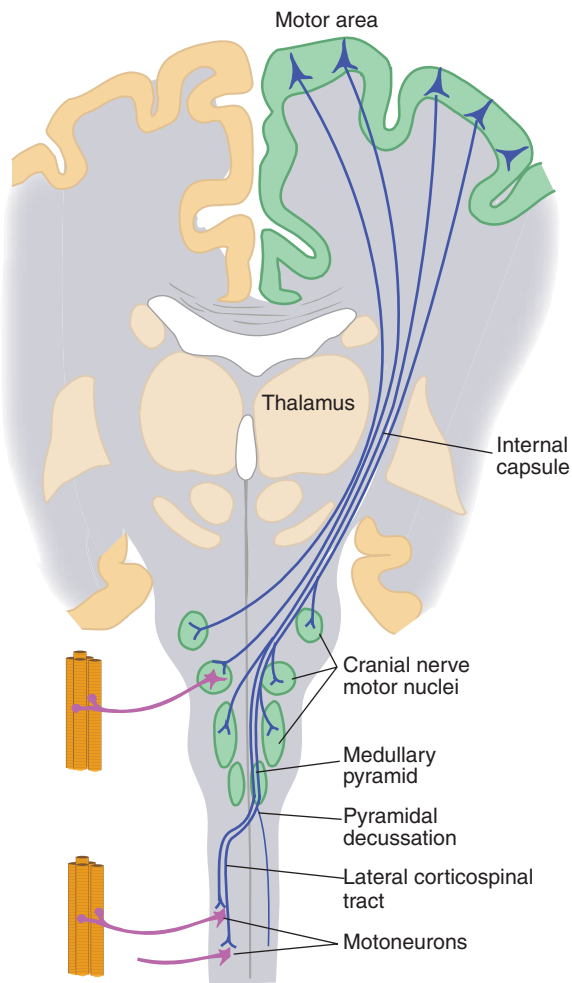


FIGURE 22.3 Direct corticobulbar and corticospinal pathways (the pyramidal tract). The corticospinal tract is mainly crossed, whereas many of the cranial nerve nuclei receive crossed and uncrossed corticobulbar fibers. Note the crossing of the corticospinal tract in the lower medulla.

Nevertheless, for practical reasons both groups are usually included in the term.

Origin of the Pyramidal Tract

A large proportion of the fibers of the pyramidal tract comes from neurons with their cell bodies in the **precentral gyrus**—that is, **area 4** of Brodmann (Figs. 22.4, and 22.5; see also Fig. 33.3). This region was called the **primary motor area (MI)**, because muscle contractions could most easily (with the weakest current) be elicited from this part of the cerebral cortex. The **somatotopical organization** of MI (Fig. 22.5) has been verified in humans with various kinds of stimulation and imaging techniques (electric and magnetic stimulation, positron emission tomography [PET], functional magnetic resonance imaging [fMRI]). It corresponds roughly to the somatotopical pattern in SI (see Fig. 14.8). We return to MI later in this chapter.

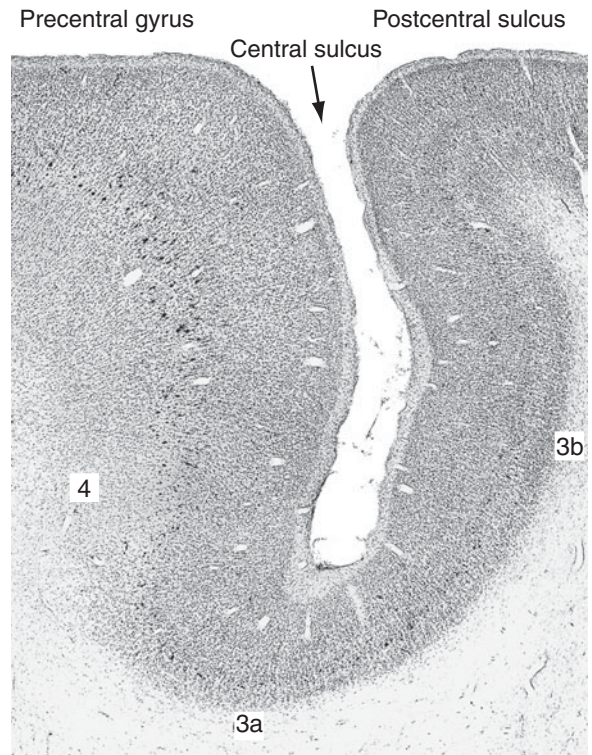


FIGURE 22.4 The central region with MI (primary motor area) and SI (somatosensory area). Photomicrograph of a section perpendicular to the central sulcus (monkey). There is a notable difference in thickness between the MI and the SI. The dark dots in the deep parts of the cortex (in layer 5) are the cell bodies of pyramidal tract cells that have been retrogradely labeled by an injection of a tracer substance (horse-radish peroxidase) in the spinal cord. There are more labeled cells in the MI than in the SI.

It was originally thought that all fibers of the pyramidal tract came from area 4 and, furthermore, only from the cells with the largest cell bodies, the **giant cells of Betz**. The number of Betz cells, however, is much too low to account for the number of axons in the pyramidal tract (about 1 million in humans). More recent studies with retrograde transport of tracer substances (Fig. 22.4) have largely clarified the origin of the pyramidal tract in various animals. In the monkey, numerous cells in area 4, in addition to the Betz cells, contribute to the pyramidal tract, as do also many cells in areas outside area 4. Although the relative contribution from various areas differ among authors, it seems that about two-thirds of all fibers arise in front of the central sulcus—that is, in area 4 and **area 6** (PMA and SMA in Fig. 22.6), whereas the rest comes from **SI** (areas 3, 1, 2), **SII**, and parts of the **posterior parietal cortex** (area 5). A considerable fraction of the fibers from SI arise in **area 3a**—that is, the part of SI adjacent to area 4 (Fig. 22.4) that receives an input from muscle spindles. Muscle contractions can also be elicited from the areas outside MI, like SI, by electrical stimulation, but the stimulation has to be more intense than in area MI. This reflects

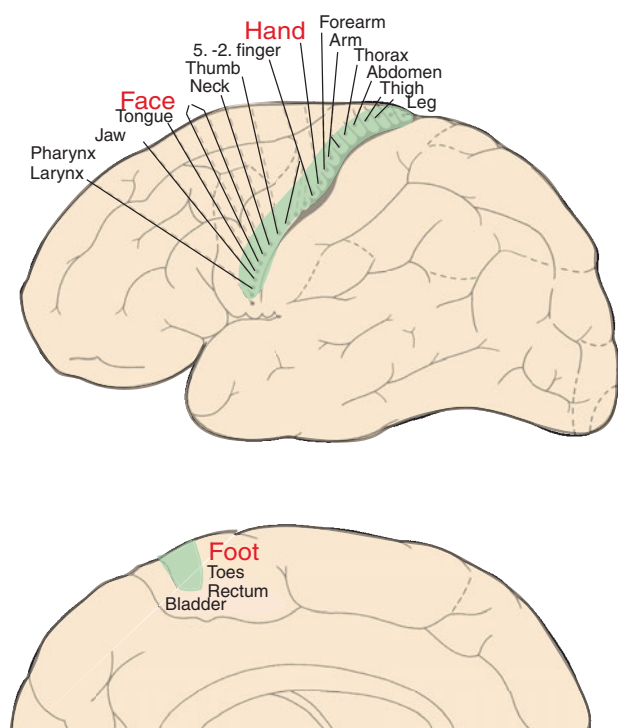


FIGURE 22.5 Somatotopic organization of the motor cortex. The points indicate the sites from which muscle contraction in a particular body part was produced by weak electrical stimulation. (Based on Foerster 1936).

that the pyramidal tract fibers from SI and the other regions outside area 4 have less direct access to the motoneurons than the neurons in area 4. The pyramidal tract fibers coming from SI and area 5 are probably more concerned with the control of sensory signal transmission than with the initiation of movements.

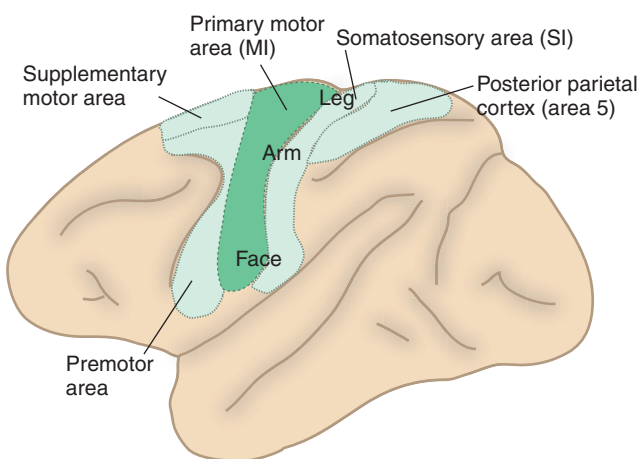


FIGURE 22.6 Cortical regions giving origin to the pyramidal tract. Based on experimental studies in monkeys with retrograde transport of tracer substances (cf. Fig. 22.4). The primary motor cortex (MI) has the highest density of pyramidal tract neurons; that is, neurons that are retrogradely labeled from the spinal cord. Lower densities of labeled neurons occur in SI, SMA, PMA, and the posterior parietal cortex.

The cell bodies of all neurons of the pyramidal tract lie in the cortical **fifth layer** (lamina V; compare Figs. 22.4 and 33.2) and are called **pyramidal cells**. The name refers to the shape of the cell body (see Figs. 1.1, 33.1, and 33.5).

Course and Crossing of the Pyramidal Tract

As mentioned, the pyramidal tract passes downward through the **internal capsule**. It occupies a posterior position (Fig. 22.7A), as judged from observations after small lesions (see Fig. 22.13) and stimulation during brain surgery of the internal capsule in humans. (It was formerly assumed that the pyramidal tract fibers were spread over a large part of the “posterior leg” of the internal capsule.) Fibers governing the muscles of the face are believed to lie most anteriorly, the leg most posteriorly. Experimental studies in monkeys indicate that descending fibers from the PMA and SMA lie in the “knee” region (genu) and the “anterior leg” of the internal capsule, respectively. The internal capsule contains many other fibers than those of the pyramidal tract; the latter constitute only a minority. For example, **sensory fibers** ascend from the thalamus to the cortex, and descending fibers from the cortex reach the thalamus, the reticular formation, and other cell groups of the brain stem. Further, in the anterior part of the internal capsule lie the **pallidothalamic fibers** (related to motor functions of the basal ganglia), and posteriorly (in the lower part) the **optic radiation** passes through on its way to the occipital lobe. Therefore, **damage** of the internal capsule—such as an infarction caused by occlusion of an artery—may produce sensory and other deficits in addition to pareses of the muscles of the opposite body half (capsular hemiplegia).

In the **mesencephalon**, the pyramidal tract fibers distribute over the middle two-thirds of the crus and mix with other descending fibers (Fig. 22.7B). As mentioned, all of the corticospinal fibers are collected within the **medullary pyramid** (Fig. 22.6D; see Fig. 6.17). At the caudalmost level of the medulla, most of the corticospinal fibers **cross** the midline and continue in the **lateral funicle** of the cord as the **lateral corticospinal tract** (Figs. 22.7E and 22.8). A small contingent continues, without crossing, downward in the ventral funicle as the **ventral corticospinal tract**. Some uncrossed fibers may also pass in the lateral funicle (Fig. 22.8).

Pyramidal-tract actions on the muscles of the **distal extremities** are mediated almost exclusively by fibers that cross in the lower medulla. This is supported by studies in humans with transcranial magnetic stimulation (TMS) of the motor cortex. The small contingent of uncrossed (and doubly crossed) fibers mentioned above can influence mainly **axial muscles**—that is, muscles of the back, the thorax, and the abdomen. This is in accordance with their termination medially in the ventral horn (Fig. 22.8; see Fig. 21.3).

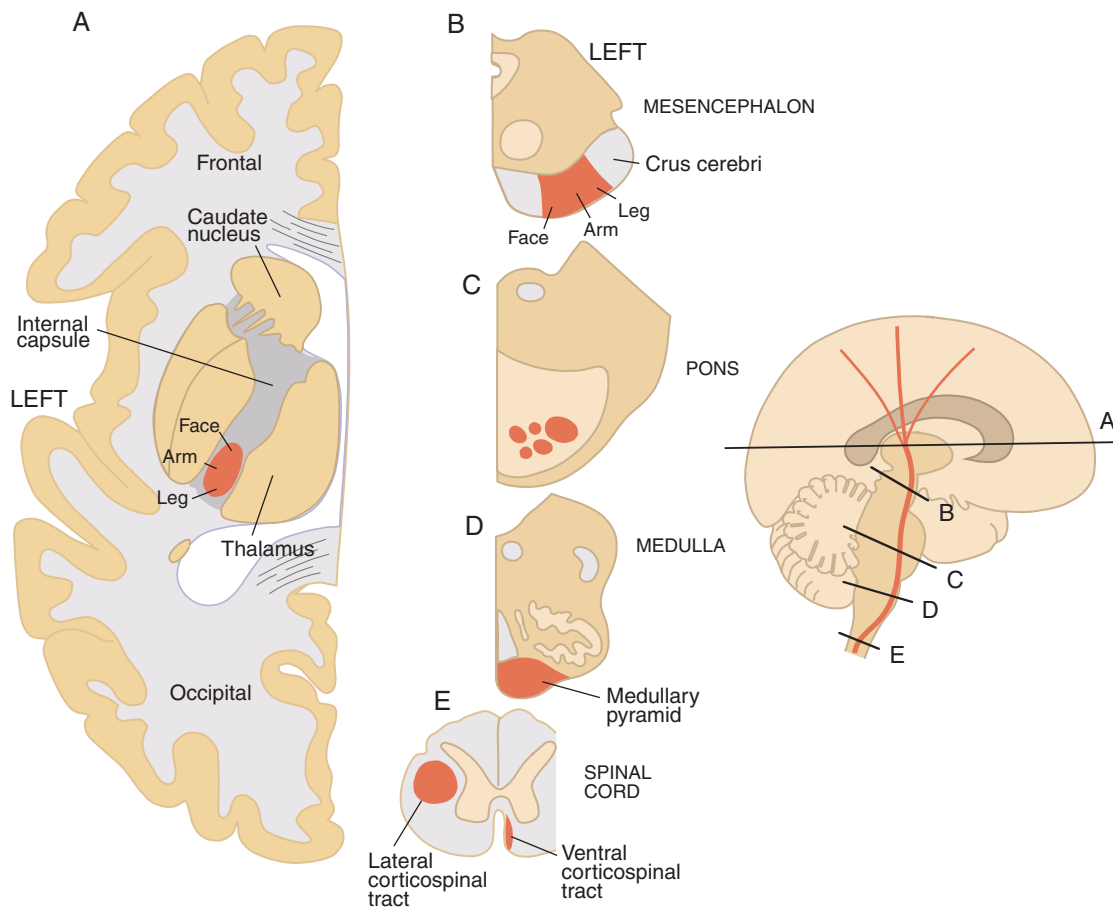


FIGURE 22.7 Position and somatotopic pattern of the pyramidal tract at various levels. **A:** Horizontal section (cf. Fig. 6.30). **B–E:** Transverse

sections. Compare with Figs. 14.2, 14.3, and 14.4 showing the positions of the somatosensory tracts at the same levels.

The pyramidal tract fibers that control the **muscles of the head** (the face, tongue, pharynx, and larynx) leave the corticospinal fibers in the brain stem to end in or close to the motor and sensory cranial nerve nuclei (Fig. 22.3). Apart from most of the facial muscles, other muscles of the head generally receive both crossed and uncrossed pyramidal tract fibers. Thus, when the pyramidal tract fibers are interrupted in the internal capsule, there are seldom clear-cut pareses of the muscles of the tongue, the pharynx, and the larynx, whereas the corner of the mouth hangs down on the opposite side of the lesion.

Individual Variations in the Position and Crossing of the Pyramidal Tract

There are considerable individual variations in both the distribution of the pyramidal tract fibers in the spinal white matter and the percentage of uncrossed fibers. Nathan and coworkers (1990) reexamined these clinically important matters and found, for example, that the corticospinal fibers are more widely distributed in the lateral funicle than is depicted in textbooks. Thus, such fibers are usually found also ventral to the level of

the central canal (Fig. 22.7E is probably reasonably accurate in this respect). Equally important for the interpretation of symptoms in patients with spinal cord injuries is that the anteroposterior location of the lateral tract shows considerable individual variations.

The majority of the corticospinal fibers cross in most individuals, and it is customary to say that there are about **15% uncrossed** fibers. The great individual variations emphasize that this is only an average number, however. Indeed, in a few individuals the pyramidal tract is completely crossed with all fibers located dorsally in the lateral funicle, or it is completely uncrossed with all fibers in the ventral funicle. As a rule, most of the uncrossed fibers continue in the ventral funicle as the ventral corticospinal tract (Fig. 22.5), whereas some join the crossed fibers from the other side in the lateral funicle (Fig. 22.8). Some fibers cross twice, and many individuals have a small crossed, ventral component. The division of uncrossed fibers between the lateral and the ventral funicles varies among individuals.

Finally, although usually not acknowledged, in about 75% of the population the pyramidal tract is **asymmetric**: the lateral and the ventral components are both larger on one side than on the other (most often on the right side).

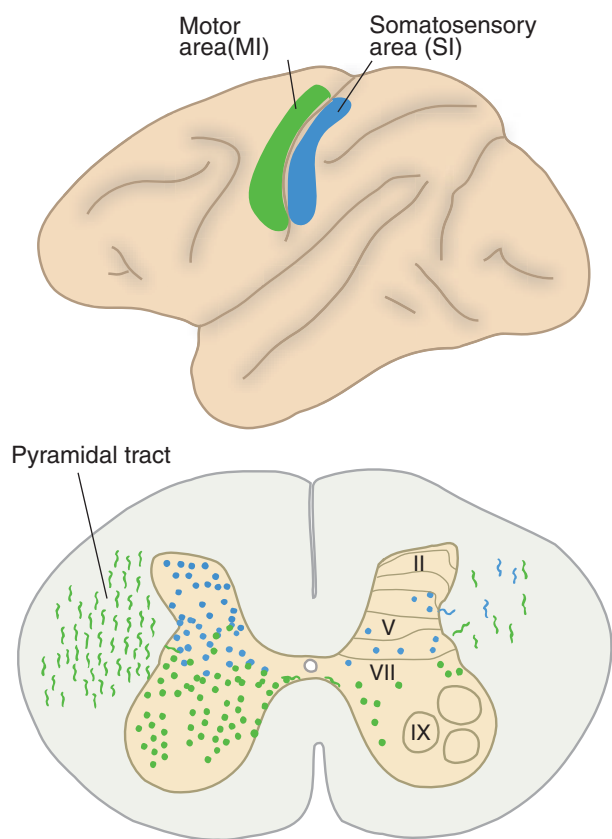


FIGURE 22.8 *Terminal regions of the pyramidal tract.* Based on experiments in monkeys with anterograde transport of radioactively labeled amino acids after injections in various parts of the motor cortex (MI) and the somatosensory cortex (SI). Corticospinal fibers from the MI end more anteriorly in the gray matter of the cord than those from the SI. The uncrossed fibers end predominantly medially in the ventral horn—that is, in contact with the motoneurons that supply axial and proximal muscles. (Based on Ralston and Ralston 1985.)

This asymmetry presumably arises because a larger proportion of the fibers from (usually) the left hemisphere cross than from the right hemisphere. Thus, the lateral (crossed) tract becomes larger on the right side than on the left, whereas the ventral (uncrossed) tract becomes smaller on the left side.

Such individual variations would influence the severity of motor symptoms after damage to the brain or spinal cord and may help explain why symptoms vary so much among patients with very similar lesions. Thus, occasionally an infarct of the internal capsule produces most severe ipsilateral pareses (on the same side), and lesions of the cord may give motor symptoms that are unexpected from their location.

Conduction Velocity and Termination of the Pyramidal Tract

The pyramidal tract fibers vary considerably in thickness, from the thickest myelinated kind to unmyelinated ones. Most are rather thin; with **conduction velocities**

between 5 and 30 m/sec. Corticospinal neurons are **glutamatergic**, exerting excitatory effects. In general, the thickest (and thus fastest conducting) fibers come from MI.

In accordance with the **somatotopic pattern** within MI (Fig. 22.5), fibers from the medial parts of the precentral gyrus (leg representation) end in the lumbosacral part of the cord, whereas fibers from more lateral parts (arm representation) end in the cervical and upper thoracic cord. Fibers from MI and SI also terminate differently in the cord (Fig. 22.8). Thus, fibers from SI end predominantly in the dorsal horn, whereas fibers from MI end in the intermediate zone and the ventral horn (laminae VII–IX).

Monosynaptic Cortico-Motoneuronal Connections

Anatomic and physiologic data show that some of the pyramidal tract fibers coming from MI end **monosynaptically** on the motoneurons. This concerns primarily motoneuron groups that control the **distal muscles** of the extremities—in particular, the intrinsic muscles of the **hand**. During evolution, the pyramidal tract has increased in size and with regard to monosynaptic connections with the motoneurons. This change has taken place in parallel with increased versatility and precision of the movements of the hand (which, in turn, depend on increased brain volume). In the rat and the rabbit, for example, the pyramidal tract is only slightly developed. It is more prominent in the cat, but there are no monosynaptic connections between pyramidal tract neurons in the motor cortex and motoneurons in the cord—all connections are with spinal interneurons. In monkeys, there is a modest proportion of monosynaptic connections, but the proportion is larger in the anthropoid apes (e.g., the chimpanzee) and still larger in humans. Even some shoulder muscles, like the deltoid, receive some monosynaptic connections in humans. The monosynaptic connections are of particular importance for the movements that require the highest degree of voluntary control—the **least automatic movements**—such as independent or **fractionated finger movements**. This relationship is witnessed by the fact that in certain subprimate mammals with an exceptional manual dexterity (and the ability to use the fingers individually), some monosynaptic motoneuronal connections are present.

Distribution and Actions of Single Corticospinal Fibers

A corticospinal neuron in MI usually has strong **excitatory** actions on some motoneurons and weaker excitatory actions or **inhibitory** actions (via interneurons) on many others. Accordingly, anatomic data from tracing of single fibers show **widespread ramifications** of a single pyramidal tract axon, enabling it to contact numerous motoneurons in different segments of the cord that

supply several muscles. This may appear surprising in light of the very discrete and precise movements the pyramidal tract is capable of producing. However, as a rule, the various muscles contacted from one corticospinal neuron have similar actions—that is, they are synergists. Further, the strength of the influence of one pyramidal tract cell probably varies greatly among the motoneurons it contacts, depending on the number and location of synaptic contacts. For example, corticospinal fibers terminate on both proximal and distal parts of the dendrites. Contacts with **inhibitory interneurons** further increase the specificity of actions. In many voluntary movements, commands from the motor cortex elicit **reciprocal inhibition** in the cord. Such varied effects of the pyramidal tract fibers fit with the motor cortex being organized to initiate **purposeful movements**, requiring coordinated activity of multiple muscles around several joints (rather than the contraction of muscles in isolation). This topic is discussed further under “Functional Organization of the Primary Motor Area.”

The Pyramidal Tract Controls Flexors More than Extensors

Experiments with transcranial magnetic stimulation (TMS) of the motor cortex show that **flexors** of the thumb have a **lower threshold** for activation than the extensors. Corresponding findings pertain to the biceps muscle (flexor) and the triceps muscle (extensor) of the upper arm. At least concerning finger movements, it seems reasonable that the flexors require somewhat more precise control than the extensors, although both certainly are necessary for manual dexterity (e.g., holding and manipulating small objects). Interestingly, fMRI studies indicate that a smaller part of MI is activated during simple finger flexions than during finger extensions. Presumably, this may be due to more monosynaptic connections to the flexor motoneurons.

The functional flexors of the ankle joint (the tibialis anterior muscle and others that dorsiflex the ankle) are more influenced by the pyramidal tract than are the plantar flexors (the triceps surae and others). This would seem to fit with clinical observations in patients with **hemiplegia**: ankle dorsiflexion is more reduced in power than plantar flexion (the foot droops). However, the typical posture of the hemiplegic arm (Fig. 22.11)—with tonic flexion at the elbow—can obviously not be explained in this way.

The Pyramidal Tract Controls Sensory Neurons in the Spinal Cord

The effects of SI (via fibers traveling in the pyramidal tract) on neurons in the dorsal horn are quite specific. Thus, projections differ from individual cytoarchitectonic subdivisions within SI: area 3b—receiving input from

cutaneous low-threshold mechanoreceptors—sends fibers primarily to laminae III and IV, which receive the same kind of sensory input (through the dorsal roots). Fibers from area 3a end deeper in the dorsal horn, where primary afferents from proprioceptors end. Physiological experiments confirm that SI influences sensory cells in the dorsal horn. Most often, SI inhibits neurons that are excited from low-threshold mechanoreceptors. As discussed in Chapter 14 (under “The Transmission of Sensory Signals Can Be Controlled from the Brain”), suppression of certain kinds of sensory information may be important, for example, during movements.

The Pyramidal Tract Fibers Open and Close Spinal Reflex Arcs

Even in humans, a large proportion of the pyramidal tract fibers end in synaptic contact with **interneurons** in laminae VII and VIII (Fig. 22.8; see also Fig. 6.12). Many—perhaps most—of these interneurons make synaptic contacts either on motoneurons or on other interneurons, which, in turn, contact motoneurons. In this manner, the pyramidal tract may mediate varied effects on the motoneurons. This is true of both the α motoneurons and the γ **motoneurons**, so that the pyramidal tract can control the sensitivity of the **muscle spindle**. Because many of the interneurons contacted by pyramidal tract fibers are intercalated in reflex arcs, the motor cortex can ensure that the various spinal reflexes are adapted to the overall aim of the movements. For example (as described in Chapter 21), the strength of the **long-latency stretch** reflex can be increased or decreased, depending on what is appropriate during motor learning. The interneurons mediating **reciprocal inhibition** during stretch reflexes (see Fig. 21.13) are also controlled by the pyramidal tract. Likewise, the **Renshaw cells** (see Fig. 21.14) are subject to supraspinal control, so that the strength of the recurrent inhibition can be increased or decreased selectively in various motoneuronal groups. A final example is the interneurons that mediate inhibition of motoneurons evoked by stimulation of **tendon organs** (in the same muscle as supplied by the motoneurons). This reflex arc can also be “opened” or “closed” by the pyramidal tract. Thus, the **autogenic inhibition** elicited by stimulation of tendon organs can be reversed to excitation during voluntary contraction of the muscle (whereas at the same time the inhibition of other muscles may be enhanced).

Function of the Pyramidal Tract as Judged from Functional Deficits after Lesions

To clarify the function of a tract, two approaches have mainly been used: studying (1) the properties of single neurons and (2) the deficits that ensue when the entire tract is eliminated. We discussed in the preceding text

some properties of single pyramidal tract neurons. Their strong (partly monosynaptic) connections with distal muscles suggest that the pyramidal tract is particularly important for delicate movements—for example, of the fingers. We discuss further the properties of pyramidal tract neurons under “Functional Organization of the Primary Motor Area.” Here we restrict ourselves to what we have learned from lesion experiments in animals and from humans with diseases that affect the pyramidal tract.

Shortly after unilateral **transection of the pyramid**, a monkey can move around apparently normally.¹ Nevertheless, to use the hand to pick up a morsel of food, for example, is almost impossible for some time, even though the hand can be used effectively for climbing. Initially, the monkey is unable to move the hand independently of the arm. Gradually, over some weeks, the monkey regains the ability to use the hand for grasping objects, but the movements are clumsy and consist only of simultaneous flexion of all fingers (a sort of scraping movement). However, the ability to move the fingers independently of each other—**fractionated movements**—does not return (even after 5 years). The precision grip, in which the thumb opposes the index finger, is permanently lost. Monkeys in which the pyramid is cut shortly after birth never develop the precision grip. Thus, no other tracts can take over this task from the pyramidal tract.

In **humans** (as in monkeys) with damage of the motor cortex or the pyramidal tract (often in the internal capsule), the most enduring symptom is difficulty with tasks that require precise, fractionated finger movements, such as writing, tying shoelaces, buttoning a shirt, and picking up small objects (like a needle). Less precise movements involving larger muscle groups are usually less severely affected. In addition, lesions of the upper motor neurons in humans produce characteristic changes of **muscle tone** and **reflexes** (spasticity), which do not occur in monkeys with lesions restricted to the pyramidal tract.² We return to the symptoms produced by lesions of the descending motor pathways in humans at the end of this chapter.

INDIRECT CORTICOSPINAL PATHWAYS

In addition to the pyramidal tract, several pathways mediate signals from higher levels of the central nervous

system to the motoneurons in the cranial nerve nuclei and the spinal cord. The nuclei giving origin to these other tracts (often included in the “extrapyramidal system”) are located in the brain stem, but several of them receive afferents from the cerebral cortex—in particular, from the motor cortex (Fig. 22.1). Thus, several **indirect pathways**—synaptically interrupted in the brain stem—transmit signals from the cortex to the motoneurons. In addition, some other brain stem nuclei with descending connections to motoneurons do not receive fibers from the motor cortex. The latter nuclei are primarily involved in control of highly automatic muscle contractions, such as those aiming at maintaining body balance and the movements of respiration.

Broadly speaking, the pyramidal tract is important primarily for the least automatic movements—that is, those requiring a high degree of conscious, voluntary attention. The other pathways are important for movements that are more automatic. This dichotomy agrees with the fact that the pyramidal tract is primarily important for the movements of the distal parts of the extremities, whereas the other pathways are more concerned with movements of the proximal parts of the extremities and of the trunk. Such a task division does not imply, however, that these two “systems”—the **direct** and the **indirect corticospinal pathways**—operate independently. Most movements, even delicate finger movements, require that the hand is moved to the right position and is kept in a specific posture. Further, even small movements of the arm displace the center of gravity, requiring postural adjustments by many other muscles.

Figure 22.9 shows the main brain stem nuclei sending fibers to the spinal cord,³ while Fig. 22.1B shows connections from the motor and somatosensory areas to the reticular formation and some other nuclei. In addition, monoaminergic spinal pathways from the locus coeruleus and the raphe nuclei are included in Fig. 22.9. The latter pathways do not participate in control of specific movements but modulate the excitability in spinal neuronal networks.

Corticoreticulospinal Pathways

The **reticular formation** is an important source of descending fibers that can influence the motoneurons. The reticular formation is discussed more completely in Chapter 26 (its functions are not restricted to motor control). **Reticulospinal** fibers arise primarily from the reticular formation of the pons and the medulla (Fig. 22.9; see also Figs. 26.1 and 26.8). These regions receive **corticoreticular fibers** from the cerebral cortex, and especially from the motor areas (areas 4 and 6). Thus, there is a **corticoreticulospinal pathway** that can mediate

1 Certainly, the pyramidal tract is more developed in humans than in monkeys. Thus, the possibility cannot be excluded that symptoms would be somewhat more pronounced in humans after a corresponding lesion (as they are far more pronounced in a monkey than in a cat).

2 In humans with lesions of the pyramidal tract, there is usually damage to other structures as well. As is well known, the fibers of the pyramidal tract are isolated only in the pyramid (only a few patients have been described with infarcts limited to the medullary pyramid). In the internal capsule, the mesencephalon (crus), the pons, and in the cord, it either is mixed with or lies in close apposition to other tracts.

3 Sensitive tracing techniques have demonstrated several minor spinal projections from the brain stem in addition to the major ones described here.

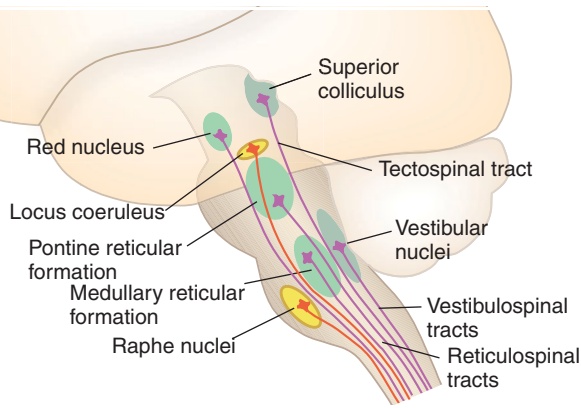


FIGURE 22.9 The major brain stem nuclei sending fibers to the spinal cord. Some of the nuclei receive afferents from the cerebral cortex (cf. Fig. 22.1). The monoaminergic nuclei are yellow.

signals from the cortical motor areas to the spinal motoneurons (see Fig. 26.8).

It is impossible to transect the reticulospinal tracts in the white matter of the spinal cord without at the same time involving other tracts. This has made it difficult to clarify the role of the reticulospinal tracts in motor control. Nevertheless, experiments with relatively selective lesions of the reticular formation indicate that the reticulospinal tracts are important for maintaining the **upright position** (posture), for **orienting movements** of the body toward external events, and for fairly crude, **stereotyped voluntary movements** of the extremities (such as extending the arm toward an object). At least in monkeys, the pyramidal tract is not necessary for such movements, as described earlier. The crude movements of the extremities are most likely mediated by pathways from the motor cortex via the reticular formation and the reticulospinal tracts (see Fig. 26.8).

Because reticulospinal neurons receive inputs from sources besides the motor cortex—such as the vestibular nuclei, the basal ganglia, and the cerebellum—many cell groups cooperate to produce the final activity of reticulospinal neurons.

More about Reticulospinal Tracts

The actions in the cord of the reticulospinal fibers are multifarious. In Chapter 15, we described reticulospinal fibers descending in the dorsolateral fascicle and ending dorsally in the spinal gray matter (see Figs. 15.2 and 15.3), with powerful effects on the central transmission of signals from nociceptors. The reticulospinal fibers that are of interest in relation to motor control terminate more ventrally in the spinal gray matter, with monosynaptic and polysynaptic effects on the motoneurons. These fibers descend in the ventral part of the lateral funicle and in the ventral funicle. They belong to at least two tracts with different sites of origin in the reticular

formation and somewhat different sites of termination in the cord.

Fibers from the **pontine** and **medullary** parts of the reticular formation (Fig. 22.9) run in different parts of the white matter: the pontine fibers are located in the ventral funicle (see Fig. 16.8), whereas the medullary fibers are in the ventral part of the lateral funicle. Together, reticulospinal fibers terminate in large parts of the ventral horn but primarily in medial parts, where the motoneurons to the axial muscles and the proximal muscles of the extremities are located. Connections are particularly ample to the motoneurons that innervate **neck muscles**, which are important for movements of the head. The medullary fibers terminate somewhat more laterally in the ventral horn (particularly in lamina VII but to some extent also in lamina IX) than the pontine fibers. The latter terminate mostly more medially in laminae VII and VIII. The **medullary reticulospinal tract** may thus be expected to have access to motoneurons innervating muscles of the **extremities**, whereas the **pontine reticulospinal tract** reaches mostly motoneurons of **axial muscles** (neck, back, and abdomen).

Many of the reticulospinal fibers send **collaterals** to both cervical and lumbar segments of the spinal cord and often to the ventral horns of both sides. Thus, such neurons are capable of influencing numerous muscles at the same time. This may be functionally meaningful in relation to adjustment of posture and body balance. Nevertheless, there is also some degree of **somatotopic** organization among reticulospinal neurons, as demonstrated with retrograde axonal transport methods. Also, electrical stimulation of the reticular formation can elicit muscle contractions restricted to smaller parts of the body. Together, these data indicate that the reticulospinal fibers can mediate both rather diffuse effects on many muscle groups and more focused influences related to specific, goal-directed movements.

Both excitatory and inhibitory effects on the motoneurons can be elicited by **electrical stimulation** of the reticular formation. **Inhibition** of reflex and voluntary movements can be produced by stimulation of a region in the lower part of the medulla (see Fig. 26.10), whereas increased reflex movements and muscle tone can be elicited from a more rostral region. The reticulospinal fibers act on both the α and γ **motoneurons**. Thus, the reticular formation also controls the sensitivity of the **muscle spindles**, and in certain situations it is probable that it acts only on the γ motoneurons and not the α motoneurons. There is even some evidence that the reticular formation may act selectively on either static or dynamic γ motoneurons.

The Tectospinal Tract

Several mesencephalic cell groups send axons to the spinal cord. A large number of fibers arise in the **superior**

colliculus and form the **tectospinal tract**. The descending fibers cross the midline shortly below the superior colliculus, and most terminate at **cervical levels** of the cord. The superior colliculus receives numerous fibers from the retina, the visual cortex, and from the so-called **frontal eye field** (area 8 in Fig. 25.7), which are of particular importance for control of conjugate eye movements. Efferent fibers from the superior colliculus do not act solely on the cord; motoneurons in the brain stem innervating extraocular muscles are also influenced (although indirectly via the reticular formation). In agreement with these anatomic data, electrical stimulation of the superior colliculus in experimental animals produces coordinated movements of the eyes and the head. The tectospinal tract is of particular importance for movements of the head and the eyes as parts of **optic reflexes**: that is, the head and the eyes are directed toward something in the visual field. (Auditory signals also reach the superior colliculus via the inferior colliculus and can thus elicit head movements.)

The superior colliculus also receives afferents from nonvisual parts of the cortex, like the **SI** and **MI**. Thus, a **corticotectospinal pathway** may perhaps play a part in voluntary movements.

Vestibulospinal Tracts

Primary sensory fibers from the vestibular apparatus terminate in the **vestibular nuclei**, which is located in the pons and medulla (see Fig. 9.7). Vestibular signals about the position and movements of the head also indirectly provide information about the position of the body and about disturbances in balance. Two tracts, issued from the vestibular nuclei to the spinal cord, can contribute to the maintenance of body balance and posture. The largest is the **lateral vestibulospinal tract**, which comes from the lateral vestibular (Deiters) nucleus and reaches all levels of the cord. The tract lies in the **ventral funicle**. The fibers exert an **excitatory** action on both α and γ **motoneurons**. Like the reticulospinal fibers, the vestibulospinal ones act primarily on motoneurons in the medial parts of the ventral horn—that is, **axial muscles** and **proximal muscles** of the extremities. Thus, the lateral vestibulospinal tract can adjust the contraction of muscles that oppose the force of gravity (antigravity muscles).

The other vestibulospinal tract is much smaller and reaches only the cervical and upper thoracic segments of the cord. This **medial vestibulospinal tract** is therefore primarily important for mediation of **reflex head movements** in response to vestibular stimuli. This conclusion is also supported by physiological experiments. Unlike the lateral vestibulospinal tract, many of the neurons of the medial tract are **inhibitory** (probably using **glycine** as a transmitter).

In contrast to the other cell groups in the brain stem sending fibers to the cord, the vestibular nuclei receive few afferents from the cerebral cortex. The vestibulospinal neurons are therefore more independent of the cerebral cortex than, for example, the reticulospinal neurons. The vestibular nuclei mediate primarily **automatic, reflex movements** and adjustments of muscle tone. Nevertheless, the activity of the vestibular nuclei may be influenced indirectly from the cortex, as they receive afferents from the reticular formation. For example, the influence of vision on automatic adjustments of posture appears to be mediated via the vestibular nuclei.

The Red Nucleus and the Rubrospinal Tract

The red nucleus (nucleus ruber) in the mesencephalon (see Figs. 6.20 and 6.21) consists of a caudal **magnocellular** part (large, motoneuron-like neurons) and a rostral **parvocellular** (small-celled) part. In monkeys the magnocellular part is quite small, and in humans it contains even fewer neurons. The large parvocellular part receives its main afferents from the **cerebellum** (especially from the dentate nucleus, receiving its main afferents from the cerebellar hemispheres). Many (most?) of the efferents from the **parvocellular** part goes to the **inferior olive** (which sends its efferent fiber to the cerebellum). By way of cerebellar pathways to the motor cortex, it appears that the parvocellular red nucleus influences movements primarily through its interplay with the cerebellum and the motor cortex, not by sending fibers to the spinal cord.

The **magnocellular** red nucleus sends fibers to the spinal cord, forming the **rubrospinal tract**. The tract crosses the midline just below the red nucleus and descends in the lateral funicle, mixed with the fibers of the pyramidal tract. The rubrospinal tract is somatotopically organized. In the cat and the monkey, the fibers terminate in largely the same parts of the spinal gray matter as the pyramidal tract and have their most marked effect on **flexor motoneurons** of distal muscles (also like the pyramidal tract). The red nucleus receives fibers from the motor cortex of the same side. A **corticorubrospinal** pathway is thus established from the cerebral cortex to the spinal motoneurons. In the cat and monkey this pathway supplements the pyramidal tract in the control of voluntary movements (although it is of greater functional importance in the cat than in the monkey). Whether it plays such a role in humans seems unlikely because the magnocellular part is so small, and the parvocellular part does not appear to project to the spinal cord (in monkey and humans). In monkeys, the ratio of rubrospinal to corticospinal fibers has been estimated to about 1:100, and this ratio is most likely even lower in humans.

Monoaminergic Pathways from the Brain Stem to the Spinal Cord

Although they are often included among the reticulospinal pathways, descending monoaminergic fibers to the cord from the **raphe nuclei** (Fig. 22.9; see also Figs. 26.6 and 26.7), the **locus coeruleus** (see Fig. 15.6), and scattered cell groups in their vicinity have distinct properties. Such monoaminergic fibers terminate in the ventral horn (in addition to in the dorsal horn). Many of the **raphespinal** fibers contain **serotonin**, whereas the **coeruleospinal** fibers contain **norepinephrine**. In addition, several **neuropeptides** coexist with the monoamines. Both these tracts end rather diffusely in the spinal gray matter and can hardly mediate information related to specific movements. More likely, they exert general, widespread **facilitatory** influences on the motoneurons, as judged from the effects on spinal motoneurons of microinjections of serotonin and norepinephrine. Thus, the excitability of most motoneurons may be enhanced, so that they react more vigorously to an input from the pathways that mediate specific motor commands (such as the pyramidal tract). At the same time, the descending monoaminergic connections to the **dorsal horn** may prevent “disturbing” signals from nociceptors from reaching consciousness. In deep **sleep** (REM sleep), all movements are suppressed, and the descending monoaminergic neurons show their lowest activity.

It is possible that these monoaminergic pathways contribute to the effects of **motivation** on the performance of voluntary movements when the muscle contractions occur with increased speed and force. This may be related to the ability of the monoamines (serotonin in particular) to alter the excitability of the motoneurons by inducing a so-called **plateau potential** (see Chapter 21, under “Muscle Cramps and Plateau Potentials”). Thus, serotonin (and other transmitters) can bring the motoneuron from a state of stable hyperpolarization with low excitability to stable depolarization (plateau potential). In this state, a brief excitatory input elicits tonic firing from seconds to minutes. Plateau potentials may also be an efficient means to control the activity of muscles used for **postural tasks** because they exhibit long periods of tonic contractions (at least in experimental animals). A brief inhibitory input suffices to terminate the tonic firing.

CONTROL OF AUTOMATIC MOVEMENTS

In this section, we treat the control of two kinds of automatic movement: the postural reflexes and locomotion. Postural control and control of movements, either automatic or voluntary, are not independent, however, since postural control is a prerequisite for proper execution of virtually every purposeful movement. Postural reflexes

were treated in more depth in Chapter 18 in conjunction with the control of body balance.

The Cerebral Cortex Coordinates Brain Stem and Spinal Networks

Brain stem and spinal networks can on their own control quite complex movement sequences; the contribution of the cerebral cortex being restricted to start and stop signals. In general, however, the cerebral cortex, the basal ganglia, and the cerebellum are needed to incorporate the activity of the lower networks in an overall plan. To achieve this, the **indirect corticospinal pathways** are instrumental in coordinating the various automatic responses initiated from the lower levels. This happens mainly by modulating the excitability of brain stem and spinal interneurons. Further, most purposeful movements challenge our balance by moving the center of gravity. When learning a sequence of movements, the higher levels make internal models that, among other things, anticipate the postural perturbations caused by the desired movements. Thus, such **anticipatory control** is an important characteristic of skilled and of harmonious movements.

Postural Reflexes

To maintain equilibrium we need rapid corrections of muscle tension in various parts of the body. Postural reflexes produce the automatic movements that help us regain equilibrium quickly—for example, when slipping on ice. It is a common experience that these compensatory movements happen so rapidly that only afterward are we aware of which movements we performed. The tasks of the postural reflexes are to maintain an appropriate posture of the body, to help regain equilibrium when it is disturbed, and to ensure optimal starting positions for the execution of specific movements. Large perturbations may require additional voluntary movements (coming later than the automatic ones). To issue the right motor commands, the brain must receive immediate and reliable information as soon as something threatens our upright position—that is, information from receptors that detect joint movements and movements of the whole body. Because our **upright position** is **labile**—with a small supporting area and a high center of gravity—constant corrections are necessary. For example, we sway a little back and forth in quiet standing, as an expression of imperceptible postural corrections (reflex responses).

Receptors that provide information used for postural control are **proprioceptors** in the legs, the spine, and the neck; **cutaneous** receptors on the sole of the foot; **vestibular receptors** in the inner ear; and photoreceptors in the **retina**. The more demanding the challenge to the

balance becomes, the more important becomes the ability to use several kinds of sensory information. The signals from the various receptors are integrated at the **brain stem** level (in the vestibular nuclei and the reticular formation) and in the **cerebral cortex**. Presumably, the sensory information is compared with an **internal model** of the motor task facing us. The sensory information serves to update the internal representations so they fit the present situation. Internal models are thought to serve as frames of reference for the continuous monitoring and adjustment of our posture. Selection of an overall motor response is done on the basis of the central processing of the sensory information. However, the final motor command depends not only on the sensory signals but also on the **context** in which they arise. Therefore, a certain sensory input can elicit very different postural responses in different situations. Mainly **reticulospinal** and **vestibulospinal** fiber tracts mediate the final postural commands.

The postural reflexes operate on a **feedback** basis: they are responses to movements that have already started. In conjunction with voluntary movements, however, commands about postural adjustments are issued in **advance**. From prior experience, the brain has determined which adjustments will be needed to keep balance during, for example, an arm movement. This is called **feedforward** or **anticipatory** control and is presumably based on the presence of internal models for well-rehearsed movements.

Development of Postural Control

Postural control is developed during childhood, starting as soon as the balance is challenged by the upright position. Although anticipatory control probably starts to develop very early, the first attempts at postural corrections depend heavily on feedback information. The corrections are large and inaccurate (ballistic). By constantly challenging the limits of its balance, the child improves both its use of feedback information and anticipatory control—improving skills by **experimentation**. This presumably goes together with establishing and refining the internal models of various skills. The development is not monotonous; there appear to be alternating periods with overreliance on one or the other strategy until the adult pattern is established around age 10 to 12. The adult pattern is characterized by flexible postural responses based on a full integration of feedback and anticipatory control strategies.

The Grasp Reflex

The grasp reflex is usually regarded as one of the postural reflexes. It is normally expressed only in infants during the first 6 months and has disappeared at 12 months. It consists of the fingers firmly grasping an

object that touches the palm. After damage of the frontal lobes (presumably, in particular, the **supplementary motor area, SMA**), the reflex may reappear in adults. The strength of the grasp reflex depends on the position of the body, reflecting that its purpose is to cling to the mother when she moves about (thus, the reflex is of less functional importance in humans than in monkeys).

Control of Locomotion and Rhythm Generators

The upright locomotion of humans requires somewhat different movements than in animals walking on four legs; not the least are that the demands on postural control are quite different. Certainly, the large size of the human brain has enabled many processes—which in lower animals are controlled from the spinal cord and the brain stem—to be controlled from the cerebral cortex. Nevertheless, with regard to basic mechanisms, the neural control of locomotion is probably very similar in humans and in other mammals. The following account is largely based on data obtained from animal experiments.

Fairly normal locomotion can be produced in animals (such as cats) even when the spinal cord is isolated from the brain stem and the brain. This can be observed on a treadmill when the paws touch the moving ground (the animal has to be supported since there are no postural reflexes). Central **rhythm generators** must, therefore, exist in the cord, able to produce rhythmic, alternating leg movements in the absence of any command signals from higher levels. The rhythm generators consist of fairly complicated spinal **networks** of interneurons with excitatory and inhibitory interconnections, eventually controlling the activity of the motoneurons. Some of the neurons within the network appear to have **pacemaker properties**; that is, they fire brief trains of action potentials with silent periods between, without receiving a rhythmically alternating input.

The rhythm generator does not depend on sensory input from the moving parts to produce the motoneuron activity typical of locomotion. Thus, the rhythmic motoneuronal activity continues even after complete paralysis of the muscles (e.g., after cutting the ventral roots). This is called **fictive locomotion**, and the pattern of motoneuronal activity is strikingly similar to that observed during normal walking in intact animals. This does not mean that sensory inputs are without significance for locomotor control, however. The activity of the rhythm generators can indeed be modified by **sensory signals** from the peripheral receptors providing information about how the movements are proceeding. **Inhibitory interneurons** can contribute to the rhythmic pattern of activity by inhibiting the antagonists when the agonists have reached their maximal activity. **Renshaw cells** (see Fig. 21.14) can probably contribute by shortening impulse trains from the motoneurons and at the same time increasing the excitability of

antagonist motoneurons. There is probably one rhythm generator for each extremity. Long **propriospinal fibers** that interconnect the forelimb and hindlimb generators ensure that their activities are coordinated.

The presence of rhythm generators in the **human** spinal cord is indicated by the occurrence of locomotion-like movements in **anencephalic infants** (born without most of the brain). In normal infants, walking movements of the legs can be elicited in the first few months (if the infant is held under the arms and the feet are made to touch the floor). This ability usually disappears, to reappear when the infant starts to crawl at about 8 months. Locomotor movements can be elicited in patients with complete **transverse lesions** of the cord if the body is supported and the feet hit a treadmill. The locomotor pattern is more normal with high than with low spinal lesions, suggesting that the network controlling human walking is not restricted to the lumbosacral cord. Presumably, the lumbosacral rhythm generators depend on cooperation with similar networks in the cervical cord (as in four-legged animals).

Central Control of Locomotor Movements

Several parts of the brain contribute to the control of the spinal rhythm generators. The most direct influence comes from parts of the reticular formation, but also the basal ganglia, the cerebellum, and the cerebral cortex contribute. Electric stimulation of a region in the mesencephalic reticular formation produces rhythmic locomotor movements in animals.⁴ This **mesencephalic locomotor region** (MLR) is situated just ventral to the inferior colliculus on the pontomesencephalic junction. The **pedunculopontine nucleus** (PPN) is another name applied to this part of the mesencephalon (although the MLR and PPN are probably overlapping but not identical). Most likely, the effects on spinal motoneurons from the MLR are mediated by reticulospinal fibers, since they are abolished by cutting the ventral and ventrolateral funicles. Further, many reticulospinal neurons are rhythmically active in pace with walking movements.

The **basal ganglia** have reciprocal connections with the PPN and appear to modulate locomotor movements without initiating them. The characteristic **gait** disturbances in **Parkinson's disease** may possibly be explained by dysfunction of these connections (see Chapter 23, under "The Efferent Connections of the Basal Ganglia"). In addition, the **cerebellum** (particularly the vermis) influences locomotor movements, and cerebellar lesions may produce an **ataxic** gait. The cerebellar effects are probably mediated by connections from the medial

(fastigial) cerebellar nucleus to the medullary reticular formation (see Fig. 24.6). Connections from the **motor cortex** (especially the pyramidal tract) are of increasing importance for the control of locomotion as the ground becomes more uneven and unpredictable—that is, as each step has to be controlled individually. Thus, after destruction of the motor cortex, cats can still walk fairly normally on an even surface but are helpless when they are required to walk along a ladder or a narrow bar.

MOTOR CORTICAL AREAS AND CONTROL OF VOLUNTARY MOVEMENTS

Motor Networks

We do not fully understand how the appropriate corticospinal neurons are selected to produce a purposeful movement. The decision to initiate a particular movement is certainly not made in the MI, and it is not likely to be caused by activity in one particular cell group. Cognitive processes are expressions of activity in **distributed networks** of cortical neurons. Likewise, many parts of the cerebral cortex and interconnected subcortical nuclei are responsible for motor control. While the final movement command issues from the primary motor cortex (M1), the **decision to move** is mediated to M1 from other parts of the cortex. Accordingly, brain-imaging methods such as PET and fMRI have shown that large parts of the cortex are activated in relation to voluntary movements. Further, when sensitive methods are used to record the electrical activity over the brain of a person asked to perform a simple flexion–extension movement of a finger, changes of activity over large parts of both hemispheres first occur. This is a slowly rising negative wave—a so-called **readiness potential**—starting about 850 msec before the movement. Approximately 60 msec before the movement, the activity is concentrated over the arm region of the motor cortex (on the side opposite the moving fingers). Studies of monkeys with implanted microelectrodes show that not only is the cerebral cortex activated before a voluntary movement; there is also increased neuronal activity in the **cerebellum**, the **basal ganglia**, the **thalamus**, and parts of the **limbic structures**.

Hierarchical Organization of Motor Areas

The regions defined today as motor in a strict sense are largely confined to the cytoarchitectonic **areas 4 and 6** of the frontal lobe (Fig. 22.10; see also Fig. 33.3). The **MI** (area 4) in the precentral gyrus has a special position because it exerts the most direct and powerful effects on the motoneurons. Further, lesions of the MI produce clear-cut **pareses**, in contrast to lesions of other cortical regions. Within area 6, two main subdivisions

⁴ Rhythmic locomotor movements can be elicited by stimulation of the so-called **subthalamic region**, which is close to the posterior hypothalamus. It is not clear, however, whether the effect is due to stimulation of passing fibers rather than neurons in the subthalamic region itself.

are usually recognized: the medially situated **supplementary motor area (SMA)**, and the lateral **premotor area (PMA)**. Besides sending fibers to the spinal cord, thus contributing to the corticospinal tract, both the SMA and the PMA send many fibers to the MI. They can therefore act on motoneurons in two fairly direct ways. Usually, however, the effect on MI has been considered most important, and, consequently, the SMA and PMA are often placed above the MI in a **hierarchy of motor areas**. The SMA and PMA—sometimes termed **supramotor areas**—are believed to instruct the MI in what to do. Several observations support this assumption—for example, that the SMA and PMA usually become active in advance of MI during voluntary movements. Clinical observations of patients with complete or partial lesions suggest that **SMA** and **PMA** are important for **sequential movements**, especially performance of rhythmic sequences (there are no pareses). A pianist suffering from such a lesion could no longer play because he was unable to keep even intervals between the key-strokes. The great Russian neuropsychologist Alexander R. Luria used the term **loss of movement melodies** to describe symptoms after lesions of area 6. Among typical

symptoms are difficulties with coordination of **bilateral movements**, such as swinging the arms in opposite directions.

The **posterior parietal cortex** and the **prefrontal cortex** are also concerned with motor control, although less directly than the MI, SMA, and PMA. In certain respects, the parietal and prefrontal areas are higher up in the hierarchy of motor areas. The posterior parietal cortex is important for the transformation of somatosensory and visual information into appropriate motor commands. The prefrontal cortex shows increased activity before self-initiated movements (i.e., movements that are not a response to external stimuli). The functions of these cortical regions are multifarious, however, and certainly not restricted to motor control. They are discussed more fully in Chapter 21.

During execution of well-rehearsed, routine movements, only relevant parts of the MI increases its activity, as judged from fMRI studies (Fig. 22.11). Apparently, “higher” motor areas are only minimally engaged; their contribution restricted to mediating the intention to move. Presumably, “motor program” is located in the motor cortex, the basal ganglia, and the cerebellum.

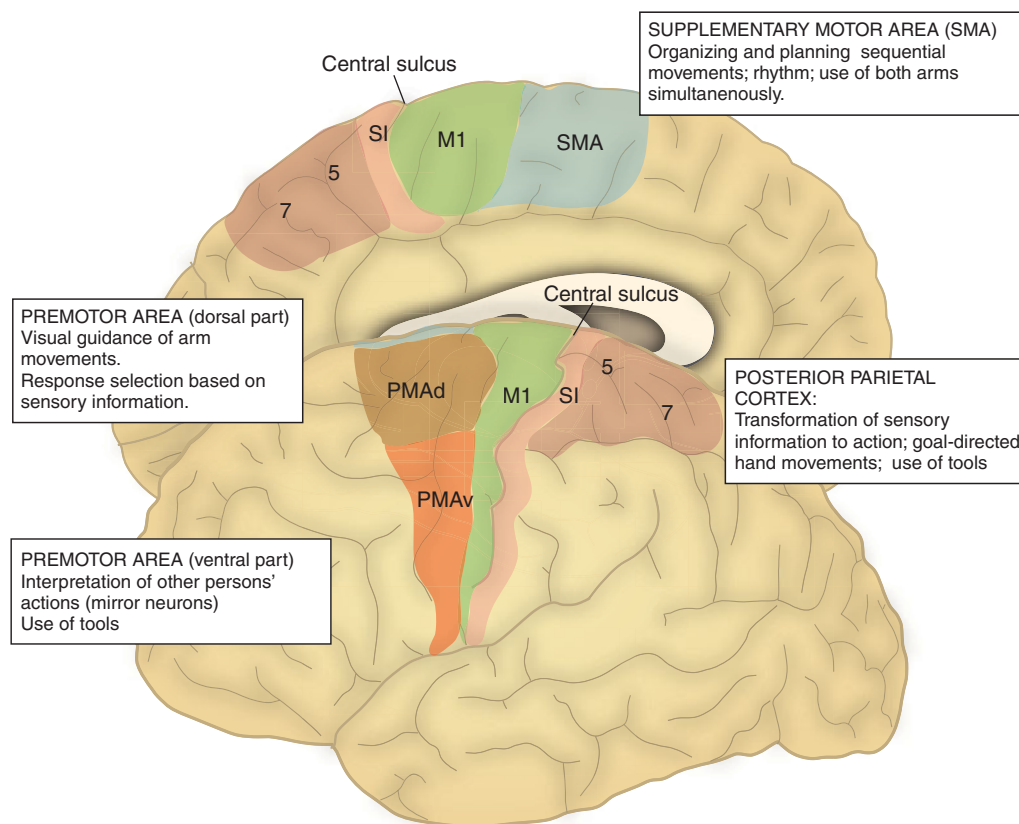


FIGURE 22.10 Areas of special importance for the control of voluntary movements. The borders of the various areas are not exact. They are partly based on cytoarchitectonic maps (Brodmann), partly on

PET and fMRI studies. Areas in the prefrontal cortex are engaged in cognitive aspects of motor control (selection of goal, choice of strategy, and so forth), but are not shown.

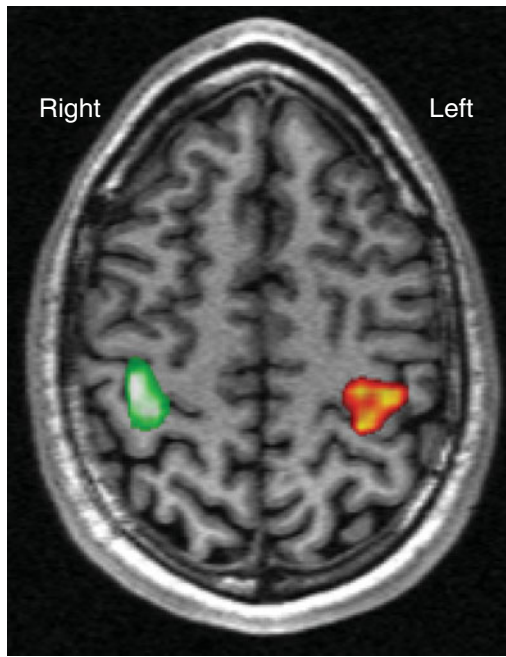


FIGURE 22.11 *Parts of the motor cortex activated by finger tapping.* T1-weighted fMRI. Areas with increased blood flow are colored (red: left motor cortex, activated by tapping the right index finger, green: right motor cortex activated by tapping the left index finger). Movements that are more complex activate additional areas. (Courtesy of Dr. S.J. Bakke, Rikshospitalet University Hospital, Norway.)

The Connections of MI

The physiologically defined MI corresponds fairly closely to Brodmann's area 4.⁵ It receives main **afferents** from **SI, SII, SMA, PMA, and the posterior parietal cortex**, in addition to afferents from “motor” parts of the **thalamus** (the ventrolateral nucleus, VL; see Figs. 6.21 and 14.16). The **cerebellum**, especially, sends important information about movement performance to MI via VL (the basal ganglia mainly influence the MI via connections to area 6).

Although a large fraction of the pyramidal tract fibers arise in MI, these nevertheless constitute only a minority of all **efferents** from MI. The motor cortex sends, for example, **feedback** connections to the cortical areas from which it receives afferents. Further, numerous fibers pass to subcortical regions involved in motor control, such as the **basal ganglia, the cerebellum**

(via the pontine nuclei), the **red nucleus**, the **reticular formation**, and the “motor” **thalamus**. This means that a copy of the motor commands that are issued to the motoneurons—an **efference copy**—reaches the basal ganglia and the cerebellum. Such information helps them to assist in the control of ongoing movements and in learning new ones. Efference copies also reach cortical regions that are responsible for the **sense of effort** (see Chapter 13, under “Perception of Muscle Force”), and regions that enable us to distinguish sensory information that arises as a result of our own movements from such that is due to external events. Efference copies are also important for updating the **body scheme** and for our perception of bodily **ownership** (see Chapter 18, under “Distributed Networks, Body Image, and Body Scheme”). Consequently, damage to the motor cortex is not identical to damage of the pyramidal tract—many neuronal groups besides the motoneurons lose crucial information.

Most if not all **pyramidal tract** fibers ending monosynaptically on the motoneurons come from MI, which explains why the threshold for eliciting movements by electrical stimulation is lower here than in any other part of the cortex.⁶ Accordingly, the movements evoked by very weak electrical stimulation of the motor cortex are mediated by the pyramidal tract. Such movements occur in the opposite body half and can be limited to a few muscles in distal parts of the extremities and the face. Increasing stimulus strength recruits more muscles, and ones that are more proximal. These effects are most likely mediated by polysynaptic pyramidal tract connections (via spinal interneurons) and by **corticoreticulospinal** pathways. Muscles that are often used simultaneously on both sides of the body, like the muscles of the back and the abdomen, can be relatively easily activated on both sides (bilaterally) by stimulation of the MI of one side. Movements of the fingers, however, can be evoked only from the opposite (contralateral) MI, which reflects that the fingers are used independently and usually differently on the two sides. The anatomic basis of this is the complete crossing of the pyramidal tract fibers that control distal muscles, as mentioned earlier. Similar conditions pertain to the **commissural fibers** that interconnect the MI of the two hemispheres: only parts of the MI representing the trunk and the proximal parts of the extremities are interconnected. The large areas representing the distal muscles are devoid of commissural connections, presumably as an expression of the independent use of the two hands (see Fig. 33.12).

⁵ It is not clear whether there is complete coincidence between the physiologically defined MI and area 4, which is defined cytoarchitectonically. Some authors maintain that the MI extends anteriorly somewhat into area 6 with the representation of axial and proximal muscles. Such disagreement may be caused by lack of clear-cut cytoarchitectonic changes when moving anteriorly from area 4 into area 6. Cytoarchitectonic borders, as depicted so confidently in maps like that in Fig. 33.3, are in reality seldom unequivocal. The considerable differences between maps published by different authors witness this point.

⁶ Near the end of the nineteenth century, electrical stimulation of area 4 in dogs produced the first firm evidence of specializations within the cerebral cortex. Before that, the existence of functional localization in the cerebral cortex was hotly debated.

Epileptic Seizures Starting in MI

The so-called **Jackson epileptic seizures** (described in Chapter 14) illustrate the somatotopic pattern within the MI (Figs. 22.11 and 22.12). The abnormal discharges of the neurons start at one site in the MI and spread out in a regular manner. Thus, the muscular cramps start in one part of the body—for example, the foot—and spread to the lower leg, then to the thigh, the abdomen, the shoulder, and so forth. Nearly always, the fits start around the mouth, in the tongue, the thumb, or the big toe. This is best explained by the fact that the cortical neurons controlling these parts occupy the largest volume of the MI and have the largest proportion of monosynaptic corticomotoneuronal connections.

Functional Organization of the Primary Motor Area

As with SI, a disproportionately large part of the MI is devoted to control of the hand (especially the thumb and the index finger) and the lips and the tongue (Fig. 22.12). In comparison, very small parts of the MI contain neurons that control muscles of the back and the abdomen. How many cortical neurons that are concerned with the motor control of a particular body part depends on the variety and precision of the movements rather than on the size of the part.

Focusing on the properties of single cells led to the view that pyramidal tract neurons were collected in groups controlling muscles around single joints (somewhat like the arrangement in the spinal cord where motoneurons are collected in groups related to single muscles; see Fig. 21.3). However, further studies have not supported this interpretation. Thus, pyramidal tract neurons controlling **motoneurons of one muscle** usually

spread over a relatively large cortical area (although respecting the rough somatotopic pattern shown in Fig. 22.5). For example, within the motor hand region, pyramidal tract neurons that control different joints of the hand are intermingled. Thus, every small patch of the motor cortex would contain neurons that control several different muscles. We also know that each pyramidal tract axon acts on more than one muscle (although with a stronger action on some than on others). Therefore, the explanation why we can do isolated movements of single joints cannot be that each pyramidal tract neuron is specified for one movement only. A particular movement is specified by the collective activity of a large neuronal **population** rather than by a narrow tuning of a few neurons. Further, such populations appear to be related to **initiation of purposeful movements** involving several joints, rather than to isolated, single-joint movements (consider how seldom our movements involve one joint only). This conclusion received recent support from experiments with **microstimulation** of the monkey motor cortex. In contrast to most previous studies, long trains of stimuli lasting for about 500 msec were used, similar to the time scale of normal reaching movements. Such stimulation evoked coordinated movements and postures that involved many joints. Notably, the movements resembled purposeful movements—such as opening the mouth and, at the same time, shaping the hand and moving it to the mouth. In other words, the motor cortex is organized to optimize **task performance**, not to control individual muscles or joints. As an example, destruction of the cat's motor cortex does not affect single-joint movements but complex, multijoint movements, such as stretching the paw toward a goal.

Several observations show that the **selection of corticospinal neurons** can be extremely precise and varied. First, one corticospinal neuron can be active when a certain hand muscle is used for a precision grip, whereas it is much less active or inactive when the muscle is used for a more crude grasping movement. Other corticospinal neurons must therefore be responsible for producing the required force from the muscle in the latter situation. Second, human subjects can quickly learn to recruit one **specific motor unit** among many others when given **biofeedback** (seeing the EMG pattern of the muscle). This suggests an almost incredible ability to focus excitation from the motor cortex on certain spinal motoneurons. Considering the precision of movements required of a violinist, a watchmaker, or a neurosurgeon, the above observations become perhaps less surprising, however.

Properties of Single Neurons in MI

Many cells in the MI become active (or increase their firing frequency) immediately before a voluntary movement

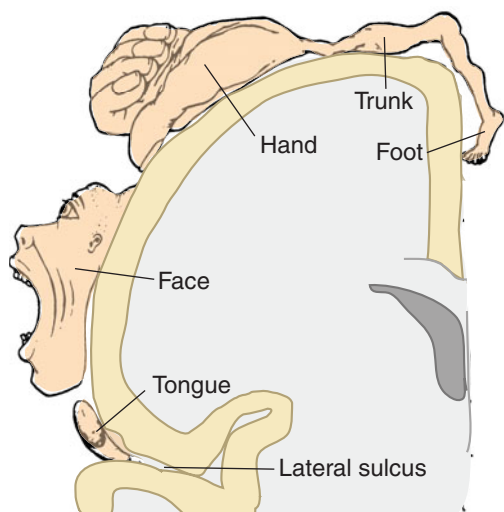


FIGURE 22.12 Relative size of the regions of the MI representing various body parts, as revealed by electrical stimulation of the exposed human MI. (Based on Penfield and Rasmussen 1950.)

starts, as shown with microelectrode recordings. In contrast, most cells in the SI become active after movement onset, indicating that they are activated by sensory signals from the moving parts but do not contribute to the initiation of the movement. Some pyramidal tract neurons (corticospinal neurons) increase their activity only in relation to the **start** of a movement, whereas others are active during the whole movement. The firing frequency of some neurons correlates with the **force** of muscular contraction. In the MI of a monkey sitting quietly waiting for a signal to move, some corticospinal neurons fire continuously. This probably means that the motor cortex not only initiates and controls movements but also determines the **readiness** of selected motoneuronal groups by keeping them slightly depolarized (below the threshold for initiation of action potentials). This enables the motoneurons to respond more quickly to a “go” signal from the motor cortex or from subcortical levels, such as the reticular formation, the vestibular nuclei, and sensory signals through the dorsal roots (e.g., from muscle spindles). At the same time as a corticospinal neuron excites motoneurons of synergistic muscles, motoneurons of antagonistic muscles are inhibited (via inhibitory interneurons mediating reciprocal inhibition).

The Supplementary Motor Area

The SMA is located in front of the MI on the medial side of the hemisphere (Fig. 22.10). It sends fibers to both the spinal cord and the reticular formation. Thus, the SMA contributes to both direct and indirect corticospinal pathways.⁷ The SMA also sends many fibers to the MI, and this pathway is probably most important for its influence on motor control. This assumption is primarily based on results obtained by microelectrode recordings in monkeys and on brain imaging in humans. **Afferents** to the SMA come from—among other sources—the **prefrontal association areas** (see Fig. 34.3).

The neuronal activity in the SMA of humans appears to increase especially in relation to somewhat complex movements. As mentioned, a series of simple flexion–extension movements of a finger increases the activity mainly in the MI hand area, not in the SMA (Fig. 22.11). When the task is to perform a series of different movements in a specified **sequence**, however, there is also increased activity in the SMA. It appears that the activity increases first in the SMA and then in the MI. Increased activity in the SMA is not related to the movement itself, as it is sufficient that the person imagines

the performance of a complex movement. Observations of the kinds discussed above have led to the suggestion that the SMA is important for organizing and planning **complex movements** and for mediating an appropriate **motor response to sensory stimuli**.

Damage of the SMA in humans usually produces severe motor impairments. Typically, the spontaneous use of the contralateral hand is abolished for weeks to some months; this condition is sometimes termed **motor neglect**.⁸ Another and more enduring problem is difficulty with the **simultaneous use of both hands** to solve a task. Monkeys with a lesion of SMA show similar symptoms. For example, when a piece of food is stuck in a hole in a Plexiglas plate, a normal monkey easily retrieves it by pushing with one finger from above and collecting the piece with the other hand from below. After damage to the SMA, the monkey pushes from above with both hands. That the **grasp reflex** is disturbed for some time after damage to the SMA may presumably be due to a disruption of normal integration of somatosensory stimuli and motor responses.

Additional Motor Areas on the Medial Wall of the Hemisphere

Several smaller motor areas have been identified adjacent to SMA on the medial wall. This concerns the so-called **presupplementary motor area** (preSMA) and the **cingulate motor area** (consisting of three small areas buried in the cingulate sulcus). These areas differ from the SMA with regard to connectivity and by being associated with somewhat different aspects of motor control. The pre-SMA seems to be particularly active in relation to the performance of **novel** movement sequences but less in relation to well-rehearsed movements. The cingulate motor area may be involved in **reward-based** movement control.

The Premotor Area

The PMA occupies the largest part of area 6 on the convexity of the hemisphere (Fig. 22.10).⁹ Like the SMA, the PMA receives its most important **afferents** from the **prefrontal cortex**, although not from identical parts. The PMA probably sends fewer **efferent** fibers to the spinal cord than the SMA but has strong connections

⁷ Spinal projections from SMA and PMA end primarily on interneurons in the intermediate part of the spinal gray matter—that is, relatively far from the motoneurons. These spinal projections may therefore serve to prepare the spinal networks for the motor commands from MI, rather than initiating movements on their own.

⁸ Some patients with lesions of SMA show the so-called **alien (anarchic) limb syndrome**. The arm makes purposeful movements, for example grasping an object, without the patient’s intention. The person recognizes the limb as his own but has apparently no control over its actions. In some patients, there is an irresistible urge to grasp and use nearby objects, even if they have no intention to do so (utilization behavior). It is unclear, however, whether some aspects of these symptoms depend on simultaneous lesions of the corpus callosum.

⁹ Assuming that area 6 of humans corresponds functionally to area 6 in monkeys, the SMA and PMA have increased in size during evolution. Thus, areas 4 and 6 are of approximately the same size in monkeys (Fig. 22.8), whereas area 6 is much larger than area 4 in humans (see Fig. 33.3).

with the **reticular formation**, the **red nucleus**, the **basal ganglia**, and the **cerebellum**. As with the SMA, however, the connections from the PMA to the MI are probably those most directly related to the motor functions of the PMA. As indicated in Fig. 22.10, the PMA consists of a dorsal (PMd) and a ventral (PMv) subdivision that differ with regard to connectivity and functional properties. Experiments in monkeys indicate that the PMA is important for the control of **visually guided movements**, such as the proper orientation of the hand and fingers when they approach an object to be grasped. The PMA thus performs **visuomotor transformations** of signals coming especially from the posterior parietal cortex. In monkeys, many cells in the PMA change their activity about 60 msec after a light signal that the monkey is trained to respond to with a certain movement. The activity of the PMA neurons continues until just before the movement starts, even when the monkey is trained to wait for many seconds after the signal before actually performing the movement. Thus, the PMA appears to hold the intention to move and the motor plan in standby until it is appropriate to start.

Monkeys with lesions of the PMA also have difficulties with moving the hand around a transparent obstacle to reach an object: They persistently use the direct approach, bumping into the obstacle. After damage to the MI, the handling of an object is clumsy and insecure, but the ability to avoid an obstacle is not lost. Connections from the **extrastriate areas** in the occipital lobe to the PMA are necessary for the ability to perform such **circumventive goal-directed movements**. Obviously, the PMA is important for the ability to adapt a goal-directed movement to altered external conditions.

Damage of the PMA often produces a peculiar tendency to continue a certain movement when first started, even though the movement is unsuccessful in achieving its goal. Thus, when the hand in one of the examples mentioned above bumps into an obstacle (in this case, a transparent plate), the monkey nevertheless repeats the same movement repeatedly. This phenomenon is called **perseveration** and occurs in humans after damage to the frontal lobes.

Mirror Neurons

In the ventral part of the PMA (PMv) and in the **posterior parietal cortex** certain neurons are active not only when the person performs a certain movement but also when she watches another person performing the same movement. Such **mirror neurons** respond particularly well to use of tools, and they may respond to the sound produced by an action. Although identified with certainty only in monkeys, mirror neurons probably exist also in the human brain, as judged from fMRI studies.

It is believed that the perceived action of the other person is automatically simulated by the mirror system without being actually carried out. The system of mirror neurons is probably involved in **learning by imitation** and in the “reading” of other persons’ **motor intentions**. Some argue that the mirror system also is responsible for mind reading in a wider sense, that is, perception of others’ intentions and the communicative content of movements (called **social cognition**). Others question this so-called “motor theory of social cognition.” As said by Jacob and Jeannerod (2005, p. 22) “. . . we grant that simulating an agent’s movements might be sufficient for understanding his motor intention, but we . . . argue that it is not sufficient for understanding the agent’s prior intention, his social intention, and communicative intention.” Certainly, identical movements may result from very different intentions and for different purposes. Another system, collecting much more varied information than the mirror system does, seems necessary for social cognition (notably association areas in the superior temporal sulcus, the amygdala, and the orbitofrontal cortex).

Motor Imagery

Do we use the same parts of the brain when we imagine a movement as when we perform it? Many studies with brain-imaging methods, such as PET and fMRI have addressed this question. Most agree that largely the same **cortical networks** are activated in both situations. This holds for the PMA, parts of the prefrontal cortex, the basal ganglia, lateral parts of the cerebellum, and the posterior parietal cortex. Some studies also show increased activity in MI, although considerably less than in relation to movement execution. In agreement with such data, magnetic stimulation of MI most easily evokes contraction of the muscles involved in the imagined movement (data are conflicting, however, whether spinal motoneurons also show similar facilitation during motor imagery). Dissimilarities concern SMA, where somewhat different subregions are active in motor imagery than in real movements. Further, inferior parts of the prefrontal cortex are active only during imagery, so perhaps this region is responsible for the **suppression** of movements. The same neuronal processes seem to underlie imagined and real movements, as indicated by the fact that they take equal time from start to end. Patients with **lesions** of the **motor cortex** can still imagine movements in the paretic side, but just as the real movements the imagined ones are slower than normal. Patients with **Parkinson’s disease** likewise exhibit similar slowing and reduced amplitude of real and imagined movements. After lesions of the **posterior parietal cortex**, however, the imagination of movements seems to be more affected than their execution.

Learning and the Motor Cortex

Brain-imaging studies show that higher association areas in the prefrontal and posterior parietal cortices are active during learning of new motor skills. As movements become more automatic, the activity decreases in these areas, presumably because of use-dependent plastic changes. We mentioned that the **pre-SMA** might be particularly engaged in learning new sequential movements. Further, animal experiments and human brain-imaging data indicate that plastic changes occur also in the MI during motor learning. Both **LTP** and **LTD** can be induced in the cerebral cortex, and motor learning appears to be associated with strengthening of **horizontal connections** within MI thereby coupling functionally related neurons. Thus, pyramidal cells in laminae II and III strengthen their synaptic couplings with neurons in other parts of the motor cortex during skill learning in rats. Further, the connections are specific for the body parts that are used in the motor performance.

In humans, the finger representation in MI increases during 1 week of intense **piano training** of a specific sequence, as judged from threshold changes to magnetic stimulation of various parts of the motor cortex. Changes were obtained both with real movements and with mental training (motor imagery), although the effect was largest with real movements. In a group of right-handed elite **badminton** players, the stimulation threshold was lower and the hand area was larger in the left than in the right motor cortex (such differences were not found in a group of recreational players).

The Posterior Parietal Cortex and Voluntary Movements

Area 5 is of particular importance for processing somatosensory information (received from SI), whereas **area 7** also receives information from visual cortical areas (see Figs. 21.9 and 21.10). Many neurons in these areas are active in relation to movements, as shown by Vernon Mountcastle (1975) and others. One kind of neuron is active before goal-directed, reaching movements, such as when a monkey stretches its hand toward a banana. Such neurons do not become active, however, in relation to a movement in the same direction but without a specific aim, or in relation to a passive movement. Other kinds of neurons increase their activity in relation to exploratory hand movements, such as when a monkey studies a foreign object. In area 7, some neurons increase their activity only when the monkey stretches the hand toward an object that it also looks at. As there are ample connections from the posterior parietal cortex to the SMA and PMA, it is likely that the posterior parietal cortex in part determines the behavior of cells in these motor areas.

Indeed, **damage** to the posterior parietal cortex also produces motor disturbances. There are no pareses, however, but rather difficulties with the execution of more complex movements. Patients with such lesions may be unable to open a door or to handle previously familiar tools like a screwdriver or a can opener. They also have difficulties with proper orientation of the hand in relation to an object, and they easily miss an object even though they see it clearly. This kind of symptom is called **apraxia** (see also Chapter 34). Interestingly, similar symptoms may occur after lesions of the frontal lobes in front of the MI, presumably reflecting the intimate connections between the posterior parietal cortex and the frontal lobes.

SYMPTOMS CAUSED BY INTERRUPTION OF CENTRAL MOTOR PATHWAYS (UPPER MOTOR NEURONS)

The term **central paresis** is used for a muscle weakness that is caused by interruption of the central motor pathways that conduct signals from the cerebral cortex (especially the MI) to the motoneurons. Only the pyramidal tract goes directly; the other pathways are indirect, with synaptic interruption in the brain stem. We have discussed that the various pathways take care of somewhat different aspects of motor control. The term **upper motor syndrome** is often used for the clinical picture resulting from interruption of the central motor pathways, to differentiate it from the **lower motor syndrome** (peripheral pareses) resulting from destruction of the motoneurons (including their axons; see Fig. 21.16).

Mechanisms underlying recovery after damage to the upper motor neurons was discussed in Chapter 11, under “Studies of Recovery after a Stroke in Humans.”

“Negative” and “Positive” Symptoms

Although pareses are present in both, peripheral and central pareses differ in other respects. In peripheral pareses, the symptoms are all **negative**, in the sense that they represent **loss of function**, like reduced or abolished muscle power, resting tone, and reflex contractions. There is also marked and rapid wasting. In central pareses **positive symptoms** also occur—that is, there are symptoms caused by hyperactivity of neurons, such as increased reflex responses and resting muscle tone. Such differences are understandable when considering that the peripheral motor neurons are still functioning in central paresis (see Fig. 21.16). The motoneurons can be activated by signals from various receptors through the dorsal roots, from spinal interneurons, and from any remaining descending pathways, even though they cannot be brought into action voluntarily. Because the

motoneurons still send signals to the muscles in a patient with central pareses, muscle wasting is modest, in contrast to that in peripheral pareses. Generally speaking, the positive symptoms occurring after damage to upper motoneurons are due to **hyperexcitability** of neuronal groups in the spinal cord, thus producing abnormal muscle contractions (e.g., on innocuous stimuli like moving or touching a limb). Because the reflex arc is intact, reflexes like the stretch reflex and the flexion reflex can still be elicited, and typically, the reflex responses are stronger than normal in patients with central pareses (**hyperreflexia**). Although the reflex responses are weak or absent shortly after a stroke, and especially after a transverse lesion of the cord (spinal shock), they recover in some days or weeks. Especially the **monosynaptic stretch reflex** (as tested with a tendon tap) becomes hyperactive on the affected side.¹⁰ In addition, most patients develop an increased reflex response to passive movements of the affected limbs—that is, to stretching of the paretic muscles. The reflex contraction depends on the **velocity of stretch**, so that slow movements typically do not elicit any contraction. In some patients, however, especially in those with spinal transection or multiple sclerosis, even slow movements or innocuous cutaneous stimuli may provoke prolonged and painful **muscle spasms**.

The term **spasticity** refers to the positive symptoms in patients with upper motor neuron lesions. Some use the term to include all positive symptoms—that is, hyperreflexia, increased muscle tone, and muscle spasms. Others define spasticity more narrowly, restricting it to the velocity-dependent increase of resistance to muscle stretch.¹¹

Mechanisms Responsible for Development of Spasticity

The development of spasticity—widely defined as the positive symptoms that occur after a lesion of the upper motor neurons—is due mainly or solely to **excitability changes in the cord** leading to exaggerated motor responses to signals from receptors, from other parts of the cord, and remaining supraspinal pathways. It is highly likely that **plastic processes**, for example, in the form of accidental collateral sprouting, contribute to excitability changes. The more precise mechanisms underlying spasticity are not known, however. To complicate matters, there is evidence that the mechanisms

differ among spastic patients with similar lesions. A priori, it seems likely that the loss of descending corticospinal fibers results in decreased activity of **inhibitory interneurons** and many studies have been performed in humans to determine the excitability of specific kinds of interneurons. As discussed earlier in this chapter, the pyramidal tract modulates reflex responses by specific actions on several kinds of spinal interneurons.

Several studies have shown reduced activity of the interneurons mediating **reciprocal inhibition** (see Fig. 21.13) in spastic patients, both after spinal cord lesions and in capsular hemiplegia. This would help explain the occurrence of hyperreflexia. For example, when testing the Achilles reflex with a tap on the tendon, a monosynaptic stretch reflex is elicited in the calf muscles (ankle plantar flexors). Normally, the reciprocal inhibition prevents a stretch reflex from being elicited also in the antagonists—that is, the ankle dorsiflexors. If the reciprocal inhibition is reduced, however, a reflex contraction of the ankle dorsiflexors can occur; in turn, this may produce a new reflex contraction in the plantar flexors, and so forth. When a single tap on the tendon elicits repeated contractions, it is termed **clonus**.

Another possible factor in spastic patients may be reduced **recurrent inhibition** of α motoneurons by **Renshaw cells** (see Fig. 21.14). When ankle dorsiflexors contract, for example, Renshaw cells inhibit the motoneurons to the ankle plantar flexors so there is less chance of eliciting an unwanted stretch reflex. Reduced activity of the Renshaw cells in such situations would contribute to the occurrence of clonus in spastic patients. Reduced recurrent inhibition has indeed been found in some spastic patients with spinal lesions, although not in hemiplegic patients. Conversely, reduced activity in inhibitory interneurons activated from **tendon organs** (Ib afferents) has been found in hemiplegic patients but not in patients with spinal lesions.

There is also evidence of reduced activity of interneurons mediating **presynaptic inhibition** of Ia afferent terminals. This would lead to a stronger than normal excitatory effect of muscle stretch (particularly rapid stretch) on the α motoneurons. This appears not always to be present in spastic patients, however. For example, one study found reduced presynaptic inhibition in the arm but not in the leg of hemiplegic patients. Reduced **postactivation depression** has been observed in patients with spasticity due to various causes. Postactivation depression means that the postsynaptic effect diminishes when action potentials in muscle spindle primary afferents follow each other with brief intervals. It is probably due to less transmitter being released. Obviously, reduction of this phenomenon might contribute to increased motoneuron excitation to a given muscle stretch.

A permanently increased **excitability of the motoneurons** themselves has not been found in patients with spasticity. The intrinsic properties of the motoneurons

10 The long-latency stretch reflex appears to be weaker than normal in spastic patients, as judged from the EMG response to carefully graded muscle stretch and vibratory stimuli. This fits with other data suggesting a transcortical route for the long-latency stretch reflex, as discussed in Chapter 21.

11 As defined by J. W. Lance (cited by Landau 1987, p. 722), “Spasticity is a motor disorder characterized by a velocity-dependent increase in tonic stretch reflexes (muscle tone) with exaggerated tendon jerks, resulting from hyperexcitability of the stretch reflex, as one component of the upper motor neuron syndrome.”

might nevertheless be changed, because **plateau potentials** may be more easily induced, as judged from animal experiments. We mentioned above (“Monoaminergic Pathways from the Brain Stem to the Spinal Cord”) that when a plateau potential is induced in a motoneuron it goes on firing for seconds to minutes without further stimulation. After a spinal transection in rats, plateau potentials disappear but return in the course of some months. Interestingly, when plateau potentials reappear, they can be induced by a variety of innocuous stimuli and appear to be closely related to the degree of spasticity. Whether lowered threshold for inducing plateau potentials plays a part in humans with spasticity remains to be determined, however.

It was formerly assumed that increased signal traffic from the muscle spindles—due to increased **γ motoneuron** firing—explained the increased stretch reflex responses in spasticity. Microneurographic recordings have not verified this, however, as the firing frequency of Ia afferent fibers was not found to be increased in spastic patients compared with normal controls.

In conclusion, the reflex hyperexcitability and increased muscle tone in patients with the upper motor neuron syndrome may arise from changes of several inhibitory mechanisms in the spinal cord, although few if any of the changes seem to be present in all patients. Possibly, varying combinations of disturbed inhibitory

mechanisms, perhaps combined with abnormally low threshold for eliciting plateau potential, produce symptoms that are indistinguishable clinically. Yet, to select the right treatment, it would be crucial to know which mechanisms are disturbed in each patient.

Other Features of the Upper Motor Syndrome

Even though central pareses can be produced by lesions located anywhere between the motor cortex and the motoneurons, the most common cause is interruption of the descending fibers by a thrombosis or bleeding in the internal capsule (Fig. 22.13). The pareses then affect the muscles of the opposite half of the body (hemipareses, hemiplegia). The term **capsular hemiplegia** is commonly used for this condition. In such patients—in contrast to those who suffer a transverse lesion of the cord—fibers descending from cell groups in the brain stem (the reticular formation and the vestibular nuclei) can still activate the motoneurons. This may explain why resting muscle tone is typically increased in some muscle groups in patients with capsular hemiplegia (Fig. 22.14). Thus, in the arm the increased tone affects the flexors so that the arm is kept flexed at the elbow, whereas the extensors of the leg are affected (interruption of motor pathways in the cord result in increased resting tone of flexors also in the leg). An important

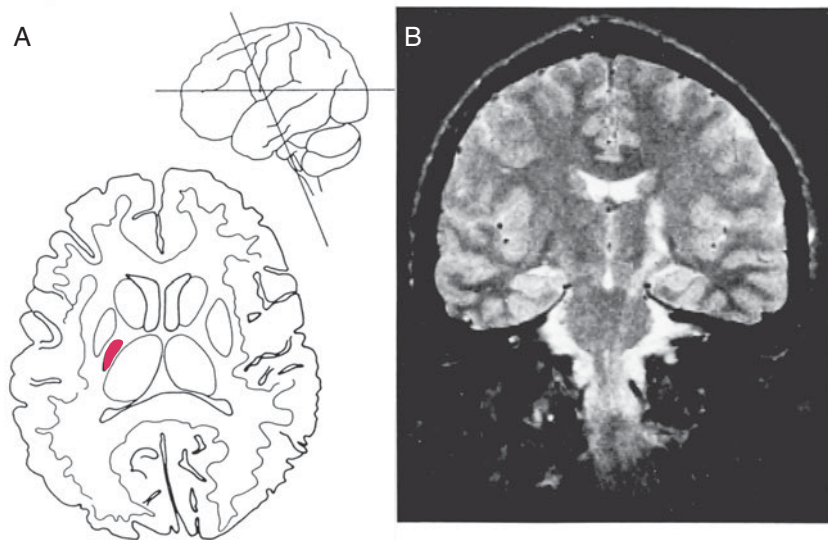


FIGURE 22.13 *Infarction in the internal capsule with degeneration of the pyramidal tract.* The patient, a 19-year-old man, suddenly became hemiplegic, probably due to occlusion of a brain artery by an embolus from the heart. **A:** Drawing of a horizontal CT showing the localization of the infarction (red) in the posterior limb of the internal capsule (cf. Fig. 6.30). Seven months after the stroke, the lesion measured 5×16 mm. **B:** Frontal MRI 4 months after the stroke, with the degenerated pyramidal tract (arrowheads) shown as whitish (cf. Fig. 22.2). It can be followed from the internal capsule through the pons and into the medulla. Immediately after the stroke, the patient had pareses on

the right side, most marked for finger movements. He recovered completely. PET studies in this patient after recovery showed that finger movements of the right hand were associated with increased activity in the left motor cortex and PMA, and bilaterally in the SMA, cingulate gyrus, and the insula (in normal controls only the motor cortex show increased activity with isolated finger movements; cf. Fig. 22.11). The patient made involuntary mirror movements of the left fingers when using the right ones. (From Danek et al. 1990. Reproduced with permission from Oxford University Press.)

feature of pareses in hemiplegia is that the **velocity** with which voluntary movements can be performed is reduced more than the isometric force (i.e., speed of movements is reduced more than strength). This is called **retardation** and concerns particularly fine finger movements and movements of the lips and tongue, whereas larger movements are less severely affected. Writing, tying, buttoning, and similar delicate movements may be impossible for a patient with capsular hemiplegia, or the movements are performed only very slowly and clumsily. This **loss of dexterity** is due to the loss of direct corticospinal fibers (the pyramidal tract), which are necessary for independent finger movements. Movements lose their rhythm and fluency. The **fatigability** is also abnormally great—that is, the muscular force drops quickly when a voluntary movement is repeated several times. The patient experiences a dramatic increase in the **mental effort** needed for voluntary movements: movements that before the stroke required no mental effort can afterward be performed only with the utmost concentration and strain.

Changes in the **contractile properties** of the paretic muscles also occur in hemiplegic patients. Thus, in the intrinsic muscles of the hand, the fatigability of type 1 muscle cells is increased and the contraction velocity of the type 2 fibers is reduced. Such changes will presumably contribute to the slowness and increased fatigability in patients with central pareses. Further, the passive,

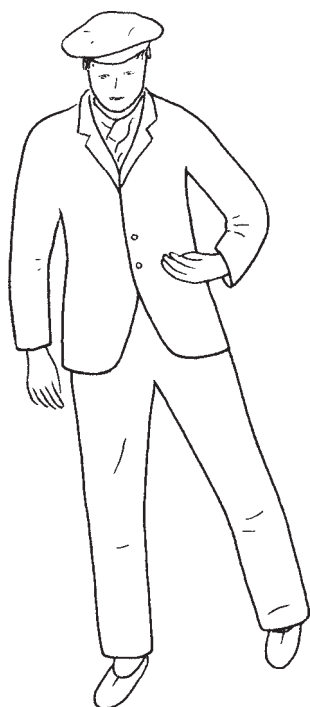


FIGURE 22.14 *Hemiplegia of the left side.* Note the characteristic position of the arm with flexion in the elbow and wrist. The paretic leg is moved laterally in a semicircle during the swing phase to keep the foot off the ground (circumduction).

viscoelastic properties of muscles can change gradually after damage to the upper motor neurons. That is, resistance to stretch may be increased without concomitant muscle contraction, and the range of joint motion may be restricted. The latter phenomenon is called **contracture** and is due to change of connective tissue elements in muscle, tendons, and joint capsules that are kept in a shortened position. Drugs used to treat spasticity such as **baclofen** (reducing the excitability of motoneurons) or **botulinum toxin** (paralyzes muscles) cannot treat this kind of reduced joint mobility.

The Plantar Reflex and Other Reflexes that Are Changed in Upper Motor Neuron Lesions

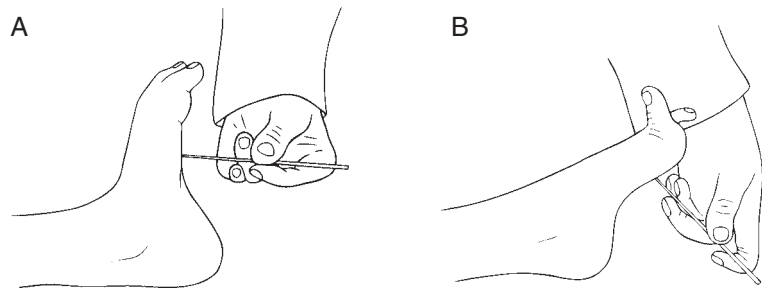
Interruption of descending central motor pathways, as in capsular hemiplegia, also produces changes of reflexes other than the stretch reflexes. The so-called **plantar reflex**, elicited by stroking with a pointed object in a forward direction along the sole of the foot (especially the lateral margin), is inverted (Fig. 22.15). Instead of the normal response, which is a flexion movement of the great toe (and the other toes), the great toe extends (moves upward). This phenomenon, with extension instead of flexion of the great toe, is called the **sign of Babinski** and is a sensitive indicator of damage of corticospinal pathways.¹² For example, increased intracranial pressure and unilateral herniation with compression of the descending motor fibers in the mesencephalon may invert the plantar reflex at an early stage (this will occur with the foot contralateral to the herniation, owing to the crossing of the pyramidal tract in the lower medulla). An inverted plantar reflex may also occur during general anesthesia and in other conditions with reduced cerebral activity. In patients with central pareses, the threshold for eliciting the plantar reflex is lowered and it can be elicited from a wider area than normally. Further, the response may include a complete flexion reflex of the lower extremity with flexion in the hip and knee and dorsiflexion in the ankle.¹³

In **newborn children**, the plantar reflex is inverted and can remain so until the age of 2 (although most infants respond with plantar flexion of the big toe by 12 months). This is most likely due to the dependence of the “normal” reflex on the integrity of the pyramidal tract, which is not fully myelinated until between 12 and 24 months after birth.

12 Studies of patients subject to cordotomy (cutting parts of the lateral funicle to achieve analgesia) indicate that the sign of Babinski occurs only if the lesion affects the pyramidal tract.

13 Electrophysiological studies indicate that Babinski’s sign is due to hyperexcitability of the reflex center in the cord, so that the extensor hallucis longus muscle is recruited together with the ankle extensors. EMG recordings and experiments with nerve blocks show that the flexors of the great toe are still active but that they are overcome by the greater force exerted by the extensors.

FIGURE 22.15 *Inverted plantar reflex in central pareses (sign of Babinski).* **A:** The normal plantar reflex is plantar flexion of all the toes when a pointed instrument is moved along the sole of the foot (from the heel to the toes). **B:** In a patient with central pareses (damage of the pyramidal tract) the big toe moves upward (dorsiflexion).



Reduction or absence of the so-called **abdominal reflex** is also typical of upper motor neuron lesions. The normal reflex response is a unilateral contraction of abdominal muscles upon stroking the skin with a pointed object. The so-called **clasp-knife reflex** or phenomenon occurs in some patients: when the patient contracts a muscle isometrically against resistance for some time, it suddenly yields. The involuntary stop of contraction is due to stimulation of high-threshold muscle afferents that inhibit the motoneurons (signals from tendon organs were formerly believed to be responsible, but this was not confirmed in animal experiments).

The “Pyramidal Tract Syndrome”

Formerly, all of the motor symptoms occurring in capsular hemiplegia were thought to be caused by damage to the pyramidal tract, and the term “pyramidal tract syndrome” is still widely used. Closer study suggests, however, that not all symptoms can be explained by damage to the pyramidal tract. Both in monkeys and humans with lesions restricted to the medullary pyramid, difficulty with fractionate finger movements is the only constantly remaining symptom after some time. Otherwise, the recovery is almost complete—a patient with an infarction limited to the right medullary pyramid even learned to play the cello afterward (see caption to Fig. 22.13). In monkeys, spasticity ensues after

lesions of the motor and premotor cortex but not after complete lesions of the medullary pyramid. To complicate matters, however, hyperreflexia and spasticity have been reported in a few patients with (most likely) pure lesions of the pyramid. Nevertheless, the weight of evidence favors the view that in capsular hemiplegia motor symptoms other than loss of dexterity arise from destruction of other corticofugal pathways than the pyramidal tract. It seems likely that interruption of the **corticoreticulospinal pathways** is important for the more severe symptoms after a capsular lesion than after one limited to the medullary pyramid, including the development of spasticity.

We should also keep in mind that a lesion of the internal capsule might interrupt tracts of importance for motor control other than the corticospinal and corticoreticular ones. Thus, many fibers acting (directly or indirectly) on the **cerebellum** and on the **basal ganglia** will usually be destroyed, and this probably contributes to the clumsiness of voluntary movements. Further, many patients with capsular hemiplegia have **sensory symptoms** in addition to the motor ones, either because the thalamus itself is affected or because the ascending fiber tracts conveying sensory signals from the thalamus to the cortex are interrupted (e.g., visual field defects). Reduced or altered cutaneous sensation and kinesthesia may therefore also contribute to the motor symptoms.

23 | The Basal Ganglia

OVERVIEW

In Chapter 22, we discussed descending pathways from the cerebral cortex, which (directly and indirectly) influence the motoneurons and thereby are crucial for the initiation and control of movements. As mentioned, the basal ganglia form a side loop to the descending motor pathways, and diseases affecting the basal ganglia lead to characteristic disturbances of voluntary movements and of muscle tone but no pareses.

Broadly speaking, the basal ganglia are intercalated in a loop of fiber connections from the cerebral cortex and back to the cerebral cortex through the thalamus. In this respect, the basal ganglia resemble the cerebellum. The basal ganglia process information from large parts of the cerebral cortex before “answers” are sent back to the cortex. Different parts of the basal ganglia assist different subdivisions of the cerebral cortex in their specific tasks. In this way, the basal ganglia are organized in several anatomically and functionally different, **parallel circuits**.

Striatum, composed of the **putamen** and the **caudate nucleus**, is the receiving part of the basal ganglia. The striatum receives excitatory (glutamate) connections from not only motor parts of the cortex but also from association areas and so-called limbic parts of the cortex (e.g., orbitofrontal cortex). In addition, the striatum receives strong connections from the thalamus. Among striatal afferents, the **nigrostriatal** pathway has a special position; it is **dopaminergic** and exerts strong modulatory control of striatal activity. Most of the striatal neurons are **GABAergic**, and send their axons to the **globus pallidus** and the **substantia nigra**. These nuclei send their GABAergic efferent connections to the **thalamus** and certain cell groups in the **brain stem**. From the thalamus, excitatory (glutamate) connections reach especially motor and prefrontal parts of the cortex, whereas the brain stem nuclei (especially the **pedunculopontine nucleus** [PPN]) influence reticulospinal pathways engaged in control of posture and locomotion.

It is noteworthy that the pathway from the striatum to the thalamus contains two inhibitory neurons in a row. Thus, increased activity of striatal projection neurons (driven from the cortex) leads to **disinhibition** in the thalamus, and hence increased excitation in the cortical areas receiving the thalamic input. This has led to the suggestion that the basal ganglia serve, as it were, to

release a brake on voluntary movements, facilitating **switching** from one movement (or mental task) to another. In addition, the basal ganglia are important for the establishment of habits and **learning** of automatic movement sequences. The tasks of the basal ganglia are not restricted to motor control, however. Even though the most obvious symptoms in diseases of the basal ganglia are related to the motor system, both clinical and experimental evidence indicates that the basal ganglia also play a role in **cognitive functions**. Further, the most ventral parts of the basal ganglia—termed the **ventral striatopallidum**—contribute to control of **motivation** and **emotions**.

The multifarious connections and tasks of the basal ganglia serve to remind us that classification of parts of the brain into rigid functional categories such as “motor,” “sensory,” and “cognitive” must not be taken literally but rather as a didactic oversimplification.

STRUCTURE AND CONNECTIONS OF THE BASAL GANGLIA

Figure 23.1 gives a simplified account of the how the basal ganglia are intercalated in a **side loop** of the direct and indirect descending motor pathways. It also shows that the basal ganglia connect with motor centers in the brain stem. Figure 23.2 gives a schematic presentation of the main connections of the basal ganglia, emphasizing the circuit cortex–striatum–globus pallidus/substantia nigra–thalamus–cortex.

On an anatomic basis the term “basal ganglia” usually includes the **caudate nucleus**, the **putamen**, and the **globus pallidus** (Figs. 23.3–23.5; see also Fig. 6.29).¹ Here we use the term “basal ganglia” of a set of functionally related cell groups rather than in a strictly topographic sense. Therefore, we also include the **substantia nigra** and the **subthalamic nucleus** because both are intimately connected with the anatomically defined basal ganglia (Figs. 23.3 and 23.4). Because of their

¹ The **claustrum** (Fig. 23.4) and the **amygdala** (amygdaloid nucleus) were also included in the term “basal ganglia” by the early anatomists. The function of the claustrum is still largely unknown, even though it is known to receive its main afferents from the cerebral cortex and to send efferents directly back to the cortex. The amygdala differs with regard to both connections and functions from the other parts of the basal ganglia, and is now usually considered a part of the so-called limbic structures (Chapter 31).

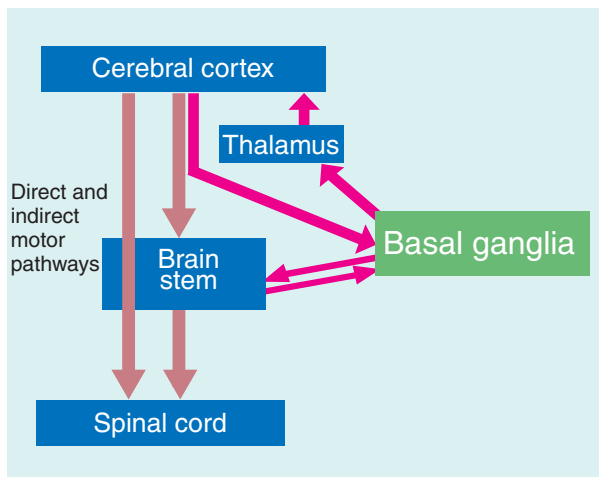


FIGURE 23.1

macroscopic shape, the putamen and globus pallidus together are called the **lentiform nucleus**. The caudate nucleus and the putamen are similarly built, with predominantly small neurons. They are also functionally related and are collectively termed the **striatum** or **neostriatum**. The neostriatum contains several neuronal types that differ with regard to where they send their axons and to which neurotransmitters they use. Most of the neurons, however, send their axons out of the striatum (projection neurons); only a minority is interneurons with axons ramifying locally within the striatum. The presence of several kinds of interneuron is consistent with the fact that the striatum is not simply a relay station but also performs considerable processing of information.

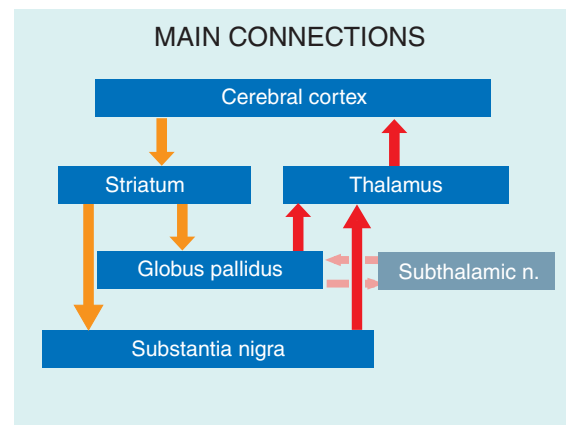


FIGURE 23.2

The **globus pallidus** has a different internal structure than the striatum, with larger, more “motoneuron”-like cells, and is also called the **paleostriatum** or **pallidum**. It consists of two parts, an **internal segment (GP_i)** and an **external segment (GP_e)** (Figs. 23.3 and 23.4). The term **corpus striatum** includes both the pallidum and the neostriatum. Phylogenetically, the caudate nucleus and the putamen developed together and are younger than the pallidum (thus the names neostriatum and paleostriatum). As we will see, these two main divisions of the basal ganglia differ also with regard to connections.

Cell groups that join the corpus striatum ventrally without sharp transitions—such as the **nucleus accumbens** and the olfactory tubercle—are now collectively termed the **ventral striatum**, and thus included in the

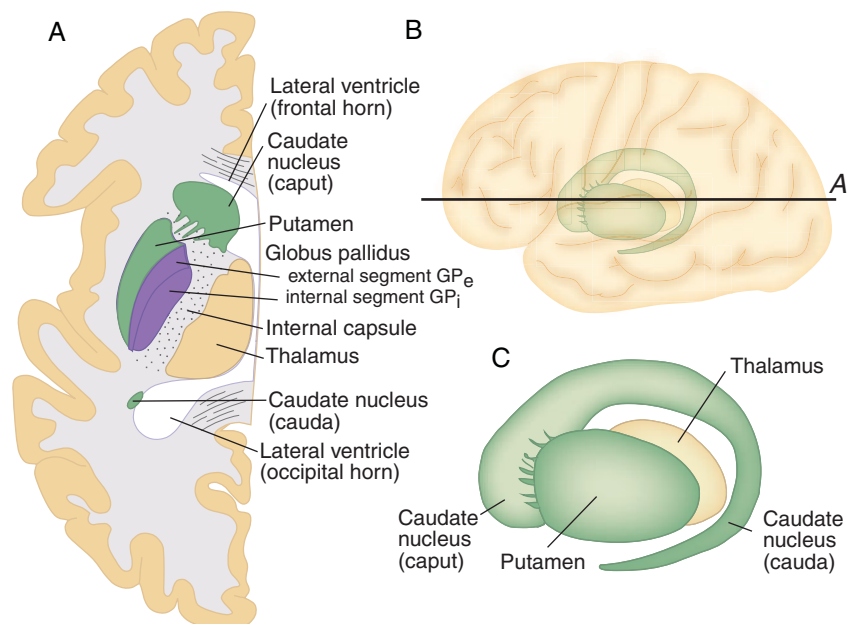


FIGURE 23.3 Shape and position of the basal ganglia. **A:** Part of a horizontal section through the hemisphere, as shown (**B**) with a line in drawing of the hemisphere (cf. Fig. 6.30 showing the whole section). **C:** Left putamen and caudate nucleus; lateral aspect.

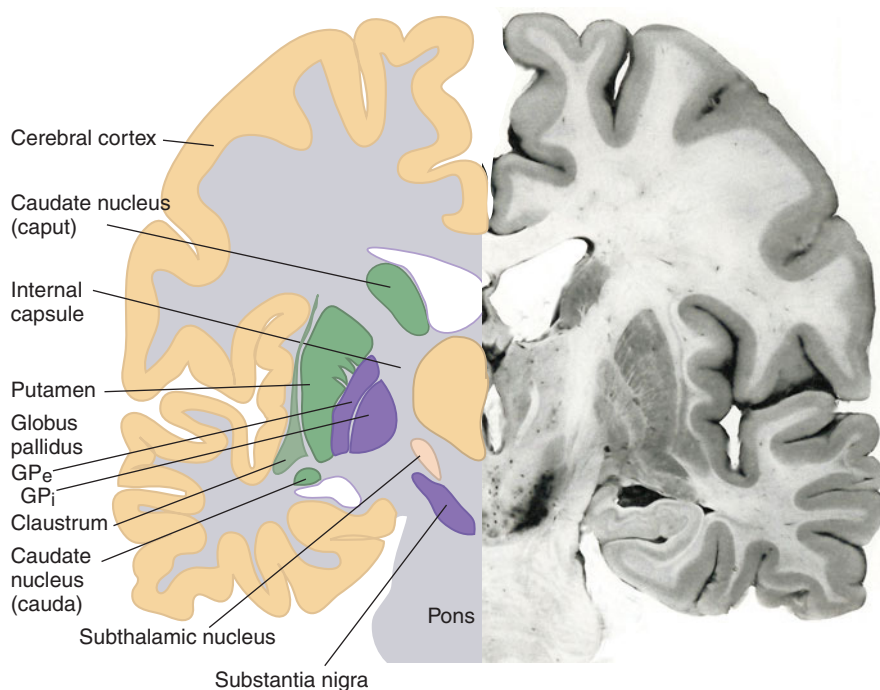


FIGURE 23.4 The basal ganglia seen in the frontal plane.

basal-ganglia concept. There is also a ventral extension of the globus pallidus, called the **ventral pallidum**. The ventral striatum differs in some respects from the rest—that is, from the **dorsal striatum**—and is treated separately in this chapter. The connections of the **ventral striatopallidum** correlate with its importance for behavior governed by emotions.

The Striatum Is the Receiving Part of the Basal Ganglia

Most of the information to be processed in the basal ganglia first reaches the striatum. It is characterized by three major sources of afferents: the **cerebral cortex**, the **intralaminar thalamic nuclei**, and **dopamine-containing cell groups in the mesencephalon** (Fig. 23.6).

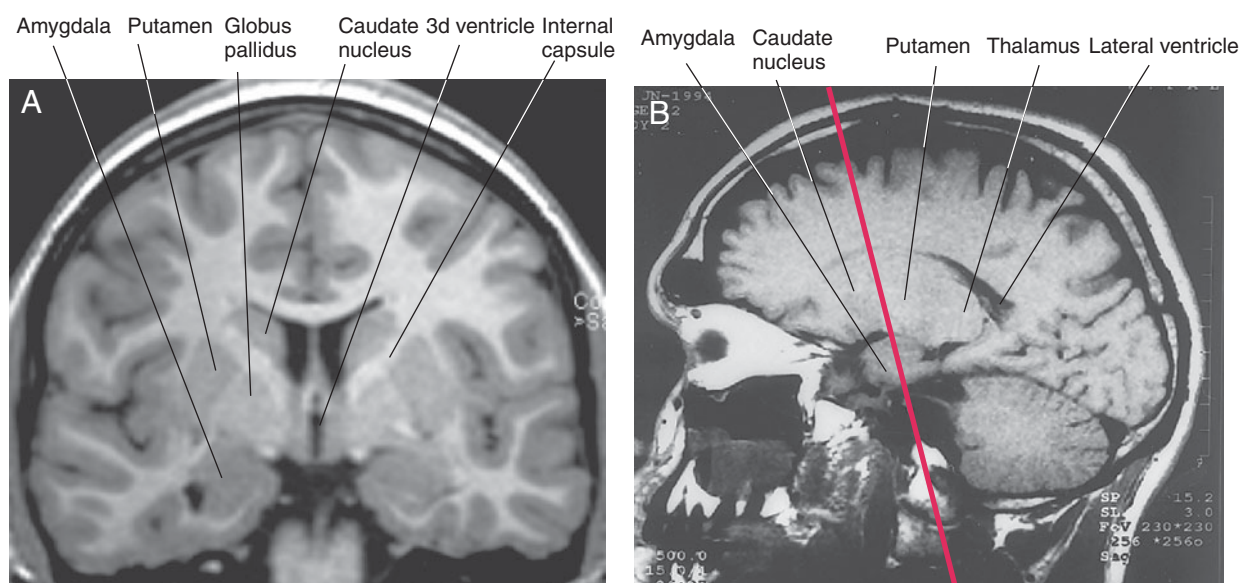


FIGURE 23.5 Magnetic resonance images (MRI) of the basal ganglia. A: Frontal plane; red line in B shows approximate position of

the section. B: Parasagittal plane. (Courtesy of Dr. S. J. Bakke, Rikshospitalet University Hospital, Oslo, Norway.)

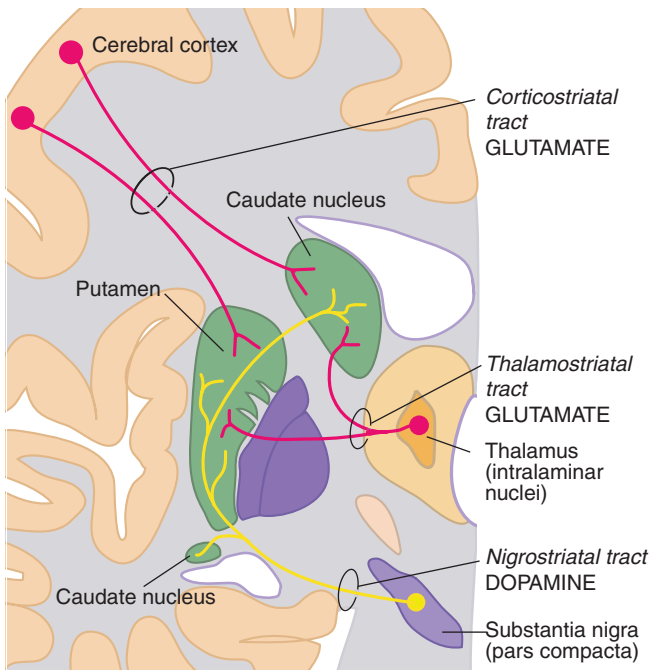


FIGURE 23.6 The main afferent connections of the striatum.

The largest contingent of afferents comes from the **cerebral cortex**. Almost all areas of the cortex send fibers to the striatum, but the caudate nucleus and the putamen receive from different parts (Fig. 23.7).

The **putamen** is dominated by somatotopically organized inputs from the **SI** and **MI**. The **caudate nucleus**, in contrast, receives fibers predominantly from the **association areas**—that is, regions that are less directly concerned with motor control than with cognitive functions and emotions. Whereas the putamen receives relatively “raw,” or unprocessed, information from sensory receptors via the SI and from upper motor neurons in the MI, the caudate nucleus receives information that is a result of integration of signals from many sources. Such information reaching the caudate nucleus may concern, for example, earlier stages in the chains of neural events leading to a decision about which movements are appropriate in the situation.

The striatal afferents from the **intralaminar thalamic nuclei** (Fig. 23.6; see Fig. 6.22) are numerous and are believed to transmit information to the striatum about stimuli that need special attention.

Dopaminergic striatal afferents to the dorsal striatum arise in the **pars compacta** of the **substantia nigra**, whereas the ventral striatum receive such fibers from more scattered dopaminergic cells in the **ventral tegmental area (VTA)** dorsal to the substantia nigra (Fig. 23.8). VTA also sends dopaminergic fibers to the prefrontal cortex.

Additional, quantitatively minor afferent contingents to the striatum come from the serotonergic **raphe nuclei** in the brain stem, among several other sources.

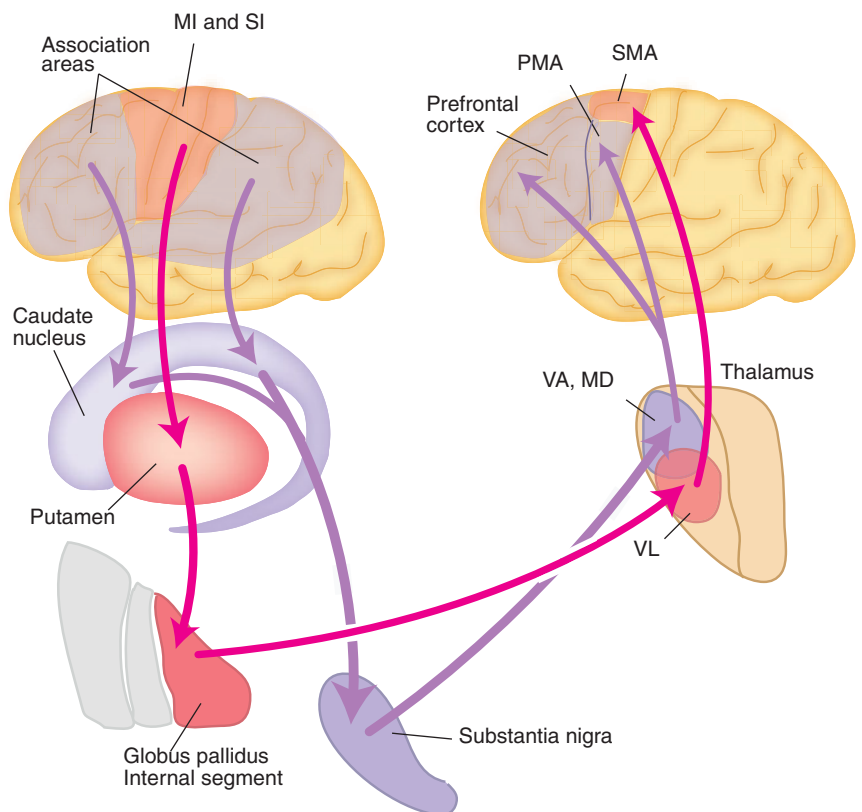


FIGURE 23.7 Parallel circuits cortex–basal ganglia–thalamus–cortex. Highly simplified. The putamen receives fibers primarily from the motor and somatosensory areas (red), whereas the caudate nucleus is dominated by inputs from the association areas in the frontal, parietal, and temporal lobes (blue). The figure further shows that sensorimotor information, after processing in the basal ganglia, ends primarily in the SMA, whereas information from the association areas reaches large parts of the prefrontal cortex.

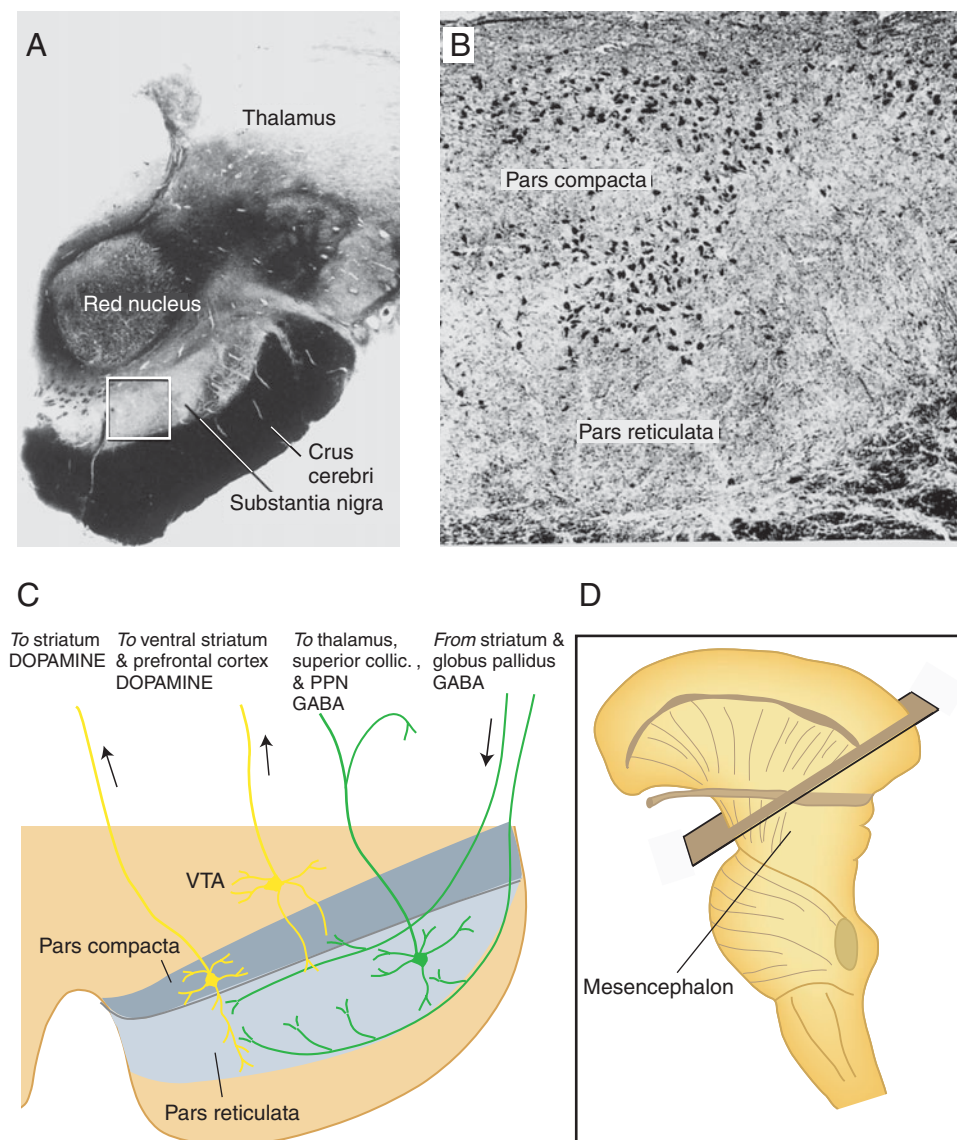


FIGURE 23.8 *The substantia nigra.* **A:** Photomicrograph of transverse section through the mesencephalon (myelin stain). Inset (**D**) shows plane and position of the section. The photomicrograph is from the section shown in Fig. 6.21. **B:** Higher magnification of framed area. The cells of the pars compacta are clearly seen as dark dots. The dark color is due to their content of pigment. **C:** The two parts of the substantia nigra differ structurally and with regard to connections. Afferent fibers to the pars reticulata contact the dendrites of the dopaminergic neurons in the pars compacta.

Thalamostriatal Connections

The thalamostriatal projections have marked effects on striatal-neuron excitability, yet their functional role is poorly understood. The main sources of thalamostriatal fibers are the **centromedian** (CM) and **parafascicular** (Pf) nuclei. In addition, projections arise in the specific thalamic nuclei. Thus, while the VL and VA send their main efferents to the cerebral cortex, they also send many fibers back to the striatum. The projection from the intralaminar nuclei has been most studied and might provide the striatum with information about events that requires special **attention**. Thus, neurons in CM integrate various kinds of sensory information and increase their activity when a person switches from a relaxed waking state to an attention-demanding task. A major input to the CM and Pf comes from the globus pallidus (internal segment) and the substantia nigra (pars reticulata) in the form of collaterals of fibers

ending in the specific thalamic nuclei. Thus, the thalamostriatal projection seems to be part of a side loop in the pathway from the basal ganglia to the cortex—handling a copy of the pallidal output and sending “answers” back to the striatum.

Efferent Connections of the Basal Ganglia: Striatal Projection Neurons

More than 70% of the striatal neurons have relatively small cell bodies and dendrites with numerous spines (Fig. 23.9A) and are called **medium spiny neurons**.² This cell type contains γ -aminobutyric acid (GABA) and sends its axon out of the striatum to the globus

² The percentage of projection neurons is higher in subprimate species (90%). This agrees with an evolutionary trend with increasing proportion of interneurons in general (e.g., in the cerebral cortex, the thalamus, and several brain stem nuclei).

pallidus and the substantia nigra. There are two main **subtypes** of this medium spiny neuron (Fig. 23.9A). One kind contains **substance P** (and usually also dynorphin) in addition to GABA; the other contains **enkephalin** in addition to GABA (a third, less characterized kind of projection neuron contains neurokinin B in addition to substance P and enkephalin). Physiological studies suggest that the two kinds of medium spiny neurons project to different targets (Fig. 23.9A): **GABA + substance P** neurons project primarily to the GP_i and/or to the **substantia nigra**, whereas **GABA + enkephalin** neurons project mainly to the GP_e .³

The two kinds of medium spiny neurons differ also with regard to expression of **dopamine receptors**: GABA + substance P neurons express predominantly D_1 receptors, whereas GABA + enkephalin neurons express mainly D_2 receptors. Because these two kinds of receptor have opposite effects, the two kinds of medium spiny neurons respond differently to dopamine. We return to this point when discussing dopamine actions in the striatum.

Striatal Interneurons: Cholinergic and GABAergic

The striatum has a complex intrinsic organization, even though the majority of neurons are “simple” projection neurons. For example, the axons of the medium spiny neurons give off recurrent collaterals in the striatum (Fig. 23.9A). Further, a number of different **interneurons** exist (i.e., their axonal arborizations remain within the striatum).

One conspicuous kind of interneuron, which constitutes about 1% of all striatal neurons, has a large cell body and smooth dendrites and contains **acetylcholine** (Fig. 23.9A). These interneurons receive excitatory synaptic influences from the cerebral cortex and the intralaminar thalamic nuclei (glutamatergic), as well as inhibitory influences by the recurrent collaterals from GABAergic projection neurons. Acetylcholine acts via **muscarinic receptors** (indirectly linked with K^+ channels) on striatal projection neurons with slow, modulatory effects. The overall effect of acetylcholine on the projection neurons is to ensure that they react with bursts of action potentials to excitatory inputs from the cerebral cortex: that is, the efficiency of signal transmission is enhanced. The cholinergic interneurons contribute—in a yet unknown way—to the symptoms in Parkinson’s disease, since anticholinergic drugs can improve the symptoms.

³ In at least apparent contradiction to such a dichotomy, tracing of single axons indicate that the majority of medium spiny neurons send collaterals to GP_e , GP_i , and the nigra. If the innervation density differs among the targets, however, a functional dichotomy may nevertheless be compatible with the observed branching of axons to several targets.

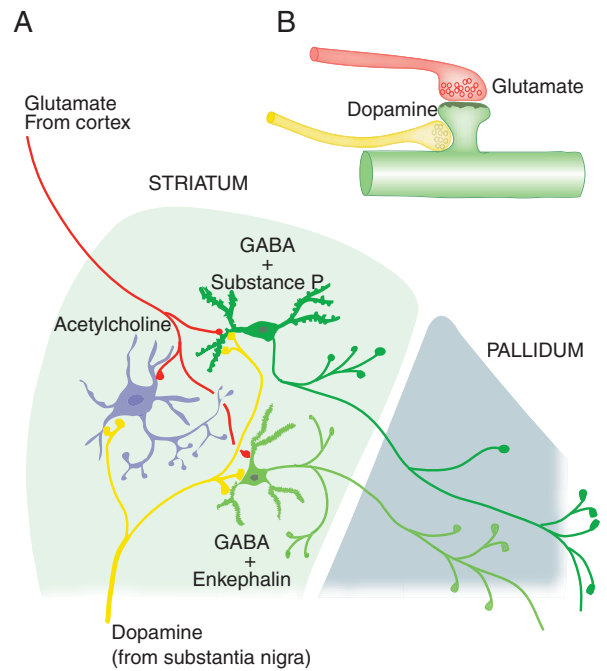


FIGURE 23.9 Three important kinds of neuron in the striatum, and the termination of dopaminergic fibers. **A:** The striatal projection neurons are relatively small and their dendrites have many spines (medium spiny neurons). All striatal projection neurons are GABAergic but two kinds can be distinguished based on their content of neuropeptides (substance P and enkephalin). Note the large, cholinergic interneurons. GABAergic interneurons are not shown. **B:** Dopaminergic synapses are often situated on spines that are contacted by glutamatergic nerve terminals from the cortex. In addition, dopaminergic fibers end without forming synapses. Therefore, dopamine presumably acts via both specific synapses and volume transmission.

GABAergic inhibition plays an important role in the functioning of the striatum, even though about 80% of all synapses are glutamatergic (mainly from the cortex and the thalamus). Thus, by blocking $GABA_A$ receptors in the striatum, the spontaneous firing rate of the projection neurons triples. The inhibition is partly due to the recurrent collaterals of the medium spiny neurons, partly to several types of fast-spiking **GABAergic interneurons**, which receive excitation from the cerebral cortex and in turn inhibit the medium spiny neurons. The recurrent inhibition has been assumed to produce lateral inhibition (see Fig. 13.4), whereas the function of the inhibitory interneurons may to regulate overall excitability in a feedforward manner, in contrast to the feedback action of the projection-neuron recurrent collaterals.⁴

⁴ The effects of GABA in the striatum are more complex than just producing inhibition by binding to $GABA_A$ receptors, however. This is because the projection neurons have a strongly polarized resting potential (-80 to -90 mV). That means that GABA *depolarizes* the projection neurons in the resting state. Only when the membrane potential has been reduced to about -60 mV (by excitatory inputs) does GABA produce hyperpolarization (inhibition).

Dopaminergic terminals of nigrostriatal neurons contact the projection neurons and the cholinergic interneurons (Fig. 23.9). In agreement with this, both cell types express mRNA for dopamine receptors.

Compartments in the Striatum: Islands, Striosomes, and Matrix

The various cell types, neurotransmitters and afferent connections are not evenly distributed throughout the striatum. First, the striatal neurons form clusters, called **islands**. Further, a **mosaic** pattern appears after staining to demonstrate **acetylcholine esterase**. Poorly stained patches called **striosomes** are embedded in a heavily stained **matrix**. The matrix can be further subdivided by visualization of various transmitters and their receptors. Cholinergic interneurons and GABAergic projection neurons are found within both the matrix and the striosomes, and the two main kinds of projection neurons do not appear to be clearly segregated, either. However, corticostriatal fibers terminating in the striosomes and in the matrix come from deep and superficial parts of layer 5, respectively. Thus, cortical information to neurons in the two compartments would be expected to differ slightly, even when coming from the same cortical area. Further, the further projections also appear to differ with regard to exact termination and patterns of arborizations. Finally, striatal afferent connections terminate in patches within the matrix. Each small part of the cortex, for example, projects divergently in many patches. These data have been taken as evidence that the striatum is organized in numerous minor compartments or **modules**, each presumably representing a functional unit. It is fair to say, however, that in spite of a wealth of data, the functional significance of striatal compartmentalization is still a matter of speculation (and the interpretation of data is made more difficult by the existence of species differences).

Efferent Connections of the Basal Ganglia: Acting on Premotor Networks in Thalamus and Brain Stem

The efferent connections of the basal ganglia may be summarized as follows. The internal segment of the globus pallidus and the substantia nigra send the information processed in the basal ganglia to premotor networks in the thalamus, the mesencephalon, and the superior colliculus. Here we use the terms premotor and **premotor networks** in a rather loose sense about neuronal groups acting either directly on motoneurons or on the motor cortex. Such premotor networks are found in the spinal cord, the reticular formation, and the thalamus. Neurons of the thalamic VL nucleus, for example, are considered premotor due to their direct projection to the motor cortex. Premotor networks organize the activity of motoneurons to produce purposeful

actions, not merely isolated movements. Thus, the effects exerted by the basal ganglia on other parts of the nervous system are mediated primarily by efferent fibers from the **pallidum** and the **substantia nigra**. These nuclei receive their main afferents from the striatum (Figs. 23.7 and 23.10). In this manner, the pallidum and nigra process information from the striatum before it is sent to premotor networks. The efferents from the striatum are **topographically organized**, so that subdivisions of the striatum are connected with specific parts of the pallidum and the nigra.

The Substantia Nigra

The substantia nigra and some of its connections have been mentioned several times, and we will also return to it when dealing with Parkinson's disease, in which the nigra plays a crucial role. A collective treatment of the main features of the substantia nigra may therefore be pertinent at this stage

The substantia nigra can be divided anatomically into two parts, the **pars compacta** and the **pars reticulata** (Figs. 23.8A and 13.11A). The compacta is richer in cells than the reticulata, whereas the latter (as the name implies) is dominated by dendritic arborizations. The reticulata also contains numerous neurons, however. The compacta neurons contain pigment (neuromelanin), which makes the nigra visible as a dark band in the cut human mesencephalon (see Fig. 23.4). The pars

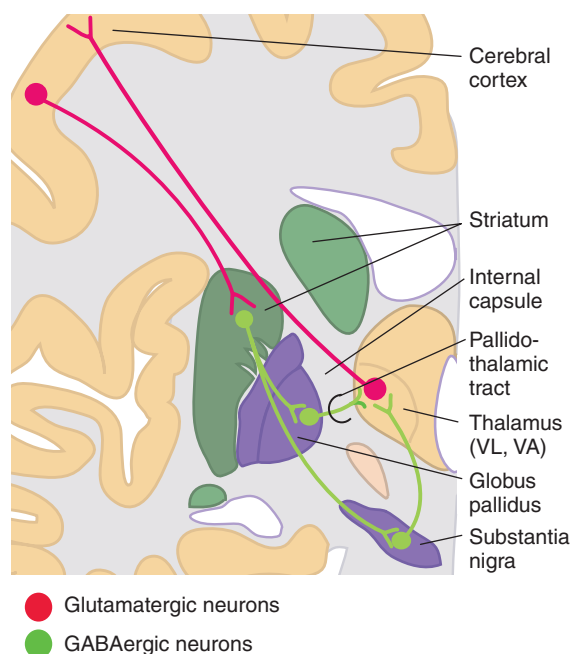


FIGURE 23.10 The loop cortex–basal ganglia–thalamus–cortex contains two inhibitory neurons in series. This means that increased signal traffic out of the striatum leads to less inhibition in the thalamus (disinhibition). Cf. text.

reticulata, located ventral to the compacta, is lighter. The dopaminergic nigrostriatal neurons are located in the pars compacta, whereas the GABAergic nigrothalamic neurons are located primarily in the pars reticulata (Fig. 23.8C).

The **efferent** connections of the **pars compacta** (Fig. 23.11A) pass primarily to the striatum (with a smaller contingents to the subthalamic nucleus and some other nuclei). This is the largest dopaminergic pathway in the brain, and nigra is the largest collection of dopamine-containing neurons. **Pars reticulata** send GABAergic fibers to the thalamus (VA, MD). In addition, it projects to the superior colliculus (Fig. 23.11A), which control coordinated eye and head movements, and to the pedunculopontine nucleus (PPN) involved control of gait and posture. The nigral neurons sending their axons to the thalamus and the superior colliculus are found in largely separate parts of the pars reticulata.

The **afferent** fiber connections of the nigra (Fig. 23.11B) arise in numerous cell groups, but quantitatively the most important input comes from the **striatum**. This projection shows some topographic organization; for example, afferents from the putamen and caudate nucleus end differently. Even though most striatonigral fibers terminate in the pars reticulata, cells in the compacta can also be influenced because their long dendrites extend into the reticulata (Fig. 23.8). GABA is the transmitter for the striatonigral fibers, exerting

inhibitory effects on the cells in the nigra. **Excitatory** afferents to the nigra arise in the subthalamic nucleus and the pedunculopontine nucleus (PPN) in the mesencephalon (glutamate). Afferents with **modulatory** effects come from the locus coeruleus (norepinephrine) and the raphe nuclei (serotonin). Additional afferents arise in the **ventral striatum** and the **bed nucleus of the stria terminalis** (BST), presumably mediating signals related to motivation, attention, and mood. An important link in the latter pathways is the **habenula** (especially the lateral nucleus). Electric stimulation of the lateral habenula effectively suppresses activity of dopaminergic neurons, presumably by activating inhibitory interneurons.

Pathways from the Globus Pallidus and the Substantia Nigra to the Cerebral Cortex

As mentioned, the **globus pallidus** consists of two parts, an **external** (GP_e) and an **internal** (GP_i) segment (Figs. 23.3 and 23.4). Both segments receive their main **afferents** from the striatum, with additional inputs from the subthalamic nucleus. Whereas the fibers from the striatum exert inhibitory actions in the GP_e , the subthalamic afferents are excitatory. The balance between these two inputs therefore to a large extent determines the activity of the GP_i neurons. The main part of the **efferents** from GP_i goes to the thalamus and the substantia nigra (pars reticulata), whereas the GP_e projects

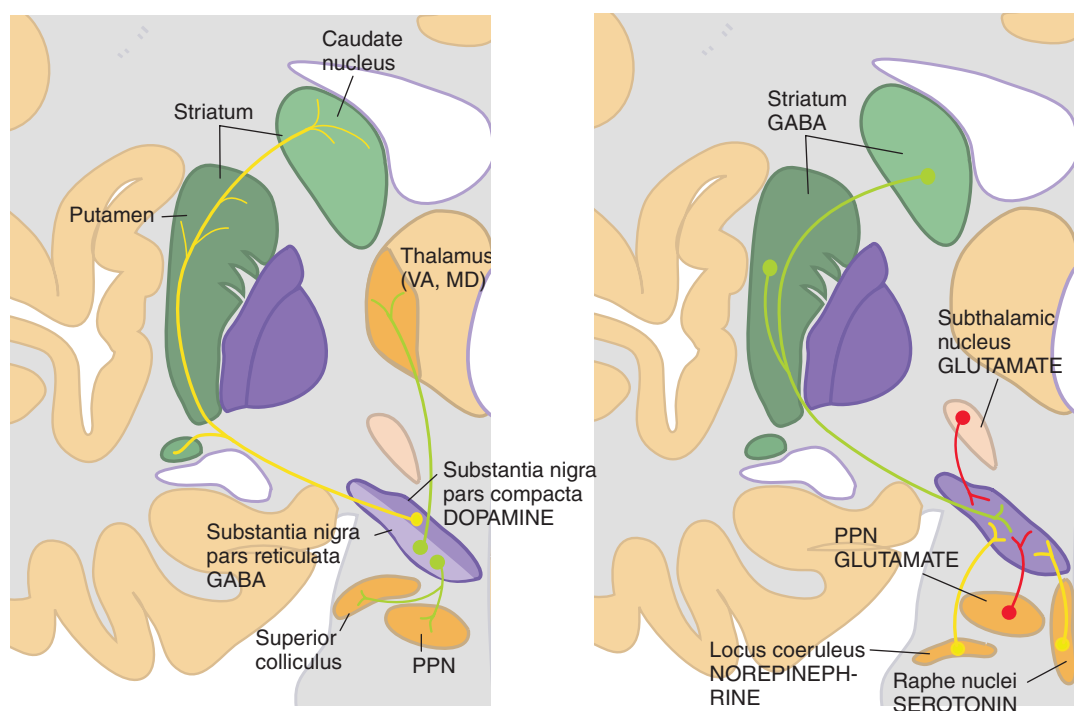


FIGURE 23.11 Main connections of the substantia nigra. **A:** Efferents. Dopaminergic neurons in the pars compacta send fibers to the striatum, whereas GABAergic neurons in the reticulata act on premotor neurons in the thalamus and the brain stem. **B:** Afferents. Inhibitory

(GABAergic) afferents come from the striatum, whereas excitatory (glutamate) fibers come from the subthalamic nucleus and the PPN. Modulatory afferents arise in the locus coeruleus (norepinephrine) and the raphe nuclei (serotonin).

mainly to the subthalamic nucleus. Many of the **pallidothalamic fibers** pass through the internal capsule (Fig. 23.10) and can therefore be damaged by lesions in this region (this should be kept in mind when analyzing the symptoms occurring after capsular infarctions, as discussed in Chapter 22).⁵ In the thalamus the pallidal fibers end in the **ventral anterior nucleus (VA)** and in parts of the **ventrolateral nucleus (VL)** (Figs. 23.3 and 13.4; see also Fig. 14.6). The VL and VA send their main efferents to the cerebral cortex but also many fibers back to the striatum. In more detail, the VA sends efferent fibers to the premotor area (PMA) and the prefrontal cortex, whereas the parts of the VL that receive fibers from the pallidum and the nigra project primarily to the supplementary motor area (SMA; see Fig. 24.16). The **substantia nigra** sends fibers to the thalamus, ending in partly different nuclei than the pallidal fibers. Thus, the efferents from pars reticulata of the substantia nigra end in parts of the VA and the mediodorsal nucleus (MD; Figs. 23.7 and 23.11A). From there, signals travel primarily to parts of the PMA and the prefrontal cortex.

Summing up, much of the information flow from the basal ganglia is directed toward the motor cortical areas. Physiological studies also indicate that the influence of the basal ganglia on motor control involves the corticospinal tract and indirect descending pathways from the cerebral cortex to the motoneurons (Fig. 23.1). It should be kept in mind, however, that the **caudate nucleus** receives afferents mainly from cortical association areas and acts primarily on the **prefrontal cortex** (Fig. 23.7), which is not directly involved in motor control but in cognitive functions, such as memory and planning of behavior. Accordingly, **lesions** of the caudate nucleus in monkeys produce symptoms similar to those seen after damage to the prefrontal cortex—among other things, reduced performance in tests requiring **spatial memory** (such as to recall where an object is located when it is out of sight).

Basal Ganglia Efferents to the Brain Stem: Pedunculopontine Nucleus and Superior Colliculus

In addition to the massive connections from the basal ganglia to the thalamus, there are also efferents reaching the **reticular formation** in the mesencephalon (Fig. 23.11A). This enables the basal ganglia to influence muscle tone and movements via reticulospinal tracts. Connections from the pars reticulata of the substantia nigra and from the GP_i to a part of the mesencephalic reticular formation called the **pedunculopontine nucleus**

(PPN) have attracted special interest. Thus, this region overlaps with the mesencephalic locomotor region (electrical stimulation of this region elicits walking movements).⁶ Conceivably, these connections are of relevance for the characteristic disturbances of muscle tone (rigidity) and locomotion seen in **Parkinson's disease** (in which there is marked cell loss in the substantia nigra but also in the PPN). Some observations furthermore indicate that the characteristic poverty of movement—**akinesia**—in Parkinson's disease might be due to increased inhibition of the PPN from the globus pallidus. For example, destruction of the PPN in monkeys has been reported to produce akinesia.⁷

Connections from the substantia nigra to the **superior colliculus** (Fig. 23.11A) participate in the control of coordinated **head and eye movements**.

The Basal Ganglia Are Arranged in Parallel Circuits

Studies with axonal tracing techniques indicate that there are four (or more) loops or **circuits** through the basal ganglia, in which the flow of information is kept at least largely segregated. This localization goes further than depicted in Fig. 23.7, which shows only the main differences between projections from association areas and the pericentral (motor and somatosensory areas) region. Four circuits are now well established: one originating in sensorimotor areas, one in association areas, one in so-called limbic cortex, and one in oculomotor cortical areas. Via the basal ganglia and the thalamus, signals are funneled back to largely separate parts of the cortex. Thus, the basal ganglia appear, at least as a general rule, to process different kinds of information in **parallel**.

One circuit arises in the **SMA, MI, and SI**; passes through the putamen; and, via the pallidum and the thalamus, ends mainly in the SMA. This circuit is probably the one most directly involved in control of movements. Recordings of single-cell activity in the putamen support this assumption. Each circuit thereby seems to be specialized for certain tasks. It should be emphasized, however, that although there probably is modest integration among the various circuits, each circuit integrates information from anatomically separated, although functionally related, cortical areas. A second circuit arises in different parts of the **prefrontal cortex** and passes through the caudate nucleus, the substantia

5 The efferent fibers from the pallidum form two bundles, the **ansa lenticularis** and the **fasciculus lenticularis**. They arise from somewhat different parts of the internal pallidal segment and fuse to form the **fasciculus thalamicus** after having traversed the internal capsule.

6 Most fibers from the GP_i to the PPN are collaterals of pallidothalamic fibers, suggesting that the thalamus and the PPN receive the same kind of information from GP_i. The PPN projects back to the pallidum and also to the subthalamic nucleus. Obviously, the PPN not only is an output channel from the basal ganglia to brain stem premotor cell groups but also acts back on the basal ganglia, influencing their internal information processing.

7 That connections from the globus pallidus to the PPN are involved in **akinesia** is further supported by the effects of stereotaxic surgery in patients with Parkinson's disease. Thus, lesions of the GP_i often improve the akinesia, whereas lesions of the thalamus do not have this effect.

nigra (to a lesser degree the pallidum), the VA, and the mediodorsal thalamic nucleus (MD in Fig. 23.3; see also Figs. 21.7 and 21.8) and back to prefrontal cortical areas. The third arises in “limbic” parts of the cortex (the cingulate gyrus, orbitofrontal areas of the prefrontal cortex, and parts of the temporal lobe cortex, which are parts of or closely connected with the limbic structures). This “limbic” circuit passes through the ventral striatum, ventral pallidum, and the MD of the thalamus and back to the cortical areas from which the circuit started. These connections are believed to be involved in regulation of mood and emotions, and to behavior directed to satisfy basal needs (such as eating and drinking), and to obtain rewards (see “The Ventral Striatum, Psychosis, and Drug Addiction,” later). In addition, a fourth circuit in parts of the frontal and parietal lobes concerned with **oculomotor control**—that is, **area 8** immediately in front of area 6 (the frontal eye field) and **area 7** (see Fig. 25.7). After synaptic interruption in specific parts of the caudate nucleus, this circuit ends primarily in area 8. This prefrontal circuit presumably enables the basal ganglia to influence cognitive functions.

While the principle of parallel processing in the basal ganglia seems well established, there is evidence that there is also considerable “crosstalk” among the circuits. Tracing experiments in animals and DWI (diffusion weighted imaging) studies humans suggest that the thalamus might be a main site for integration in the basal ganglia circuits.

All the circuits are influenced by the **subthalamic nucleus** (Fig. 23.4). Indeed, subdivisions of this small nucleus project with topographic order to the dorsal and ventral pallidum. This might explain why electric stimulation of the subthalamic nucleus (**deep brain stimulation**) aimed at improving motor symptoms in Parkinson’s disease can give side effects such as mood changes (depression or mania), cognitive decline, and personality changes. Affections of circuits involving the prefrontal cortex are especially likely to produce such side effects.

Transmitters and Synaptic Actions in the Cortex–Basal Ganglia–Cortex Loop: Disinhibition

To understand the processing going on within the basal ganglia and their effects on other parts of the brain, we need to know the transmitters and the synaptic actions of all the neurons involved. The conditions turn out to be extremely complex. Numerous neuroactive substances have been found in the striatum, although only a few can be correlated with anatomic and physiologic data.

The **corticostriatal** and **thalamostriatal** fibers are excitatory due to release of glutamate (Fig. 23.12). As discussed in the preceding text, the **striatopallidal** and **striatonigral** fibers contain GABA and have inhibitory effects. The neurons in the globus pallidus are also

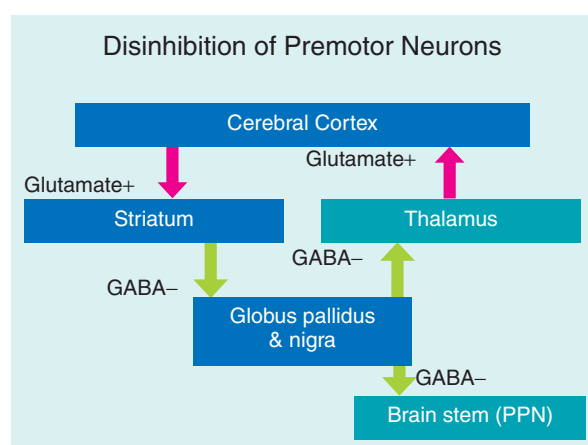


FIGURE 23.12

GABAergic—that is, the **pallidothalamic** fibers from the GP_i and the **pallidosubthalamic** fibers from the GP_e are inhibitory. The **nigrothalamic** fibers, arising in the pars reticulata, are also GABAergic and thus inhibitory. The **thalamocortical** fibers (from the VL and VA, as from other thalamic nuclei) are excitatory (glutamate).

The flow of information from the cerebral cortex through the basal ganglia and back is obviously less straightforward than if all involved synapses were excitatory. Notably, there is a chain of two inhibitory neurons from the striatum to the **premotor** neurons in the thalamus and the brain stem (Fig. 23.12). Increased cortical input to the striatum would lead to decreased activity of the pallidal neurons, because excitation of striatal neurons produces increased inhibition in the pallidum. In turn, this would increase the activity of thalamocortical neurons (because they would receive less inhibition from the pallidum). Thus, excitatory impulses from the cortex would eventually produce **disinhibition** of the thalamocortical neurons (and other “premotor” neurons in the reticular formation and the superior colliculus receiving fibers from the substantia nigra).

Functional Significance of Disinhibition in Premotor Neuronal Groups

Recordings of **single-cell** activity in the basal ganglia in awake monkeys are compatible with the aforementioned considerations. At rest, most striatal neurons are “silent”—that is, they do not produce action potentials—whereas the pallidal neurons and neurons in the pars reticulata of the substantia nigra fire with a **high, regular frequency**.⁸ This would presumably keep the

8 This is not quite true for the GP_i neurons, however, because they fire with a somewhat lower and more uneven frequency than GP_i neurons, and furthermore tend to fire in bursts.

premotor neurons in the thalamus and the brain stem in a state of inhibition when the animal is not moving. Commands from the cortex to the basal ganglia in relation to the preparation or execution of movements would release the **premotor neurons** from this inhibition. Indeed, electrophysiological experiments show that increased striatal activity reduces the activity of many pallidal and nigral neurons, followed by increased firing of thalamocortical neurons.

It has been proposed that the disinhibition of premotor neurons by the basal ganglia is a **gating mechanism** to control the access of other inputs (e.g., sensory) to the motor cortex. As the connections of the basal ganglia are topographically organized at all levels, this would be a specific and focused gating rather than a diffuse one, varying with the nature of the motor task. Such focused effects might serve to reinforce wanted movements while suppressing unwanted ones.

The Subthalamic Nucleus Regulates Pallidal and Nigral Activity

While efferents of the internal pallidal segment (GP_i) end primarily in the thalamus (and the nigra), the GABAergic fibers from the GP_e are directed toward the subthalamic nucleus (Fig. 23.13). The subthalamic nucleus also receives excitatory afferents from the **motor cortex**, thus constituting a cortical input to the basal ganglia in addition to the major corticostriatal pathway. Most of the **efferents** from the subthalamic nucleus go back to both segments of the pallidum and to the pars reticulata of the substantia nigra. As mentioned, the **subthalamopallidal** fibers exert excitatory actions.

The efferents of the subthalamic nucleus are **topographically** organized, as shown with axonal transport methods. Thus, different neuronal populations project to the substantia nigra and the pallidum, and there are differences with the regard to projections to minor parts of the pallidum. Fibers from the subthalamic nucleus and the striatum converge on the same neurons in the GP_i. These data suggest that the activity of the GABAergic neurons in GP_i are determined largely by the sum of synaptic influences from the striatum (–) and the subthalamic nucleus (+). Because GP_i neurons inhibit thalamocortical neurons, increased subthalamic activity would be expected to produce reduced excitation of the motor cortex—that is, inhibition of voluntary movements. Thus, the subthalamic nucleus is believed to control or **stop ongoing movements**, rather than to select and initiate movements. Considering the somatotopical organization of its connections, the subthalamic nucleus presumably exerts specific actions rather than a diffuse inhibition of all motor activity.

Loss of influence from the subthalamic nucleus would produce **disinhibition** of thalamocortical neurons.

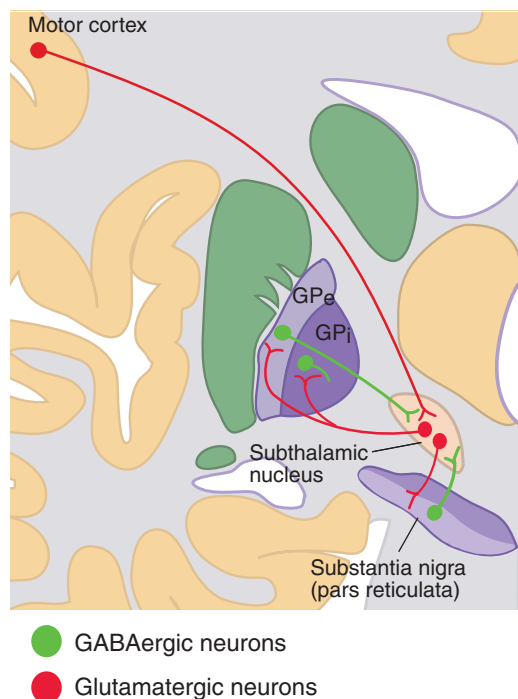


FIGURE 23.13 *Main connections of the subthalamic nucleus.* Inhibitory afferents arise in the globus pallidus and substantia nigra, whereas excitatory connections come from (among other sources) the motor cortex. Glutamatergic neurons in the subthalamic nucleus excite the GABAergic neurons in the globus pallidus and the substantia nigra. Especially the connections from GP_e to the subthalamic nucleus and from there to GP_i have been postulated to be of special significance for the symptoms in Parkinson's disease.

Thus, the violent involuntary movements of the opposite body half after destruction of the subthalamic nucleus—**hemiballismus**—might be caused by hyperactivity among the thalamocortical neurons. In a yet unknown way, disturbed activity of subthalamic neurons (abnormal high-frequency firing and oscillatory activity) appears to be crucial to the symptoms in **Parkinson's disease**. Indeed, “switching off” the subthalamic nucleus by lesion or by electric stimulation has produced marked symptomatic relief in many patients.

Actions of Dopamine in the Striatum

The connections and transmitters described so far are concerned with fast transmission of specific information, mediated by glutamate and GABA. In addition, modulatory transmitters have important roles in the functioning of the basal ganglia. This concerns **dopamine** in particular. Indeed, the most studied basal ganglia connection is the dopaminergic **nigrostriatal pathway**, which is the most massive dopaminergic pathway in the central nervous system (CNS). Many dopaminergic nerve terminals are strategically placed on spines close to the corticostriatal nerve terminals

(Fig. 23.9B).⁹ This is a pathway of great clinical interest because several diseases are related to disturbed dopamine-mediated synaptic transmission (Parkinson's disease, schizophrenia, Tourette's syndrome, and others).

Striatal neurons express, not unexpectedly, **dopamine receptors**. Dopamine alters the response of striatal neurons to specific inputs from the cerebral cortex and the thalamus. In more detail, however, actions of dopamine on neuronal excitability in the striatum are multifarious and not fully understood.

There are two main kinds of dopamine receptor—**D₁** and **D₂**, with subtypes of each. Common to the **D₁-like** receptors (D₁ and D₅) is that they increase the synthesis of intracellular cyclic AMP, whereas **D₂-like** receptors (D₂–D₄) have the opposite effect. Dopamine receptors are present both pre- and postsynaptically in the striatum, and dopamine influences, among other things, several kinds of ion channel. The actions of the D₁ receptor is most studied, whereas more remains to be known about the D₂ receptor.

Although there is, as mentioned previously, physiological evidence that the major effect on striatal projection neurons of D₁-receptor activation is excitatory and that of D₂-receptor activation is inhibitory, the actions of dopamine is not properly described by this simple dichotomy. One factor complicating the analysis is that dopaminergic neurons fire in two characteristic modes, **tonic** and **phasic**, with either a sustained or transitory rise in dopamine concentrations. There is some evidence that phasic release of dopamine acts on D₁ receptors (excitatory, LTP induction) whereas tonic release activates D₂ receptors (inhibitory, LTD induction). Because the two main types of medium spiny neurons express different dopamine receptors (Fig. 23.14), this would imply the neuronal target of dopamine depends on the firing mode of the nigrostriatal neurons. Another factor making it difficult to generalize dopamine actions is that the effect of dopamine depends on the **state** of the postsynaptic neuron. For example, D₁ receptors may produce depolarization or hyperpolarization, depending on the membrane potential of the postsynaptic neuron. In one state, the neuron is depolarized and fire bursts of action potentials; in the other state it is hyperpolarized and inactive. In the inactive state, activation of D₁ receptors depolarizes the cell (by closure of K⁺ channels) so that it reacts more easily with burst to an excitatory input from the cortex. If the neuron is in the active state, however, D₁ receptor activation closes a Na⁺ channel that keeps the neuron depolarized by leakage of cations into the cell. Closing this channel would

stabilize the membrane potential, perhaps to avoid excessive activation by glutamate from the cortex.

In conclusion, dopamine probably serves to keep the membrane potential in the range where the postsynaptic neuron is apt to fire in **bursts**—that is, in a state suited for efficient signal transmission. Some data suggest that D₁ receptor activation enhances the activity of neurons that receive a strong and **focused excitatory input** (from the cerebral cortex), while reducing the activity of neurons receiving weak inputs. This would help in focusing striatal activity, in accordance with other data indicating that the basal ganglia assists in the **selection of behavior**, such as the choice of specific movements.

The “Indirect Pathway,” the Subthalamic Nucleus, and Parkinson's Disease

The signal pathway we described above—cortex-striatum-GP_i-thalamus-cortex—are often said to establish a

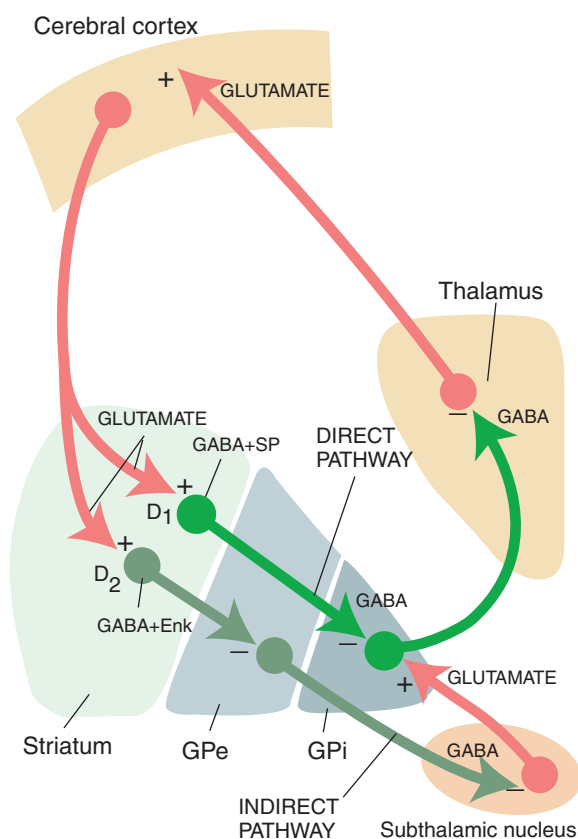


FIGURE 23.14 “Direct” and “indirect” pathways through the basal ganglia. One pathway goes directly from the globus pallidus to the thalamus; the other goes indirectly via the subthalamic nucleus. Excitation of striatal projection neurons from the cerebral cortex would produce disinhibition in the thalamus via the direct pathway, while producing inhibition via the indirect pathway. Disturbance of the balance between these two pathways is postulated to explain some of the symptoms in Parkinson's disease. However, anatomic data speak against that the division between the direct and indirect pathway is as sharp as shown in this figure.

⁹ Dopamine depletion in the striatum is associated with loss of dendritic arborizations and spines (in Parkinson's disease and in animal experiments). Therefore, dopamine probably has a direct growth-promoting effect on neurons, or it protects against the harmful effects of strong glutamatergic excitation (or both).

direct signal pathway to distinguish it from an indirect pathway (Fig. 23.14). The latter involves a side-loop connecting the GP_e to the subthalamic nucleus, which projects back to the GP_i. Findings of increased activity in the subthalamic nucleus in animal models of Parkinson's disease led to great interest in the "indirect" pathway because it offered a possible explanation of how loss of striatal dopamine can lead to subthalamic hyperactivity. As described in the preceding text, it has been assumed that striatal medium spiny neurons containing GABA and substance P project mainly to the GP_i and express D₁ receptors, whereas neurons containing GABA and enkephalin project mainly to GP_e and express D₂ receptors (Fig. 23.9). These assumptions led to the postulation of two parallel signal pathways out of the striatum: the direct and the indirect. In conjunction with data showing that D₁ receptors excite striatal projection neurons whereas D₂ receptors inhibit them, this could explain the subthalamic hyperactivity. The model predicts that release of dopamine in the striatum excites striatal neurons that project to the GP_i while inhibiting neurons projecting to the GP_e. This would increase inhibition in the GP_i but reduce it in the GP_e, thereby evoking increased inhibition of subthalamic neurons. Because of reduced firing, the subthalamic excitatory influence on the GP_i would be diminished. Thus, release of dopamine in the striatum would—via both the direct and the indirect pathways—reduce excitation in the GP_i. Thus would give less inhibition (disinhibition) of thalamic and nigral neurons from GP_i—that is, the excitation of cortical motor areas would increase. In this way, dopamine would facilitate movements via both pathways. In case of **dopamine depletion** in the striatum, the GP_e would receive stronger inhibition from the striatum (because the GABA + enkephalin neurons are no longer inhibited by dopamine), and this was postulated to explain the increased subthalamic activity.

Even though subthalamic dysfunction undeniably is of crucial importance in Parkinson's disease, the clear division between a direct and an indirect pathway may not be tenable. One problem is, as mentioned, that many striatal neurons send branches to several targets (not only to GP_i or GP_e). Moreover, the scheme in Fig. 13.14 is highly simplified because it leaves out several fiber connections. For example, the subthalamic nucleus does not only project to the GP_i but also to the GP_e. Finally, the segregation of D₁ and D₂ receptors in the striatum may not be as sharp as depicted in Fig. 13.14. Thus, studies with sensitive techniques indicate that many striatal projection neurons express both D₁ and D₂ receptors, although apparently the expression of one receptor type dominates. Finally, experimental studies in Parkinsonian monkeys have not been able to confirm the postulated reduced activity in GP_e neurons (explaining the subthalamic hyperactivity).

What Activates the Nigrostriatal Neurons?

In light of the dominating effects of dopamine in the striatum, it is of particular interest to know under which conditions the nigrostriatal neurons are activated. Clues may come from the fact that many of the nuclei sending fibers to the nigra change their activity in relation to **arousal, motivation, and emotionally** driven behavior. This concerns the PPN, the locus coeruleus, the raphe nuclei, and the ventral striatum and other cell groups in the basal forebrain.¹⁰ Accordingly, physiological studies show that striatonigral neurons do not change their firing in relation to movements but in relation to stimuli that are **unexpected** or are judged to be of particular **salience** for the behavior of the animal at the moment. This may concern stimuli signaling reward or punishment, although they must have a fairly high intensity to activate dopaminergic neurons.

In general, dopamine assists in establishing associations between stimuli and their reward value. This kind of **learning** underlies motivated behavior based on prior experience. As mentioned, dopaminergic neurons in the midbrain fire either in bursts or tonically. Selective reduction of burst firing in mice (by genetic manipulations), support the idea that a brief release of dopamine acts as a signal for associative learning, especially related to potentially rewarding or dangerous events.

THE VENTRAL STRIATUM

The term ventral striatum is used for rather diffusely distributed cell groups in the basal part of the hemispheres (the basal forebrain; see Chapter 31, under "The Basal Forebrain," for a discussion of its constituting parts). The ventral striatum merges with the dorsal striatum without sharp boundaries. The **nucleus accumbens** represents a fairly distinct part of the ventral striatum, and connects the caudate nucleus and the putamen ventrally (Fig. 23.15). A similar diffuse cell group, the **ventral pallidum**, merges with the dorsal well-defined parts of the globus pallidus (see Fig. 31.8). The **ventral striatopallidum** is used as a collective term.

Connections of the Ventral Striatum

Although the ventral striatopallidum has no sharp boundaries toward other cell groups in the basal forebrain, its

10 The **habenula** receives afferents from these nuclei and the hypothalamus and projects to the substantia nigra pars compacta and the VTA. It seems to play a particular role in controlling the activity of dopaminergic neurons (and other monoaminergic neurons). Signals from the habenula appear to occur especially in situations when an expected reward is not given. It has therefore been suggested to constitute a pivotal link in a "circuit of disappointment" (Fritz A. Henn, personal communication). Indeed, the lateral habenula shows altered activity in **depression** and is the target of **deep brain stimulation** for depressive disorders.

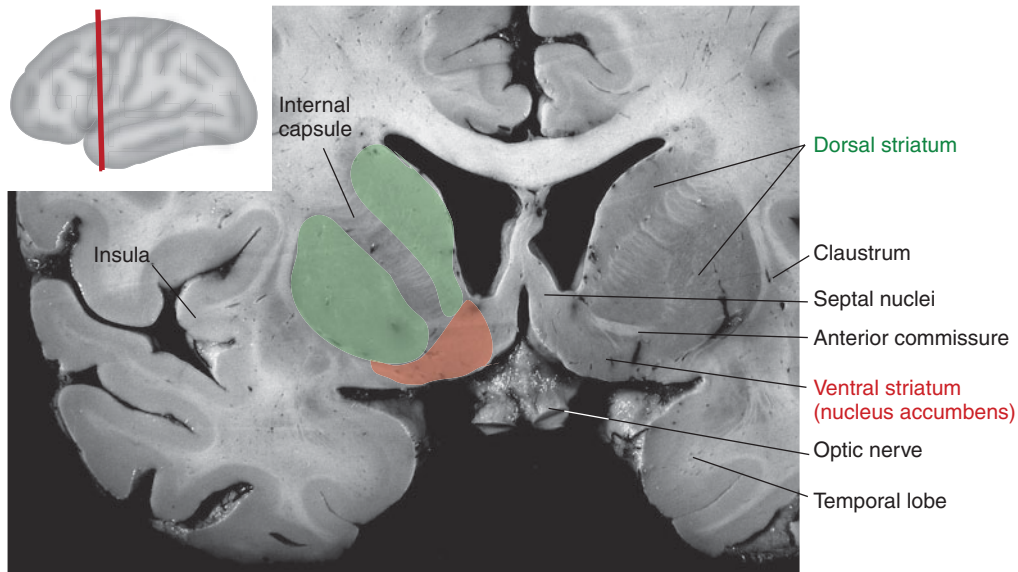


FIGURE 23.15 *The ventral striatum with the nucleus accumbens.* The septal nuclei, which belong to the basal forebrain, are also indicated.

Photo of a frontal section through the brain at the level of the anterior commissure (cf. Fig. 6.26).

relationship to the basal ganglia is witnessed by the similarities of their connections. We mentioned one of the **circuits** through the basal ganglia: from parts of the prefrontal cortex and cingulate gyrus to the ventral striatum, from there to the ventral pallidum, then to the mediodorsal thalamic nucleus (MD), and finally back to the prefrontal cortex. There is probably a further differentiation of connections through the ventral striatum (several circuits). Thus, it receives **afferents** with some topographic localization from the hippocampal formation, the amygdala, the orbitofrontal cortex, and parts of the temporal lobe (all these sources are either parts of the limbic structures or closely connected with them). The nucleus accumbens also send **efferent** fibers to the hypothalamus and the mesencephalic reticular formation (PPN).

Like the dorsal striatum, the ventral striatum receives many **dopaminergic fibers**; those projecting to the ventral striatum are located mainly dorsomedially to the substantia nigra, in the **ventral tegmental area (VTA)**. Since the dopaminergic neurons in VTA project primarily to the ventral striatum, the prefrontal cortex and other cell groups that are closely linked with limbic structures, the term the **mesolimbic dopaminergic system** is now widely used. To use the word “system” for this is hardly justified, however, as there is little reason to assume that these widespread dopaminergic projections are functionally homogeneous. **Afferents to the VTA** have been traced from the prefrontal cortex, the nucleus accumbens, and the PPN (and other nuclei).

The “Mesolimbic Reward System”

A popular theory suggests that the dopaminergic mesolimbic connections constitute a “**reward pathway**”

and that the pleasurable feelings evoked by, for example, narcotic drugs are caused by release of dopamine in the nucleus accumbens. Indeed, much evidence points to dopamine as the neurotransmitter most directly involved in the pleasurable effects of drugs of abuse. Nevertheless, many other parts of the brain than the nucleus accumbens are activated by stimuli evoking reward-motivated behavior, and there are substances that produce reward behavior without activating the mesolimbic dopaminergic neurons. Today, several other transmitters than dopamine are under intense scrutiny in addiction research—such as glutamate, acetylcholine, serotonin, and several neuropeptides. As said by the American neuropharmacologist Ann Kelley (2002, p. 448) in connection with nicotine addiction: “Thus, repeated exposure of the brain to drugs with abuse potential sets in motion a cascade of activity involving dopamine, glutamate and acetylcholine signaling.” Further, opinions on what drives the mesencephalic dopaminergic neurons has changed from solely focusing on reward to a more general activation by unexpected or highly salient stimuli requiring a change of behavior, as discussed above.

The Ventral Striatum, Psychosis, and Drug Addiction

The dopaminergic projection to the ventral striatum (especially to the nucleus accumbens) has attracted much interest, because **antipsychotic drugs** appear to bind with particularly high density in the ventral striatum. Such drugs are dopamine antagonists with preferred binding to D₂ receptors. Neurons in the nucleus accumbens of experimental animals change their pattern of activity during development of drug **addiction**,

no matter whether the drug is amphetamine, cocaine, or morphine. The addictive behavior is reduced by lesions of the nucleus accumbens or by removing its dopaminergic innervation. Thus, after giving a dopamine antagonist, the experimental animals stop self-administration of cocaine (they could easily obtain an intravenous dose by a movement). Further, dopaminergic activity is increased in paranoid **psychoses** elicited by amphetamines or cocaine (see also Chapter 5, under “GABA Receptors Are Influenced by Drugs, Alcohol, and Anesthetics,” “Nicotinic Addiction,” and “Drugs Altering Monoamine Activity in the Brain”).

However interesting these observations are, they provide only limited insight. By focusing on one transmitter and one part of the brain, one may even give the impression that there is a simple biologic explanation to complex mental phenomena. Indeed, many parts of the brain other than the nucleus accumbens show altered activity in the conditions discussed here (see Chapter 34 for some comments on mental illness and the cerebral cortex). In drug addiction, for example, changes of neuronal activity occur in the locus coeruleus (norepinephrine) and in the intralaminar thalamic nuclei, amygdala, and parts of the basal forebrain adjoining the nucleus accumbens. It is not clear whether the nucleus accumbens or the ventral tegmental area is the primary target of narcotic drugs in the brain; neither is it known how such drugs alter the properties of dopaminergic neurotransmission.

FUNCTIONS OF THE BASAL GANGLIA

In spite of enormous research activity during recent years, the functions of the basal ganglia are still far from fully understood. One reason is that recent research has made us realize that the basal ganglia participate in much more than just motor control; they also provide important contributions to cognitive and emotional processing. Further, their role in various aspects of learning attracts increasing interest.

Movement Planning and Learning

Taking the **connections** of the basal ganglia as a starting point (leaving out the ventral striatum for the time being), it is noteworthy that their major output goes to the SMA, PMA, and the prefrontal cortical areas. The properties of these areas suggest that the basal ganglia would be important in the **planning** phase of a movement, such as when several single-joint movements have to be put together to produce a complex movement, or when sensory stimuli or stored information has to be translated into an adequate motor response. Observations of symptoms in monkeys with lesions of the globus pallidus indicate that learned movements are slower than

normal, whereas the manner in which the task is performed is not significantly altered. Therefore, neither the movement command nor the movement program appears to be located within the basal ganglia themselves. There is also evidence that the basal ganglia participate when movements are **learned by repetition** and not by gaining insight into the nature of the task and, furthermore, that the basal ganglia enable **automatic** performance of well-rehearsed movements by the use of motor programs located elsewhere in the CNS. We also discussed earlier the role of the basal ganglia (and dopamine in particular) with regard to **associative learning**.

Motivation

Some experimental evidence shows that the basal ganglia contribute to the linking of **motivation** and **emotions** to the execution of movements. Thus, recording the activity of single cells in the striatum indicates that many respond best when a stimulus is linked with memory of an event that has a particular significance for the animal. For example, certain cells in the substantia nigra are active just before a rapid eye movement, but only when the movement is directed toward a target whose location the animal must remember in order to obtain a reward. As mentioned, the nigrostriatal dopaminergic fibers appear to provide information about the relevance of a stimulus. Further, motivation (expectation of reward) influences strongly how sensory information is processed in the striatum.

Shift of Behavior

A central theory, trying to unite several lines of investigation, proposes that the basal ganglia are important for the **selection** and **shift of behavior** in particular. This appears to concern both movements and cognitive functions. In response to unexpected changes in the environment, it is often necessary to switch attention quickly from one target to another. At the same time, the behavior might need to be shifted on the basis of a decision of what is appropriate in the new situation. Thus, a new behavior is selected, and the current behavior must therefore be stopped.¹¹ The **evaluation** (giving priority to a certain behavior) probably requires cooperation between the prefrontal cortex and limbic structures, while the **execution** of the behavioral switching may depend on the basal ganglia (regardless of whether it concerns change of movements or thoughts).

11 Because the activity of most GP_i neurons *increases* in relation to a movement (thus increasing inhibition of thalamocortical neurons), one would think that the basal ganglia were concerned primarily with stopping ongoing movements rather than helping to initiate new ones. Such stopping is of course a prerequisite for starting anew.

Interval Timing

Finally, the basal ganglia seem to be involved in **interval timing**, underlying the ability to judge intervals and **duration**. This is the basis of our ability to form **temporal expectations** and predictions about ongoing and future events (the expected time of movement sequences and cognitive processes). Striatal neurons may monitor oscillatory patterns in large areas of the cortex. Thus, human functional magnetic resonance imaging (fMRI) studies of persons performing timing tasks show altered activity in the striatum and the substantia nigra, in addition to in areas such as the prefrontal cortex, the cingulate gyrus, SMA, parietal cortex, and the insula. There is some evidence that these regions function as a network in relation to interval timing. However, also other parts of the brain may be of importance in this respect, notably the **cerebellum** (see Chapter 24, under “The Timing Theory: Does the Cerebellum Perform a Basic Operation Used in All Its Functions?”). It should be emphasized, however, that it is not settled whether the “sense of time” is represented by specific modules and networks, or whether it is an ubiquitous property of neuronal activity in itself.

Symptoms after Focal Lesions of the Basal Ganglia in Humans

Studies of patients with destruction (vascular lesions in particular) confined to parts of the basal ganglia have given limited insight in their normal functions. A meta-analysis of 240 such patients found that **dystonias**—that is, abnormal positions of body parts due to persistently increased muscle tone—were most frequently encountered. Parkinsonism or choreatic movements occurred in only very few patients. Motor symptoms were most frequent, after lesions comprising the putamen and the globus pallidus, which is in agreement with the connections of these parts. Most frequent among behavioral changes was **apathy**, with loss of initiative, spontaneous thought, and emotional responses (**abulia**). Such symptoms were most frequent after lesions affecting the caudate nucleus, in agreement with its extensive connections with the prefrontal cortex. Nevertheless, on further analysis, many observations could not be fitted into the current theories of basal ganglia functions, and the symptoms varied considerably, even among patients with very similar localization of their lesions.

DISEASES OF THE BASAL GANGLIA

As the functions and internal operations of the basal ganglia are still incompletely known, it should come as no surprise that we have only partial explanations of

the relationships between certain symptoms and the elements responsible within the basal ganglia.

As mentioned, the most obvious symptoms in diseases of the basal ganglia are motor ones. As a rule, there is a mixture of symptoms due to **loss of neuronal activity** (compare with pareses after destruction of central motor pathways) and symptoms due to abnormally **increased neuronal activity** (compare with spasticity occurring after lesions of central motor pathways). The most frequent diseases of the basal ganglia impede the initiation of movement and lead to **akinesia**. When started, the movements are slower and of smaller amplitude than normal (**bradykinesia**). Because movements are difficult to initiate, there is also a conspicuous paucity of movements, usually implied in the term “akinesia” but sometimes termed **hypokinesia**. In addition, there are more or less pronounced **involuntary** movements (**dyskinesia**).

The diseases of the basal ganglia fall into two broad groups: those characterized by akinesia and rigidity (Parkinson’s disease is the most common example), and those dominated by dyskinesia (e.g., Huntington’s disease). The first group is also characterized by loss of dopamine effects in the basal ganglia, whereas the second group shows signs of dopaminergic hyperactivity. Some authors use the terms **hyperkinetic** and **hypokinetic** disorders to distinguish the two main kinds of basal ganglia diseases. The hypokinetic disorders are usually combined with rigidity and tremor (i.e., signs of neuronal hyperactivity), whereas the hyperkinetic disorders are often combined with muscular hypotonia.

The most frequent disease affecting the basal ganglia is **Parkinson’s disease**, mentioned several times already in this chapter. Typically, voluntary movements are hard to initiate (akinesia) and they are slower and smaller than normal (bradykinesia). In addition, there is an increased muscular resting tone in the form of **rigidity** (cf. Chapter 21) and **resting tremor**—that is, involuntary, rhythmic, alternating movements. Patients with Parkinson’s disease have regularly been found to have a pronounced **neuronal loss** in the **substantia nigra** and a corresponding decrease of dopamine in the striatum.

In **Huntington’s disease**, in which there is a marked cell loss in the striatum, the most pronounced symptom is involuntary, jerky, often “dance-like” movements (chorea).

Violent, large involuntary movements of one side of the body, called **hemiballismus**, occur typically after damage to the subthalamic nucleus of the opposite side. Other diseases are also dominated by sudden, involuntary muscular contractions, occurring with uneven intervals, called **tics**. One example is **Tourette’s syndrome**, believed to be caused by dysfunction of the basal ganglia. The symptoms can be partially relieved by dopamine antagonists.

Parkinson's Disease

This disease of unknown etiology usually starts during the fifth or sixth decade of life. The syndrome includes, as mentioned, **akinesia**, **bradykinesia**, **tremor**, and **rigidity**. In addition, there are disturbances of **postural reflexes**. Notably, the body balance during walking is disturbed, so that the patient appears to be “running after his center of gravity.” The steps are typically very short, which is due to the bradykinesia (the movements are not only slower than normal but also reduced in amplitude). When being pushed from behind, the patient has difficulty stopping and continues to move forward. The normal pendulum movements of the arms during walking are absent. There is also a conspicuous loss of facial expression (the face becomes like a mask). There are disturbances of the **autonomic nervous system**, such as increased salivation and secretion from sebaceous glands of the skin. We do not know how the basal ganglia would influence the autonomic nervous system, and the autonomic symptoms perhaps may be caused by a more general disturbance of monoamine metabolism (not only dopaminergic cell groups are affected in Parkinson's disease).

The **tremor** is typical, with its frequency of 3 to 6 per second, and it is most pronounced at rest. When the patient uses the hand, the tremor disappears or is reduced in amplitude. The increased muscle tone, the **rigidity**, is different from the spasticity that occurs after lesions of central motor pathways. The resistance to passive movements is equal in extensors and flexors and is independent of whether the muscle is stretched slowly or rapidly. There is no clear-cut increase in the strength of the monosynaptic stretch reflex (such as the patellar reflex), whereas the long-latency stretch reflexes are increased, and this may perhaps contribute to the difficulties with balance and locomotion. The rigidity is apparently not caused by hyperactivity of γ motoneurons but by descending influences that increase the excitability of the α motoneurons so that they fire continuously—perhaps like the situation in a nervous person who is unable to relax his muscles. The **plantar reflex** is normal in patients with Parkinson's disease, indicating that there is no damage to the central motor pathways. It is indeed an old clinical observation that the tremor and rigidity disappear if a patient with Parkinson's disease has a capsular hemiplegia, and corresponding observations are made in experimental animals. The last central link in the pathways mediating the tremor and rigidity must therefore be pathways from the cerebral cortex to the motoneurons.

The most pronounced structural change in the brain of patients with Parkinson's disease is a profound loss of pigmented (melanin-containing) neurons in the pars compacta of the **substantia nigra** (Fig. 23.8). These are

the dopamine-containing neurons. As a consequence of the loss of dopaminergic nigrostriatal fibers, the dopamine content of the striatum is reduced. Symptoms first appear, however, when the dopamine content of the striatum is reduced by 80% to 90%. Another striking change is loss of 20% to 30% of the spines on striatal projection neurons. As mentioned, this may be due to loss of a possible growth-promoting effect of dopamine. Another possibility is that spine loss is due to glutamatergic overactivity in corticostriatal synapses. The latter explanation is suggested by the observation that loss of dopamine indeed increases activity in glutamatergic synapses, and by the close proximity of dopaminergic and glutamatergic synapses on the spines (Fig. 23.9B).

Can We Explain the Symptoms in Parkinson's Disease?

Increased firing of neurons in the subthalamic nucleus is the most striking electrophysiological finding in Parkinson's disease. There is also increased activity in the GP_i , presumably due to the increased excitatory input from the subthalamic nucleus. As discussed earlier (“The “Indirect Pathway,” the Subthalamic Nucleus, and Parkinson's Disease”), it is not clear how loss of dopamine in the striatum leads to subthalamic hyperactivity. Even if we ignore that problem, the relationship between the subthalamic hyperactivity and the symptoms is not obvious. True enough, increased GP_i activity would increase thalamic inhibition and thus reduce excitation in the motor cortical areas, and this fits with positron emission tomography (PET) studies that show reduced blood flow in the SMA in Parkinsonian patients. Although reduced thalamocortical activation of motor areas thus helps to explain bradykinesia/akinesia, it is nevertheless puzzling that **thalamotomy** (a stereotaxic lesion of VL/VA) mainly improves rigidity and tremor with less effect on the akinesia and bradykinesia, and neither is voluntary movement made more difficult by the surgery. Further, thalamic infarcts affecting the VL/VA do not produce Parkinson-like symptoms, even though thalamocortical activation of motor areas must be severely reduced. Like thalamotomy, **pallidotomy** improves rigidity and tremor; in addition, it improves the akinesia. Because loss of dopamine in the striatum produces such devastating motor disturbances, it seems paradoxical that thalamotomy and pallidotomy—which eliminate the major effects of the basal ganglia on other parts of the brain—do not worsen control of voluntary movements. It may be mistaken, however, to try to explain the symptoms simply in terms of more or less excitation or inhibition at stations in the cortex–basal ganglia–cortex loop. There is evidence, for example, that thalamic neurons (both in the lateral nucleus and in the intralaminar nuclei) in Parkinsonian patients

show various forms of abnormal, partly **rhythmic** firing. Further, their activity—in contrast to that in normal persons—shows no modulation in relation to movements. In addition, the hyperactivity in the subthalamic nucleus takes the form of high-frequency oscillations (which appear to be reduced by dopaminergic medication). We furthermore know that hyperpolarization of thalamocortical neurons can induce **oscillatory** activity under certain circumstances (cf. Chapter 26, under “Thalamocortical Neurons and EEG”). Perhaps the abnormal inhibition from the GP_i in some way releases the thalamic neurons from normal control. As proposed by Marsden and Obeso (1994), the Parkinsonian symptoms may arise mainly because the basal ganglia outputs become “noisy” and thus disrupt the activity of their targets. Some neurons fire rhythmically while others become very difficult to activate. The oscillatory (20 Hz) firing of large populations of neurons may be abnormally synchronized. Presumably, such states are labile, with rapid shifts between oscillatory activity and inactivity. Neuronal networks might shift from one functional state to another in milliseconds. Conceivably, such shifts produce the rapid fluctuations in symptoms sometimes observed in Parkinson’s disease. The motor system may be better off when the disruptive basal ganglia outputs are removed (be it by thalamotomy, pallidotomy, or deep brain stimulation)—leaving it in “peace” to compensate for the loss of basal ganglia inputs.

Therapeutic Approaches to Parkinson’s Disease

The finding of reduced striatal dopamine led to attempts to alleviate the symptoms by giving the patients dopamine, to substitute for the loss of dopaminergic neurons. For various reasons, a precursor of dopamine—**levodopa** (L-dopa)—has to be used. This is converted to dopamine in the brain. This treatment has a beneficial effect on the symptoms, particularly on the very troublesome bradykinesia. Even though L-dopa has proved to be a very helpful drug, the first hopes turned out to be unrealistic. The drug does not affect the course of the disease, and the therapeutic effect decreases as the disease progresses. There probably must be a certain number of remaining dopaminergic neurons in the nigra for L-dopa to be converted to dopamine in the striatal dopaminergic boutons. More profound changes of the striatal neurons and their ability to react to dopamine may also occur. One might think that when the brain is unable to produce enough dopamine itself, it would not be able to produce dopamine from a supplied precursor, either. However, enzymes converting L-dopa to dopamine are present not only in dopaminergic neurons in the nigra but also in glial cells and in other types of neurons.

The L-dopa treatment has **side effects**, as one might expect when giving a drug that acts not only in the striatum but also in many cell groups in the brain.

For example, altered dopamine functioning in the hypothalamus might explain side effects such as nausea, loss of appetite, and reduced control of blood pressure. Other side effects are sleep disturbances and changes of mood. In some cases, the treatment can precipitate a psychosis that might be due to actions of dopamine on the prefrontal cortex limbic structures. Long-term treatment may provoke motor symptoms that are different from those caused by the disease, such as chorea-like and **athetoid dyskinesias** (athetosis is used of slow, “wormlike,” involuntary movements of the fingers and toes).

Stereotaxic surgery has now been replaced by **deep brain stimulation** (DBS) of the subthalamic nucleus or the GP_i. Such treatment gives considerable improvement to many patients (the effect is better on positive than on negative symptoms). It appears that improvement occurs regardless of whether the subthalamic nucleus is destroyed or stimulated electrically. Although this might seem to support the hypothesis that DBS removes a disturbing influence (rather than normalizing basal ganglia functioning), the mechanism whereby DBS achieves its effect is poorly understood.

Replacement of lost dopaminergic neurons by **transplantation** has now been tried for some years in patients with Parkinson’s disease, with marked improvement in at least some patients. The basis for this approach is animal experiments—pioneered by, among others, the Swedish neurobiologist Anders Björklund—showing that transplantation can eliminate virtually all symptoms produced by lesions of the substantia nigra in adult rats. The transplanted cells must be embryonic or transformed stem cells to avoid rejection, and are injected into the striatum of the recipient. The surviving cells synthesize and release dopamine. Neurons for transplantation can be developed from stem cells in culture, and this approach might be used in the future (thus avoiding ethical and practical problems with the use of human embryos).

Huntington’s Disease

This disease is dominantly inherited and usually starts in the fourth decade of life. It is caused by expansion of trinucleotide (CAG) repeats in the **huntingtin gene** (for mechanisms of neuronal death in neurodegenerative diseases, see Chapter 10, under “Common Molecular Mechanisms in Neurodegenerative Diseases”). The disease has a steadily progressing course, and is characterized by rapid, jerky, involuntary movements of the face, arms, and legs. In advanced stages, the patient is never at rest. The dance-like quality of the movements led to the terms **chorea** and **choreatic movements** (Greek: *chorea*, dance). The most prominent pathological change involves the **striatum**, with a marked loss of GABAergic projection neurons.

Observations suggest that in **early stages** of the disease, mostly neurons projecting to the GP_e degenerate (the “indirect pathway”; see earlier). This would produce reduced inhibition of GP_e neurons and, by that, increased inhibition of the subthalamic nucleus. (Animal models support that reduced activity of the subthalamic nucleus may be crucial for development of choreatic movements.) This in turn leads to reduced excitation of GP_i , and thereby to less inhibition (disinhibition) of thalamocortical neurons in VL and VA. Thus, the choreatic movements might be causally related to increased excitatory input to motor cortical areas.

In **later stages** of the disease, the striatal neurons projecting to the GP_i also die (as do many neurons in other parts of the brain). This leads to reduced inhibition of GP_i , with subsequent increased inhibition of thalamocortical neurons (as in Parkinson’s disease). This is taken to explain why bradykinesia develops in later stages of Huntington’s disease, while the choreatic movements continue. This is not obviously logical, however, if the effects of both the direct and indirect pathways are mediated by the GP_i . Another explanatory model focuses on dopaminergic hyperactivity in the striatum, caused by loss of GABAergic inhibition in the substantia nigra. Although there are problems also with this model, it fits with the observation that **L-dopa** worsens the choreatic movements of patients with Huntington’s disease. The disease also leads to mental deterioration with **dementia**, which probably may be explained by cell loss occurring also in the cerebral cortex (particularly the frontal lobes).

The genetic defect is located on the short arm of **chromosome 4**, and this makes it possible to decide whether a person is a carrier of the disease long before the symptoms occur. This is of importance regarding the choice whether or not to have children. However, to provide individuals with such knowledge as long as there is no effective treatment poses obvious ethical questions.

Tourette’s Syndrome

This is a peculiar and multifarious condition with multiple **tics** (quick involuntary movements that are repetitive at irregular intervals) associated with involuntary vocalization as central manifestations. It has a strong

heritable component but responsible genes have not been identified. The condition is probably caused by a functional disturbance of the basal ganglia and the prefrontal cortex, although the mechanisms are not known. Volumetric studies with MRI indicate that the caudate nucleus is smaller than normal, and fMRI studies show altered activity in the striatum and related cortical regions. It has been postulated that the tics are caused by abnormal activity in small groups of striatal projection neurons (cf. **striosomes** and **matrix** discussed in the preceding text). Involvement of both motor and limbic circuits has been proposed to explain both the motor and associated behavioral and emotional symptoms. Dopaminergic hyperactivity may play a role in the disease, but other transmitters, such as serotonin and GABA, have also been implicated.

The disease starts during childhood, usually between the ages of 5 and 10 years, and it affects boys more frequently than girls. Often the symptoms diminish before or during adolescence. Many patients have a repertoire of **repetitive behavior**, such as touching others, repeating their own words, or echoing the words or movements of others. Vocalizations (vocal tics) commonly include explosive cursing or compulsive utterance of obscenities, which interrupt normal speech. Many patients describe that an inner tension builds up, and this is temporarily relieved by the tic. Some people with Tourette’s syndrome appear to have unusual energy and creativity. As children, they often show hyperactive behavior. This may lead to serious social problems, especially when, as often happens, the disease is misdiagnosed as a behavioral disorder of social origin.

Treatment with **dopamine antagonists** (especially haloperidol, a D_2 -receptor antagonist) may reduce the involuntary movements and other troublesome symptoms. However, the patient’s mental energy and self-perception may be severely affected by the treatment. As vividly described by Oliver Sacks in *The Man Who Mistook His Wife for a Hat*, this may be experienced by some patients as worse than the symptoms. Since 1999, **deep-brain stimulation** has been used in some patients with severe symptoms and poor response to conventional treatments. Beneficial results have been reported with different sites of stimulation (the GP_i , the **accumbens**, and the intralaminar thalamic nuclei).

24 | The Cerebellum

OVERVIEW

The cerebellum and the basal ganglia have in common that they are involved in motor control without being responsible for the initiation of movements, and both are built into “side loops” of the motor cortical areas and the central motor pathways. Both contribute to motor learning, although in different ways. Although the major fiber connections of the cerebellum suggest a motor function, some connections are compatible with a cerebellar contribution to cognitive functions. A brief account of the cerebellum was given in Chapter 6.

Damage to the cerebellum produces characteristic symptoms primarily with respect to the execution of **voluntary movements**. Our normal movements are always **coordinated**—that is, the various muscles participating contract at the proper time and with the proper force. This is a prerequisite for our ability to hit a nail with a hammer, for writing even and rounded letters, for the words to follow each other with proper loudness and rhythm during speech, and so forth. After cerebellar damage, such coordination is lacking, and the movements become uncertain and jerky.

The cerebellum **receives information** from many sources. Sensory signals come from the skin, joints, muscles, vestibular apparatus, and eye. In most instances, this sensory information appears to be related to aspects of movement, such as signals from muscle and joint receptors about positions and ongoing movements. Information is also coming from other parts of the central nervous system (CNS), especially from the cerebral cortex—primarily from cortical areas treating information about movements or involved in planning or initiation of movements. The cerebellum **sends information** primarily to cell groups that give origin to the central motor pathways, like the motor cortical areas and the reticular formation of the brain stem.

A striking feature of cerebellar organization is that the number of fibers leading to the cerebellum is much larger than the number of fibers leading out; the relationship is about 40:1 in humans. The high degree of **convergence** shows that considerable integration and processing of information takes place in the cerebellum before an answer is issued.

The **afferent fibers** end in segregated regions of the cerebellar cortex, and the efferents from these regions are also largely segregated. As a general rule, the cerebellum sends signals to same regions from which it

receives afferents. In contrast to the cerebral cortex, the cerebellar cortex has no association and commissural connections: different parts of the cerebellum do not “talk” to each other. Consequently, the cerebellum consists of many functional units or **modules**, each carrying out a specific task. The cerebellum can be subdivided in three main parts based on connections: **vestibulocerebellum**, **spinocerebellum**, and **cerebrocerebellum**. Within each of these major divisions numerous small modules have been identified. The cerebrocerebellum—receiving large number of afferents from the cerebral cortex via the pontine nuclei—comprise about 90% of the human cerebellar volume. The spinocerebellum receives afferents from the spinal cord informing about sensory events and the state of premotor networks and act back by way of reticulospinal and vestibulospinal pathways.

Owing to the structural **homogeneity of the cerebellar cortex**, all modules have the same kind of machinery and, presumably, perform the same kind of information processing. Among several possibilities, the cerebellum may be specially designed for accurate **timing** of events, not only for movements but also for certain cognitive tasks carried out by the cerebral cortex. Further, the cerebellum is involved in aspects of **motor learning**.

SUBDIVISIONS AND AFFERENT CONNECTIONS OF THE CEREBELLUM

The macroscopic anatomy of the cerebellum is described in Chapter 6 (see Figs. 6.1, 6.13, and 6.32), but some points bear repetition here. Figure 24.1 shows how the cerebellum resides in a “side loop” of the motor cortical areas and the central motor pathways. Figure 24.2 shows how all afferent and efferent connections pass through the cerebellar peduncles. The thickness of the arrows indicates the relative number of afferent and efferent fibers. The external part of the cerebellum is a thin, highly convoluted sheet of gray matter, the **cerebellar cortex** (Fig. 24.3). The folds are arranged transversely and form numerous **folia**. If the cerebellum were unfolded completely in the anteroposterior direction, it would measure 2 m! The cerebellar cortex is thinner and of a simpler build than the cerebral cortex, consisting only of three layers (Fig. 24.2). The total number of neurons is nevertheless much higher than in the cerebral cortex due to the dense packing of tiny neurons in the granular layer. All signaling out of the cerebellum is

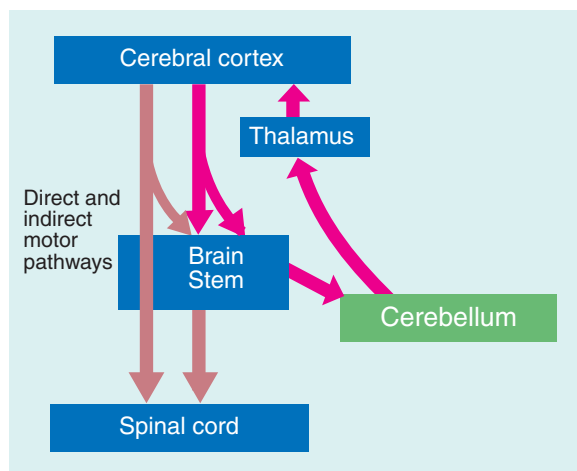


FIGURE 24.1

mediated by the axons of the **Purkinje cells** (Figs. 24.2 and 24.12). The underlying white matter contains the afferent and efferent fibers of the cerebellar cortex. In addition, masses of gray matter, the **intracerebellar nuclei**, are embedded in the white matter in the central parts of the cerebellum (Fig. 24.3). These nuclei are relay stations in the efferent connections of the cerebellar cortex—that is, the majority of the Purkinje cell axons terminate in the intracerebellar nuclei.

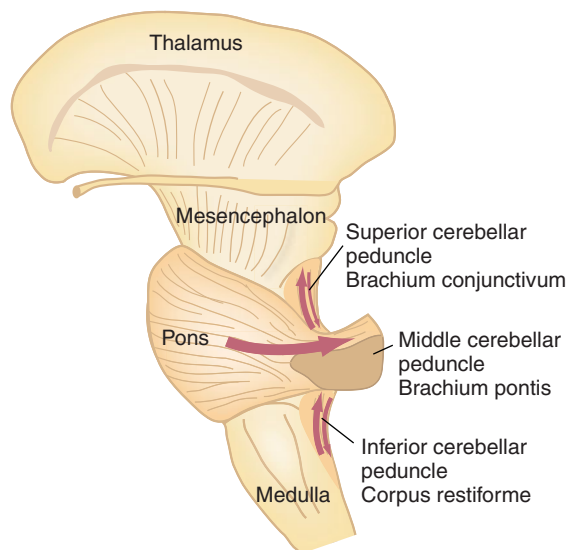


FIGURE 24.2 The cerebellar peduncles contain the afferent and efferent connections of the cerebellum. The brain stem, as viewed from the left. Arrows indicate the peduncle used by the main afferent and efferent connections on entering or leaving the cerebellum. The thickness of the arrows indicates the magnitude of the various connections. Note the massive afferent pathway from the pontine nuclei that enter through the middle cerebellar peduncle (the brachium pontis). The largest number of efferent fibers leaves the cerebellum through the superior cerebellar peduncle (the brachium conjunctivum).

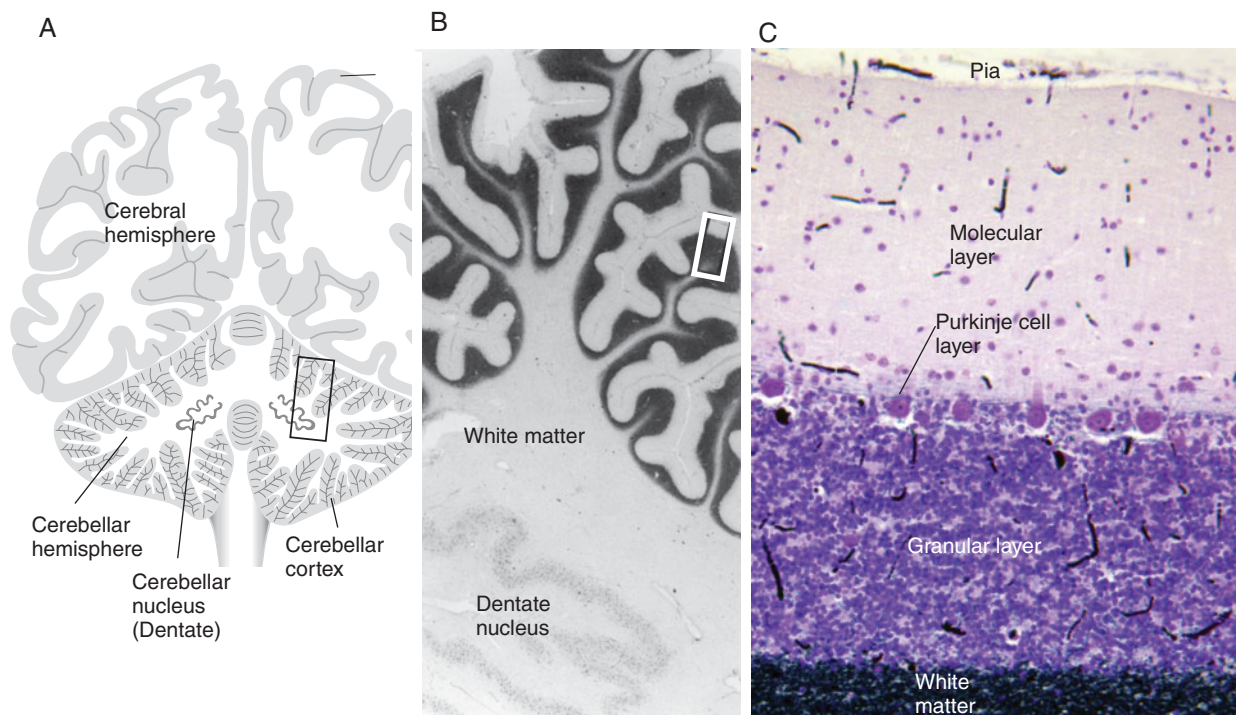


FIGURE 24.3 The cerebellum, the cerebellar cortex, and an intracerebellar nucleus. **A:** Frontal section through the cerebrum and the posterior fossa with the cerebellum. Note the folding of the cerebellar surface to form thin folia. Cf. Fig. 6.32. **B:** Photomicrograph from the

area corresponding to the frame in **A**. Thionine staining. **C:** Higher magnification of framed area in **B**, showing the three layers of the cerebellar cortex. Cf. Fig. 24.12 showing the structural elements of the cerebellar cortex in detail.

The Cerebellum Consists of Three Functionally Different Parts

The structure of the cortex is the same all over the cerebellum. Therefore, various parts of the cerebellum differ in function because of differences in fiber connections: the subdivisions of the cerebellum receive afferents from and act on different parts of the nervous system. Indeed, a subdivision of the cerebellum based on **functional differences** corresponds closely with a subdivision based on differences in **afferent connections** (Fig. 24.4). Further, such a subdivision corresponds also with one based on the cerebellar phylogenetic development from lower to higher animals.

The most primitive part of the cerebellum—the part occurring first during phylogeny—is the small **flocculonodular lobe** (Fig. 24.4; see Figs. 6.13 and 6.32). It consists of the **nodulus** in the midline (a part of the vermis) connected laterally with a thin stalk to the **flocculus**. The size of this part of the cerebellum varies little among mammals. The flocculonodular lobe receives afferents primarily from the vestibular apparatus and the vestibular nuclei and is therefore called the **vestibulocerebellum**. The rest of the cerebellum is called the **corpus cerebelli**. This comprises all of the **vermis** (except the nodulus), which forms a narrow zone on both sides of the midline, and the large lateral parts called the **cerebellar hemispheres**. The medialmost part of the

hemispheres, bordering the vermis medially, is called the **intermediate zone** (Fig. 24.4). This zone cannot be distinguished from the rest of the cerebellum on a macroscopic basis but only on the basis of fiber connections. A deep, transversely oriented cleft, the **primary fissure**, divides the corpus cerebelli. The parts of the corpus cerebelli in front of and behind the primary fissure are called the **anterior lobe** and the **posterior lobe**, respectively. The anterior and posterior parts of the vermis constitute the phylogenetically oldest parts of the corpus cerebelli. The midportion of the vermis and the hemispheres are younger and are therefore called the **neocerebellum**. The anterior and posterior portions of the vermis and the adjoining parts of the intermediate zone of the corpus cerebelli receive afferents primarily from the spinal cord and are therefore also termed the **spinocerebellum**. The hemispheres receive their main input from the cerebral cortex (synaptically interrupted in the pontine nuclei) and are therefore termed the **cerebrocerebellum** or the **pontocerebellum**.

With regard to the **efferent connections**, the three main subdivisions of the cerebellum act on the parts of the CNS from which they receive their afferents; that is, the vestibulocerebellum sends fibers mainly to the vestibular nuclei, the spinocerebellum influences the spinal cord, and the cerebrocerebellum acts on the cerebral cortex.

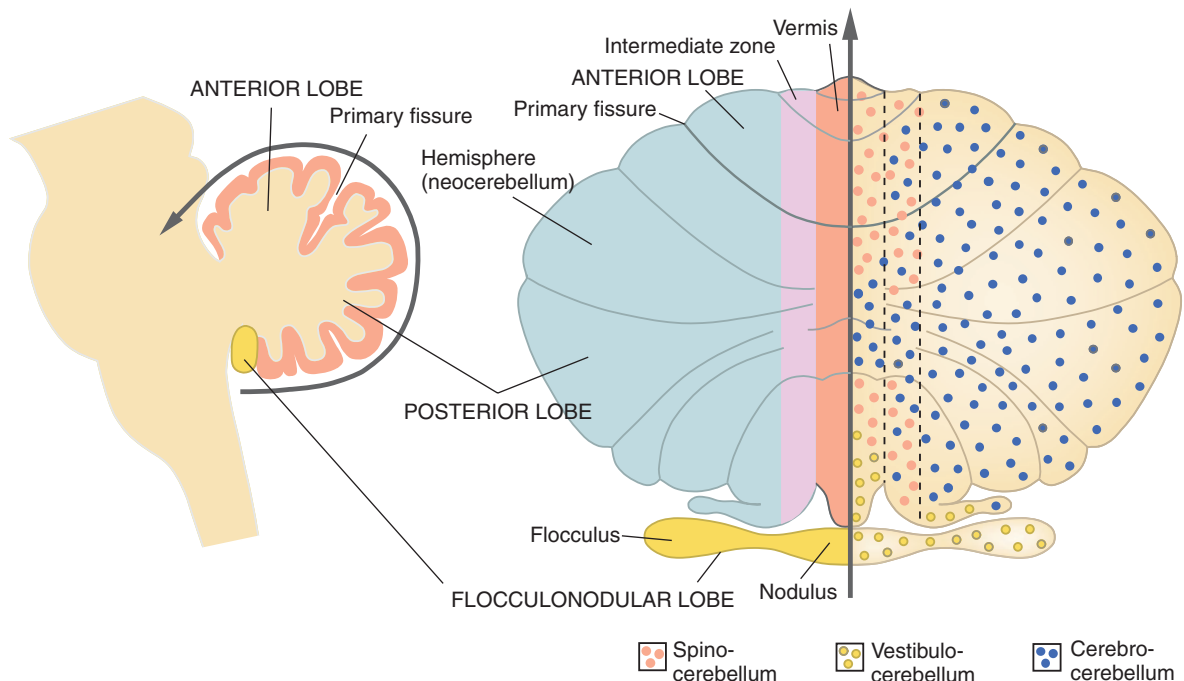


FIGURE 24.4 Main cerebellar subdivisions. **Left:** A sagittal section through the cerebellum and the brain stem. **Right:** The human cerebellum unfolded and seen from the dorsal aspect. The left half shows the macroscopic subdivisions of the cerebellum into lobes, as well as the border between the vermis and the hemisphere. Compare with

Fig. 6.32. The right half shows the terminal regions of the three major afferent contingents to the cerebellum (vestibular, spinal, and cerebrocortical). Because the intermediate zone receives information from both the spinal cord and the cerebral cortex, there is no sharp border between the spinocerebellum and the cerebrocerebellum.

Afferents from the Labyrinth and the Vestibular Nuclei

Primary vestibular afferents bring sensory signals from the vestibular apparatus in the inner ear. They enter the cerebellum through the inferior cerebellar peduncle (Fig. 24.1) and end in the flocculonodular lobe and adjoining parts of the vermis (Fig. 24.5). Although most of the primary vestibular afferents end in the **vestibular nuclei**, many neurons in the latter send axons to the cerebellum. In this manner, the cerebellum receives vestibular information also via **secondary vestibular afferents**. The vestibular input provides the cerebellum with information about the position and movements of the head. **Efferents** from the flocculonodular lobe end in the vestibular nuclei and can thereby influence the body equilibrium (via the vestibulospinal tracts) and eye movements via the medial longitudinal fascicle (Fig. 24.5 and see Fig. 18.9).

Afferents from the Spinal Cord

Several pathways bring signals from the spinal cord to the cerebellum (Fig. 24.6). Some of these pathways go uninterrupted from the cord to the cerebellum and are called **direct spinocerebellar tracts**, whereas others are synaptically interrupted in brain stem nuclei and are

therefore termed **indirect spinocerebellar tracts** (not shown in Fig. 24.6). The spinocerebellar tracts originate from neurons with their cell bodies located in different lamina of the spinal cord (Fig. 24.7). They are therefore influenced by different kinds of **sensory receptors** and spinal **interneurons** and bring different kinds of information to the cerebellum. The spinocerebellar axons end mostly in the spinocerebellum of the same side (as their cell bodies). Some pass uncrossed; other fibers cross twice (first in the spinal cord and then back again in the brain stem). The fibers are located in the **lateral funicle** as they ascend (Fig. 24.7). The tracts are **somatotopically** organized, so that signals from different body parts are kept segregated. The somatotopic pattern is maintained within the cerebellum (Fig. 24.8) so that the leg is represented anteriorly within the anterior lobe, with the arm and the face represented successively more posteriorly. In the posterior lobe, the arrangement is the reverse, with the face represented anteriorly.

Direct Spinocerebellar Tracts

Functionally, the direct spinocerebellar tracts consist of two main groups. One group of tracts conveys information from **muscle spindles**, **tendon organs**, and **cutaneous low-threshold mechanoreceptors**. Physiological studies show that many of the neurons of the direct spinocerebellar tracts are activated monosynaptically by primary

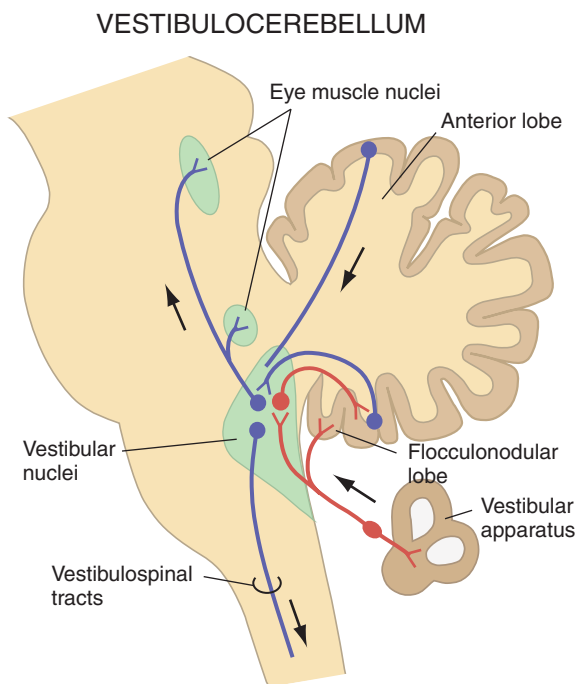


FIGURE 24.5 Main connections of the vestibulocerebellum. Afferents are shown in black and efferents in blue in a schematic of a sagittal section through the brain stem. Also shown are primary and secondary vestibulocerebellar fibers and the projection back to the vestibular nuclei. In addition to afferents from the vestibulocerebellum, the vestibular nuclei receive cerebellar afferents from the anterior lobe vermis and from the fastigial nucleus (not shown).

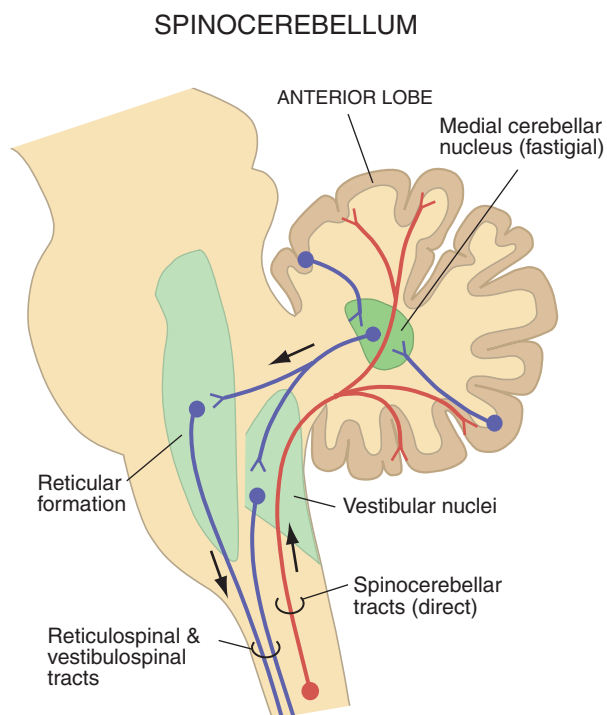


FIGURE 24.6 Main connections of the spinocerebellum. The spinocerebellum can influence spinal motoneurons by both the reticulospinal and vestibulospinal pathways.

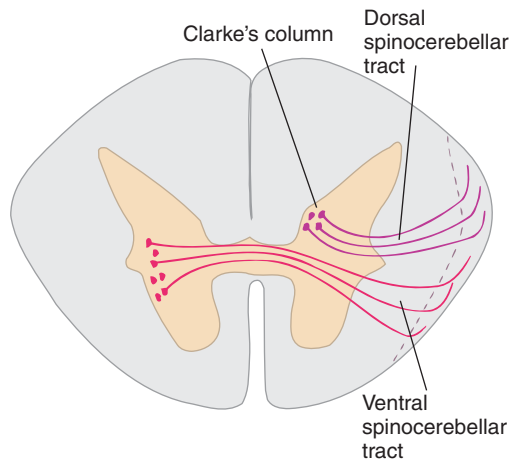


FIGURE 24.7 *The dorsal and ventral spinocerebellar tracts.* Transverse section through the thoracic cord, showing the position of the cell bodies in the gray spinal matter and the tracts in the white matter. Most of the fibers of the ventral spinocerebellar tract cross twice (once in the cord and once in the brain stem), so that most spinocerebellar signals reach the cerebellar half on the same side as the cell bodies are located.

afferent (sensory) fibers. The tracts conduct very rapidly and appear to give precisely timed information about movements.

Another group of direct spinocerebellar tracts does not convey signals from receptors but provides information about the level of activity among specific groups of **spinal interneurons**. As a rule, such interneurons are intercalated in spinal reflex arcs and between descending motor pathways and motoneurons forming **premotor networks**. Information about their activity is therefore presumably highly relevant for cerebellar operations.

Together, the direct spinocerebellar tracts provide the cerebellum with information about the activity both *before* and *after* the motoneurons—that is, about the

commands issued to the motoneurons and the movements they produce. The cerebellum can probably judge whether the command led to the desired result. When, for example, an unexpected increase of external resistance to a movement reduces the velocity compared with what was intended, the cerebellum would be informed immediately. By means of its connections to the spinal cord, the cerebellum can then help to adjust the firing of the motoneurons to the new situation, so that the correct velocity is regained.

Indirect Spinocerebellar Tracts

There are several indirect spinocerebellar tracts, but here we will only mention the largest one, which is synaptically interrupted in the **inferior olive** in the medulla (see Fig. 6.17). Neurons at all levels of the cord send fibers to the inferior olive of the opposite side. The **spino-olivary** fibers end in parts of the inferior olive that project to the spinocerebellum of the opposite side; thus, like the direct spinocerebellar tracts, the spino-olivocerebellar tract conveys information mainly from one side of the cord to the cerebellar half of the same side. This pathway, too, is precisely **somatotopically** organized. The kind of information conveyed by the spino-olivocerebellar tract appears to be different from that conveyed by the other spinocerebellar tracts. We return to the inferior olive later in this chapter, because it has a unique role among the cell groups that send fibers to the cerebellum.

The Dorsal, Ventral, and Some Other Spinocerebellar Tracts

Originally, two direct spinocerebellar tracts were described, differing with regard to the position of their fibers in the lateral funicle of the cord. The fibers of one of the tracts are located dorsally, and the tract was called the dorsal spinocerebellar tract; the fibers of the ventral spinocerebellar tract are located more ventrally (Fig. 24.7).

The **dorsal spinocerebellar tract** comes from a distinct cell group with fairly large cells at the base of the dorsal horn, forming the **column of Clarke** located in the spinal segments from T_1 to L_2 only. The axons of the cells in the column of Clarke ascend on the same side and enter the cerebellum through the inferior cerebellar peduncle (Fig. 24.1). Primary afferent fibers entering the cord through spinal nerves below the L_2 ascend in the dorsal columns before entering the lowermost part of the column of Clarke. The dorsal spinocerebellar tract conveys information only from the trunk and the lower extremities and ends in the corresponding parts of the spinocerebellum. Another cell group, the **external cuneate nucleus**, located laterally in the medulla oblongata, mediates the same kind of information from

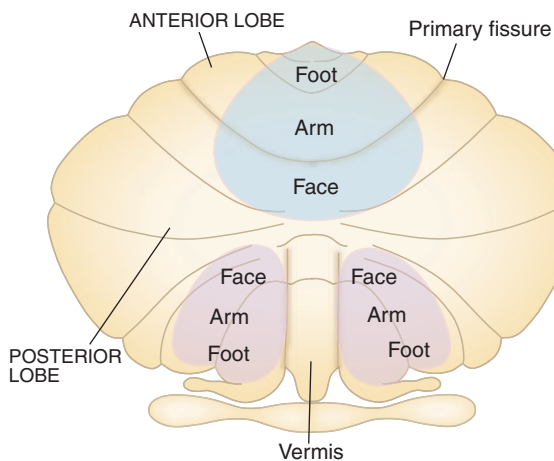


FIGURE 24.8 *Somatotopic localization of the cerebellar cortex.* Based on experiments with electrical stimulation of peripheral nerves and of somatotopic subdivisions of the SI and the MI.

the upper extremities. Functionally, this nucleus corresponds to the column of Clarke, and primary afferents from the arms ascend in the dorsal columns to end in the nucleus. The cells of the external cuneate nucleus send their axons to the arm regions of the spinocerebellum of the same side (Fig. 24.8), forming the **cuneocerebellar tract**. As mentioned, primary afferent fibers from muscle spindles and tendon organs end **monosynaptically** on the neurons of the column of Clarke (and the external cuneate nucleus). In addition, the column of Clarke receives **polysynaptic** information from the same kind of receptors, as well as from joint and cutaneous receptors. Indeed, recent physiological studies show that most of the column-of-Clarke neurons **integrate** information from various receptor types activated by a specific movement. Thus, the dorsal spinocerebellar tract appears to inform primarily about **complex joint movements**, for example, those that involve the whole extremity. This is another example that sensory relay nuclei do not merely transmit signals unchanged from receptors; they always process information in some way.

The **ventral spinocerebellar tract** originates from cells located mainly laterally in lamina VII (Fig. 24.7; see Fig. 6.14)—that is, in the lamina containing the largest number of interneurons. Most of the axons cross at the segmental level in the cord to the lateral funicle of the opposite side. However, after having reached the cerebellum through the superior cerebellar peduncle (Fig. 24.1), many of the fibers cross once more. The cell bodies of the neurons giving origin to the ventral spinocerebellar neurons are found only below the midthoracic level of the cord and have a rostral counterpart conveying information from higher levels of the cord, the **rostral spinocerebellar tract**. These tracts convey information about the activity of the spinal interneurons, as mentioned above. Physiological experiments indicate that they inform primarily about the activity of **inhibitory interneurons**.

In addition to the spinocerebellar tracts mentioned so far, other cell groups in the spinal cord also send fibers to the cerebellum. Among such cell groups is the **central cervical nucleus**, located in the upper cervical segments. The fibers end in the anterior folia of the anterior lobe and transmit signals primarily from receptors around the **cervical joints**.

An additional indirect spinocerebellar pathway is relayed in the **lateral reticular nucleus**, located in the medulla just lateral to the inferior olive. This **spinoreticulocerebellar** pathway, too, is mainly uncrossed and appears to convey information about the activity of certain groups of spinal **interneurons**. Because most of the spinal interneurons are strongly influenced by descending **motor pathways** (e.g., the pyramidal tract), the ascending tract to the lateral reticular nucleus probably informs about their activity as well. The lateral reticular

nucleus also receives afferents from sources other than the spinal cord, notably the red nucleus, the vestibular nuclei, and the motor cortex. Many cells, for example, are strongly influenced by tilting of the head, which stimulates **vestibular receptors**.

Finally, information from sensory receptors can reach the cerebellum not only through some of the spinocerebellar tracts but also through fibers sent to the spinocerebellum from the **dorsal column nuclei** and the **trigeminal nuclei**.

Afferent Connections from the Cerebral Cortex

In humans, by far the largest number of cerebellar afferent fibers arises in the **pontine nuclei** (Fig. 24.9; see Fig. 6.18). The **pontocerebellar tract** ends primarily in the cerebellar hemispheres, which constitute the major part (90%) of the cerebellum in humans. The vast majority of afferents to the pontine nuclei arises in the cerebral cortex and forms the **corticopontine tract** (Fig. 24.8).¹ The main task of the pontine nuclei therefore is to process information from the cerebral cortex and forward it to the cerebellar cortex. The corticopontine tract is uncrossed, whereas most of the pontocerebellar fibers cross; thus, the cerebral cortex of one side acts mainly on the cerebellar hemisphere of the opposite side.

The **corticopontine tract** runs in the internal capsule and then in the crus cerebri (see Fig. 6.20), where it occupies a large part; of the approximately 19 million fibers in the crus of the human brain, the corticopontine fibers constitute the majority (the pyramidal tract, in comparison, contains only about 1 million fibers).

A large fraction of the corticopontine fibers arise in the **MI** and **SI**. There are also substantial contributions from the supplementary motor cortex (**SMA**) and premotor cortex (**PMA**) and from areas 5 and 7 of the **posterior parietal cortex** (Fig. 24.9). Contributions that are more modest come from parts of the **prefrontal cortex**. In various ways, all of these areas are active during or before movements. Presumably, the cerebellum thus receives information about movement **planning** and about the **motor commands** that are sent out from the cortex; in response, it can modulate the activity of the motor cortex so that the movements are performed smoothly and accurately. The pontine nuclei also receive afferents from the **visual cortex** (mainly extrastriate areas), and physiological experiments indicate that these fibers inform primarily about moving objects in the visual field. Such connections may be important for

¹ Other **precerebellar nuclei**—that is, brain stem nuclei that send their efferents to the cerebellum—also receive afferents from the cerebral cortex. This concerns mainly the **reticular tegmental nucleus**, located just dorsal to the pontine nuclei (participating in the control of eye movements, see Chapter 25), and the **lateral reticular nucleus**. In quantitative terms, these pathways play minor roles compared with the corticopontocerebellar pathway.

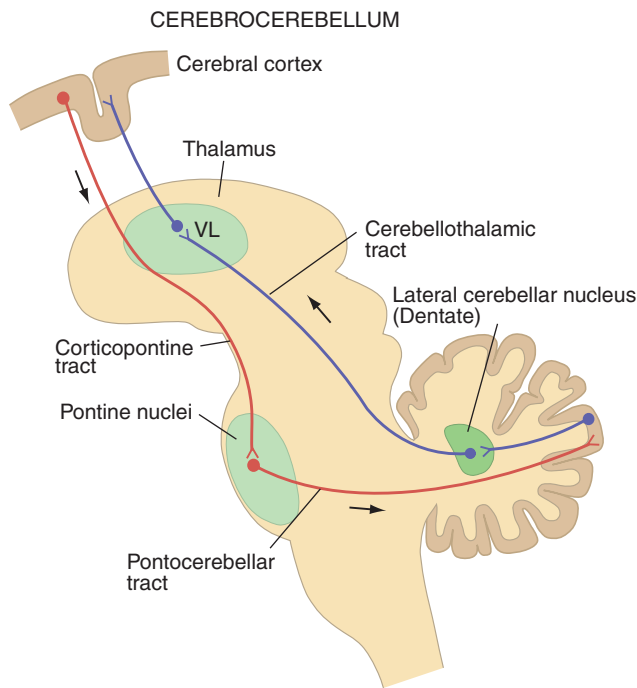


FIGURE 24.9 *Main connections of the cerebrocerebellum.* The ascending connections to the cerebral cortex are synaptically interrupted in the dentate nucleus and in the thalamus. The projection from the cerebellar cortex to the cerebellar nuclei is GABAergic and inhibitory, whereas the other links in the cerebrocerebellar pathway are excitatory. Compare Fig. 24.17 showing the crossing of the cerebrocerebellar connections.

the execution of **visually guided movements**.² The pontine nuclei also receive connections from parts of the **hypothalamus** and **limbic structures**, notably the **mammillary bodies** and the **cingulate gyrus**. The information conveyed from these regions is probably not directly related to movement. The cingulate gyrus is concerned, for example, with attention and with error-detection during motor, sensory, and cognitive processes. These connections and those from the prefrontal cortex may form the basis of cerebellar contributions to certain **cognitive tasks** (see later in this chapter, “The Cerebellum and Cognitive Functions”). In addition, “limbic” corticopontine connections might contribute to the well-known influence of motivation and emotions on movements.

Topographic Organization of the Cerebrocerebellar Pathway

The corticopontine projection is **topographically organized**, so that different cortical regions project to largely different parts of the pontine nuclei (Fig. 24.10). The fibers coming from the MI and SI are **somatotopically organized**. Further, the pontocerebellar projection shows

a topographic organization—for example, upholding the somatotopic pattern in the connections from MI and SI (Fig. 24.7). The connections from the cerebral cortex to the pontine nuclei exhibit an extreme **divergence**: that is, a small part of, let us say MI, influences small clusters of neurons in widespread parts of the pontine nuclei. Presumably, each pontine neuron receives converging inputs from specific combinations of cortical cell groups, thus integrating various kinds of information before forwarding messages to the cerebellum. The next link—the pontocerebellar tract—shows a marked **convergence** (as well as divergence). In this way, the corticopontocerebellar pathway seems to produce numerous specific combinations of cerebrocortical inputs in the cerebellar cortex. These and other data indicate that there is a functional localization within the cerebellar hemispheres, so that smaller parts take care of specific tasks.

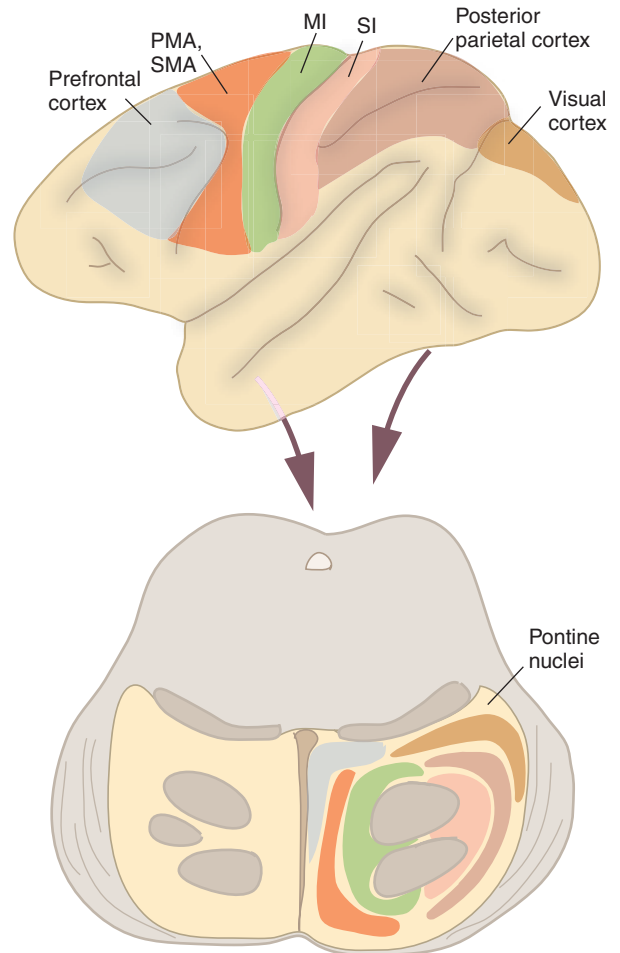


FIGURE 24.10 *The corticopontine pathway.* Areas of the cerebral cortex that are connected with the cerebellar hemispheres by means of the pontine nuclei are shown in different colors. In the pontine nuclei, fibers from each cortical region terminate in its own territory. Note the lamellar arrangement, resembling the skins of an onion.

² There are modest connections to the pontine nuclei from the **frontal eye field**, involved in control of rapid eye movements, and from area 9 in the dorsolateral part of the prefrontal cortex. Some fibers arise in parts of the **auditory cortex**, whereas few if any pontine fibers come from the rest of the temporal lobe.

The Intermediate Zone Is a Meeting Place for Signals from the Cord and the Cerebral Cortex

The intermediate zone (Fig. 24.4) is mainly defined on the basis of its efferent connections (projecting to the interposed intracerebellar nuclei; Fig. 24.11). It also has special features with regard to afferents, however (Fig. 24.11). Whereas the lateral parts of the hemispheres are strongly dominated by inputs from the cerebral cortex, and the vermis is dominated by spinal inputs, the intermediate zone receives connections from both the cerebral cortex and the spinal cord (Fig. 24.4). Animal experiments indicate that the cortical input to the intermediate zone comes primarily from the MI and SI. Single neurons in the intermediate zone can be activated from both the cerebral cortex and the spinal cord. For example, in the “arm region” of the anterior lobe, intermediate-zone neurons receive converging input from the arm region of the MI and SI and from sensory receptors in the arm. Perhaps the cerebellum in this case **compares** copies of the motor commands sent from the cerebral cortex with the signals from the periphery providing information about the actual movement that was produced by the command (signaled by the spinocerebellar tracts).

THE CEREBELLAR CORTEX AND THE MOSSY AND CLIMBING FIBERS

Before discussing the efferent connections of the cerebellum, we need to know something about the cerebellar

cortex. Here, the vast amount of information provided by all of the afferents is processed. To some extent, different kinds of information are integrated, and then “answers” are issued to various motor centers of the brain and spinal cord. As mentioned, the cerebellar cortex has the same structure all over (it cannot be subdivided into cytoarchitectonic areas, differing also in this respect from the cerebral cortex), and it lacks association fibers that interconnect different regions. The structural arrangement of the neuronal elements is strictly **geometric**, so the individual elements can be distinguished fairly easily. This helps explain why the structure and internal connections of the cerebellar cortex are much better known than that of the cerebral cortex.

The Cerebellar Cortex Consists of Three Layers

The superficial, outermost layer is the **molecular layer** (Figs. 24.2 and 24.12). It contains mainly dendrites and axons from cells in the deeper layers and only a few cell bodies. The middle layer is dominated by the large Purkinje cells, arranged in a monolayer, and is called the **Purkinje cell layer**. The deepest, lowermost layer is the **granular layer**, named so because it is packed with tiny **granule cells**. The axons of the granule cells ascend through the Purkinje cell layer into the molecular layer, where they divide at a right angle into two branches running parallel with the surface of the cortex (Figs. 24.12 and 14.13). These branches are called **parallel fibers** and run in the direction of the long axis of the folia. The parallel fibers form numerous synapses with the Purkinje cell dendrites. The **Purkinje cell dendritic tree** is unusual: first, it has an enormously rich branching pattern; second, the dendritic tree is compressed into one plane, forming an espalier oriented perpendicular to the long axis of the folia and the parallel fibers. This arrangement ensures that each parallel fiber forms synapses with many Purkinje cells (the parallel fibers can be several millimeters long). At the same time, an enormous number of parallel fibers contact each Purkinje cell: it has been estimated that each Purkinje cell receives about 200,000 synapses. Considering that there are approximately 100 billion granule cells but only 30 million Purkinje cells, each Purkinje cell would integrate signals from about 3000 granule cells.³

In addition to granule cells and Purkinje cells, the cerebellar cortex contains **inhibitory interneurons** (Fig. 24.13) that serve to limit the activity of the Purkinje cells and probably increase the spatial precision of the

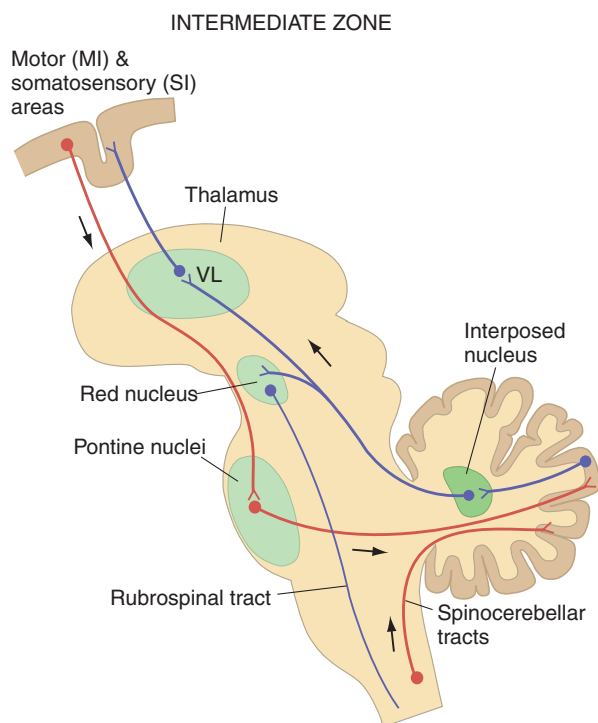


FIGURE 24.11 Main connections of the intermediate zone. Both the spinal cord (via the red nucleus) and the cerebral cortex (via the thalamus) can be influenced by the intermediate zone.

³ Although older quantitative studies agreed that the number of Purkinje cells in the human cerebellum is about 15 million, a study with an improved stereological method estimated the number to about 30×10^6 million (Andersen et al. 1992). The same study estimated the number of granule cells to about 100×10^9 , while the number of neurons in the dentate nucleus was 5×10^6 .

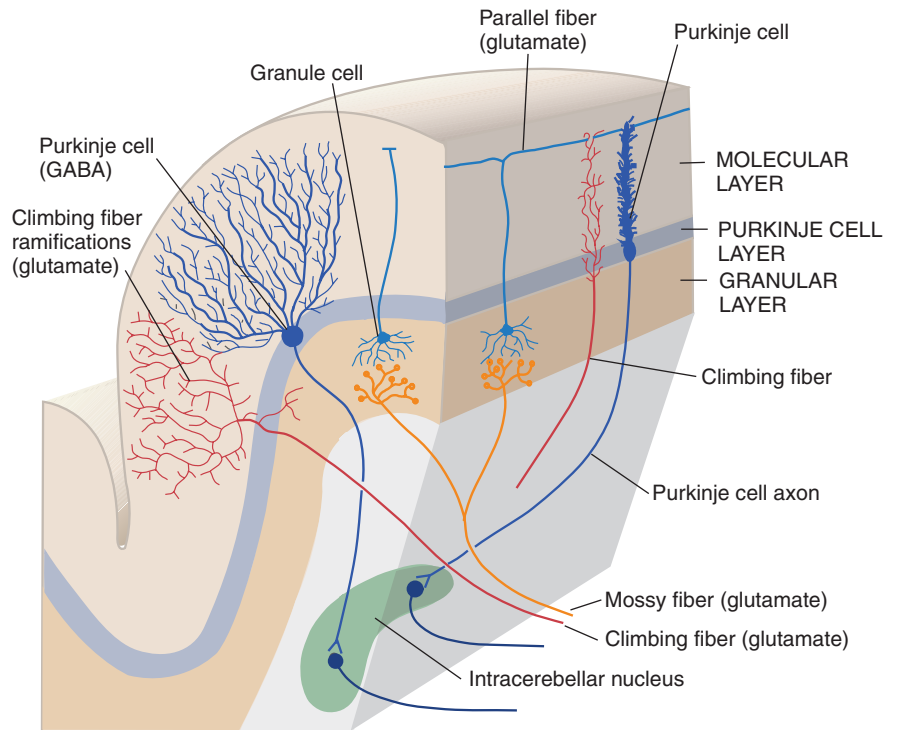


FIGURE 24.12 *Structure of the cerebellar cortex.* The three layers and the main cell types are shown schematically in a piece of cerebellar folium. The Purkinje cell dendrites are arranged perpendicular to the long axis of the folium and the parallel fibers. The two main kinds of afferent fibers (mossy and climbing fibers) are also shown. The mossy fibers end on the granule cells, whereas the climbing fibers enter the molecular layer to end on the Purkinje cell dendrites.

incoming signals (cf. inhibitory interneurons in sensory systems; see Fig. 13.4).

As mentioned, the **Purkinje cells** are the only ones that send their axons out of the cerebellar cortex and thus constitute the efferent channel. The Purkinje cells

contain γ -aminobutyric acid (**GABA**), and they inhibit their target cells, as shown physiologically. The **granule cells** have an excitatory action on the Purkinje cells, releasing **glutamate**.

The Cerebellar Cortex Contains Three Kinds of Inhibitory Interneurons

All cerebellar interneurons contain GABA (some may also contain glycine, another inhibitory neurotransmitter). One main type of interneuron is the **stellate cell**, located in the molecular layer (Fig. 24.13). It receives afferent excitatory input from the granule cells (parallel fibers), and its axons form synapses with the Purkinje cell dendrites. Another kind of interneuron, the **basket cell**, is located close to the Purkinje cell layer. Basket cells are also contacted by parallel fibers, whereas their axons end with synapses around the initial segment of the Purkinje cell axons—a location that enables the basket cells to inhibit the Purkinje cells very efficiently. The axonal branches of the basket cells are arranged perpendicular to the long axis of the folia, so that they inhibit Purkinje cells lateral to those that are being activated by parallel fiber excitation. Activation of a group of granule cells would lead to a narrow band of excitation of the Purkinje cells along the folium, flanked by a zone of basket-cell mediated inhibition on each side. Thus, it appears to be a kind of lateral inhibition, which is common in sensory systems to increase the spatial precision. Correspondingly, the extent of the cerebellar cortical region activated by each mossy fiber is reduced. The third kind of inhibitory interneuron, the **Golgi cell**,

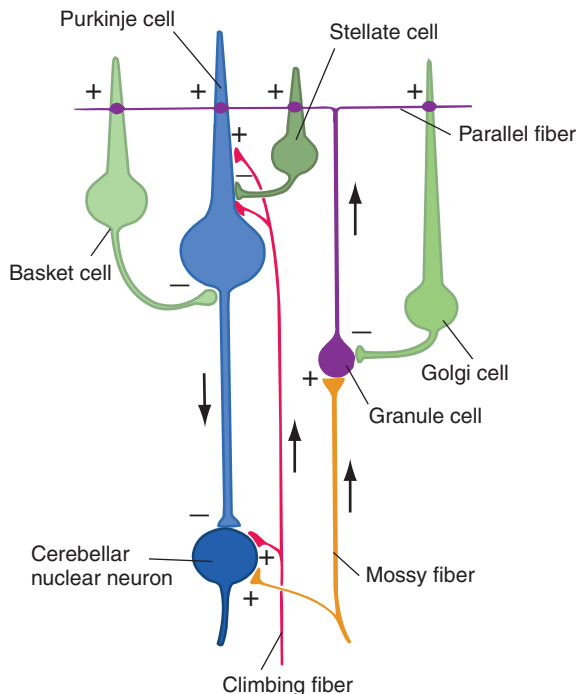


FIGURE 24.13 *The cerebellar cortex.* Schematic of the main cell types and their synaptic arrangements. The three types of GABAergic interneurons are colored green. (Redrawn from Eccles et al. 1967.)

is located in the granular layer. The dendrites of the Golgi cells extend upward into the molecular layer and are therefore contacted by parallel fibers (like the stellate cells and the basket cells). The axonal branches form synapses with the dendrites of the granule cells and thus reduce the excitation received by the Purkinje cells from the granule cells.

Afferents to the Cerebellar Cortex Are of Two Main Kinds: Mossy and Climbing Fibers

The afferent fibers to the cerebellar cortex fall into two categories, which differ in how the fibers end in the cerebellar cortex. Both kinds have an **excitatory** synaptic action, most likely mediated by glutamate. The **climbing fibers** all come from the **inferior olive**, whereas afferents from nearly all other nuclei end as **mossy fibers** (such as the vestibulocerebellar, the spinocerebellar, and the pontocerebellar fibers).⁴

The **mossy fibers** conduct signals relatively rapidly and end in the granular layer, establishing synapses with the granule cell dendrites (Figs. 24.12 and 24.13). One mossy fiber branches extensively and contacts a large number of granule cells, each of which, in turn, contacts many Purkinje cells. Thus, each mossy fiber influences many Purkinje cells, but the excitatory effect on each is weak, so that many mossy fibers must be active together to provide sufficient excitation (via the parallel fibers) to fire a Purkinje cell (as mentioned, the parallel fibers excite the Purkinje cells). A typical feature of the mossy fibers is that they transmit action potentials with a high frequency and make the Purkinje cells fire so-called **simple spikes** with a frequency of 50 to 100 per second.

All the **climbing fibers** cross to the opposite side and end very differently from the mossy fibers: the fibers ascend directly to the molecular layer and divide into several branches, each “climbing” along a Purkinje cell dendrite (Figs. 24.12 and 24.13). As they climb, they form numerous synapses with the dendrites. Each Purkinje cell receives branches from only one climbing fiber (i.e., from only one cell in the inferior olive). Each olivary cell, however, innervates more than one Purkinje cell, as the number of Purkinje cells is higher than the number of olivary cells. Because each climbing fiber forms so many synapses with a Purkinje cell, the total excitatory action is strong. Thus, even a single action potential in a climbing fiber elicits a burst of action potentials in the Purkinje cells—called **complex spikes**.

In contrast to mossy fibers, the firing frequency of the climbing fibers is very low under natural conditions (often less than one signal per second). Even maximal stimulation does not bring the firing frequency above 10/sec. Accordingly, recordings of Purkinje cell activity in an animal at rest shows continuous firing of simple spikes with high frequency, interrupted now and then by complex spikes.

Acute **destruction** or inactivation of the inferior olive demonstrates its importance for motor control. Movements become uncoordinated, similar to the effect of removing the whole cerebellum. Accordingly, physiologic studies show that inactivation of the olive (by injection of a local anesthetic) produces a marked disinhibition of the Purkinje cells. This increases the inhibition of the intracerebellar nuclear cells so that the cerebellar output to other parts of the brain is virtually eliminated.

In conclusion, the Purkinje cells receive excitation from both mossy and climbing fiber afferent inputs but with very different spatial and temporal characteristics of their actions.

Connections of the Inferior Olive

The climbing fibers cross and end in the cerebellar nuclei and the cerebellar cortex with an extremely precise topographic order. One subdivision of the olive receives **afferents** primarily from the **cord** via several precisely organized spino-olivary tracts, and project to the spinocerebellum. Another part receives afferents from the **superior colliculus** and project to the midportion of vermis (the “oculomotor vermis”). The superior colliculus receives signals from the retina, from the visual cortex, and the SI. These parts of the olive contribute to the control of eye and head movements. A small olivary subdivision receives afferents from the **pretectal nuclei** (see Fig. 27.19) and project to the flocculonodular lobe. The pretectal nuclei receive signals from the retina, and its projection to the olive is of importance for adaptation of the **vestibuloocular reflex** (see Fig. 25.4; see also later, “Examples of the Cerebellar Role in Motor Learning”).

The main part of the human inferior olive (the principal nucleus) projects to the cerebellar hemispheres and receives afferents from various **mesencephalic nuclei**, notably the parvocellular **red nucleus**. These mesencephalic nuclei receive afferents from the cerebral cortex (especially M1, SMA, and PMA) and can therefore mediate cortical information to the cerebellar hemispheres. In addition, the mesencephalic nuclei receive strong connections from the cerebellar nuclei, thereby establishing a loop: cerebellum–red nucleus–inferior olive–cerebellum. The functional role of this rather massive pathway is unknown. Finally, the inferior olive receives GABAergic fibers from the cerebellar nuclei (of the opposite side).

4 In addition to the climbing fibers and mossy fibers, demonstrated with the Golgi method a long time ago, a third type of cerebellar afferent has been demonstrated by using the histofluorescence method (visualizing fibers containing catecholamines). Such fibers come from the **raphe nuclei** and from the **locus coeruleus** and contain serotonin and norepinephrine, respectively. They appear to end rather diffusely in both the granular and molecular layers. Direct fibers from the **hypothalamus** also end in this manner.

Mossy and Climbing Fibers Mediate Different Kinds of Information

The great differences between the mossy and climbing fibers with regard to both structural and physiological properties strongly suggest that they convey different kinds of information and thus play different parts in cerebellar functions. Because of their ability to vary their signal frequency over a wide range, the **mossy fibers** are presumably well suited for providing precisely graded information about movements (the muscles involved, as well as the direction, speed, and force of movements), localization and characteristics of skin stimuli, details concerning motor commands issued from the cerebral cortex, and so forth. Such assumptions also fit with the physiological properties of spinocerebellar fibers, known to end as mossy fibers (less is known about the corticopontocerebellar pathway in this respect).

The **climbing fibers**, because of their low range of firing frequency, are less likely to provide precisely graded information. Recordings of the firing of the Purkinje cells in response to climbing fiber activity also suggest that the climbing fibers have a unique functional role. Thus, as mentioned, a single action potential in a climbing fiber is sufficient to trigger a burst of Purkinje cell action potentials (complex spikes). This would suggest an all-or-none action rather than a graded one. Several theories of cerebellar functions postulate that the climbing fibers inform about **errors** in the execution of a movement (giving an “error signal”) when the movement does not correspond to what was intended, and there is some experimental support for this hypothesis. Some studies show that the firing frequency of climbing fibers increases in relation to a perturbation of an ongoing movement, whereas the firing frequency is unrelated to, for example, the direction and velocity of the movement. In experiments with walking cats, the firing frequency of climbing fibers leading from the forelimb increases when the foot meets an obstacle, so that the walking pattern has to be changed. In monkeys learning a new movement, there is increased climbing-fiber activity from the relevant body parts. When the movement is well rehearsed—that is, the learning phase is over—the climbing fiber firing frequency does not increase during execution of the movement (no more error signals?).

A specific role of the climbing fibers during **motor learning** has been postulated based on this and other kinds of experiments. The climbing fiber input is thought to alter for a long time (days, perhaps years) the responsiveness of the Purkinje cells to mossy fiber inputs. This appears to happen only on simultaneous activation of a Purkinje cell by specific sets of climbing and mossy fibers, leading to a change of Purkinje cell excitability, so that the following mossy fiber signals have less effect than previously. This phenomenon,

long-term depression (LTD, see Chapter 4, under “Different Kinds of Synaptic Plasticity”) depends on a rise in Ca^{2+} concentration in the Purkinje cell dendrites, which is produced by the binding of glutamate to amino-methylisoxazole propionic acid (AMPA) and metabotropic glutamate receptors. Most likely, however, several cellular mechanisms underlie cerebellar plasticity.⁵

The Inferior Olive and Rhythmic Movements

As mentioned, the climbing fibers arise in the inferior olive—a large, folded nucleus in the ventral medulla (see Fig. 6.17). A characteristic property of olivary neurons is that their membrane potential shows spontaneous fluctuations that facilitate **rhythmic firing** with a frequency of 5 to 10 Hz. Llinás and Sugimori (1992) propose that the inferior olive functions as a **pace-maker** for movements, by alerting specific combinations of premotor neurons in the cord. Indeed, making the olivary neurons to fire rhythmically by systemic administration of the alkaloid **harmaline** produces rhythmic muscular contractions—tremor—in large parts of the body with a frequency of 10 Hz. Further, olivary neurons fire rhythmically in pace with licking movements in the rat, which occur with a frequency of 6 to 8 Hz. Another peculiarity of olivary neurons is that they are **electrically coupled** (nexus). This enables large assemblies of neurons to fire **synchronously**. The GABAergic fibers from the cerebellar nuclei, mentioned earlier, can switch off the electric coupling so that the synchronously firing neuronal assemblies become much smaller. Conceivably, this relates to how the cerebellum organizes the activation of muscle groups at specific moments during a movement.

The Mossy Fibers and Fractured Somatotopy in the Cerebellar Cortex

The American neurophysiologist Welker (1987) using very precise, micromapping methods, described a novel aspect of the somatotopic localization within the cerebellum. The mossy fibers, formerly thought to end with a fairly diffuse somatotopic pattern, were shown to end in numerous, discrete **patches** in the cortex, each patch being defined by its sensory input from a specific minor part of the body. Each patch is usually less than 1 mm in diameter (in the rat and cat). Thus, the leg region within the posterior lobe consists in reality of a **mosaic** of patches. A salient feature is that adjacent patches can receive inputs from body parts that are widely separated.

⁵ The hypothesis that LTD is the cellular basis of cerebellar plasticity (and thus learning) is not supported by all available data, however. For example, knockout mice lacking certain glutamate receptors exhibited reduced motor learning in spite of retained ability to produce LTD in the cerebellar cortex.

Further, the same body part is usually represented in several widely separated patches. This arrangement of the mossy fibers was termed **fractured somatotopy** by Welker. It presumably is a means to integrate various inputs sharing relevance for a certain movement. How this pattern of mossy fiber inputs is coordinated with the climbing fiber inputs is not clear. As discussed later (Fig. 24.18), the climbing fibers terminate in narrow sagittal strips or zones, and one such strip is often related to one body part only.

EFFERENT CONNECTIONS OF THE CEREBELLUM

As previously mentioned, the three main subdivisions of the cerebellum act largely on the parts of the nervous system from which they receive afferent inputs. The vast majority of the Purkinje cell axons end in the cerebellar nuclei (corticonuclear fibers). The neurons of these nuclei forward the information to the various targets of the cerebellum.

The Cerebellar Nuclei and the Corticonuclear Connections

The cerebellar nuclei are located in the deep white matter of the cerebellum, just above the roof of the fourth ventricle (Figs. 24.2 and 24.14). In humans, there are four nuclei on each side. Close to the midline, under the vermis, lies the **fastigial nucleus** (medial cerebellar nucleus); then follow two small nuclei; and most laterally lies the large, folded **dentate nucleus** (lateral cerebellar nucleus). The two small nuclei have specific names in humans (the **globose** and the **emboliform** nuclei) and correspond to the anterior and posterior **interposed nuclei** in animals.

The **corticonuclear connections** are precisely, topographically organized, so that, as a rule, fibers from the anterior parts of the cerebellar cortex end in anterior parts of the nuclei, fibers from medial parts of the cortex end medially, and so forth. There is in addition a marked **longitudinal localization**, with the vermis sending fibers to the fastigial nucleus, the intermediate zone to the interposed nuclei, and the hemispheres to the dentate nucleus (Fig. 24.15). There is also somatotopic localization within each of the nuclei so that different parts influence movements in different parts of the body. Overall, signals from different parts of the cerebellum are kept segregated through the nuclei and further on to other parts of the brain. Thus, each of the nuclei sends efferent fibers to a separate target region, as shown in a very simplified manner in Figs. 24.6, 24.9, and 24.11.

Direct Projections from the Cerebellum to the Vestibular Nuclei

Parts of the vestibular nuclei correspond in certain respects to the cerebellar nuclei. Thus, the Purkinje cells of the **vestibulocerebellum** send their axons directly to the vestibular nuclei as **corticovestibular** fibers (Fig. 24.5). These fibers end primarily in vestibular nuclei that send ascending connections to the nuclei of the external ocular muscles (the medial longitudinal fasciculus; see Fig. 18.8) and, to a lesser extent, in parts of the nuclei sending fibers to the spinal cord. The vestibular nuclei also receive direct projections from Purkinje cells of the vermis of the **anterior** and the **posterior lobes**—that is, outside the vestibulocerebellum as defined here. These fibers end primarily in the lateral vestibular nucleus (nucleus of Deiters; see Figs. 18.7 and 18.8), which projects to the spinal cord. Thus, the cerebellar vermis

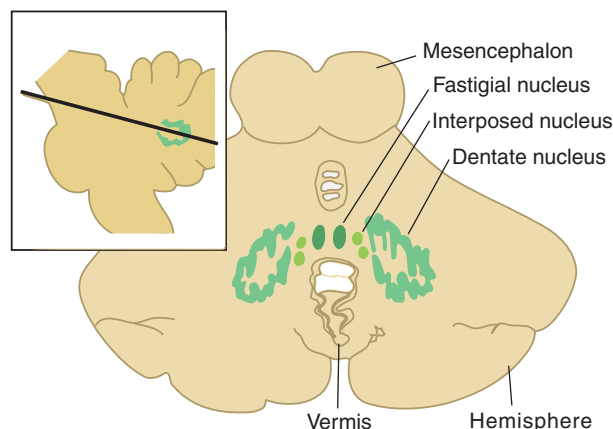
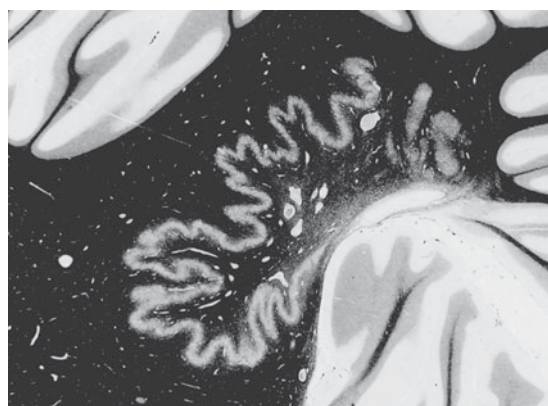


FIGURE 24.14 *The cerebellar nuclei.* **Left:** Drawing of an oblique section through the cerebellum and the brain stem. **Right:** Photomicrograph of a myelin-stained section placed slightly more



dorsally than the drawing. Therefore, only the dentate and the interposed nuclei are seen in the photomicrograph.

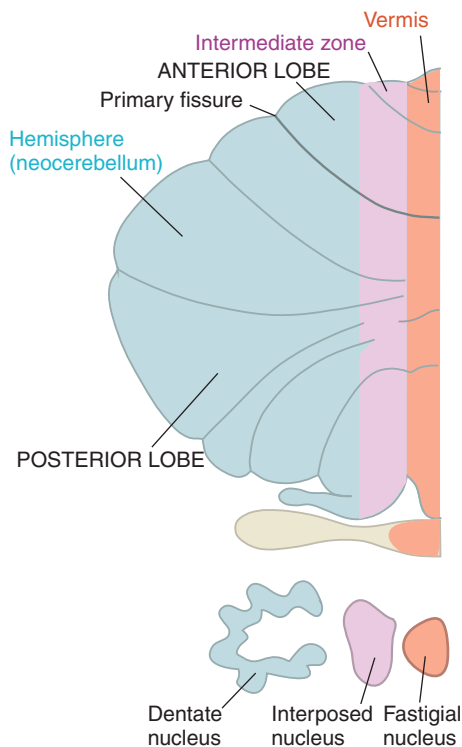


FIGURE 24.15 *Sagittal arrangement of corticonuclear connections.* Purkinje cells in the hemispheres project to the dentate nucleus, whereas Purkinje cells in the intermediate zone and the vermis project to the interposed and fastigial nucleus, respectively. In addition, Purkinje cells in the anterior and posterior vermis project to the vestibular nuclei.

can influence spinal motoneurons via the lateral vestibulospinal tract, besides its effects mediated by means of the fastigial nucleus to the reticular formation and the lateral vestibular nucleus. The vestibulocerebellum thus contributes to the control of eye movements, whereas the vermis via the vestibular nuclei primarily controls posture and equilibrium.

Cerebellar Nuclear Neurons Are Spontaneously Active

The nuclear cells fire with a high frequency even in an animal sitting quietly. When neurons fire without any obvious excitatory input, they are said to be **spontaneously active**. Indeed, in vitro studies of cerebellar slices show that the nuclear neurons have intrinsic properties that depolarize the membrane even in the absence of an excitatory input (**pacemaker** properties). Because all Purkinje cells are inhibitory (GABA), a continuous firing of the nuclear cells is a prerequisite for the information from the cerebellar cortex to be passed on (increased Purkinje cell activity leads to reduced nuclear cell firing). In addition to their tendency for spontaneous depolarization, the nuclear cells receive some excitatory inputs, namely the spinocerebellar and olivocerebellar projections that give off collaterals to the cerebellar nuclei.

Increase or decrease in the firing frequency of the Purkinje cells immediately causes change in the activity of the nuclear cells. A prerequisite for this is synchronous firing of many Purkinje cells with axons converging on a few nuclear cells. Then even minute changes of the activity of each Purkinje cell changes the signals issued from the cerebellum to its target nuclei. Indeed, there is evidence that **synchronous firing** of assemblies of functionally related Purkinje cells is a fundamental feature in cerebellar functioning.

In conclusion, the outputs of the cerebellar nuclei reflect with high temporal precision even very weak inputs to the cerebellar cortex. This is presumably important for, among other tasks, the cerebellar role in control of rhythm, as we discuss later in this chapter.

Organization of Efferent Connections from the Cerebellar Nuclei

The fibers from the **dentate nucleus** leave the cerebellum through the superior cerebellar peduncle (Fig. 24.2). They cross the midline in the mesencephalon, and some fibers end in the red nucleus of the opposite side. Most fibers continue rostrally, however, to end in the thalamus. Here, the dentate fibers end primarily in the **ventrolateral nucleus, VL** (some also reach the VA) (Fig. 24.16). These nuclei also receive fibers from the basal ganglia, but they end in different parts than the cerebellar fibers (as shown schematically in Fig. 24.16), in agreement with physiological studies showing that the basal ganglia and the cerebellum do not influence identical parts of the cortex. Signals from the cerebellar hemispheres pass primarily to the **MI**, whereas the basal ganglia via the thalamus act mainly on premotor areas and the prefrontal cortex. In addition, the cerebellum influences parts of the **SMA** and **PMA** that apparently differ from the parts influenced from the basal ganglia. There is also anatomic and physiological evidence of connections from the dentate nucleus (via the thalamus) to area 9 in the dorsolateral **prefrontal cortex**. Even though such connections most likely are modest compared to those reaching the motor areas, they are suggested to be of decisive importance for the cerebellar influence on cognitive tasks.

As mentioned, the efferents from the intermediate zone reach the **interposed nuclei**. These send their efferents both to the contralateral **thalamus** (to the VL mainly, like the dentate) and the **red nucleus** (Fig. 24.11). This enables the interposed nuclei to influence motoneurons via both the rubrospinal tract and the pyramidal tract. Because the rubrospinal tract is crossed, the interposed nucleus (and the intermediate zone) acts on the body half of the same side. The human rubrospinal tract is, however, most likely too small to be of much functional significance (cf. Chapter 22, under “The Red Nucleus and the Rubrospinal Tract”).

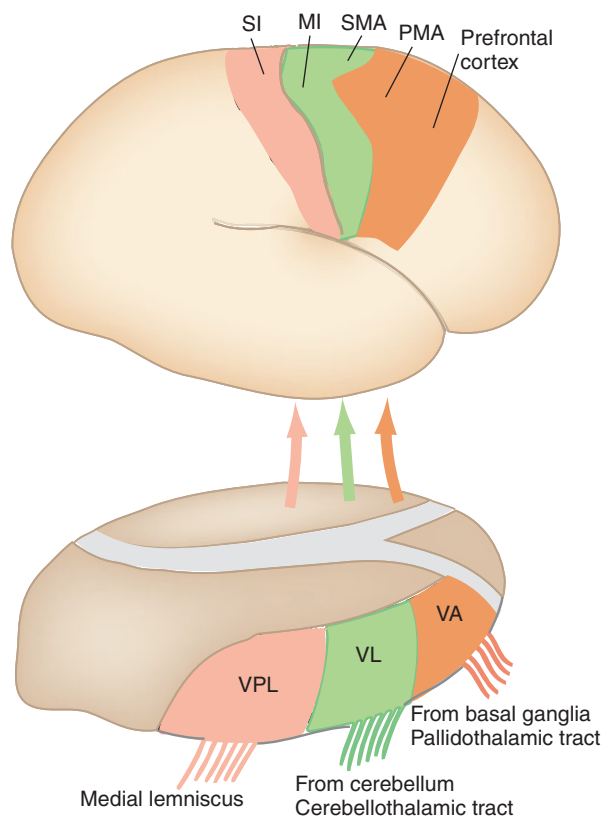


FIGURE 24.16 *Thalamocortical connections.* Schematic of the arrangement within the ventral thalamic nucleus of afferents from the somatosensory pathways, the cerebellum, and the basal ganglia and their further projections to the SI, MI, and the premotor cortex.

The **fastigial nucleus** (receiving Purkinje cell axons from the vermis) sends its efferents to both the **vestibular nuclei** and the **reticular formation** (Fig. 24.6). Thus, motoneurons can be influenced via the vestibulospinal and the reticulospinal tracts. On the basis of what we discussed in Chapter 22, the cerebellum via the fastigial nucleus can influence posture and relatively automatic movements like locomotion. This is supported by the results of animal experiments and by clinical observations. In addition, connections from the fastigial nucleus to the reticular formation mediate cerebellar influences on autonomic functions, as shown in animal experiments.

Sagittal Zones and Modules in the Cerebellum

We have so far described that the main subdivisions of the cerebellum—such as the flocculonodular lobe, the vermis, and the hemispheres—differ with regard to connectivity. Further, we have seen that the lateral parts of the hemispheres, the intermediate zone, and the vermis differ with regard to their efferent connections (Fig. 24.15). This organization of the cerebellum into three longitudinal zones, each projecting to separate parts of the cerebellar nuclei, was first described by the Norwegian neuroanatomists Jan Jansen and Alf Brodal

in the 1940s. But the localization within the cerebellum is far more sophisticated than what could be revealed by the fairly primitive methods 70 years ago. Each of the three **longitudinal zones** of Jansen and Brodal can thus be further subdivided, as shown by the Dutch neuroanatomist J. Voogd and coworkers. Figure 24.17 shows the zones of the intermediate zone as an example.

The zonal pattern is especially sharp within the **olivo-cerebellar** and **corticonuclear** projections. The neurons located within a particular small part of the olive send their fibers to a narrow longitudinal zone, whereas the neighboring zones receive climbing fibers from other parts of the olive (Fig. 24.17). The Purkinje cells of the zones also send their axons to different parts of the cerebellar nuclei. The parcellation of the cortex into sagittal zones in fact goes further than shown in Fig. 24.17. Physiological studies show that within the anterior lobe each zone consists of several **microzones**, which differ in the information they receive. Apart from the parallel

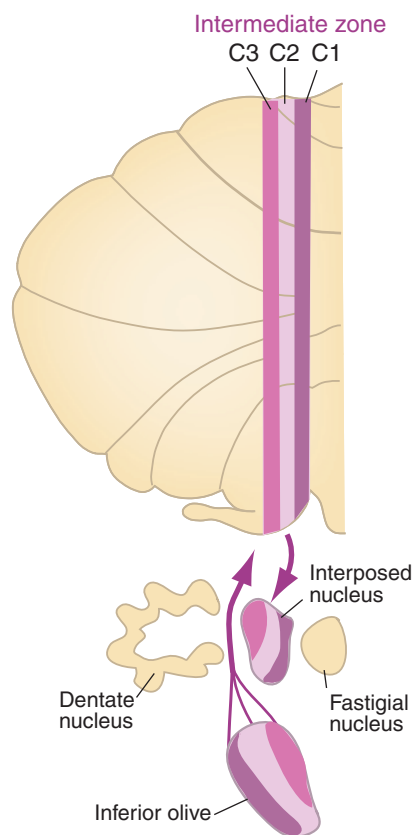


FIGURE 24.17 *Zonal organization of the cerebellar cortex.* Very simplified illustration of the precise topographic pattern of cerebellar efferent and afferent connections. The cortex can be divided into longitudinal zones projecting to different parts of the cerebellar nuclei and receiving afferents from different parts of the inferior olive. Here, the intermediate zone is shown as an example. The intermediate zone is divided into three zones termed C1–C3 differing with regard to olivary afferents and nuclear target region. Each zone, or rather parts of a zone, control certain muscular groups and their coordination in goal-directed movements.

fibers, extending for a few millimeters perpendicular to the longitudinal zones, there are no association fibers interconnecting different parts of the cerebellar cortex. Thus, the cerebellar cortex and the cerebellar nuclei consist of numerous compartments, or **modules**, that function (at least largely) independently of each other. Some data suggest that each module takes care of a specific motor task. As mentioned, in this respect the cerebellar cortex differs from the cerebral cortex, which is characterized by extensive association connections and cooperation among different regions.

CEREBELLAR FUNCTIONS AND SYMPTOMS IN DISEASE

We finally discuss the function of the cerebellum, with special reference to the motor disturbances that occur in humans and experimental animals after damage to the cerebellum. (To some extent, the function of an organ is easier to discover when it does not work properly.) Because both the ascending fibers from the cerebellum to the cerebral cortex and the descending fibers from the cerebral cortex to the spinal cord are crossed, the cerebellar hemisphere exerts its influence on the body half of the same side (Fig. 24.18). Consequently, with diseases of the cerebellum, the **symptoms** occur on the **same side as the lesion**.

Clinically, only certain motor symptoms can with certainty be referred to cerebellar lesions, even though a multitude of symptoms have been produced in experimental animals after lesions of the cerebellum.⁶ Modern neuroimaging techniques, however, provide the opportunity to localize accurately even fairly small lesions in living subjects and, furthermore, to study cerebellar neuronal activity (indirectly) in healthy and diseased subjects performing specific tasks. Such studies have demonstrated change of cerebellar activity in relation to performance of tasks that were unexpected on the basis of clinical observations. We return to this below when considering the cerebellar role in cognitive functions.

To judge from the cerebellar connections as we have discussed them here, it is clear that the cerebellum **influences movements** by acting on the neuronal groups that give origin to the central motor pathways (the pyramidal tract, the reticulospinal tracts, and the vestibulospinal tracts). These pathways, of course, control the activity of spinal and cranial nerve motoneurons. The three main subdivisions of the cerebellum (Fig. 24.4) act on different neuronal groups, and lesions restricted to each of

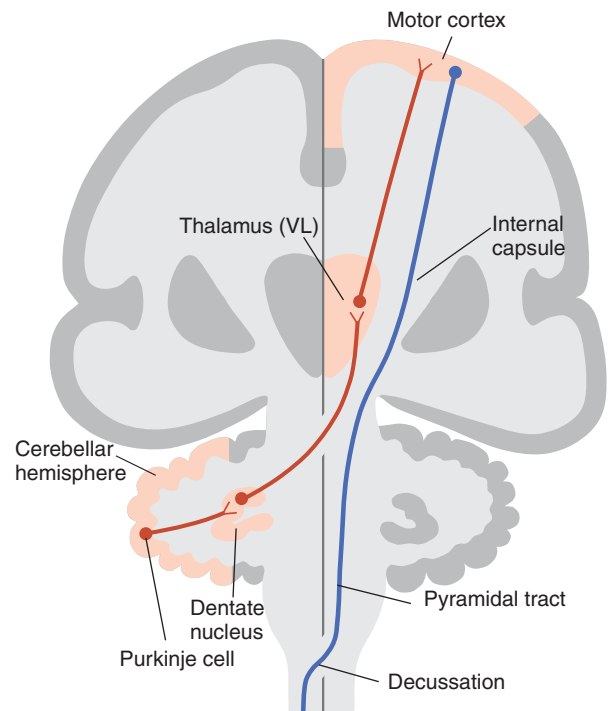


FIGURE 24.18 Unilateral lesions of the cerebellum produce symptoms on the same side. Ascending connections from the dentate nucleus cross in the mesencephalon on their way to the thalamus. The next link passes to the motor cortex, where pyramidal tract neurons are influenced. Since the pyramidal tract crosses (in the lower medulla), the cerebellar hemispheres control muscles in the ipsilateral body half.

the subdivisions give different symptoms. On this basis, one usually distinguishes three cerebellar syndromes:⁷ the **flocculonodular syndrome**, the **anterior lobe syndrome**, and the **neocerebellar syndrome**. The existence of three distinct syndromes is most clear-cut in experimental animals. In humans, the neocerebellar syndrome is most often seen, which is not surprising because the hemispheres (the neocerebellum) make up the major part of the human cerebellum.

The Flocculonodular and Anterior Lobe Syndromes

Isolated damage to the flocculonodular lobe in monkeys produces disturbances of the equilibrium—that is, unsteadiness in standing and walking. When the body is supported, movements of the extremities can be performed normally, however. Sometimes, similar symptoms occur in humans with a special kind of tumor in the posterior cranial fossa, most often a so-called medulloblastoma arising from the nodulus. The animals also exhibit **nystagmus** with the quick phase to the side of the lesion. (Nystagmus is movements in which the eyes move slowly in one direction—as when tracking a moving object with the gaze—and rapidly back.)

⁶ This is probably explained by the fact that in humans the lesions that give clear-cut symptoms most often are large and not confined to particular functional or anatomic cerebellar subdivisions. Further, they are usually combined with lesions in other parts of the CNS (e.g., in parts of the brain stem). It is therefore often difficult to decide whether a particular symptom is caused by the cerebellar lesion.

⁷ A syndrome is a constellation of symptoms that occur together.

Eye movements may also be disturbed in humans with cerebellar lesions that affect the vestibulocerebellum.

Damage to the **anterior lobe** in experimental animals primarily produces a change of **muscle tone**. In decerebrate animals, the decerebrate rigidity increases, as do the postural reflexes. This fits with the observation that electrical stimulation of the anterior lobe reduces the decerebrate rigidity (as mentioned, the Purkinje cells inhibit the cells of the cerebellar nuclei, whereas the nuclear cells have excitatory actions on reticulospinal and vestibulospinal neurons).⁸ In addition, some Purkinje cells in the anterior lobe vermis send axons directly to the vestibular nuclei, and removal of this inhibitory action would also tend to increase the activity of the vestibulospinal neurons and thus the decerebrate rigidity. In **humans**, it is doubtful whether lesions of the anterior lobe produce increased muscle tone. More marked is **gait ataxia** (unsteadiness of walking) in patients with damage that mainly affects the anterior lobe vermis and the intermediate zone (this occurs in cerebellar degeneration caused by alcohol abuse). The anterior lobe vermis, by means of its efferent connections to the fastigial nucleus and from there to the reticular formation, must therefore be assumed to have a role in coordination of the half-automatic movements of walking and postural adjustments. Similarly, selective lesions of the fastigial nucleus in monkeys cause difficulties with walking, sitting, and maintaining the upright position.

Cerebellar Lesions and Eye Movements

As mentioned, lesions of the flocculonodular lobe can cause nystagmus. This may manifest itself as **spontaneous nystagmus** (i.e., nystagmus occurring in a person at rest with no kind of stimulation) or only when the patient tries to keep the gaze in an eccentric position (paralysis of gaze nystagmus). Conceivably, these symptoms are due to the loss of Purkinje cells that normally inhibit the vestibular nuclei (especially the medial nucleus) sending fibers to the nuclei of the extrinsic eye muscles (see Fig. 18.8). In addition, the patients with lesions of the flocculonodular lobe may have difficulties with slow **pursuit movements** (tracking a moving object with the gaze). Pursuit movements may also be impaired after lesions restricted to lateral parts of the **pontine nuclei** or the **cerebellar hemispheres** (impaired to the side of the lesion). Finally, lesions of the cerebellar hemisphere may cause so-called **saccadic dysmetria**—that is,

the rapid eye movements overshoot the target and are followed by several correcting movements before the gaze is finally fixed. The role of the cerebellum in the control of eye movements is further discussed in Chapter 25, under “The Cerebellum Controls Both Saccades and Pursuit Movements.”

The Neocerebellar Syndrome

The neocerebellum plays a different functional role than the phylogenetically older parts of the cerebellum: it is primarily concerned with the coordination of the (least automatic) voluntary movements. This stands to reason, since the cerebellar hemispheres send their main output to the MI (via the dentate nucleus and the thalamus) and thus influence the neurons of the pyramidal tract (Figs. 24.9 and 24.18). After removal of one cerebellar hemisphere in a monkey, the voluntary movements become uncertain on the same side of the body: they become **uncoordinated** or **ataxic**. The same effect can be produced by cooling the dentate nucleus (by the use of a cooling electrode) in a monkey that is performing a well-rehearsed movement (as soon as the cooling is reversed, the movements again become normal). Movements that were performed quickly and smoothly become unsteady and jerky by the cooling. The monkey misses repeatedly when trying to grasp an object, even though it knows perfectly well where it is and what is demanded. Sometimes the hand is moved too far in relation to the object, sometimes too short. The movements tend to be **decomposed**; that is, instead of occurring simultaneously in several joints, they take place in one joint at a time, and the **velocity** is uneven—sometimes too high and sometimes too low. Selective damage or transient “uncoupling” of the dentate nucleus in monkeys indicates that the cerebellar hemispheres are particularly important when movements must take place in **several joints** at the same time. Picking up a raisin, for example, became impossible because the monkey could no longer coordinate the movements of the joints of the wrist, thumb, and index finger. Precise movements of one finger at the time could be done normally, however. Difficulties with hitting an object when trying to grasp it with the hand can probably be attributed to the same basic defect; that is, difficulties with coordinating the movements of the wrist, shoulder, and elbow joints.

Ataxia of this kind is also the most prominent symptom in **humans** with damage to the cerebellar hemispheres. For example, difficulty with the **precision grip** similar to that described in monkeys has recently been observed in patients with unilateral infarcts of the cerebellar hemispheres. The increase of force when grasping an object is slower than normal and the adjustment of the grip force is deficient when grasping and lifting at the same time. In clinical neurology, the various elements

8 Electrical stimulation with electrodes surgically implanted at the cerebellar surface has been used in patients with neurological disorders such as **epilepsy** and **cerebral palsy**. The theoretical basis is the inhibitory action of the Purkinje cells with subsequent reduction of abnormally increased neuronal excitability and muscle tone. Even though some report favorable results with such stimulation, there is no agreement as to whether the effect is due to the cerebellar stimulation or to some other factor.

of ataxia have particular names, such as **dysmetria** (movement is not stopped in time), **asynergia** (decomposition of complex movements), **dysdiadochokinesia** (reduced ability to perform rapidly alternating movements of, for example, the hand), and **intentional tremor** (tremor arising when trying to perform a movement, such as grasping an object). Speech is also often disturbed in cerebellar diseases. It has been called **speech ataxia**, to emphasize that it also appears to be caused by incoordination (in the respiratory muscles, the muscles of the larynx, and others), making the strength and velocity of the speech uneven.

All the elements of ataxia have been attributed to a fundamental defect in control of the force and of the exact timing of the **starting** and **stopping** of movements.⁹ As mentioned, the **temporal** aspect appears to be central to the cerebellar contribution to motor control. This is evidenced by patients with cerebellar damage who are unable to perform sequences of finger movements in a particular **rhythm**. The movements are done by each finger without a fixed temporal relation to movements of the other fingers.

In acute damage to the cerebellar hemispheres in humans, the muscle tone often appears to be reduced when tested by passive stretch (the symptom is transient). This is called **cerebellar hypotonia**. The underlying mechanism is not clear, although experiments in anesthetized animals suggested that it is caused by reduced γ motoneuron activity. Recent experiments in awake animals and in humans, however, show that the muscle-spindle sensitivity to stretch is not significantly altered by cerebellar lesions (even when they include the dentate and interposed nuclei).

The Timing Theory: Does the Cerebellum Perform a Basic Operation Used in All Its Functions?

Even though the study of cerebellar symptoms provides reasonable insight into the functions of the cerebellum, we are far from understanding how the cerebellum performs its tasks. The striking uniformity and strictly geometric structure of the cerebellar cortex has led to comparison with a **computer**, which can perform the same kinds of computations on various kinds of information. The subdivision of the cerebellum into numerous, apparently independent units or modules may fit such a concept. Several theories have been put forward to explain how the cerebellum operates. None of them has so far been universally accepted, however, and there is disagreement with regard to interpretation of some experiments said to support one theory or the other.

⁹ It is not certain, however, that this suffices to explain why the symptoms are most marked when movements in two or more joints must be coordinated (such as moving the index finger quickly to the tip of the nose from a position with the arm stretched out).

The German neuroscientist Valentin Braitenberg proposed more than 50 years ago that the cerebellum functions as a kind of **clock**, measuring temporal intervals with great accuracy. The theory was subsequently modified to emphasize the cerebellar role in the control of **movement sequences** and perhaps other sequential behaviors. The theory is based on, among other things, the regular arrangement of the parallel fibers (Figs. 24.12 and 24.13). Action potentials conducted along the parallel fibers will excite the Purkinje cells in a fixed temporal sequence. We discussed that a timing function might be crucial in the cerebellar contribution to motor control, and recent data suggest that the cerebellar timing function may also be used in nonmotor tasks. Thus, patients with cerebellar damage were impaired not only in their ability to reproduce a certain rhythm by tapping their fingers but also in discriminating different sound rhythms—that is, not only impaired execution but also **perception of rhythm**. They had no problems with discriminating sounds of different intensities, however, which suggest that the defect is specific to discrimination of **temporal intervals**. Reduced ability to judge the velocity of visual stimuli has also been reported in patients with cerebellar lesions.

The **inferior olive** and the climbing fibers may have a crucial role in the timing function of the cerebellum. Thus, the olivary neurons fire rhythmically, and neurons that activate Purkinje cells within a narrow sagittal zone fire synchronously. Further, there is experimental evidence that the firing rhythm of olivary neurons and the rhythm of certain movements is correlated (see also the earlier section, “The Inferior Olive”).

Another aspect of timing is the judgment of **duration**, for example, how long time has passed since I started a particular action (mental or physical)? Whether the cerebellum plays a role for this time function as well is not clear (see also Chapter 23, under “Interval Timing”).

The Cerebellum and Motor Learning

Animal experiments indicate that long-lasting changes in synaptic efficacy may take place in the cerebellar cortex during motor learning. Much interest is devoted to theories that consider the cerebellum to be a learning machine. The cerebellum may help automation of movements and perhaps also of certain cognitive functions. There is considerable evidence that plastic changes occur in the cerebellum during motor learning. For example, the activation of the cerebellar hemispheres (as measured with fMRI) is higher when a new sequence of movements is learned than when the automated movement is performed afterward. We discussed the possible role of the **climbing fibers** in motor learning, and LTD as a likely cellular mechanism (see the earlier section, “Mossy and Climbing Fibers Mediate Different Kinds of Information”). Recent studies found that patients

with cerebellar lesions have impaired ability to learn **conditioned responses** (shown for the blink reflex and the withdrawal reflex in the leg). It has not been directly shown, however, that LTD is induced in the cerebellum during motor learning.

Because learning takes place in the cerebellum in conditioned responses, it is tempting to assume that corresponding changes occur in the cerebellar hemispheres when humans learn complex **voluntary movements** (such as playing a musical instrument). Indeed, some experiments support this assumption. Thus, patients with cerebellar lesions (or lesions of the inferior olive) show reduced motor learning capacity when wearing prismatic glasses while throwing darts. Both patients and controls missed systematically to one side of the target immediately after putting on the prisms (because the aiming follows the direction of the gaze). During subsequent repetitions, the control persons improved their performance, while no improvement occurred in the cerebellar patients. Conversely, the controls overshot the target after removal of the prisms, although this did not occur in the cerebellar patients.

Although there is a strong case for a cerebellar role in motor learning, this should not be taken to imply that motor learning involves *only* the cerebellum. Indeed, there is good evidence that motor learning involves synaptic changes in distributed networks, including at least motor cortical areas and the basal ganglia (cf. Chapter 22, under “Learning and the Motor Cortex,” and Chapter 23, under “Movement Planning and Learning”).

More about the Cerebellum and Motor Learning

Structural changes of the cerebellum during motor learning have been found in rats that were trained in an **acrobatic task**. After 30 days, the trained rats had significantly more synapses in the molecular layer than the untrained controls had. The increase was in both parallel fiber synapses and climbing fiber synapses with Purkinje cells.

A much-studied example of cerebellar plasticity is adaptation of the **vestibulo-ocular reflex** (VOR) (see Fig. 25.4), as first shown by the Japanese neurophysiologist Masao Ito. The VOR ensures that when the head moves in one direction, the eyes move in the opposite direction with exactly the same speed. This makes it possible to keep the gaze fixed on a stationary object even though the head moves. The magnitude of the reflex response to a certain head movement (that is, the gain of the reflex) needs to be adjusted when, for example, the head grows and alters its proportions. By means of relay stations (see Fig. 25.4), signals from the retina provide information about **retinal slip** (the image is not kept stationary on the retina but moves). Climbing fibers ending in the flocculus provide such signals and thus tell the cerebellum that the velocity of the eye movement is incorrect.

Information about head movements from the vestibular apparatus is provided by mossy fibers, which also end in the flocculus. The sensitivity of the vestibulo-ocular reflex can be altered experimentally in a short time, as shown by making experimental animals wear **prismatic glasses** that displace the image on the retina. The most drastic experiment is when the movement of the surroundings appears to be the opposite of the real movement, leading to a complete reversal of the reflex response. Destruction of the **cerebellar flocculus** prevents adaptation of the reflex. (It is disputed, however, whether the change in synaptic efficacy—that is, the learning—is caused by changes in the cerebellum or elsewhere.)

Certain **conditioned responses** are examples of possible cerebellar participation during motor learning. Especially the so-called **nictitating membrane reflex** (a part of the blink reflex) in rabbits has been investigated. When a jet of air hits the eye the nictitating membrane moves together with the eyelid. This is an unconditioned reflex, in which the trigeminal nerve is the afferent link; the reflex center is in the brain stem involving the sensory trigeminal nucleus, the reticular formation, and the facial nucleus; and the efferent link is the facial nerve to the muscles around the eye. If the jet of air is regularly preceded by a tone (conditioning stimulus), the rabbit will eventually react with a nictitating membrane movement even when the tone is presented alone. The signal pathway for the conditioned response is much more complicated than that for the unconditioned reflex (e.g., the auditory pathways and parts of the cerebral cortex are involved). What is interesting in this connection, however, is that after destruction of the cerebellum, the reflex can no longer be conditioned (i.e., only the unconditioned response occurs, and the animal can no longer be trained to react to the tone only). Damage to the cerebellum after having made the response conditioned abolishes the conditioned (but not the unconditioned) response. It is sufficient to remove a small part of the cerebellar “face area” of the intermediate zone to get these effects (this is a further example of the cerebellar functional localization). After establishing the conditioned response, it can be evoked by electric stimulation of pontocerebellar mossy fibers.

The Cerebellum and Cognitive Functions

Many observations now suggest that the cerebellar functions are not restricted to motor ones. While this topic has attracted much interest, and numerous neuroimaging studies have been performed in normal persons and cerebellar patients, findings are partly contradictory. It is therefore too early to draw conclusions as to the importance of the cerebellum for mental tasks that are usually considered the domain of the cerebral cortex.

We mentioned that the cerebellum might be important for the **perception of rhythm**, not only the proper execution of rhythmic movements. Further, reduced ability to rapidly **shift the attention** from one kind of stimulus to another was found in six patients with lesions of the cerebellar hemispheres. The patients' ability to maintain focused attention was not reduced, and neither was their perception of the stimuli. These and other studies have been taken as evidence that the cerebellum participates in a purely cognitive task—that is, the switching of attention.¹⁰ Many neuroimaging studies demonstrate (indirectly) that blood flow increases (or decreases) in parts of the cerebellum during the performance of various cognitive tasks. For example, persons trying to solve a **pegboard puzzle** showed increased blood flow in the dentate nucleus. In such studies, one seeks to eliminate the possibility that the increase is caused by the movements alone (e.g., moving the pegs). Another study reported increased blood flow in the cerebellar hemispheres in persons trying to find **verbs** going with nouns. Further, increased blood flow in the hemispheres was observed in persons playing **imaginary tennis**, and in persons **counting silently**. Finally, the

cerebellum may also participate in other kinds of learning than learning of movements. Thus, measurements of regional cerebral oxidative metabolism with PET in humans show that the posterior parts of the cerebellar hemispheres increase their neuronal activity when learning a **tactile recognition task** (more than during just tactile recognition of an object).

As mentioned, interpretation of associations between cerebellar change of activity and the performance of cognitive tasks is not straightforward. For example, it is difficult to eliminate the possibility that the cerebellar contribution concerns aspects of execution rather than earlier stages in the processing. When trying to assess the importance of the cerebellum for cognitive functions, we should remember that rather sophisticated tests are required to reveal cognitive defects in cerebellar patients while their motor impairments are obvious and incapacitating. This might be illustrated by a study of children who had their cerebellum partly removed before the age of 3 (because of tumors). These children showed no significant signs of disturbed cognitive development, even though their motor acquisition of motor skills was clearly subnormal. The problems with interpreting clinical data become obvious, however, when adding that among the children, those who had received radiation therapy scored below normal on both cognitive and motor tests.

10 An fMRI study of normal persons challenged this interpretation, however. Thus, altered cerebellar activity occurred only in relation to a motor response associated with the shift of attention.

25 | Control of Eye Movements

OVERVIEW

For the eye to provide useful information to higher visual centers, the picture must be held stationary on the retina. Further, the eyes must be positioned so that the most salient part of a visual scene falls on the central part of the retina with the highest visual acuity. Finally, to sample enough information, the eyes must be moved quickly from one point of salience to another. The control system must therefore be able to move the eyes quickly and precisely to make the image fall on the macula; such movements are called **saccades** (or saccadic movements). In addition, the control system must move the eyes so the retinal image is stationary even if the head or the object is moving. The latter are called **slow-pursuit movements**. The extraocular muscles responsible for moving the eyes receive their nerve supply from the nuclei of the **third, fourth, and sixth cranial nerves**. The control system coordinates the activity of the α motoneurons in these nuclei. **Premotor networks** interconnecting areas in the cerebral cortex, the brain stem, and the cerebellum carry out this task. To enable coordinated activity, the nuclei of the extraocular muscles are interconnected by numerous fibers forming the **medial longitudinal fascicle**. The control system uses **sensory information** from the retina, which informs about whether the retinal image is stationary or slipping, from the **vestibular apparatus** about the movements of the head, and from **proprioceptors** in the eye muscles about the movements of the eyes in the orbit. All this sensory information is integrated and transformed into a **motor signal** specifying the total activity of the extraocular muscles at any time. For control of **horizontal eye movements**—especially saccades—the most important premotor area, the **paramedian pontine reticular formation (PPRF)**, lies close to the abducens nucleus on each side. The PPRF sends fibers to the abducens and oculomotor nuclei and coordinates their activities. A corresponding premotor area for **vertical eye movements** is found in the mesencephalic reticular formation close to the oculomotor nucleus.

There is an important difference between the control of eye muscles and of other muscles subjected to precise voluntary control (e.g., intrinsic hand muscles): the nuclei of the extraocular muscles—in contrast to spinal α motoneurons—receive no direct fibers from the cerebral cortex. The central control is exerted via premotor

networks in the brain stem. At least two regions of the cerebral cortex are closely involved in the control of eye movements (Fig. 15.7): the **frontal eye field** is primarily related to initiation of saccadic movements, whereas several smaller areas in the **parietotemporal region** are mainly involved in the control of pursuit movements.

The third, fourth, and sixth cranial nerves and their nuclei are treated in Chapter 27, together with the light reflex and accommodation reflex mediated by the intrinsic eye muscles. Chapter 16 gives a brief account of the structure of the eye.

MOVEMENTS OF THE EYES AND THE EYE MUSCLES

Horizontal, Vertical, and Rotatory Movements of the Eye

The eye is a sphere, lying in the orbit, surrounded by fat. It can rotate freely in any direction around its center, whereas translatory movements are prevented. To describe the rotatory movements of a sphere, we define **three axes**, passing through the center and oriented perpendicular to each other. For convenience, we describe the movements as taking place in three planes: a **frontal**, **sagittal**, and a transverse or **horizontal** plane. A movement in the horizontal plane takes place around a vertical axis, and the anterior part of the eye—and therefore the gaze—moves from side to side. We perform **horizontal eye movements** when looking to one side; when looking to the left, the left eye rotates laterally and the right eye rotates medially. A movement in the sagittal plane takes place around a transverse axis, and the anterior part of the eye moves up and down. Such movements, directing the gaze up and down, are called **vertical eye movements**. Movements in the frontal plane take place around a sagittal axis, and the eye rotates without any horizontal or vertical movement. For practical reasons only, such movements around a sagittal axis are called **rotatory eye movements** (strictly speaking, all eye movements are rotations around the center of the eyeball).

The Extraocular Muscles and Their Actions

The six extraocular (extrinsic) muscles (Fig. 25.1) ensure that the **visual axes** of the eyes (see Fig. 16.2) can be directed precisely toward any point in the visual field. The scheme in Fig. 15.2 shows the main movements produced by each of the extraocular muscles if they

were acting alone (this is a theoretical situation, because in reality they always work in concert). Most of the muscles produce combinations of vertical, horizontal, and rotatory movements. Further, the actions of each of the muscles change with the position of the eye because this changes the position and direction of the line of pull. The extraocular muscles (Fig. 15.1; see also Fig. 27.16) all **attach** to the sclera and originate from the wall of the orbit (a brief account of the structure of the eye bulb is given in Chapter 16). We analyze the **actions** of the extraocular muscles in relation to the above-defined three axes through the center of the eyeball. We then need to know the direction of the force exerted by the muscles in relation to the axis. We must furthermore know whether the muscle insertion in the sclera is anterior or posterior to the **equatorial plane** of the eye, a frontal plane dividing the eye in an anterior and a posterior half.

There are **four straight** and **two oblique extraocular muscles** (Fig. 25.1, see also Fig. 27.16). Figure 25.2 shows schematically the actions of each muscle if it acted alone. Most muscle produce movements that are composed of horizontal, vertical, and rotatory components. We can nevertheless simplify their functions by stating that two muscles produce predominantly horizontal movements (the medial and lateral rectus muscles), two produce predominantly vertical movements (the superior and inferior rectus muscles), whereas two muscles produce mainly rotatory movements (the superior and inferior oblique muscles). The straight ones (the rectus muscles) come from the posterior end of the orbit and run forward to insert in front of the equatorial plane. This means that the **lateral rectus** muscle pulls the front of the eye (the cornea) laterally, whereas the **medial rectus** muscle pulls it medially (these two muscles thus produce pure horizontal movements). Correspondingly, the

superior rectus muscle pulls the eye (the cornea) upward, whereas the **inferior rectus** muscle pulls it downward. These two therefore produce vertical movements. Because the superior and inferior rectus muscles run anteriorly in a lateral direction, however, they do not only produce vertical movements but also some horizontal movement (in the medial direction). Thus, when the superior rectus muscle acts alone, it produces an upward movement combined with a (smaller) medial—that is, an oblique movement. In addition, the muscle produces a small medial rotation of the eye (around the sagittal axis). The **superior oblique** muscle has a more complicated course than the other extraocular muscles (Fig. 15.1; see also Fig. 27.16). It originates posteriorly in the orbit and runs forward medially. Just behind the anterior margin of the orbit, it bends sharply around a small hook of connective tissue and continues in a posterolateral direction to insert posterior to the equatorial plane. The muscle has actions around all three axes: it rotates the eye around the sagittal axis so that the upper part moves medially and furthermore directs the gaze downward and laterally (Fig. 15.2). The **inferior oblique** muscle originates from the bottom of the orbit in its anteromedial part and runs, like the superior oblique, posterolaterally to insert behind the equator (Fig. 15.1). It directs the gaze laterally and upward and rotates the eye around the sagittal axis with its upper part laterally.

These considerations and the scheme in Fig. 15.2 concern movements starting from a **neutral position** of the eye—that is, when viewing a distant object straight ahead. Changing the position of the eye in the orbit also changes the action of the muscles (for some muscles the change is small, for others quite marked). For example, the superior rectus is a pure elevator when the eye is 23 degrees abducted. The oblique muscles perform pure vertical movements when the eye is about 50 degrees abducted (cf. Fig. 27.16 showing the line of action of the superior oblique). This is basis for the test of the extraocular muscles in Fig. 25.3, designed to test each muscle in isolation (as far as possible).

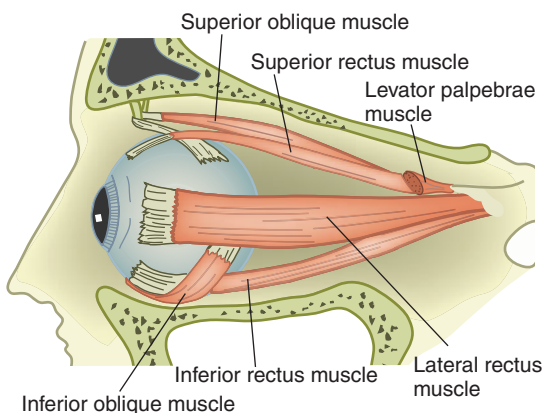


FIGURE 25.1 The extraocular muscles seen from the lateral aspect. The lateral wall of the orbit is removed. The straight muscles insert in front of the equatorial plane of the eye, whereas the oblique muscles insert behind it (see also Fig. 27.16, which shows the orbit and the eye muscles from above).

Natural Eye Movements Are Conjugated

Virtually every natural eye movement is a combination of the various movement directions described in the preceding text. By combining proper amounts of vertical and horizontal movements, any oblique movement can be produced. Further, all natural eye movements are **conjugated**—that is, the two eyes move together to ensure that the image always falls on corresponding points of the two retinas (see Fig. 16.3). **Double vision (diplopia)** results if the eye movements do not occur in conjugation. This is a typical symptom of pareses of the extraocular muscles.

Almost all eye movements require a complicated cooperation of numerous muscles, with activation of synergists

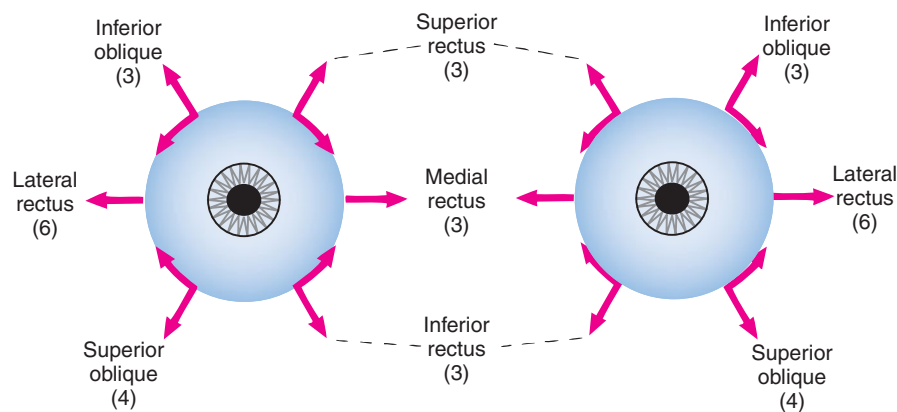


FIGURE 25.2 Actions of the extraocular muscles. These are as observed when movements start from a neutral position of the eye. Arrows indicate direction of the action (but not the force). Except for the medial and lateral recti, all muscles rotate the eye, in addition to their other actions. Starting positions away from the neutral may alter the actions of the muscles. For example, the superior rectus is a pure elevator when the eye is 23 degrees abducted (cf. Fig. 15.3).

and inhibition of antagonists. When, for example, we look to the left, we activate the left lateral rectus and inhibit the left medial rectus, whereas we activate the right medial rectus and inhibit the right lateral rectus. This is the simplest possible example, with a purely horizontal movement. In most other situations, the cooperation between various muscles becomes much more complicated and requires an extensive, sophisticated neural network for control. Electromyographic (EMG) recordings in awake human subjects with their eyes open show that there is some activity in virtually all of the extraocular muscles. The tension produced by each muscle varies, of course, with the position of the eyes.

The Eye Muscles Are Built for Precise Control

The structure of the extraocular muscles reflects their use in extremely delicate and precisely controlled movements. The muscle fibers are very thin in comparison

with ordinary skeletal muscle fibers, and the **motor units** are among the smallest in the body (only 5–10 muscle fibers per motoneuron). Consequently, the nerves to the extraocular muscles contain many nerve fibers: the abducens nerve in humans (supplying only one muscle) contains around 6000 axons.

The extraocular muscles are required both to hold a certain tension for a long time (static position holding) and to produce extremely fast movements. Accordingly, the maximal **speed of contraction** is high in comparison with other skeletal muscle fibers. Further, the maximal **firing frequency** of the motoneurons—occurring during saccadic movements—is unusually high. Patients with a **paralysis of the lateral rectus** muscle demonstrate that there is **constant activity** in extraocular muscles. Even though the only muscle capable of active abduction is paralyzed, a small abduction nevertheless occurs when the patient attempts to look to the affected side. This abduction is caused by the relaxation of muscles that adduct the eye, primarily the medial rectus (an example of reciprocal inhibition).

The extraocular muscles are composed of a mixture of fibers with fast and with slow **twitch** contractions. In addition, there are muscle fibers that do not produce twitches but only a slow graded contraction (they receive multiple end plates along the length of the muscle and occur only in eye muscles). Their function is unknown, but presumably, they contribute to static position holding.

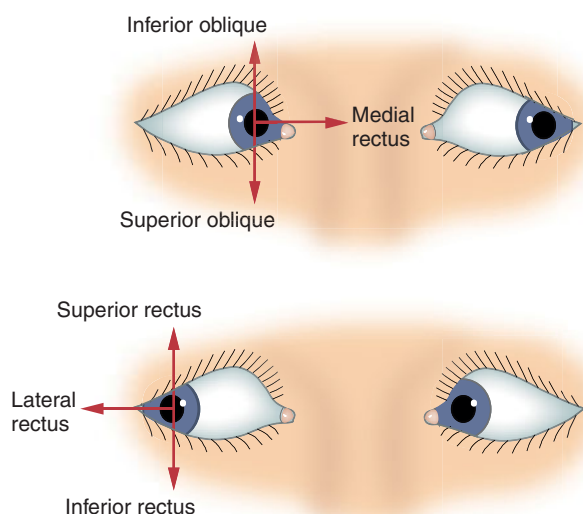


FIGURE 25.3 Scheme for testing the eye muscles. Based on testing in extreme positions in which the muscles produce relatively pure vertical or horizontal movements. See text for further discussion.

Eye Muscles, Proprioception, and Efference Copy

There are **muscle spindles** in the extraocular muscles (100–150 altogether in the six human extraocular muscles). Apart from the general rule that muscles subject to precise control have a high density of muscle spindles, their functional role in the extraocular muscles is not clear. Thus, signals from extraocular muscle spindles are apparently not consciously perceived—that is, they do not contribute to the awareness of eye position and movements (in this respect differing from muscle spindles elsewhere). Nor do signals from extraocular

muscle spindles evoke stretch reflexes. There is evidence that muscle-spindle signals, providing information about the length and the change of length of the extraocular muscles, are integrated in the cerebral cortex with signals providing information about movements of the head from the labyrinth and with signals from the retina. Experiments in monkeys with transection of the ophthalmic nerve (assumed to contain most of the muscle spindle afferents from the extraocular muscles) did not disturb saccadic and pursuit movements, although they did produce instability of the eyes and slow pendular movements in darkness. It seems that the control system receives enough information from the **efference copy** (i.e., the copy of the motor commands sent to the extraocular muscles).

Presumably, information from the eye muscle proprioceptors contributes to the long-term calibration of the efference copy. When there is a mismatch between the command and the feedback (informing about the true movement of the eye) this may be used to change the motor program. Such mismatch arises, for example, when the dimensions of the eye and the eye muscles change as we grow. Thus, proprioceptive information is presumably necessary to ensure fusion of the retinal images (avoid strabismus with double vision).

BRAIN STEM AND CEREBELLAR CONTROL OF EYE MOVEMENTS

Kinds of Eye Movement

As mentioned, the control system for eye movements must ensure that the gaze can be moved quickly from one point of fixation to another by **saccadic movements**, but also that the gaze can be kept stationary on an object when the object or the head moves by **pursuit movements**. The eye movements may be either **voluntary**, as when we follow a moving object with the eyes, or **reflex**, as when the head moves (a **vestibulo-ocular reflex**) or the surroundings move (an **optokinetic reflex**). The vestibulo-ocular reflex and nystagmus are discussed in Chapter 18.

The following is a highly schematic description of eye movements. Different **premotor networks** control each of the kinds of movement, although they all converge on the motoneurons of the extraocular muscles. The separation of the different central networks is less clear-cut than formerly believed, however. Thus, recent studies show that several regions, for example in the cerebral cortex and the cerebellum, participate in the control of both saccadic and pursuit movements.

1. **Saccades** are conjugated movements that change the visual axis of the eyes from one point of fixation to another with maximal speed. Saccades can (at least in a certain sense) be **voluntary**, as when we look at a stationary

landscape and fix the gaze at one point for a moment and then move on (with a saccade) to another point of fixation. They can also occur **reflexly**, as part of vestibular or optokinetic nystagmus. When awake, we perform saccades all the time, often several per second. In this way, we constantly **scan** the visual scene to provide maximal information. Because the visual acuity declines so rapidly outward from the macula, scanning is necessary to use the full analyzing capacity of the visual system.

2. **Smooth-pursuit movements** are performed when we follow a small moving object to keep the image stationary on the central part of the retina. As a rule, we use pursuit movements both of the eyes and of the head when looking at a moving object. One might think that the movements of the head would elicit conjugated eye movements in the opposite direction of the head because of the vestibulo-ocular reflex. This does not occur, however, because the vestibulo-ocular reflex is suppressed during such smooth-pursuit movements. This kind of smooth movement is **voluntary** in the sense that it requires that our attention be directed to something in the visual field; that is, the gaze is voluntarily fixed. Vestibular and optokinetic stimuli can also elicit reflex slow movements that stabilize the retinal image. These are involuntary in the sense that they occur also without paying attention to something in the visual field.

3. **Optokinetic reflex movements** are movements intended to stabilize the retinal image when the whole visual field moves relative to the head. The stimulus is movement of the image on the retina (**retinal slip**). When looking out from a train this kind of movement occurs. Then a slow movement (following the scene moving outside the window) is interrupted by a quick, opposite movement—resetting the eyes to their original position—when the eyes cannot follow the scene anymore. This movement pattern is called **optokinetic nystagmus**. The optokinetic reflex can be suppressed voluntarily only by fixating on an object that is stationary (or moves with a different speed) in relation to the movement of the rest of the visual scene.

4. **Vestibulo-ocular reflex movements** (VORs) are eye movements elicited by movements of the head (Fig. 25.4). The eyes move with the same velocity as the head but in the opposite direction, thereby ensuring that the retinal image is stationary. When the eye cannot move further, a fast movement occurs in the opposite direction, resetting the position of the eye; then the slow movement starts again. These alternate movements are called **vestibular nystagmus**. The quick phase of vestibular nystagmus and saccades share many characteristics, but are not identical. The VOR enables us to see an object sharply even when we move around. When the head rotates, the stimulus originates in the semicircular ducts. Even though the VOR has been most studied in relation to rotations in the horizontal plane (Fig. 15.4), the

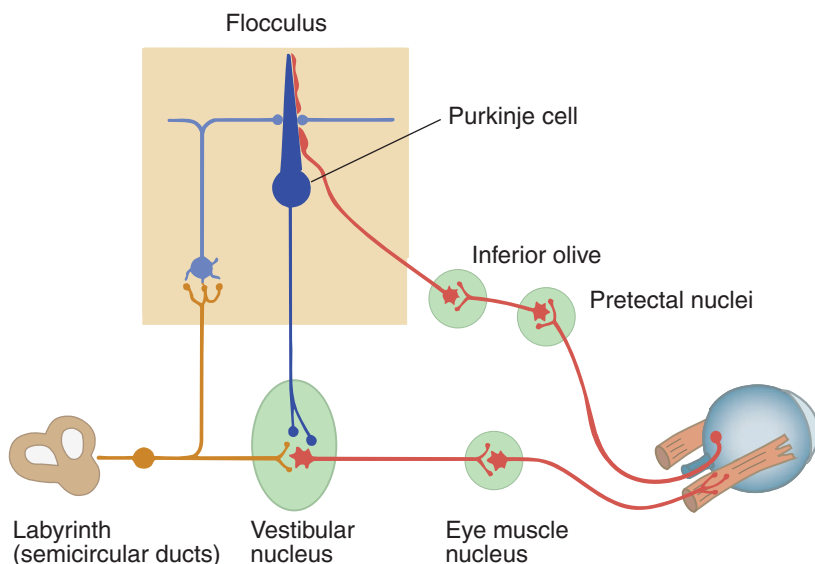


FIGURE 25.4 Main structural elements of the vestibulo-ocular reflex. Only excitatory connections are shown, even though there are inhibitory neurons in the vestibular nuclei that influence the motoneurons of the antagonists. The reflex arc consists of three neurons from the semicircular duct to the extraocular muscles. The cerebellar flocculus receives signals from the labyrinth and from the retina, and the output of the Purkinje cells can adjust the sensitivity of the vestibular neurons, if necessary, to avoid retinal slip. (Based on Ito 1984.)

reflex is three-dimensional in the sense that all directions of head rotation elicit specific compensatory eye movements.¹

5. **Vergence** movements change the visual axes of the eyes in relation to each other when the point of fixation moves away from or toward the eyes. This is necessary to keep the image on corresponding points of the retina. Vergence movements are a prerequisite for fusion of the two images and for stereoscopic vision. **Convergence of the visual axes**, which takes place when an object is approaching the eyes, depends primarily on the activity of the medial rectus muscles, with some contribution also from the superior and inferior recti (Fig. 15.2). **Accommodation** and **pupillary constriction** accompany convergence movements.²

More about Voluntary Saccades and Scanning

When **reading** we fixate a point on the line for an average of 250 msec (60–500) before the gaze is moved on by a saccade. How far the gaze moves before reaching a new point of fixation varies greatly. There is a tendency to fixate on long “content” words rather than on short

“functional” words. Native readers of English perceive about 4 letters to the left and 15 to the right of the point of fixation.

A woman with inborn **ophthalmoplegia** (inability to move the eyes) had surprisingly small problems and was able to live a normal life. She apparently used quick head movements to compensate for the lack of saccadic eye movements (Gilchrist 1997), and was thereby able to scan the visual scene with sufficient speed and accuracy.

The Cerebellum Can Adjust the Vestibulo-Ocular Reflex to Changing External Conditions

The magnitude of the reflex response (not the response itself) to a certain rotational stimulus depends on signals to the vestibular nuclei from the cerebellum (Fig. 25.4). The Purkinje cells of the vestibulocerebellum receive primary vestibular fibers (ending as mossy fibers) that provide information about direction and velocity of the head movement. In addition, the same Purkinje cells receive information, via the inferior olive and climbing fibers, about whether the image is stationary or slips on the retina. A retinal slip indicates that the velocity of the compensatory head movement is too high or too low. The cerebellum is then capable of adjusting the excitability (the gain) of the neurons in the vestibular nuclei—that is, in the reflex center of the vestibulo-ocular reflex. Such adaptive change of **gain of the reflex** is presumably needed continuously during growth and in situations of muscular fatigue. Experiments in which animals wear optic prisms that deflect the light so that it appears to come from another direction than it really does, show the remarkable capacity for adaptation (learning) in this system.

1 The VOR described here—the **rotational VOR** or RVOR—is not the only vestibulo-ocular reflex. Translatory (linear) accelerations of the head (stimulating the sacculus and utricle) also elicit compensatory eye movements (**translational VOR**, or TVOR; **otolith-OR**, or OOR). In real life, both kinds of head movement occur, and the different sensory inputs from the labyrinth must be integrated centrally to yield a motor command that ensures a stable retinal image.

2 **Gaze holding** is sometimes included among the kinds of eye movement. It is the ability to stabilize the eye position (and the image on the retina) after a shift of gaze. Premotor neurons controlling gaze holding appear to rely on a partly separate premotor network, although located close to the paramedian pontine reticular formation (PPRF) and the rostral interstitial nucleus of the medial longitudinal fasciculus (riMLF).

Brain Stem Centers for Control of Eye Movements

Signals informing about desired eye position, actual position, retinal slip, and position of the head are integrated in the **reticular formation** close to the eye muscle nuclei. From these **premotor** neuronal groups commands are sent to the α motoneurons. By combining anatomic data on the fiber connections with physiological results (obtained by single-cell recordings, electrical and natural stimulation, and lesions) the preculomotor networks have been described in detail.

A “center” for **horizontal eye movements** has been identified in the **paramedian pontine reticular formation (PPRF)**. PPRF lies close to, and sends fibers to, the abducens nucleus (Fig. 25.5). It also sends fibers to the parts of the oculomotor nucleus that contains the motoneurons of the medial rectus muscle. Together with the lateral rectus (innervated by the abducens nucleus), the medial rectus participates in horizontal movements. In addition, there are so-called **internuclear neurons** in the abducens nucleus that send axons to the medial rectus motoneurons of the opposite side (Fig. 25.6). This premotor network ensures simultaneous activation of the lateral rectus on one side and the medial rectus on the other, along with inhibition of the antagonists.

A **lesion** in the region of the **PPRF** reduces horizontal conjugate movements to the side of the lesion. Especially marked is the reduction in saccadic movements. A unilateral lesion of the **medial longitudinal fasciculus** between the abducens and the oculomotor nucleus produces

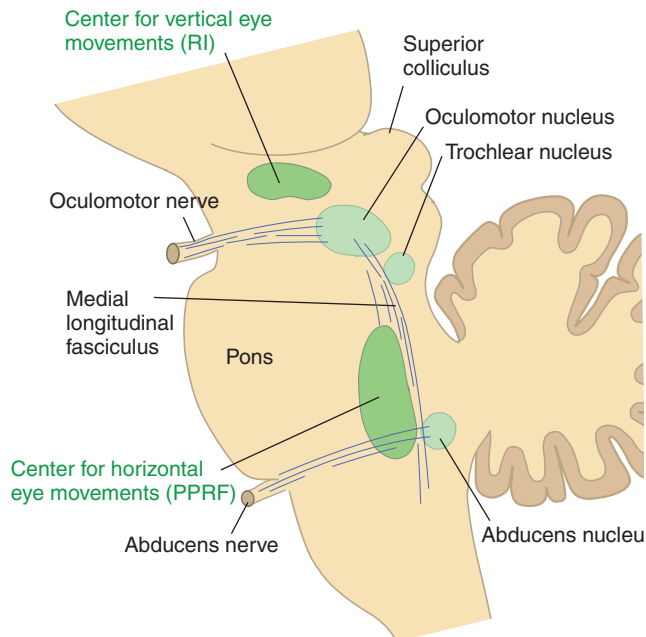


FIGURE 25.5 Centers for vertical and horizontal eye movements. Sagittal section through the monkey brain stem. (Based on Büttner-Ennever 1988.)

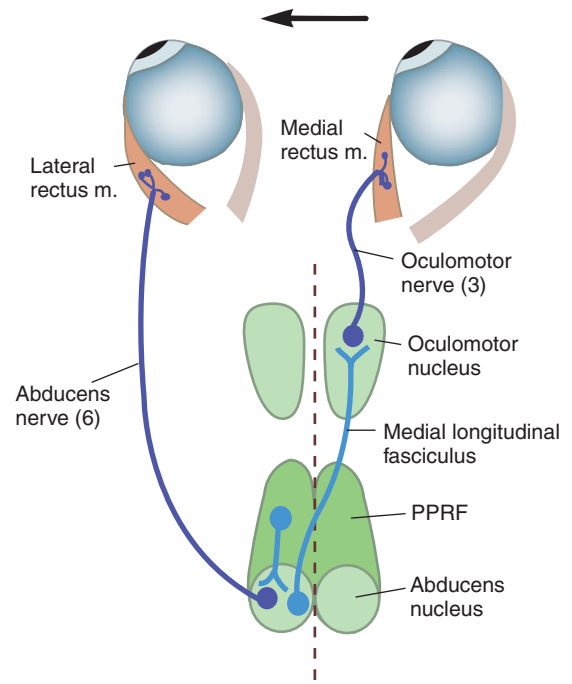


FIGURE 25.6 Control of horizontal conjugate eye movements. Highly simplified scheme to show some of the connections responsible. Only excitatory connections are included. The figure also shows how interruption of the medial longitudinal fascicle produces a paralysis of the gaze in the medial direction (internuclear ophthalmoplegia).

so-called **internuclear ophthalmoplegia** with abolished ability to adduct the eye on the same side (the medial rectus muscle). This may be understood based on the diagram in Fig. 25.6. However, **vergence** movements are possible even though the medial rectus is responsible also in that case. Thus, pathways other than the medial longitudinal fasciculus are responsible for the activation of the medial rectus muscle during vergence movements.

A center for **vertical** and rotatory eye movements has been identified in the reticular formation close to the oculomotor nucleus. This region includes a nucleus called the **rostral interstitial nucleus of the medial longitudinal fasciculus (riMLF or RI)** and probably another small cell group (the **interstitial nucleus of Cajal**). This region receives **afferents** from the vestibular nuclei, the pretectum (and thereby indirectly from the superior colliculus), and the frontal eye field. Physiologic studies indicate that there are monosynaptic connections from the region of the riMLF to the trochlear nucleus and to the motoneuron groups within the oculomotor nucleus that produce vertical eye movements.

The PPRF and the region of the RI are interconnected, indicating that they do not operate independently of each other. The fact that most natural eye movements are neither purely horizontal nor purely vertical but combinations of both strongly suggests that the centers

for horizontal and vertical movements must coordinate their activity.

Properties of Neurons in the PPRF

The PPRF receives signals directly and indirectly from the vestibular nuclei, the superior colliculus, and the frontal eye field. It appears to function as the last premotor station for the initiation of conjugate horizontal saccadic movements. In agreement with this, there are neurons in the PPRF that are active immediately before and during a saccade to the same side, whereas they are inactive during slow-pursuit movements and fixation. These so-called **excitatory burst neurons** establish excitatory connections with the motoneurons responsible for the saccade. There are also **inhibitory burst neurons**, located close to the PPRF, with a pattern of activity similar to that of the excitatory burst neurons. Monosynaptically they inhibit the motoneurons in the abducens nucleus of the opposite side. The activity of both kinds of burst neurons appears to be controlled by a third kind of neuron in the PPRF, which is tonically active. Such **omnipause neurons** probably inhibit the burst neurons tonically, except in relation to saccades. Thus, the omnipause neurons are silent immediately before and during saccades, regardless of the direction and amplitude of the movement. Recent studies indicate that the omnipause neurons receive direct connections from both the superior colliculus and the frontal eye field.

Afferent Connections of the Ocular Premotor Networks

Many kinds of information must be integrated and translated into motor commands to achieve the extremely precise control of eye movements. **Visual signals** do not pass directly from the retina to the eye muscle nuclei but are mediated by several brain stem nuclei, which also receive impulses other than visual information. This concerns the **superior colliculus** and the **pretectal nuclei** (situated rostral and ventral to the superior colliculus; see Fig. 27.19). These nuclei receive afferents from the retina and from the parts of the cerebral cortex that are involved in control of eye movements (the frontal eye field, the posterior parietal cortex, and parts of the extrastriate visual areas; Fig. 25.7). Many neurons in the superior colliculus are active immediately before a saccade and most likely have an important role in the organization of conjugated saccades. This is mediated by connections from the superior colliculus to the reticular formation and from there to the eye muscle nuclei. In addition, the superior colliculus is important for integration of somatosensory and visual information. This is needed, for example, to rapidly direct the gaze to a somatosensory stimulus (involving simultaneous and coordinated movements of the eyes and the head).

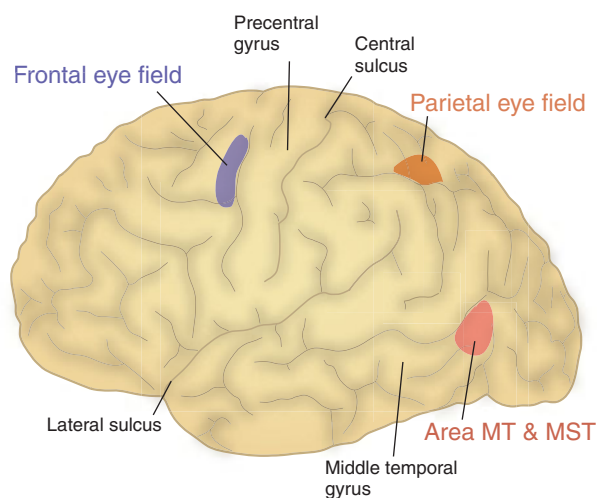


FIGURE 25.7 Cortical areas involved in control of eye movements. Areas on the medial wall of the hemisphere are not shown. Size and position of the areas are only approximate. (Based on Müri 2006.)

We have previously mentioned connections from the **vestibular nuclei**, which, by means of the **medial longitudinal fasciculus**, reach all the cranial nerve nuclei of the eye muscles (see Fig. 18.8A). These connections are both crossed and uncrossed. Some of the vestibular neurons sending their axons into the medial longitudinal fasciculus are excitatory; other neurons are inhibitory (GABA). The connections are precisely organized, enabling the activation of specific synergists and the inhibition of their antagonists (with regard to the intended eye movement). Other efferent fibers from the vestibular nuclei end in cell groups situated near the eye muscle nuclei, which send their axons to the latter. The **perihypoglossal nuclei** are examples of such preoculomotor cell groups. These nuclei (the most prominent one is the prepositus hypoglossus nucleus) lie around the hypoglossal nucleus in the medulla. In addition to afferents from the vestibular nuclei, the perihypoglossal nuclei receive afferents from other structures involved in eye movement control, such as parts of the reticular formation and the cerebellum. The direct and indirect impulse pathways from the vestibular nuclei to the eye muscle nuclei mediate the **vestibulo-ocular reflexes** (Fig. 15.4) and are probably involved also in **optokinetic reflexes**. In the latter case, signals from the retina reaching the vestibular nuclei (synaptically interrupted in several nuclei) must be involved rather than signals from the labyrinth.

Proprioceptive signals from muscle spindles of the extraocular muscles reach the eye muscle nuclei from the **mesencephalic trigeminal nucleus** (see Fig. 27.13). As mentioned, proprioceptive signals are probably necessary for the long-term correction and adaptation of the mutual position of the eyes in fixation.

The Cerebellum Controls Both Saccades and Pursuit Movements

Cerebellar lesions can impair both saccadic and slow-pursuit movements. The middle part of the **vermis** (lobules VI–VII, vermal oculomotor region) is particularly concerned with the performance of **saccades**, while the **flocculus** and adjoining parts of the posterior lobe (the **paraflocculus**) seem most important for **pursuit** movements. After cerebellectomy, monkeys are unable to perform pursuit movements, whereas saccades can be performed although with reduced precision and velocity. Clinical observations support the role of the cerebellum in control of pursuit movements also in humans (the deficits observed in patients with cerebellar lesions resemble the dose-dependent effects of alcohol on pursuit movements). The distinction between the vermal oculomotor region and flocculus/paraflocculus is not absolute, however, because both contain neurons related to either saccades or pursuit movements. In accordance with this, animal experiments and observations in humans indicate that the vermal oculomotor region controls both precision of saccades and the velocity of pursuit movements.

Recent studies strongly suggest that lateral parts of the **pontine nuclei** are necessary for voluntary pursuit movements and for the slow phase of optokinetic nystagmus.³ This is based on recordings of single-cell activity and lesion experiments in monkeys, as well as in a few patients with small brain-stem infarctions. The relevant parts of the pontine nuclei send efferents to both oculomotor-related cerebellar regions, and they receive afferents from (among other areas) the **middle superior temporal area (MST)**, which is related to visual analysis of moving objects. Parallel pathways seem to exist for the control of pursuit movements, however—for example, the **pontine reticulotegmental nucleus (PRN)**, located immediately dorsal to the pontine nuclei, also relays signals from “cortical oculomotor areas” to the cerebellum. The fact that the PRN is involved also in saccadic movements underscores that the networks controlling different kinds of eye movement are not entirely separate.

CORTICAL CONTROL OF EYE MOVEMENTS

There are no direct connections from the cerebral cortex to the eye muscle nuclei. Activation of the extraocular muscles from the cortex in conjunction with voluntary eye movements is mediated via other brain stem cell groups,

among them the premotor “gaze centers” discussed in the preceding text. Recent functional magnetic resonance imaging (fMRI) and positron emission tomography (PET) studies in humans indicate that a distributed cortical **network** controls eye movements. This network includes discrete regions in the frontal and parietal lobes, and at the temporo-occipital junction. While each area appears to be most important for one kind of movement, all areas alter their activity in relation to both saccades and pursuit movements (based on human fMRI studies). Interestingly, areas for control of eye movements (especially saccades) overlap areas related to **shift of attention**. This seems logical because a voluntary saccade is a motor expression of a shift of attentional focus.

Saccades

With regard to specializations among cortical areas, clinical observations indicate that the **frontal eye field (FEF, or area 8 of Brodmann; Fig. 25.7, see also Fig. 33.3)** is of particular importance for **voluntary saccades**. Electrical stimulation of the frontal eye field elicits conjugated eye movements to the opposite side. The effect is mediated by fibers descending in the internal capsule to brain stem premotor cell groups, such as the **superior colliculus**, the **pretectal nuclei**, and the **PPRF**, which, in turn, activate motoneurons in the eye muscle nuclei. A unilateral **lesion** of the frontal eye field makes the patient unable to move the gaze voluntarily to the side opposite the lesion (e.g., when asked by the examiner to move the gaze). This is called a **gaze paralysis**. Correspondingly, the activity in the FEF increases more during voluntary than during reflex-evoked saccades. Indeed, the ability to move the gaze laterally is not completely lost after a lesion of the FEF. Thus, smooth-pursuit movements occur when an object is brought into the visual field and then moved slowly laterally—that is, the patient is able to follow the object with her gaze. Imaging studies show additional activation of the **SMA**, **dorsolateral prefrontal cortex**, and **posterior parietal cortex** (see Fig. 21.13) in relation to voluntary saccades. An area close to the frontal eye field, the **supplementary eye field (SEF)** is particularly active when saccades are coordinated with hand movements (based on single-unit recording in behaving monkeys). It is also active when saccades are made to a remembered target location—that is, in the absence of visual or other information about where the target is.

The **superior colliculus** and the **vermal oculomotor area** of the cerebellum both receive information from the frontal eye field and the posterior parietal cortex, and both are necessary for proper execution of saccades. They are not necessary for their initiation, however, because lesions of neither the cerebellum nor the superior colliculus abolish saccadic movements.

3 Voluntary pursuit movements and the slow phase of optokinetic nystagmus appear to be controlled by largely the same neuronal groups because both kinds of movement are disturbed after lesions of the middle superior temporal area (MST), the pontine nuclei, or the flocculus/paraflocculus.

Pursuit Movements

With regard to cortical control of **smooth-pursuit movements**, several small areas in the parietal lobe, and at the temporo-occipital junction are of particular importance (Fig. 15.7; see also Figs. 16.25 and 34.2). This notion is based on, among other things, single-cell recordings. The **middle temporal area** (area MT, V5) is important for perception of movement. Neurons in MT respond to movement of the image on the retina (retinal slip), which is a strong stimulus to elicit pursuit movements of the eyes. The **middle superior temporal area** (area MST) lies close to area MT and is of special interest because it contains many neurons that respond preferentially to moving visual stimuli in a specific direction. The subcortical cell groups intercalated between these cortical regions and the eye muscle nuclei are not fully clarified. One important pathway seems to go from the cortex to the **pontine nuclei** (see Fig. 24.10), then to the **cerebellum**, further to the **vestibular nuclei** and the **reticular formation**, and thence to the eye muscle nuclei. In addition to in the parietal and temporal areas, neurons with activity related to pursuit movements occur in the frontal eye field and the supplementary eye field. Accordingly, impairment of pursuit movements has been reported in humans with frontal lesions including the supplementary eye field.

Gaze Fixation

The ability to **fix the gaze** is a prerequisite for voluntary slow-pursuit movements. A specific fixation center in the cerebral cortex has not been found; as with most other tasks carried out by the cerebral cortex the task is solved in a distributed network. In monkeys, single

neurons that change their activity in relation to gaze fixation are present in the frontal eye field and in the posterior parietal cortex. PET studies in humans, however, indicate that frontal lobes are particularly important for the ability actively to fix the gaze. This includes the **frontal eye field**, the **anterior cingulate gyrus**, and parts of the **prefrontal cortex**. In general, these regions become active in conjunction with **directed attention**, and lesions in humans produce an impaired ability to fix the gaze.

Often the tendency to fix the gaze on an object is not voluntary (cf. the term fixation reflex), at least not in the strict sense (even though fixation depends on the person being conscious). **Optokinetic nystagmus** is an expression of the strong tendency to fix the gaze. When a screen with alternating black and white vertical stripes is moved horizontally in front of a person, nystagmus occurs. The gaze is fixed automatically on one of the stripes and follows it until it leaves the visual field. Then a quick movement resets the eyes to the starting position, and the gaze is fixed again on another stripe. This sequence repeats itself as long as the screen moves.

Miniature Eye Movements

Closer study shows that even during active fixation, the eyes are not completely still. Several kinds of **miniature movement** occur, such as a slow drift of the eyes, miniature saccades, and tremor. The amplitudes of these movements are very small, however, and are not normally perceived. Nevertheless, the phenomenon can be demonstrated by fixating a square pattern for about 20 seconds and then move the gaze to a white surface. The afterimage of the square pattern is then seen to move, because the eyes are not completely still (Ilg 1997).

V

THE BRAIN STEM AND THE CRANIAL NERVES

THE definition of the term brain stem varies. The widest definition—used in Chapter 6—includes the medulla oblongata, the pons, the mesencephalon, and the diencephalon. In this part of the book, and most often in clinical contexts, we restrict it to the **medulla**, the **pons**, and the **mesencephalon**, which together form a macroscopically distinct part of the brain. Which definition is used is of no great importance because the brain stem is a topographically and embryologically defined unit; it does not represent a functional “system.” Neuronal groups within the brain stem take part in virtually all the tasks of the central nervous system.

Functionally, the brain stem may be said to have two levels of organization. On the one hand, most of

the cranial nerves and their nuclei represent “the spinal cord of the head.” On the other hand, many neuronal groups in the brain stem represent a superior level of control over the spinal cord and the cranial nerve nuclei. Examples are the vestibular nuclei and various premotor nuclei in the reticular formation. In addition, other neuronal groups—usually included in the reticular formation—exert ascending influences on the thalamus and the cerebral cortex related to consciousness, arousal, and sleep. This section deals with both levels of organization. Chapter 26 deals with the reticular formation, which represents the superior level of control. Chapter 27 describes most of the cranial nerves and their nuclei.

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The Reticular Formation: Premotor Networks, Consciousness, and Sleep

OVERVIEW

The reticular formation extends from the lower end of the medulla to the upper end of the mesencephalon. At all levels, it occupies the central parts and fills the territories not occupied by cranial nerve nuclei and other distinct nuclei and by the large fiber tracts. The **raphe nuclei** (serotonin) and the **locus coeruleus** (norepinephrine) are often included in the reticular formation, and are treated in this chapter. Although the reticular formation consists of many functionally diverse subdivisions, they all share some anatomic features. Thus, the neurons have wide dendritic arborizations and their long axons give off numerous collaterals. The afferent and efferent connections show only a rough topographic order. Efferent connections reach most parts of the central nervous system (CNS, from the cord to the cerebral cortex), while afferents bring all kinds of sensory information. These features show that the reticular formation is built for **integration**. Accordingly, the reticular formation attends primarily to tasks involving the nervous system and the organism as a whole. Subdivisions of the reticular formation form **premotor networks** that organize several complex behaviors. These behaviors include control of body **posture**, **orientation** of the head and body toward external stimuli, control of **eye movements**, and coordination of the activity of the **visceral organs**. These tasks fit with the reticular formation being phylogenetically old and present even in lower vertebrates. In addition, parts of the reticular formation (especially in the upper pons and mesencephalon) send ascending connections to the thalamus and the cerebral cortex. These connections form the **activating system of the brain stem**. The integrity of this system is a prerequisite for **consciousness**, and is closely linked to control of **awareness** and **attention**. Further, parts of the reticular formation, including cholinergic cell groups in the upper pons, are concerned with regulation of **sleep**.

STRUCTURE AND CONNECTIONS OF THE RETICULAR FORMATION

Structure and Subdivisions

The reticular formation extends from the lower end of the medulla to the upper end of the mesencephalon, where it gradually fuses with certain thalamic cell groups (Fig. 26.1).¹ At all levels it occupies the central parts and fills the territories not occupied by cranial nerve nuclei and other distinct nuclei (such as the dorsal column nuclei, the pontine nuclei, and the colliculi) and by the large fiber tracts (such as the medial lemniscus and the pyramidal tract). The reticular formation received its name from the early anatomists because of its network-like appearance in microscopic sections (Figs. 26.2 and 26.3). It is built of cells of various forms and sizes that appear to be rather randomly mixed. Between the cells, there is a wickerwork of fibers passing in many directions (Fig. 26.2). These fibers are partly axons and dendrites of the neurons of the reticular formation and partly afferent axons from other sections of the CNS. The reticular neurons have typically very long and straight **dendrites**, so that they cover a large volume of tissue (Fig. 26.3). This distinguishes the reticular neurons from those found in specific nuclei of the brain stem, such as the cranial nerve nuclei (see the hypoglossal nucleus in Fig. 26.3).

Medial Parts Are Afferent and Lateral Parts Are Efferent

A more detailed analysis of the reticular formation makes clear, however, that it consists of several **subdivisions**, among which the cells differ in shape, size, and arrangement even though the borders between such subdivisions are not sharp (Fig. 26.1). It is especially

¹ Some authors include certain thalamic cell groups—especially the intralaminar nuclei—in the term “the reticular formation of the brain stem.”

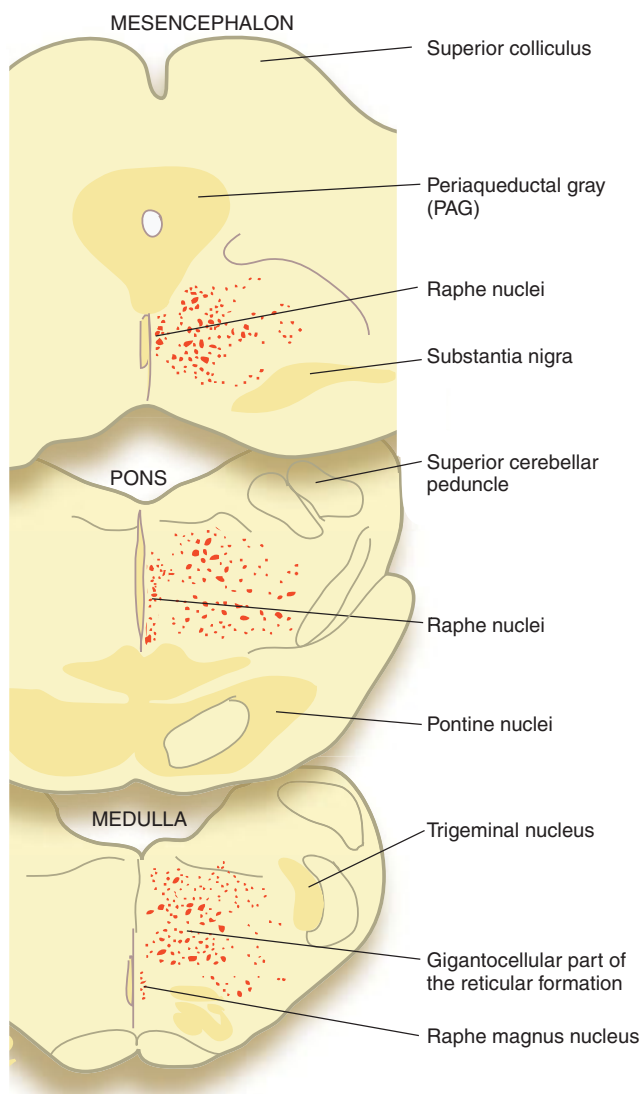


FIGURE 26.1 *The reticular formation.* Transverse sections through various levels of the brain stem (cat) showing the position of the reticular formation. The size of the red dots indicates the size of the neurons, which varies considerably among subdivisions of the reticular formation. (Based on Brodal 1957.)

important that such cytoarchitectonically defined subdivisions also differ with regard to fiber connections, neurotransmitters, and functions. In the pons and the medulla, approximately the **medial two-thirds** of the reticular formation consists of many large cells, in part so-called **giant cells** (Figs. 26.1 and 26.2). The **lateral one-third** contains almost exclusively **small cells**. Tract-tracing methods have shown that the medial part sends out many long, ascending and descending fibers, whereas the lateral small-celled part receives most of the afferents coming to the reticular formation. In general, therefore, we may say that the lateral part is **receiving**, whereas the medial part is **efferent** (executive). The efferents convey the influence of the reticular formation to higher

parts, such as the thalamus, and lower parts, such as the spinal cord.

The Reticular Formation Is Built for Integration

Studies with the Golgi method and with intracellular tracers give evidence of how complexly the reticular formation is organized. The long ascending and descending efferent fibers give off numerous **collaterals** on their way through the brain stem (Fig. 26.4). As can be seen in Fig. 26.3, the collaterals run primarily in the transverse plane. Most dendrites of reticular cells have the same preferential orientation, so the reticular formation appears to consist of numerous **transversely oriented disks** (Fig. 26.3). The numerous collaterals of the axons from the cells in the medial parts ensure that signals from each reticular cell may reach many functionally diverse cell groups (such as other parts of the reticular formation, cranial nerve nuclei, dorsal column nuclei, colliculi, spinal cord, certain thalamic nuclei, and hypothalamus).

The Raphe Nuclei and the Locus Coeruleus: Common Features

The **raphe nuclei** (raphe, seam) together form a narrow, sagittally oriented plate of neurons in the midline of the medulla, pons, and mesencephalon (Figs. 26.1, 26.5, and 26.6). In many ways, these nuclei have similarities with the reticular formation proper and are often considered part of it. The **locus coeruleus** is small group of only about 15,000 strongly pigmented neurons located under the floor of the fourth ventricle (Fig. 26.7). It has clear borders except ventrally where it merges gradually with the adjoining reticular formation, which also contains many norepinephric neurons. This is one reason the locus coeruleus is often included in the reticular formation.

A characteristic feature of these two nuclei is that they contain only a small number of neurons, while their axons have extremely widespread ramifications, reaching virtually all parts of the brain and the spinal cord. Their synaptic actions are **modulatory**, although the effects vary among different targets due to different distribution of receptors. The raphe nuclei contain mainly **serotonergic** neurons, whereas the locus coeruleus neurons contain **norepinephrine** (cf. Chapter 5, under “Biogenic Amines”). Another special feature is that they send fibers directly to the cerebral cortex, that is, without synaptic interruption in the thalamus, as is typical of most cortical afferent pathways from lower levels. These features are shared with other cell groups, notably the dopaminergic neurons in the mesencephalon and the cholinergic cell groups in the pons and the basal forebrain (cf. Chapter 5, under “Modulatory Transmitter “Systems”).

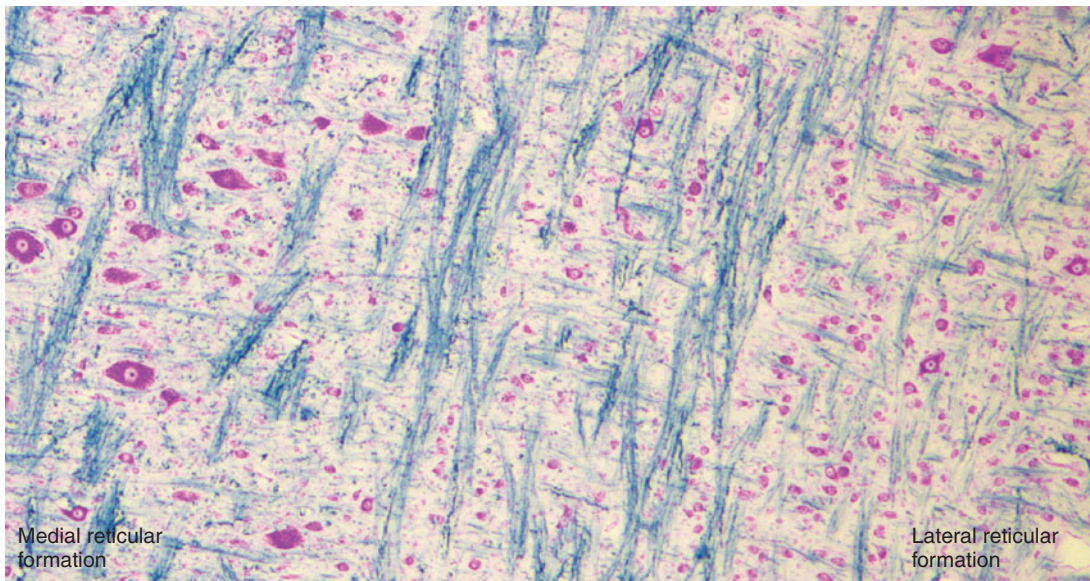


FIGURE 26.2 *The reticular formation.* Photomicrograph of section of the medulla, stained to show myelinated fibers (blue) and cell bodies (red). The cell bodies are distributed diffusely without clear nuclear borders and with small fiber bundles coursing in various directions.

The photomicrograph shows the transition between the large-celled medial division and the small-celled lateral division. Magnification, $\times 300$.

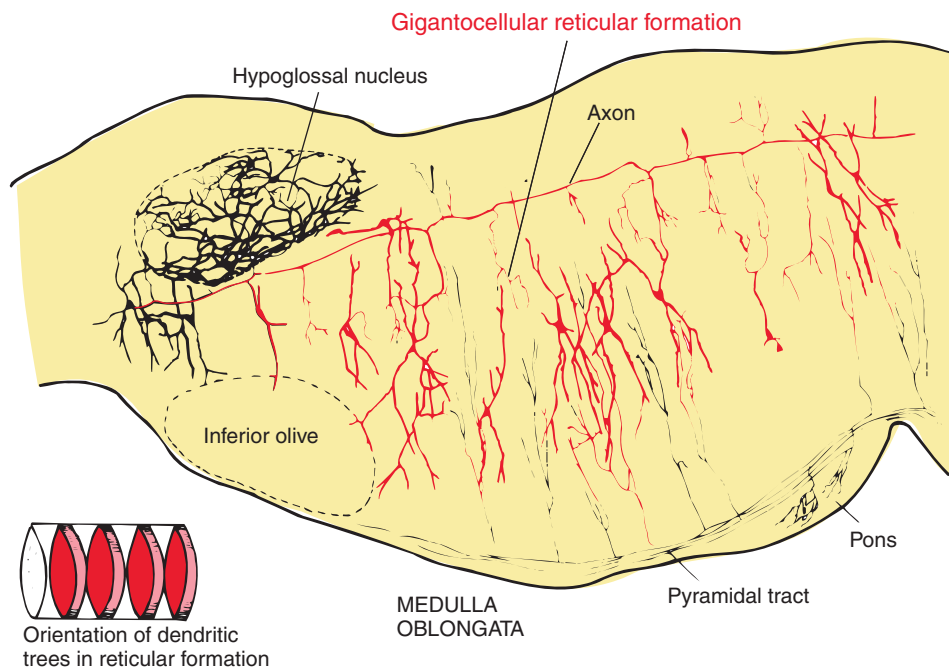


FIGURE 26.3 *Orientation of dendrites in the reticular formation.* Sagittal section through the medulla (rat). The long, straight dendrites are typical of the neurons of the reticular formation, in contrast to the neurons of a cranial nerve nucleus (the hypoglossal) and other specific

brain stem nuclei. A long axon with numerous collaterals extending ventrally in the transverse plane is also shown. Collaterals of the pyramidal tract fibers enter the reticular formation. (Modified from Scheibel and Scheibel 1958.)

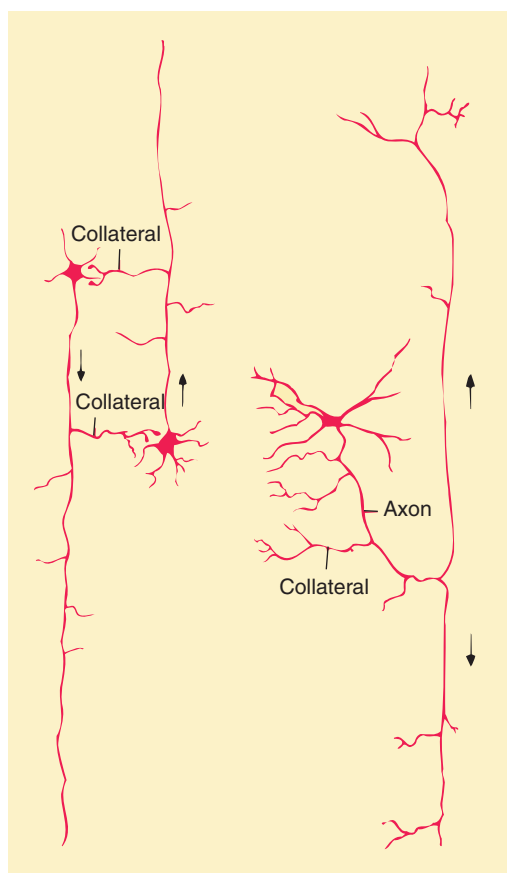


FIGURE 26.4 *Neurons in the medial part of the reticular formation with long ascending and descending axons. Right:* Example of a neuron with a bifurcating axon, with one ascending and one descending branch. Both branches give off collaterals as they course through the reticular formation. *Left:* Two neurons with ascending and descending axons, respectively. In this way, neurons with ascending axons can influence the activity of neurons with descending axons.

In other chapters, we discuss the effects of the monoamines serotonin and epinephrine on **sensory-information** processing (see Chapter 15, under “Central Control of Transmission from Nociceptors and Pain Sensation,” and Chapter 19, under “Modulation of Taste Cell Sensitivity”) and on **movements** (Chapter 22, under “Monoaminergic Pathways from the Brain Stem to the Spinal Cord”). We give examples of the importance of these monoamines for **emotions** and **motivation** in Chapter 5 (“Monoamine Oxidase, Serotonin Transporters, and Behavior”). They are of crucial importance for brain **plasticity** and **learning**, and influence cardiovascular control and other functions governed by the **autonomic nervous system**. Finally, serotonin and norepinephrine participate in regulation of **attention** and **sleep**. The serotonergic raphe neurons and the norepinephric locus coeruleus neurons have in common that they are active during wakefulness and less active during sleep (we return to this, under “Neuronal Groups and Transmitters Controlling Sleep”). They differ, however, in their response to sensory stimuli: only the norepinephric neurons increase their activity in relation to salient stimuli (like the dopaminergic neurons in the substantia nigra).

The Raphe Nuclei

Based on cytoarchitectonic and connectional differences, several raphe nuclei have been identified (Fig. 26.6) even though their borders are far from sharp. Together, the raphe nuclei receive **afferents** from many sources, such as the cerebral cortex, the hypothalamus, and other

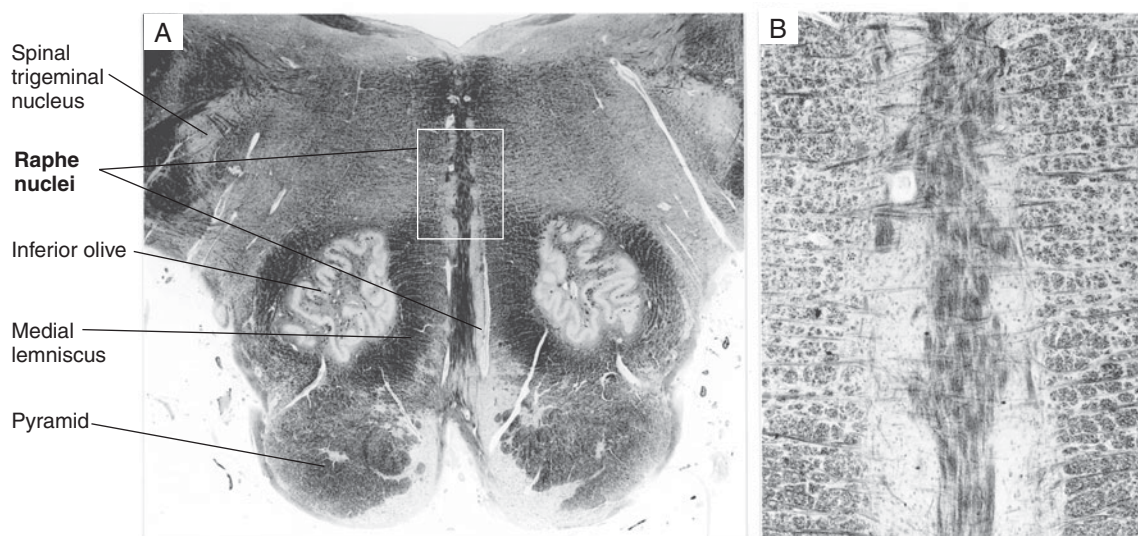


FIGURE 26.5 *The raphe nuclei. A:* Transverse section of the upper medulla, stained to visualize neuronal cell bodies and myelinated axons. The raphe nuclei are seen as narrow light stripes on each side

of the midline. *B:* Larger magnification from framed area in A. The cell bodies of raphe neurons are seen as small dots.

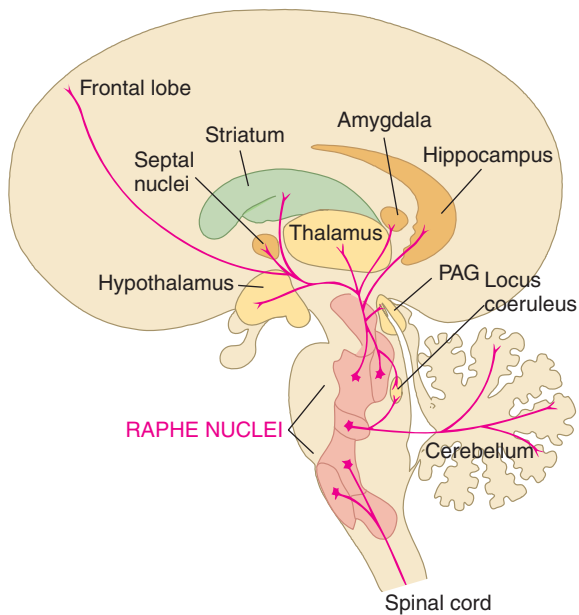


FIGURE 26.6 *The raphe nuclei and their efferent connections.* Schematic midsagittal section through the brain, showing the various subdivisions of the raphe nuclei and some main connections. The most rostral nuclei send their fibers rostrally to the thalamus, cortex, and other cell groups, whereas the caudal nuclei project to the spinal cord. Together, the raphe nuclei supply large parts of the central nervous system with serotonergic fibers.

parts of the reticular formation. In spite of the extensive distribution of the **efferent** fibers, the different raphe nuclei have largely different targets. Thus, the caudal raphe nuclei send their efferents to the spinal cord, whereas the rostral nuclei send fibers upstream (Fig. 26.6). The actions of **serotonin** depend on the kinds of receptor that are present postsynaptically in the target nuclei (see Chapter 5, under “Monoamine Receptors”). In addition, most raphe neurons contain **neuropeptides** (substance P and thyrotropin-releasing hormone, [TRH]), which presumably contribute to the synaptic effects of efferent fibers from the raphe nuclei. Serotonin also influences the **cerebral circulation**. A final peculiarity of the raphe nuclei is that they send fibers ending in close relation to the **ependymal cells** (which cover the interior aspect of the brain ventricles). Presumably, such fibers contribute to the regulation of transport processes through the ependyma.

Various kinds of behavioral changes have been reported after **lesions** of the raphe nuclei, such as aggression and increased motor activity (to destroy the raphe nuclei in isolation, however, is, virtually impossible).

The Locus Coeruleus

In spite of containing so few neurons, the **efferent** fibers of the locus coeruleus reach virtually all parts of the central nervous system. It sends, for example, direct fibers to the cerebral cortex, the hypothalamus, the basal ganglia,

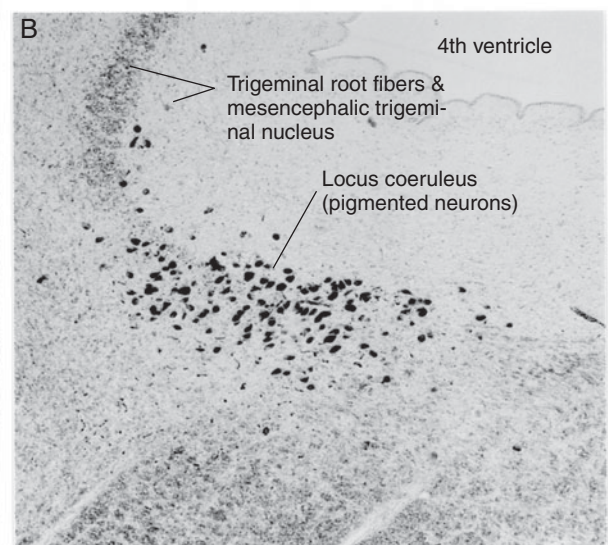
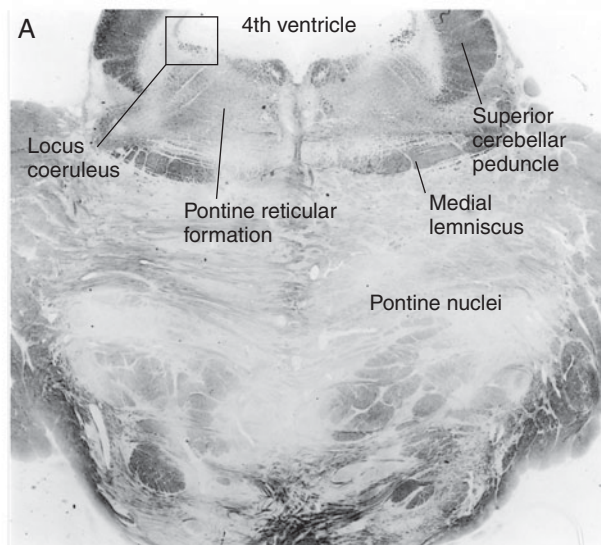


FIGURE 26.7 *The locus coeruleus.* **A:** Transverse section of the upper pons (approximately the same level as in Fig. 3.18). The locus coeruleus is positioned just below the floor of the fourth ventricle. **B:** Larger magnification from framed area in A. The norepinephrine-containing

neurons in the locus coeruleus are heavily pigmented. This nucleus and other norepinephric cell groups in the vicinity have neurons with highly branched axons that end rather diffusely in most parts of the central nervous system.

the hippocampus, and other limbic structures. Direct fibers also reach the spinal cord but only parts of the brain stem (especially sensory nuclei). The dense norepinephric innervation of the reticular formation and the motor nuclei must therefore originate from the scattered norepinephric neurons outside the locus coeruleus.

The **afferent** connections come mainly from a few regions. The locus coeruleus receives direct connections from the **cingulate gyrus** and the **orbitofrontal cortex** (in monkeys and presumably in humans as well). Subcortical afferents seem from recent studies to arise in the medullary part the reticular formation and in the amygdala. The medullary connections probably provide the locus coeruleus with integrated sensory information, while fibers from the amygdala might signal the emotional value of sensory information. **Single neurons** in the locus coeruleus respond preferentially to novel, “exciting” sensory stimuli. Because norepinephrine increases the response to specific stimuli and improves the signal-to-noise ratio of postsynaptic neurons, the locus coeruleus is believed to play a particular role in mediating **arousal** and **shifts of behavior**. One example is the facilitatory effect of norepinephrine on **spinal motoneurons** (see Chapter 22, under “Monoaminergic Pathway from the Brain Stem to the Spinal Cord”). Norepinephrine also has a special role in relation to synaptic **plasticity**. For example, it is especially concentrated in the cerebral cortex during sensitive periods of development.

Possible Tasks of the Raphe Nuclei and the Locus Coeruleus

The modulatory neurotransmitters—including serotonin and norepinephrine—influence virtually all brain functions. This is not surprising, considering the widely branching axons of the neurons containing these neurotransmitters. In view of the small and relatively homogeneous nuclei giving origin to the axons, it seems nevertheless possible that the multifarious actions are subject to a common aim or “plan.” However, in spite of several unifying theories, none has obtained general acceptance. We here present just a few possible unifying concepts. One of the theories concerning **serotonin** views the raphe nuclei as especially important for **homeostatic control** (another theory tries to collect all serotonin actions under the heading of motor control). One of the facts favoring homeostatic control as a common theme is that many raphe neurons are **chemosensitive** and measure the CO_2 level in the blood (thus indirectly monitoring the pH of the nervous tissue). Chemosensitive neurons in the medulla participate in control of respiration (normalization of CO_2). Actions of serotonin on spinal motoneurons can perhaps have something to do with the fact that muscular activity is a major source of CO_2 production. Serotonergic actions

on pain transmission might be viewed in the perspective of pain as part of a homeostatic response, as discussed in Chapter 15. Further, chemosensitive neurons in the rostral raphe nuclei can increase wakefulness and alter cerebral circulation. They also seem to mediate signals that evoke anxiety associated with high blood CO_2 levels (a single inhalation of air with 35% CO_2 provokes acute anxiety).

An overarching function of the **locus coeruleus** may be to increase **arousal** and **attention** in response to salient sensory information. However, the task may be more specific than this, according to a theory put forward by Aston-Jones and Cohen (2005). During wakefulness, the locus coeruleus neurons alternate between phasic and tonic firing. The phasic state is proposed to optimize ongoing actions (help to maintain the focus on the present task). Tonic activity, on the other hand, allows shift of attention away from the ongoing activity to enable exploratory behavior to select a new (and more rewarding) behavior. Instruction to **shift behavior** appears to reach the locus coeruleus from parts of the prefrontal cortex (the cingulate and the orbitofrontal cortex). The gyrus cinguli, for example, seems to monitor errors during execution, and helps decide whether actions produce the expected results.

The Efferent Connections of the Reticular Formation

The reticular formation sends fibers to (and thereby acts on) four main regions: the **thalamus**, the **spinal cord**, **brain stem nuclei**, and the **cerebellum**. The cell groups giving off ascending axons are located somewhat more caudally than those emitting descending axons (Fig. 26.8). By means of the numerous collaterals of both the ascending and descending fibers in the reticular formation (Figs. 26.3 and 26.4), the two kinds of cell groups can influence each other. Further, there are also many interneurons connecting different parts of the reticular formation. Thus, a close cooperation is possible between the parts of the reticular formation that act on the cerebral cortex and those that act on the spinal cord. Collaterals of ascending and descending axons mediate actions on brain stem nuclei.

Parts of the reticular formation form **premotor networks**. Chapter 25 deals with premotor networks in the pons and mesencephalon controlling and coordinating the activity of the eye muscle nuclei. Other premotor networks in the reticular formation control rhythmic movements such as locomotion (Chapter 22) and respiration. Further, premotor networks coordinate the activity of the widely separated motoneurons responsible for the cough reflex and the vomiting reflex.

The **descending** fibers run in the ventral part of the lateral funicle and in the ventral funicle (Fig. 26.9), and are related to **motor control**. Such reticulospinal fibers end primarily on interneurons, which, in turn, can

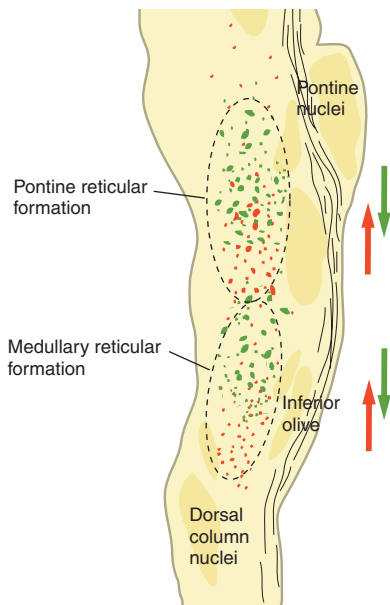


FIGURE 26.8 *Position of efferent reticular cell groups.* Drawing of sagittal section through the brain stem (cat). Neurons sending axons to higher levels (the thalamus) are red, while neurons with descending axons are green. The cell groups sending descending fibers are located somewhat more rostrally than the regions sending ascending fibers, providing opportunity for mutual influences by collaterals (as shown in Fig. 26.4).

influence motoneurons. The ventral reticulospinal tracts are discussed in Chapter 22 (“More About Reticulospinal Tracts”). The **ventral reticulospinal tracts** are both crossed and uncrossed and mediate both inhibitory and excitatory effects on spinal motoneurons. The reticulospinal neurons are further characterized by their axonal branching pattern, with collaterals given off at several levels of the spinal cord. Thus, each neuron can influence muscles in different parts of the body. As discussed in Chapter 22, the ventral reticulospinal tracts are of particular importance for **postural control**, for the **orientation of the head and body** toward external stimuli, and for **voluntary movements of proximal body parts**. Connections from the **superior colliculus**—that mediate sensory information to the reticular formation—are crucial for orienting movements toward novel stimuli. Combined anatomic and physiologic studies of single cells have shown the importance of a tecto-reticulospinal pathway for such movements. The **dorsal reticulospinal** fibers concern primarily **control of sensory information**. Many of these fibers are monoaminergic and arise partly in the raphe nuclei and adjoining parts of the reticular formation (see Fig. 15.3).

The **ascending fibers** from the reticular formation end in the **intralaminar thalamic nuclei** (Fig. 26.9), unlike the specific sensory tracts that end in the lateral thalamic nucleus. Some fibers also end in the **hypothalamus**. We discuss the functional significance of the ascending

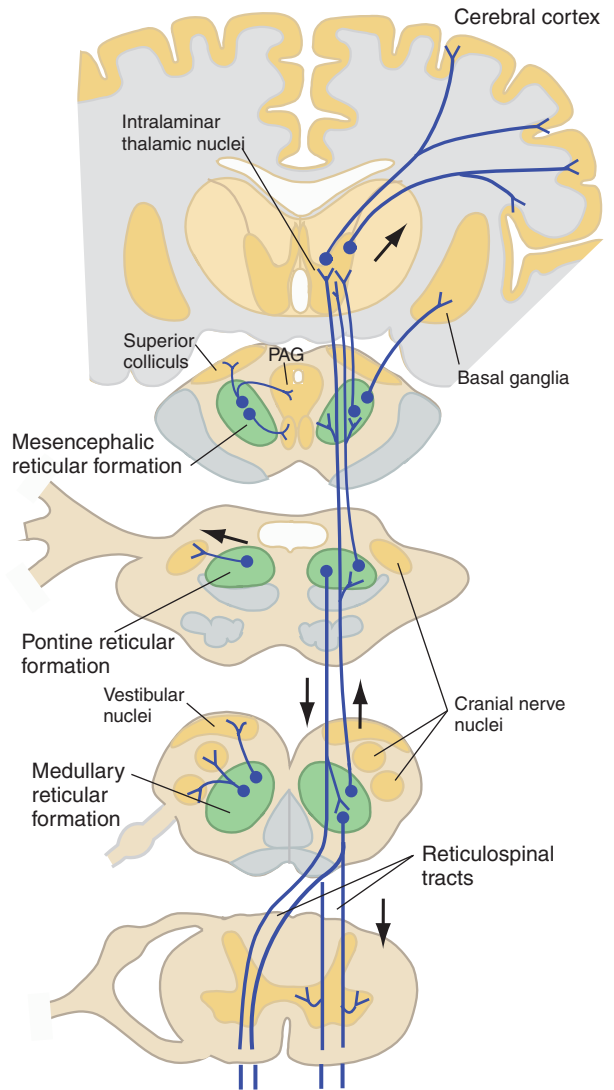


FIGURE 26.9 *Efferent connections of the reticular formation.* Various subdivisions of the reticular formation send fibers to higher levels, such as the thalamus, the basal ganglia, and the cerebral cortex. Descending fibers end in the spinal cord. In addition, the reticular formation sends fibers to the cranial nerve nuclei and other brain stem nuclei, such as PAG and the superior colliculus, and the vestibular nuclei. Most of the long connections are both crossed and uncrossed, although this is not shown in the figure. Numerous shorter fibers interconnecting subdivisions of the reticular formation are not shown.

reticular connections later in this chapter; suffice it to say here that they are of particular importance for the general level of activity of the cerebral cortex, which, in turn, concerns consciousness and attention.

The Reticular Formation Receives All Kinds of Sensory Information

Various cell groups send fibers to the reticular formation (Fig. 26.10). **Spinoreticular fibers** are discussed in Chapter 14. These fibers ascend in the ventral part of

the lateral funicle together with the spinothalamic tract, but diverge in the lower medulla. Among other destinations, the fibers end in the parts of the reticular formation that give off long ascending axons to the thalamus. This provides a **spino-reticulothalamic pathway** that is anatomically and functionally different from the major sensory pathways. Some of the spinoreticular fibers end in areas containing neurons that send axons back to the spinal cord, thus establishing **feedback** loops between the reticular formation and the cord.

In addition to spinal neurons that send their axons only to the reticular formation, many secondary sensory neurons send **collaterals** to the reticular formation. This concerns many of the fibers of the **spinothalamic tract**,

which, presumably, mediate **nociceptive** and **thermoceptive** signals to the reticular formation. Collaterals of ascending axons from the sensory (spinal) trigeminal nucleus supply the same kind of information from the face.² **Visceral sensory** signals reach the reticular formation by collaterals of ascending fibers from the **solitary nucleus** (which receives afferents from the vagus nerve, for example).

The **superior colliculus** sends fibers to parts of the reticular formation, as mentioned. These connections make it possible for **visual signals** to influence the reticular formation because the superior colliculus receives visual information directly from the retina and from the visual cortex. In addition, the superior colliculus receives somatosensory information from the cortex and integrates visual and somatosensory stimuli enabling orientation toward external stimuli.

Auditory signals reach the reticular formation by collaterals of ascending fibers in the auditory pathways. **Vestibular** signals come from the vestibular nuclei.

The preceding description of afferents indicates that signals from virtually all kinds of receptors can influence neurons of the reticular formation. This is verified by physiological experiments. Electrodes placed in the reticular formation can record potentials evoked by stimulation of receptors for light, sound, smell, and taste. Furthermore, stimulation of peripheral nerves carrying signals from cutaneous receptors and proprioceptors and of visceral nerves evokes activity. Whenever a receptor is stimulated, the signals reach not only the cortical areas important for the perception of the stimulus but also the reticular formation.

Afferents from Subcortical Nuclei and the Cerebral Cortex

Corticoreticular fibers arise mainly in the cortical areas that give origin to the pyramidal tract (Fig. 26.10). They end preponderantly in the regions of the reticular formation that send axons to the spinal cord. As discussed in Chapter 22, the **corticoreticulospinal pathway** is of special importance for the control of voluntary and automatic movements.

As discussed in Chapter 23, the **basal ganglia** (the substantia nigra) send efferents to the mesencephalic reticular formation (the locomotor region, or PPN). Fibers from the **hypothalamus** ending in parts of the reticular formation serve to coordinate the activity of different peripheral parts of the autonomic system. Limbic structures, notably the **amygdala**, also send fibers to the reticular formation. Such connections probably mediate emotional effects on autonomic and somatic motor

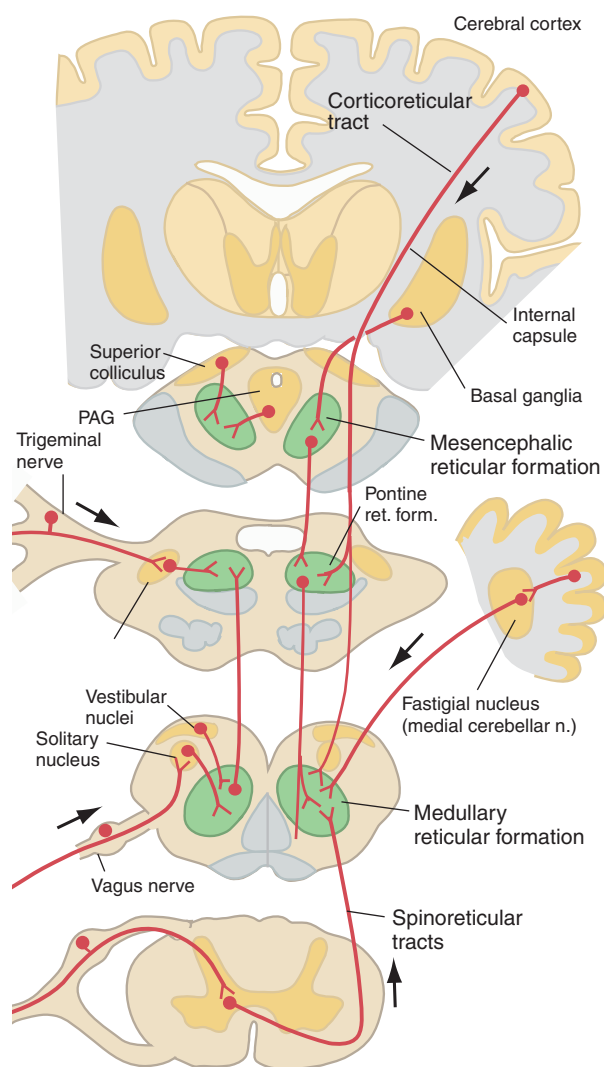


FIGURE 26.10 *Afferent connections of the reticular formation.* Several spinoreticular tracts mediate various kinds of sensory information. In addition, the reticular formation is influenced from sensory cranial nerve nuclei and other brain stem nuclei, such as the PAG, the superior colliculus, and the vestibular nuclei (not shown). The descending fibers from the cerebral cortex are links in indirect corticospinal pathways (cf. Fig. 22.1).

2 The medial lemniscus does not give off collaterals to the reticular formation. Information from low-threshold mechanoreceptors must therefore reach the reticular formation by means of spinoreticular neurons.

functions. The periaqueductal gray (PAG) is discussed in Chapter 15 in relation to suppression of nociceptive signals, but this mesencephalic complex of smaller nuclei has a wider specter of functions; for example, it helps initiate defensive reactions to external threats or other kinds of stress. This happens by way of its efferent connections to premotor networks in the reticular formation. The reticular formation receives afferents from the **cerebellum** (primarily the fastigial nucleus; see Fig. 24.6). This is an important pathway for the cerebellar influence on α and γ motoneurons. In addition, this pathway presumably mediates cerebellar influence of the autonomic nervous system. Thus, electric stimulation of the vermis can elicit changes in autonomic functions.

FUNCTIONS OF THE RETICULAR FORMATION

It follows from the preceding discussion of its efferent connections that the reticular formation can act on virtually all other parts of the CNS. We considered its effects on the spinal cord in Chapters 14 and 22, and we add more here on this point. In addition, we discuss the effects on the cerebral cortex, which are of particular importance. Parts of the reticular formation that are involved in the control of eye movements are discussed in Chapter 25, whereas the parts involved in the control of autonomic functions are described briefly in Chapter 28.

“The Activating System”

Electrical stimulation of parts of the reticular formation alters several functions mediated by the spinal cord, such as muscle tone, respiration, and blood pressure. In addition, the general activity of the cerebral cortex, which is closely related to the level of consciousness, can be altered by stimulation of the reticular formation. The **activating system of the brain stem** was therefore introduced as another name for the reticular formation. It should be emphasized, however, that these widespread effects are not served by the reticular formation as a whole but by relatively specific subregions. Further, stimulation of the reticular formation can also produce **inhibitory** effects on several processes. Therefore, the term “activating system” is not synonymous with “the reticular formation” but pertains to effects obtained from specific subregions only (to be discussed further later in this chapter).

There is continuous activity in the reticular formation maintained by a constant inflow of signals from various sources. The level of consciousness reflects the tonic activity in specific parts of the reticular formation. In particular, such activity is essential for the conscious awareness of sensory stimuli and for adequate behavioral responses to them. When a novel stimulus

catches our attention, this is mediated by the reticular formation. At the same time, premotor networks in the reticular formation produces the motor responses that ensure automatic orientation of the head and the body toward the source of the stimulus. The motor apparatus is mobilized, together with alterations of respiration and circulation.

Actions on Skeletal Muscles

Animal experiments during the 1940s led to the identification of two regions within the reticular formation that influence muscle tone. Stimulation of a region in the medulla that sends particularly strong connections to the spinal cord (Fig. 26.9) could inhibit stretch reflexes and movements induced by stimulation of the motor cortex. This region was therefore called the **inhibitory region** (Fig. 26.11). A region with opposite effects, the **facilitatory region**, was found rostral to the inhibitory one. It extends rostrally into the mesencephalon and is located somewhat more laterally than the inhibitory region. The distinction between inhibitory and facilitatory regions is less sharp than originally believed, however. In several places, single neurons with inhibitory or facilitatory actions on muscles are intermingled.

The actions in the cord concern not only α but also γ **motoneurons**, so that the reticular formation can control the sensitivity of the muscle spindles. The activity

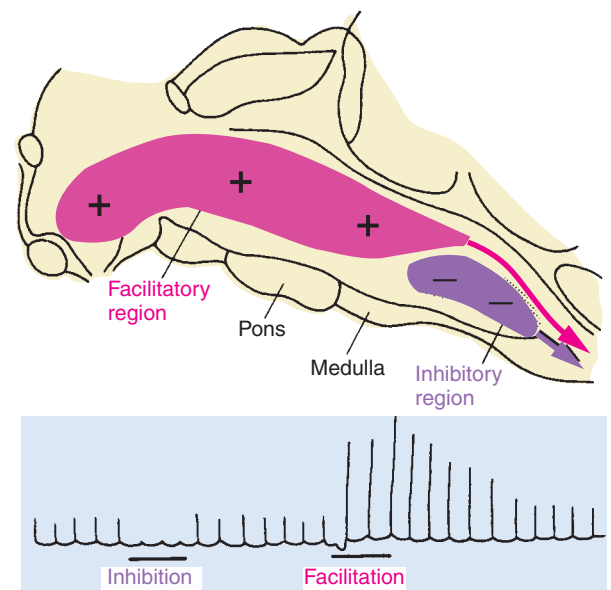


FIGURE 26.11 Facilitatory and inhibitory regions of the reticular formation. Schematic sagittal section through the brain stem (cat). The diagram at the bottom shows the amplitude of the patellar reflex (measured with EMG). In the period marked “Inhibition,” the inhibitory region was stimulated electrically and the reflex response is almost abolished. In the period marked “Facilitation,” the facilitatory region was stimulated and the patellar reflex response is markedly enhanced. (Modified from Kaada 1950.)

of the γ motoneurons is increased by stimulation of the facilitatory region and inhibited by the inhibitory region. The muscle tone can therefore be up- and down-regulated by altering the balance between the influences from the two regions. Since the cerebral cortex, the cerebellum, the basal ganglia, and other regions send fibers to the reticular formation, it follows that muscle tone can be influenced from these parts as well. It should be recalled, however, that cell groups other than the reticular formation also have direct access to the motoneurons, notably the vestibular nuclei with their strong facilitatory effects on the tone of muscles maintaining the upright position (see Chapter 21 for a discussion of muscle tone).

In addition to the rather diffuse effects on muscle tone obtained by electrical stimulation of the reticular formation, its role in the control of certain kinds of **voluntary** and **reflex movements** were discussed in Chapter 22. No principal difference exists, however, between the influence of motoneurons in eliciting a movement and in maintaining a (static) posture. Based on the present state of knowledge of the organization of the reticular formation, the emphasis is more on its control of specific motor tasks than on its diffuse effects on the muscle system as a whole. For example, certain subregions have a particular role in controlling the **rhythmic locomotor movements**, whereas other regions are devoted to controlling **eye movements**, and some are concerned primarily with **orienting movements** of the head and the body in response to optic and vestibular stimuli. In such functions, parts of the reticular formation—often consisting of extensive premotor networks—collect the relevant information about, for example, the position of the head and body, and ensure through their output signals the coordinated activity of all the muscles that are necessary to produce a proper response. Common to premotor networks is their control of the activity of large groups of muscles.

Effects on Respiration and Circulation

Microelectrode studies have shown that neurons with respiratory movement-related activity lie in several regions of the reticular formation, even though they are concentrated in the **ventrolateral medullary reticular formation**, which is often termed the **ventral respiratory group (VRG)**. This region contains many premotor neurons that control (monosynaptically and polysynaptically) the rhythmic activity of motoneurons innervating the diaphragm and other respiratory muscles. The **rhythm generator** itself consists most likely of a small neuronal network in the rostral part of the VRG, the so-called **pre-Bötzinger complex**. Thus, animal experiments indicate that this small region is both necessary and sufficient to elicit rhythmic respiratory movements (this network has properties in common with other

rhythm generators—e.g., those producing locomotor movements). Normally, however, a much wider network participates in respiratory control, including neurons in the **pontine reticular formation**. The VRG receives sensory signals from the thoracic cage and the **lungs** about the degree of expansion, and from **chemoreceptors** about blood pH and CO₂ content. Such information modulates the activity of the rhythm generator, without being necessary for the maintenance of breathing.

The respiratory rhythm generator is unstable shortly after birth in rats (but not in pigs and cats). A theory proposes that similar immaturity—so that the breathing rhythm is easily disturbed or abolished—lies behind **sudden infant death syndrome**.

Parts of the reticular formation also receive all necessary information for **cardiovascular control**. Thus, these regions control blood pressure, the blood volume distribution among the organs, the stroke volume, and the heart rate. Rather extensive networks in the **rostral ventrolateral medulla (RVLM)** are responsible for coordinating the necessary adjustments of vascular resistance and cardiac output. The effects are mediated by reticulospinal fibers acting on preganglionic sympathetic neurons and fibers to brain stem preganglionic parasympathetic neurons (with fibers passing in the vagus nerve).

The Reticular Formation and the Relationship between Mental and Bodily Processes

Variations in the activity of the reticular formation are reflected in virtually all aspects of the nervous processes and in the activity of the endocrine organs controlled by the hypothalamus. Such interactions may help explain the intimate correlation of mental and bodily processes. There is much evidence to suggest that our mental state influences the activity of parts of the reticular formation. The thought of a forthcoming, unpleasant event or the memory of an embarrassing or agonizing situation may suddenly make a drowsy person alert and tense (increased muscle tone), produce sweating, increase the heart rate, and so forth. Every doctor who routinely tests reflexes knows that apprehension and anxiousness is accompanied by increased reflex responses and increased muscular tone. An exaggerated patellar reflex at the start of a consultation becomes “normal” as the patient relaxes. Another everyday example is the difficulty in falling asleep when one is preoccupied with distressing thoughts.

All of the above-mentioned effects covary with altered activity in parts of the reticular formation that is closely linked with neuronal activity in the thalamus, cerebral cortex, limbic structures, and the hypothalamus. Thus, stimulation of certain cortical areas can increase the activity of reticular neurons followed by desynchronization of the EEG (unanesthetized animals become attentive).

These effects are mediated by direct and indirect connections from the cortex to the brain stem. There are also connections from the hypothalamus and limbic structures (such as the amygdala) to the reticular formation, which may be of particular importance because emotions are most effective in causing activation. That insomnia is related to increased activity in the reticular formation is suggested by the fact that most drugs used against **insomnia** reduce the activity of certain groups of reticular neurons. **General anesthesia** abolishes the transmission through central parts of the reticular formation (resulting in lack of activation of the EEG and unconsciousness), whereas the transmission in the specific sensory pathways is less affected.

We also know that **bodily processes influence our mental state**. For example, a treatment that leads to muscle relaxation usually also reduces mental tension. Again, the effects are probably mediated by a reduction of the activity of the reticular formation. At the same time, there are alterations of respiration, blood pressure, heart rate, and other autonomic functions such as sweat secretion, peristaltic movements of the bowel, and secretion of gastric juice. There is evidence that some of the autonomic expressions of anxiety (e.g. palpitations) may by themselves serve to maintain and increase the anxiety.

We return to the interactions between the mind and the body in Chapters 30 (under “Hypothalamus and Mental Functions”) and 32 (under “Amygdala and Emotions”).

CONSCIOUSNESS

What Is Consciousness?

“Consciousness” eludes attempts at precise scientific definitions. Yet, we all “know” what it is to be conscious. The American philosopher John Searle (2000, p. 559) gives this definition: “Consciousness consists of inner, qualitative, subjective states and processes of sentience or awareness. Consciousness, so defined, begins when we wake in the morning from a dreamless sleep and continues until we fall asleep again, die, go into a coma, or otherwise become ‘unconscious.’” A salient feature of consciousness is **unity** or wholeness: we perceive ourselves as a whole, and all the different bits and pieces of information analyzed by specialized brain systems are incorporated into a unitary experience. (Certain brain lesions, however, can dramatically change this; e.g., a stroke patient can deny that his paralyzed leg is a part of his body.) Further, consciousness cannot be separated from its **content**, including subjective sensations, moods, motives, inner images, and thoughts. Yet, we know that specific sensory information is not sufficient for conscious experiences to arise: the brain must be in a conscious state that depends on the integrity of the

ascending activating system, and probably specific activity patterns in the thalamocortical system. As put by Searle (2000, p. 574): “Only the already conscious subject can have visual experiences.”

Most people would accept that consciousness is a product of neuronal activity and, as such, an emergent property of the brain. It cannot be singled out as a separate “part”; rather, consciousness represents a particular state of the brain. This in itself does not explain the phenomenon, and consciousness cannot be fully understood by a **reductionistic** approach, studying in ever more detail the constituent parts of the brain. The brain—consisting of billions of neurons—produces something that is more than the sum of its parts, and even the most detailed knowledge of synaptic couplings, transmitters, and receptors does not suffice to explain consciousness and other mental phenomena. A “consciousness center” or “module” somewhere in the brain is incompatible with our present knowledge about the brain. That unconsciousness follows interruption of the ascending activating system in the mesencephalon does not imply that the locus of consciousness is somewhere in the mesencephalon (similarly, the fact that a car cannot run without a battery does not imply that the battery is the car).

Neurobiological Basis of Consciousness

For higher mental functions, the cerebral cortex is essential, and the cerebral cortex is certainly necessary for consciousness. It would nevertheless be misleading to state that the cerebral cortex alone “owns” the property of consciousness. In general, consciousness depends on coordinated activity in a network comprising the brain stem, the thalamus, and the cerebral cortex. Isolated activity in any one of the parts of the network is not sufficient. For example, activity in specific cortical areas responsible for (conscious) analysis of sensory information or control of cognitive processes cannot produce conscious experience on their own. We do not know the neurobiological mechanisms producing the conscious state.

In Chapter 16, we discuss whether specific parts of the cortex can be linked to conscious visual experiences (“How Are Data from Different Visual Areas Integrated?”). A “building block theory” of consciousness has some experimental support (e.g., from the study of persons with blindsight) but seems unable to explain important aspects. Much recent research has focused on what has been termed the “hypothesis of unified field consciousness.” This implies that consciousness might not be separable into different domains—as visual, auditory, somatosensory, emotional, and so forth. Much information now points to **synchronized activity**—binding vast assemblies of neurons in a coherent state—as a prerequisite for

conscious experience.³ In particular, synchronization within an extensive **frontoparietal network** may be characteristic of a conscious state. Indeed, conditions with unconsciousness (such as coma, deep sleep, general anesthesia, and the vegetative state) show reduced metabolic activity in the frontoparietal network.

Only few of our mental processes—in spite of their dependence on the integrity of the cerebral cortex—take place in “the searchlight of consciousness.” There is in fact much experimental evidence suggesting that of our “conscious” or “voluntary” behavior often reflects processes that do not come to full awareness.⁴

The Study of Consciousness: Electroencephalography

Direct recording from the surface of the cerebral cortex of animals shows that the patterns of electrical waves differ among parts of the hemispheres. In humans, too, direct recordings from the exposed cortex can be done during brain surgery (**electrocorticography**). The electrical waves are conducted through the skull and the soft tissues of the scalp and can therefore be recorded with electrodes placed on the skin of the head. With this method, **electroencephalography** (EEG), the electrical signals are dispersed and attenuated on their way through the skull and soft tissues, so that the origin of the signals within the hemisphere can be only roughly determined. Nevertheless, the method has been of great importance both for research and diagnostically since its introduction around 1930.

When analyzing the EEG, one can discern various waveforms and patterns. The **α waves** are relatively slow, with a frequency of 8 to 12/sec (Fig. 26.12). They occur typically in an awake person who is relaxed and resting with her eyes closed. When the person opens her eyes or starts to think about a mental task (Fig. 26.12), the EEG immediately changes to a pattern with more irregular waves with a higher frequency and lower amplitude. These are called **β waves**. The change of wave pattern

from α to β waves is called **desynchronization**, or **activation** of the EEG. During sleep, other wave forms and patterns occur that are typical of specific stages of sleep. Various neurological diseases can be diagnosed because they are associated with characteristic changes of the EEG. To some extent, the EEG can also help localize the disease process. An EEG is particularly important in the diagnosis of **epilepsy**.

The EEG varies considerably with **age**; most marked are the changes during the first couple of years. There are also individual variations. Further, the EEG pattern changes during **sleep** and during **general anesthesia**. **Hyperventilation**, which leads to a reduced carbon dioxide concentration in the blood, changes the excitability of cortical neurons, and this is reflected in the EEG. Finally, several **drugs** affect the EEG.

Thalamocortical Neurons Have Two States of Activity

We do not fully know how the various waveforms in the normal EEG arise, but it is clear that they depend in large measure on different activity states in thalamocortical neurons. Thalamocortical neurons (functioning as relay cells for transmission of sensory and other signals to the cortex) have unique membrane properties owing to the presence of a special kind of Ca^{2+} channel that opens only when the cell is hyperpolarized to a certain level. In addition, massive connections from the cerebral cortex to the thalamus—**corticothalamic** connections—play an important part in the rhythmic firing of thalamocortical neurons.

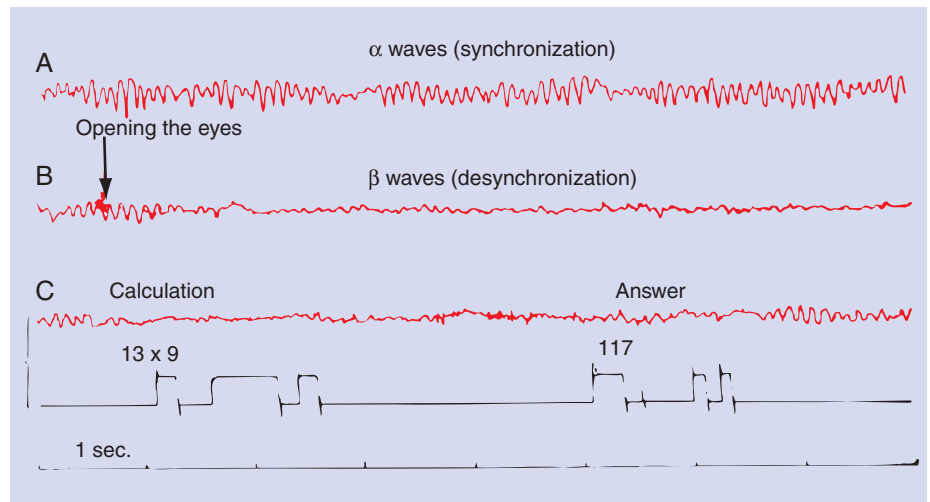
Thalamocortical neurons generate action potential in two different patterns or modes: they may fire in **bursts** (that is, they fire 2 to 8 action potentials with a high frequency followed by a pause), or they fire **single spikes** with a varying overall frequency. The bursting pattern is associated with synchronization of the EEG (in states of drowsiness and so-called slow-wave sleep), whereas the single-spike firing occurs together with desynchronization (during attentiveness and during rapid-eye-movement sleep). It appears that these different functional states of the thalamocortical (relay) neurons determine whether they transmit signals to the cortex. Only in the single-spike mode do the cells transmit to the cortex the information they receive from, for example, peripheral receptors.

Switching between bursting and single-spike firing can be produced by modulatory influences from the reticular formation (see later, “Multiple Pathways and Transmitters Are Responsible for Cortical Activation”). Further, the **reticular thalamic nucleus** (see Fig. 33.8)—consisting of GABAergic neurons with local actions in the thalamus—has a central role in the switching (cf. Chapter 33, under “Inhibition in the Thalamus: Interneurons and the Reticular Thalamic Nucleus”).

3 EEG recordings in humans during watching of ambiguous pictures support the relevance of synchronization for conscious experience. The picture could be interpreted either as meaningless or representing a face. When persons reported that they saw a face the EEG showed synchronization of widely separated cortical areas, whereas synchronization did not occur when they saw no meaning in the picture.

4 A woman suffering a stroke that damaged the midportion of the corpus callosum was asked to identify objects with the right and left hand (without vision). The damaged part of the corpus callosum connects the somatosensory areas. Therefore, in this case somatosensory information could not be transferred between the hemispheres. When holding a rubber in her right hand she was able to name it correctly. This is as expected, because the sensory information passes from the right hand to the left hemisphere that is responsible for speech in most people. When she held the rubber in her left hand, however, she did not notice anything or she did not know what it was. In this case, the information went to the “mute” right hemisphere. Nevertheless, she was afterwards able to pick out a rubber from a bag with her left hand when asked to find the same object she just held. Thus, the right hemisphere was able to identify the object, remember it, and use the “hidden” information in conscious behavior (Gazzaniga 1993).

FIGURE 26.12 *Electroencephalography (EEG).* Three traces are shown, all of them recorded over the occipital lobes. **A:** The person is in a relaxed state, with his eyes closed. There are α waves typical of synchronization. **B:** At the arrow, the person opens his eyes, and the α waves are replaced by faster, irregular waves with smaller amplitude, called β waves. The EEG is desynchronized. **C:** Desynchronization produced by solving a mental task. The person is asked to make a calculation (13×9). After the calculation is completed (at 117), the slower waves return.



A Prerequisite for Consciousness: Effects of the Reticular Formation on the Cerebral Cortex

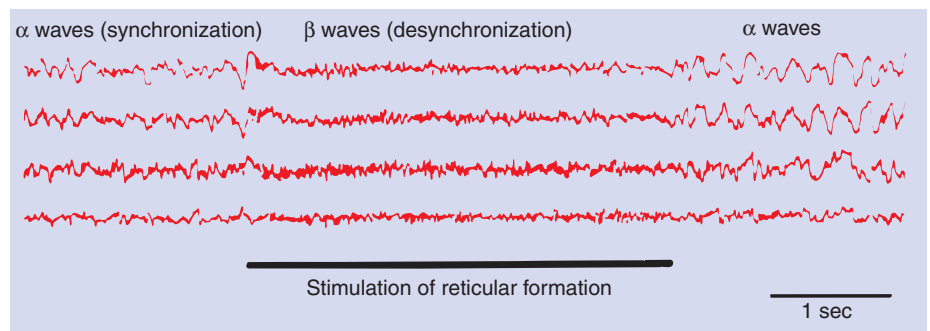
The effect of the reticular formation on the cerebral cortex was brought into focus by a paper published by the neurophysiologists G. Moruzzi (Italian) and H.W. Magoun (American) in 1949. Suddenly, new perspectives were opened on important aspects of brain functioning, especially consciousness, sleep, and wakefulness. Based on their animal experiments, Moruzzi and Magoun formulated a new concept: the **ascending activating system of the brain stem**. Later studies confirmed their hypothesis that (parts of) the reticular formation is necessary for maintenance of normal wakeful consciousness. The reticular formation determines the **level of consciousness**, with its many variations from tense alertness to drowsiness and sleep.

Electrical stimulation of the reticular formation in anesthetized animals produces certain changes of the electrical activity in the cortex, as recorded by EEG. Under anesthesia (without stimulation), relatively slow α waves dominate (Fig. 26.13). On electric stimulation of the reticular formation, the slow waves of the EEG are replaced by β waves with a faster and more irregular

rate and with lower amplitude. Together with this **desynchronization** of the EEG occur signs of increased attention and alertness—that is, an **arousal** is produced. Such an arousal can also be produced in unanesthetized animals with implanted electrodes. For example, a cat lying still with no obvious interest in its surroundings becomes suddenly alert and attentive when the reticular formation is stimulated with a high-frequency train of impulses. Marked reduction of the activity of the reticular formation is associated with unconsciousness, as shown experimentally in animals with selective lesions.

As mentioned, the reticular formation can have either activating or inhibiting effects on functions mediated by the spinal cord, depending on which region is stimulated (Fig. 26.11). Similar findings were made with regard to the ascending effects on the cerebral cortex, even though so far we have discussed only the activating effects. It appears that inhibition—that is, **synchronization** of the EEG—is evoked most easily from the caudal parts, whereas activation or desynchronization is evoked from the rostral parts (corresponding to the effects on skeletal muscles). We return to this below when discussing sleep.

FIGURE 26.13 *Electroencephalographic (EEG) recordings from the classic studies of Moruzzi and Magoun (1949).* The four tracings are recorded over different parts of the cerebral cortex of a cat. The thick line at the bottom shows the period of electrical stimulation of the reticular formation. The stimulation produces desynchronization of the EEG (activation), which returns to a synchronized pattern when the stimulation stops.



In humans with prolonged periods of unconsciousness after **head injuries**, there is often damage of the **mesencephalic reticular formation**. The lesion can be surprisingly small and yet produce deep unconsciousness. This fits with animal experiments showing that interruption of the ascending reticular connections in the mesencephalon produces loss of consciousness, in spite of normal conduction in the large sensory pathways (medial lemniscus, spinothalamic tract, and visual and auditory pathways).

Control of Sensory Information and Focusing of Attention

The main task of the ascending activating system is probably to **focus our attention** on certain stimuli or internal events, rather than to produce a diffuse awareness (if such a state is conceivable). To achieve this, it is necessary to prevent irrelevant stimuli from entering consciousness. Together with other mechanisms, inhibition from the reticular formation can ensure that, for example, we do not notice that someone is talking to us while we are absorbed in a book. In Chapters 14 and 15, we discussed how the central transmission of sensory signals is controlled from higher levels of the CNS, usually so that signals from certain kinds of receptor or parts of the body are inhibited. For example, descending connections to the spinal cord from the raphe nuclei and various (other) parts of the reticular formation suppress the central conduction of signals from nociceptors. Further, the central transmission of visual, auditory, and other sensory impulses is controlled by the reticular formation. Thus, even though it is not alone in this capacity, the reticular formation plays an important part in eliminating sensory signals that are considered irrelevant, so that our attention can be focused on salient signals.

Pathways and Transmitters Responsible for Cortical Activation

What are the pathways used by the reticular formation to influence consciousness, attention, and sleep? All of the major specific sensory pathways (the spinothalamic tract, the medial lemniscus, and the visual and auditory pathways) can be interrupted without affecting consciousness or the activation of the EEG produced by stimulation of the reticular formation. If these sensory pathways are left intact but the ascending connections of the reticular formation are interrupted by a cut in the mesencephalon, the animal becomes unconscious. Electrical stimulation of the reticular formation can no longer activate the EEG, even though stimulation of peripheral receptors evokes potentials in the cortical sensory regions. Thus, the sensory signals reach the cortex, but they are restricted to the sensory regions, and,

most importantly, they are unable to arouse the animal. Pathways other than the major sensory ones must therefore be responsible when the reticular formation activates the EEG over major parts of the cerebral hemisphere and produces behavioral changes indicating increased attention.

Connections from the reticular formation to the **intralaminar thalamic nuclei** are likely candidates. Thus, electrical stimulation of these nuclei can produce activation of the EEG similar to that seen after stimulation of the reticular formation itself. The intralaminar thalamic nuclei send widespread efferents to the cerebral cortex (cf. Chapter 33, under “The Intralaminar Thalamic Nuclei”). Therefore, a reticulothalamocortical pathway probably is important for the actions of the reticular formation on the cerebral cortex. Many of the reticulothalamic fibers to the intralaminar nuclei are **cholinergic** and come from a few small cell groups in the **dorsal part of the pons**.⁵ In addition to this indirect reticulothalamocortical pathway, there are direct projections to the cerebral cortex from **monoaminergic** cell groups in the brain stem—usually considered parts of the reticular formation—such as the **raphe nuclei** (serotonin), the **locus coeruleus** (norepinephrine), and the **ventral tegmental area** in the mesencephalon (dopamine). These nuclei also project to the thalamus, however. Stimulation of each of these cell groups can produce synchronization of the EEG, even though they behave differently in other respects. For example, their activities differ with regard to sleep, as discussed below. Norepinephric neurons increase their firing rate shortly before and during periods with cortical activation and focused attention. Similarly, **serotonergic** raphe neurons and **histaminergic** neurons in the hypothalamus (tuberomammillary nucleus) are more active during wakefulness than during sleep.⁶ As with the other modulatory inputs, activation of histaminergic fibers can bring thalamocortical neurons from a state of burst firing to single-spike firing.

Experiments in rats with selective elimination of various neurotransmitter systems suggest the following (very simplified) specializations with regard to tasks requiring focused attention: the **cholinergic** connections increase the precision of the performance, the **norepinephric** ones reduce the effect of distracting stimuli, the **dopaminergic** ones increase the speed of execution, and the **serotonergic** ones limit the frequency of impulsive response errors (see also “Possible Tasks of the

5 The largest pontine cholinergic cell groups are the **pedunculopontine nucleus** (PPN) and the **lateral dorsal tegmental nucleus** (LDT). The PPN consists of several subdivisions, however, and many neurons are glutamatergic; moreover, a part of the PPN has important connections with the basal ganglia and other parts of the brain stem.

6 Histamine-receptor antagonists, commonly used for allergy and motion sickness, block the activating effect of histamine, and this might explain why they have sleepiness as a side effect.

Raphe Nuclei and the Locus Coeruleus” earlier in this chapter).

To summarize, at least five cell groups, using as many transmitters, cooperate in the control of consciousness and attention. They exert their effects partly in the thalamus and partly in the cerebral cortex. Most likely, each of the cell groups and transmitters influence different aspects of wakefulness.

Coma, Vegetative State, and the Locked-in Syndrome

For **clinical purposes**, the definition of consciousness is practical: a condition of wakefulness in which the person responds appropriately to stimuli and, by his behavior, demonstrates awareness of himself and his surroundings. Patients with diseases or damage of the brain can exhibit states of consciousness ranging from full awareness to coma. A person in a **coma** appears to be sleeping, but cannot be awakened by any kinds of sensory stimulus. Purely reflex movements may be evoked, however (e.g., the withdrawal reflex). A coma may have several different causes but it is always a serious condition. The prognosis is poorer the longer a comatose state lasts. After acute head trauma, persistent unconsciousness suggests brain stem involvement, typically of the **mesencephalic reticular formation**.

After a few weeks, some comatose patients enter a **persistent vegetative state** in which they show some signs of wakefulness: they may open their eyes upon strong stimulation and after some time even spontaneously. They may briefly fix the gaze on a person or an object. They show no other signs of being aware of their surroundings, however, and they do not talk. Functional magnetic resonance imaging (fMRI) studies in such patients show activity patterns compatible with unconsciousness. For example, in 15 patients noxious stimuli activated expected parts of the brain stem, thalamus, and SI but not the rest of the “pain network,” including the insula, the cingulate gyrus, and the SII. Nevertheless, doubt exists as to whether some patients in the vegetative state can perceive and respond adequately to external events. Thus, a patient showed activation (measured via fMRI) of relevant motor networks when asked to imagine playing tennis. Further, speech areas were activated when the patient was presented with spoken sentences. It is not clear, however, whether these observations mean that the patient was in fact conscious, or whether it just shows how much of cerebral processing that can occur without entering consciousness.

On rare occasions, a patient may appear unconscious yet be fully awake. This occurs typically after brain stem infarctions that damage the ventral parts of the pons with the pyramidal tract and corticobulbar tracts on both sides. The sensory pathways located more dorsally are spared, as is also the ascending activating

system. This condition is called the **locked-in syndrome**. The patient is unable to move or talk but is otherwise fully conscious and mentally intact.

SLEEP

The necessity of sleep and its contribution to mental and physical health may seem self-evident. Yet, the more specific functions of sleep and its neurobiological basis are not fully understood. Indeed, control of sleep and its various phases has turned out to be very complex, involving many neuronal groups and neurotransmitters. Here we provide only a brief and simplified treatment of this topic.

Sleep Phases

Sleep consists of several phases that can be distinguished based on differences in the EEG. The transition from alertness to drowsiness changes the EEG in the direction of synchronization. When a subject is falling into a deep, quiet sleep, the α waves disappear altogether and are replaced by **irregular slow waves** with greater amplitude (**slow-wave sleep**). After an initial light **phase I**, the sleep becomes gradually deeper, until **phase IV**. To waken a person in sleep phase IV requires relatively strong stimuli, whereas only weak stimuli are necessary in phase I. **Phase V** is special because the EEG is desynchronized, and there are conjugated movements of the eyes, much like a person looking at moving objects. Because of these **rapid eye movements** (REM), this phase is called **REM sleep** or **paradoxical sleep**. The eye movements appear to relate to the content of the dream. Thus, patients suffering from neglect of the left visual hemifield (after damage to the right hemisphere) have conjugated eye movements during REM sleep only to the right side; that is, the visual field they attend to when awake. **Muscle tone** is generally reduced, with occasional muscular twitches, and there are changes of **blood pressure** and **heart rate**. Dreaming occurs—at least mainly—during REM sleep. The various phases of sleep follow each other with the same order throughout the night. Usually, the first REM phase occurs after about 1.5 hours of sleep and lasts for about 10 minutes. Thereafter the REM phases return at intervals of 1 to 2 hours, and become gradually longer up to about 30 minutes. When waking up (or being wakened) just after a REM phase, a person remembers the content of the dream vividly.

The cerebral cortex is as active during REM sleep as when awake, but is partly uncoupled from the **thalamus**, which in the awake state delivers a constant stream of information about the external world. The EEG pattern during sleep is produced by complex interactions between neuronal firing patterns in the thalamus and

the cortex. **Inhibitory interneurons** in both places have important roles in producing the synchronized firing of thalamocortical neurons characteristic of sleep.

Why Do We Sleep?

One hypothesis assigns a **homeostatic** function to sleep. The energy stores of the brain need rebuilding, the synaptic vesicles need refilling with neurotransmitters, and so forth. Another theory emphasizes that sleep is necessary for the sake of **plasticity**. Growth of neurons and establishment (and pruning) of synaptic contacts proceed presumably better when the neurons are not required to participate in task solving and production of goal-directed behavior. Useful connections need consolidation while other should be weakened or removed. The plasticity hypothesis would imply that sleep is important for **learning** and **memory**, a notion that has received experimental support. Thus, there is evidence that brain plasticity is increased during sleep. In agreement with this, even a brief nap immediately after a learning session can improve the subsequent performance.

Animal experiments suggest that the **consolidation** of newly learned material takes place predominantly in specific sleep phases. The earliest studies focused on the phases with dreaming (REM-sleep), while the phases with slow-wave sleep now seem to be equally important.⁷ For example, human EEG studies found increase in slow waves over cortical regions involved in a preceding learning session (e.g., over the posterior parietal cortex after a spatial-task training session). Further, slow-wave increases were associated with improved performance when tested afterwards. Consolidation of newly learned material is believed to depend on a dialog between the hippocampus and the cortex, and this process appears to go on during slow-wave sleep (but not during REM sleep), as witnessed by correlated firing in the hippocampus and the prefrontal cortex during slow-wave sleep.

Yawning

Most mammals yawn but we are not sure why. The hypothalamus, the basal ganglia, and the reticular formation have been implicated in the control of yawning (in addition, a number of cranial nerves and spinal nerves are responsible for the execution). One hypothesis proposes that yawning serves to increase alertness in situations where we are passive but need to pay attention (lectures may be an example). This assumption is based on the observation that yawning activates parts of the reticular formation and is immediately followed

by EEG desynchronization. The yawning movement evokes sensory inputs via (among others) the trigeminal nerve.

“**Contagious**” yawning—that is, yawning elicited by the sight or the sound of someone else yawning—may be an expression of our ability to imagine the state of mind of other people. It does not occur in animals that are unable to recognize their own mirror image, and not in children younger than the age of 2. An fMRI study showed change of activity in parts of the parietal cortex and the cingulate gyrus in contagious yawning (regions otherwise implicated in empathy). Regions thought to contain mirror neurons were not activated, however.

Pathologic yawning occurs in various diseases, for example in migraine. It also occurs as a side effect of **dopaminergic drugs** and **serotonin-reuptake inhibitors**. The importance of dopamine is supported by the observation that yawning is reduced in patients with Parkinson’s disease (with loss of dopamine). Increased yawning has been described as an early sign in infarctions of the upper pons, and in patients with **amyotrophic lateral sclerosis (ALS)**. In both cases, it may be due to loss of descending inhibitory control of brain stem “yawning centers.”

Neuronal Groups and Transmitters Controlling Sleep

As mentioned, the neural basis of sleep and its various phases is not fully understood and has turned out to be very complex. Several brain regions and neurotransmitters are important. Lesions of the **hypothalamus**, for example, can lead to increased or reduced amounts of sleep (see Chapter 30, under “Hypothalamus and Sleep”), but it is evident from animal experiments that the neuronal groups most directly involved in sleep control are located in the **brain stem**, especially in the **pons**. In agreement with this, fMRI in humans shows increased blood flow in the dorsal pons during REM sleep. Initially, after the discovery of the activating system, it was assumed that sleep was simply the result of reduced activity of the activating system—that is, a purely passive process. Further studies showed, however, that sleep could be induced by electrical stimulation of the lower parts of the reticular formation and that lesions of the same region prevented sleep (insomnia).

For **induction** of sleep, activity in **cholinergic neurons** in the **dorsolateral pons** is crucial (the pedunculopontine nucleus, and some smaller cell groups). These neurons project to the thalamus to influence the activity of the large sensory relay nuclei (the geniculate nuclei and the VPL). In addition, they project to neurons in the nearby reticular formation, which, in turn, project to cholinergic neurons in the hypothalamus and the basal nucleus (among other targets). The effects of acetylcholine in the thalamus are mediated by **muscarinic** receptors.

⁷ There is some evidence that REM sleep may be most important for consolidation of emotional and procedural memories, whereas slow-wave sleep is more involved in spatial and declarative memories.

The pontine cholinergic neurons fire in **bursts** (bursting neurons) ahead of the eye movements in REM sleep. Their close relation to REM sleep is further shown by the observation that microinjection of acetylcholine in the dorsolateral pons increases REM sleep for several days.

The roles of **serotonin** and **norepinephrine** in control of sleep are less clear, probably because they are difficult to study in isolation. Thus, cholinergic and monoaminergic neurons lie partly intermingled in the dorsolateral pons. It seems established, however, that the monoamines exert their main effect on sleep by modulating the activity of the pontine cholinergic neurons. Mostly, the monoamines **inhibit** the **cholinergic neurons**, thus increasing wakefulness. This conclusion is based on the fact that, among other things, the monoaminergic neurons fire during wakefulness and are “silent” during REM sleep.⁸ Further, microinjection of serotonin hyperpolarizes (inhibits) the bursting neurons in the dorsolateral pons. Finally, **drugs** that enhance the synaptic effect of serotonin (such as reuptake inhibitors) reduce REM sleep in humans.

Spinal **motoneurons** are strongly inhibited from the brain stem during REM sleep. This prevents the execution of movements that are part of the dream (small, “miniature” movements nevertheless occur during dreaming). The inhibition is partly initiated by the pontine cholinergic neurons, although the major control most likely is exerted by **norepinephric** neurons close to the locus coeruleus (the **subcoeruleus nucleus**). Reticulospinal neurons in the medial medullary reticular formation mediate the effects on the motoneurons. After destruction of descending norepinephrine-containing fibers, animals still have REM sleep but behave as if they were acting out their dreams with orienting movements, more complex exploratory behavior, and attack or flight.

Other neurotransmitters—GABA, dopamine, and several neuropeptides—are involved in control of sleep by influencing the activity of the pontine cholinergic neurons. In addition, glutamate and glycine act by way of pathways that are more indirect. Thus, many drugs acting on the brain would be expected to influence sleep and wakefulness.

Narcolepsy

Narcolepsy is a genetically linked disease with sudden, irresistible attacks of REM sleep (or components of REM sleep). Starting usually between the ages of 10 and 30, the disease affects equally often men and women. The patients experience increased sleepiness during

daytime, whereas night sleep is fragmented. The disease seems to be due to degeneration of a cell group in the lateral hypothalamus. The neurons of this group have widespread axonal ramifications and release neuropeptides called **hypocretins (orexins)**. Hypocretin-containing neurons appear especially to target modulatory cell groups, such as the locus coeruleus, raphe nuclei, dopaminergic neurons in the mesencephalon, and the tuberomammillary nucleus (histamine). Because hypocretins seem to be excitatory, they would increase arousal by increasing norepinephrine release from the locus coeruleus neurons, as confirmed in animal experiments.

Patients with narcolepsy also have increased concentration of **muscarinic** receptors within the brain stem (cf. the increased REM sleep caused by injection of acetylcholine in the dorsolateral pons, mentioned above). Further, there is evidence that monoamine metabolism is deficient. Both drugs that block muscarinic receptors and drugs that increase synaptic concentration of monoamines (such as amphetamine and tricyclic antidepressants) are reported to reduce the narcoleptic attacks. Possible connections between the loss of hypocretin neurons in the hypothalamus and the alterations of muscarinic receptors and monoamine metabolism in the brain stem are unknown.

Dreaming

Why we dream has been the subject of much speculation and many theories. REM sleep appears to occur in all mammals, even in the earliest species that developed about 140 million years ago.⁹ This fact alone suggests that REM sleep has an important function. Further support for its biological significance comes from other observations. Thus, the proportion of REM sleep increases after a period with REM-sleep deprivation, and it is much more difficult to prevent the occurrence of REM sleep than the other phases of sleep. **Newborn infants** have approximately 8 hours of REM sleep per day and a special sleep rhythm. It consists of periods of 50 to 60 minutes of sleep usually starting with REM sleep rather than slow-wave sleep. By the age of 2, REM sleep lasts only about 3 hours, and the sleep pattern is largely as it is in adults. The reason that REM sleep is so dominating during infancy (and before birth) is unknown. One possibility is that REM sleep is necessary for neuronal growth and development of connections, as discussed earlier. The dominating view today is that dreams contribute to the processing and consolidation of newly learned material, along with the integration of new experiences with older ones. Animal experiments indicate that storage of new material is

8 Raphe neurons increase their activity and serotonin release during wakefulness, and reduce their activity during sleep. Yet, destruction of the raphe nuclei or blockage of the serotonin synthesis produces insomnia. To reconcile these at least apparently conflicting observations, it has been postulated that serotonin released during waking gradually activates sleep-promoting neurons in the anterior hypothalamus (by acting at the gene level). This might serve as a homeostatic mechanism to initiate sleep after some time of wakefulness.

9 Whales (Cetacea) appear not to have REM sleep. In these mammals, the two hemispheres alternate to exhibit slow-wave sleep. Perhaps constant vigilance is a necessity for adaptation and survival.

reduced by prevention of REM sleep a certain period after the learning situation. During REM sleep in certain animals, a particular pattern of electrical activity—**theta rhythm**—occurs in the hippocampus. Since the hippocampus plays a crucial role in learning, this has been taken to support the relationship between REM sleep and learning.

The view that dreams are psychologically meaningful, relating in a disguised form to inner conflicts and life events, is the basis for their central place in the

psychoanalysis since Freud. Rather than emphasizing the role of dreams in learning, psychoanalytic tradition would stress their importance for the elaboration of inner conflicts that are not consciously accessible. These two possibilities might not be mutually exclusive, however. Perhaps a common purpose for all dreams is to help the individual develop **coping strategies**, as this requires the integration of new with old experiences, as well as ensuring that inner conflicts do not block learning and appropriate behavior.

27 | The Cranial Nerves

OVERVIEW

We usually count 12 cranial nerves, even though neither the first (the olfactory nerve) nor the second (the optic nerve) are true nerves. These two and, in addition, the cochlear nerve and the vestibular nerve are dealt with in Chapters 16 to 19.

A brief survey of the cranial nerves is given in Chapter 6. The prenatal development of the cranial nerve nuclei is treated in Chapter 9 (see under “Cranial Nerves and Visceral Arches”).

The cranial nerves connect the brain stem with structures in the head, neck, and in the thoracic and abdominal cavities. The cranial nerves are not as regularly built as the spinal nerves because some are purely motor, others are purely sensory, and some are mixed (like the spinal nerves). The cranial nerves contain four main fiber types. **Somatic efferent** fibers supply skeletal (striated) muscles, while **visceral efferent** fibers supply smooth muscles and glands and belong to the **parasympathetic** part of the autonomic nervous system. **Somatic afferent** fibers conduct sensory signals from the skin and mucous membranes of the face, from muscles and joints, and from the vestibular apparatus and the cochlea, while **visceral afferent** fibers bring sensory signals from the visceral organs. In early embryological development, the **cranial nerve nuclei** form longitudinal columns (cf. columnar arrangement of motoneurons in the cord), each column giving origin to only one of the four kinds of fiber. The columns are arranged so that, in general, motor cranial nuclei (somatic and visceral efferent) lie medially in the brain stem while sensory nuclei (somatic and visceral afferent) lie laterally. Later in development, the columns break up into discrete smaller nuclei but their mediolateral position and fiber composition remain the same (with some exceptions). The cranial nerve nuclei and the cranial nerves are links in various **brain stem reflexes** (e.g. the blink reflex and the vomiting reflex). The somatic motor nuclei receive innervation from the motor cortical areas, partly as collaterals of **pyramidal tract** fibers, partly as **corticobulbar** fibers destined only for the brain stem. Somatic sensory nuclei convey signals to the sensory areas of the cerebral cortex by joining the large **ascending sensory tracts** (e.g. the spinothalamic tract and the medial lemniscus).

A fair knowledge of the position of the various cranial nerve nuclei, the course of the nerves, and their main functions serve as a necessary basis for a topographic diagnosis of brain stem lesions.

GENERAL ORGANIZATION OF THE CRANIAL NERVES

Before dealing with specifics for each of the cranial nerves (Fig. 27.1), we discuss some features that are common to them all. Like the spinal nerves that connect the spinal cord with the body, the cranial nerves connect the brain stem with the peripheral organs. Several structural features are shared by the spinal cord and the brain stem, and thus also by the spinal and cranial nerves. Nevertheless, the brain stem is less regularly built and more complex in its organization than the spinal cord and the cranial nerves are not as schematic in their composition as the spinal nerves. Most of the cranial nerves, for example, lack a distinct ventral (motor) root and a dorsal (sensory) root. Some of the cranial nerves are purely sensory, others are purely motor, and others are mixed. Like the spinal nerves, several of the cranial nerves contain autonomic (preganglionic) fibers supplying smooth muscles and glands. Finally, some also contain afferent fibers from visceral organs.

The Cranial Nerves Can Contain Four Different Kinds of Nerve Fibers

The cranial nerves can contain the following kinds of fibers:

1. **Somatic efferent** fibers innervating skeletal (striated) muscles
2. **Visceral efferent** fibers supplying smooth muscles and glands and belonging to the parasympathetic part of the autonomic nervous system
3. **Somatic afferent** fibers with sensory signals from the skin and mucous membranes of the face, from muscles and joints, and from the vestibular apparatus and the cochlea
4. **Visceral afferent** fibers with sensory signals from the visceral organs

The **efferent fibers** of the cranial nerves have their cell bodies in brain stem nuclei corresponding to the columns of spinal motoneurons (see Fig. 21.3) and the intermediolateral cell column of the cord. We use the terms **somatic efferent** and **visceral efferent** cranial nerve nuclei of these cell groups (Figs. 27.2 and 27.3). The afferent fibers have their cell bodies in **ganglia** close to the brain stem, corresponding to the spinal ganglia. The central process of the ganglion cells enters the brain stem and ends on neurons in nuclei corresponding to

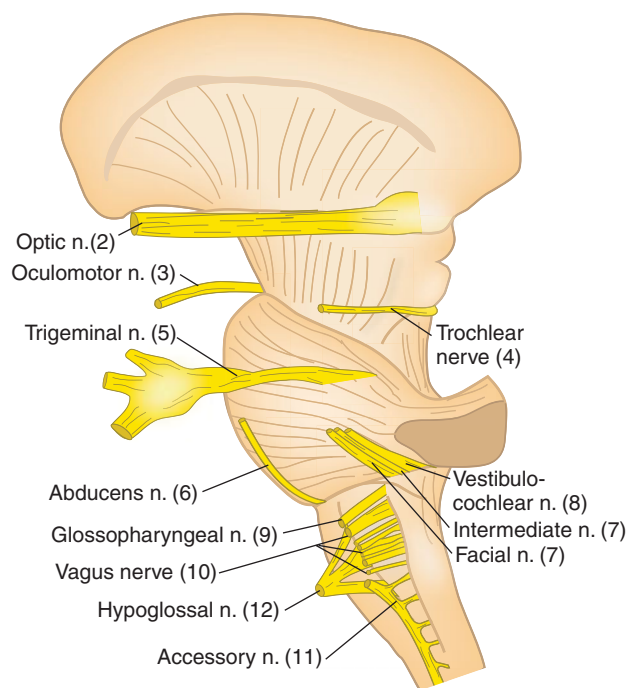


FIGURE 27.1 The brain stem and the cranial nerves, as viewed from the left side. See also Figs. 6.14 and 6.15.

the spinal dorsal horn and the dorsal column nuclei (Figs. 27.3 and 27.4). We use the terms **somatic afferent** and **visceral afferent** cranial nerve nuclei for such groups. From such afferent (sensory) nuclei, signals are conducted centrally to the thalamus and the cortex, and via brain stem interneurons to the somatic and visceral efferent (motor) cranial nerve nuclei (as links of brain stem reflex arcs; Fig. 27.4).

A Broad View on the Position of the Cranial Nerve Nuclei

In early embryonic life, all neurons giving origin to a particular kind of fiber lie together in one longitudinal column in the brain stem. During further development, these columns break up into smaller groups, and some of them move away from the original column. Nevertheless, the tendency for cranial nerve nuclei of the same category to form columns can be recognized also in the adult brain (Fig. 27.2). The demonstration of this regular pattern has been of value in understanding the functions of the cranial nerves. Further, knowledge of this pattern aids us in learning about the cranial nerves

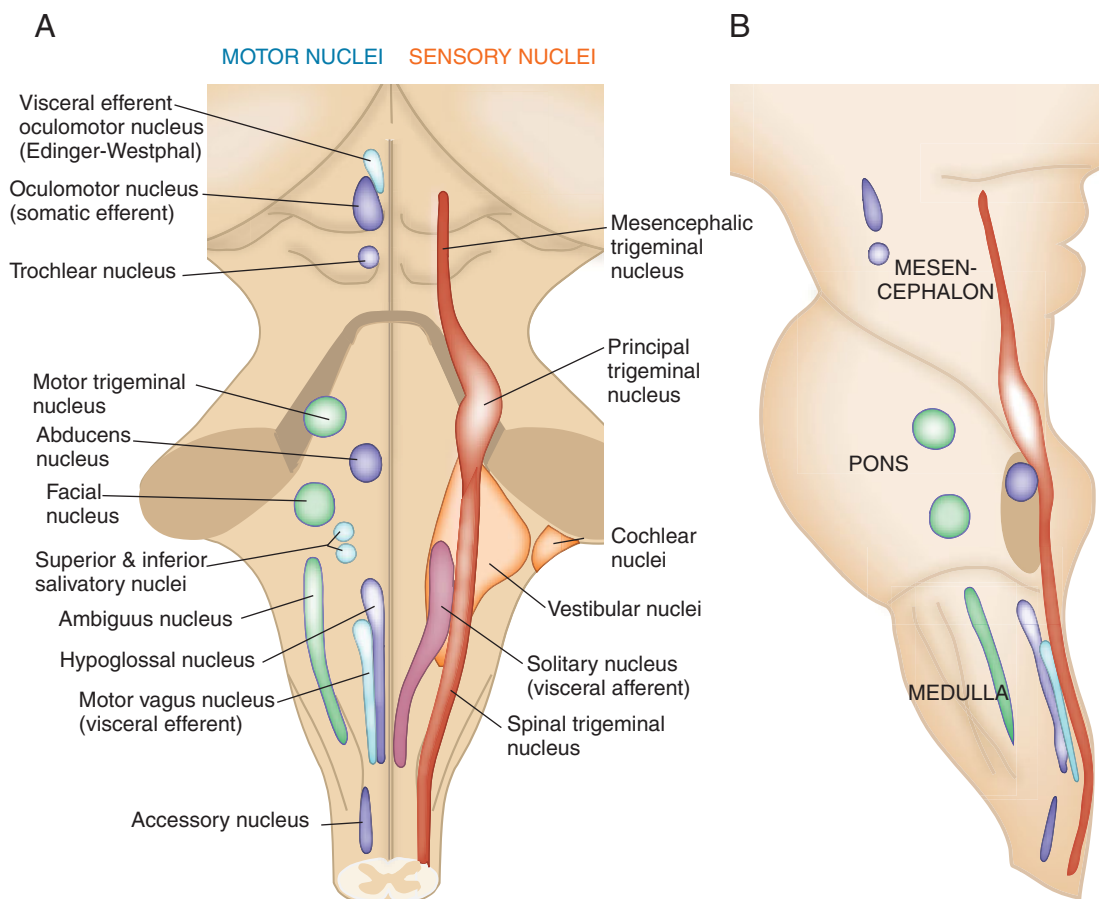


FIGURE 27.2 Position of the columns of the cranial nerve nuclei. A: Schematic of the brain stem, as viewed from the dorsal side. The nuclei belonging to the same kind (e.g., somatic efferent, visceral afferent)

are shown in the same color. They form more or less continuous columns. B: The position in the brain stem of some nuclei as seen from the left side. The positions of the various nuclei are only approximate.

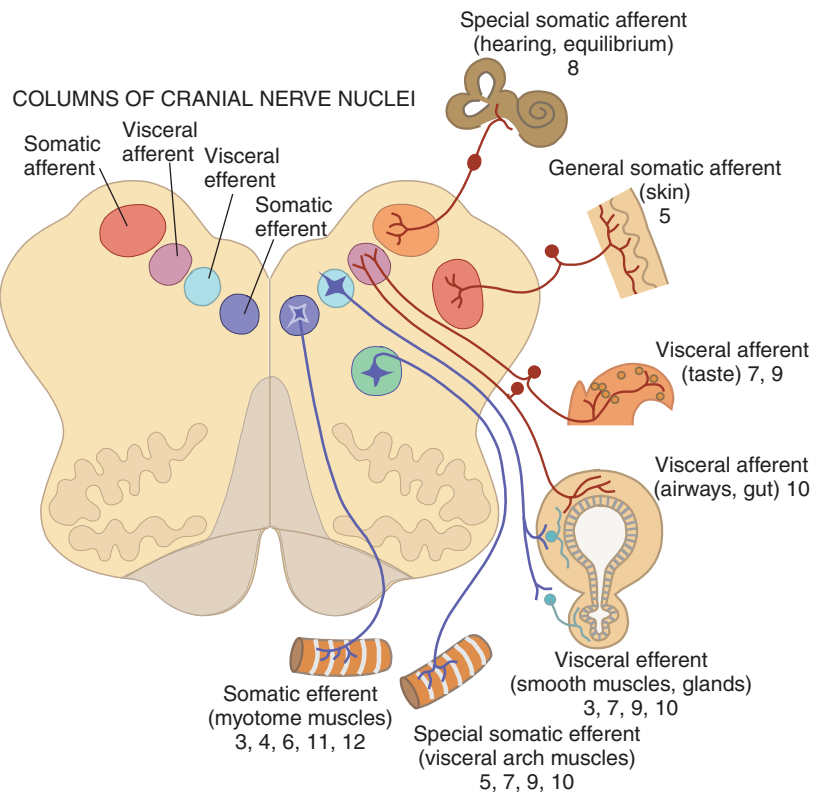


FIGURE 27.3 Position of the columns of the cranial nerve nuclei. Schematic cross section of the medulla (cf. Figs. 27.2 and 27.5). As a rule, the motor nuclei are located medial to the sensory nuclei, as shown schematically on the left side. The right half shows how the various kinds of fibers are distributed among the cranial nerves. Note that one kind of fiber usually distributes to several cranial nerves. Some of the peripheral organs innervated by the nerves are also shown to the right.

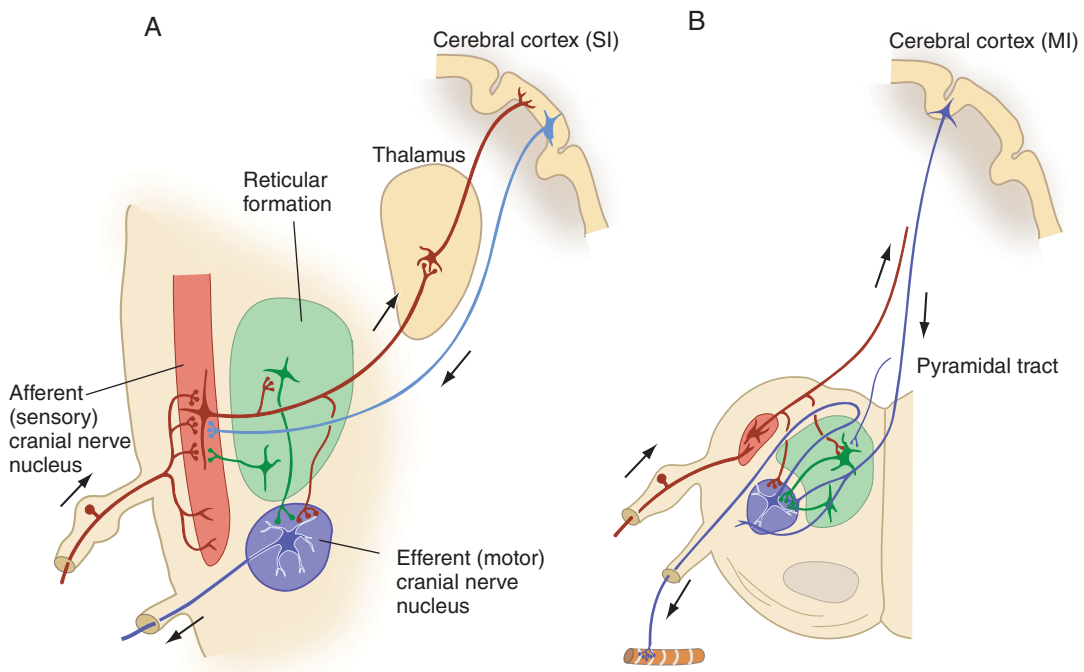


FIGURE 27.4 Main features of the organization of the cranial nerve nuclei. **A:** Sensory nucleus (e.g., the trigeminal nucleus). The efferent fibers of the nucleus ascend to the thalamus of the opposite side, and from there the next neuron projects to the cerebral cortex. The fibers destined for the thalamus give off collaterals on their course through the reticular formation and the motor cranial nerve nuclei. Thus, a

reflex arc with reflex center in the brain stem is established. Descending fibers from the cerebral cortex influences the sensory nucleus. **B:** Motor nucleus (e.g., the facial nucleus) sending its efferent fibers out of the brain stem to striated muscles. The neurons of the nucleus are influenced by the cerebral cortex, by the reticular formation, and by collaterals of the ascending fibers from the sensory nuclei.

and their nuclei. It is helpful at the outset to remember the following general rule, evident from Figs. 27.2 and 27.3: the efferent (motor) nerve nuclei lie medially in the brain stem, whereas the afferent (sensory) nuclei are located laterally. Further, in most cases the nuclei are located at about the same rostrocaudal level as their nerves leave the brain stem. In clinical neurology it is important to know the approximate mediolateral and rostrocaudal level of each nucleus (Fig. 27.5).

Further on the Position of the Cranial Nerve Nuclei

The **somatic efferent** nuclei are in early embryonic life all arranged in a column close to the midline, but later some move away in a ventrolateral direction (Figs. 27.3 and 27.5). The nuclei remaining in the medialmost column are termed **general somatic efferent** and comprise (from caudal to rostral) the **nucleus of the accessory nerve** (11), the **nucleus of the hypoglossal nerve** (12), the **nucleus of the abducens nerve** (6), and the **nucleus of the oculomotor nerve** (3). (In the following, for practical reasons we use abbreviated names, such as the accessory nucleus and the hypoglossal nucleus.) All of these nuclei innervate **myotome muscles**—the muscles that are developed from the segmentally arranged somites of early embryonic life. The somatic efferent nuclei that have moved away from the medial column all innervate **branchial muscles**—the striated muscles developed from the branchial (visceral) arches (facial and masticatory muscles, and muscles of the pharynx and larynx). We call these nuclei **special somatic efferent** (although they most commonly are termed “special visceral efferent”).¹ This group comprises the **ambiguus nucleus** (9, 10), the **facial nucleus** (7), and the **motor trigeminal nucleus** (5).

The **visceral efferent** (parasympathetic) column of cranial nerve nuclei is located immediately lateral to the somatic efferent column (Figs. 27.2, 27.3, and 27.5) and comprises the (dorsal) **motor nucleus of the vagus** (10), the small **inferior** and **superior salivatory nuclei** (9, 7), and the **parasympathetic oculomotor nucleus of Edinger-Westphal** (3). The **visceral afferent fibers** all end in one long nucleus, the **solitary nucleus**, which is located lateral to the visceral afferent column (Figs. 27.2, 27.3, and 27.5).

¹ Both anatomically and functionally, however, “the special visceral efferent neurons” are more similar to the somatic efferents. First, branchial (visceral) arch striated muscles are among those subject to the most precise voluntary control (the mimetic muscles of the face and the muscles of the larynx) and should therefore not be mixed up with visceral (smooth) muscles. Second, the neurons are structurally like α motoneurons; that is, they are larger than the preganglionic, parasympathetic neurons. In addition, the visceral arch neurons contain the neuropeptide CGRP that occurs in spinal motoneurons but not in preganglionic parasympathetic neurons. Therefore, we find it preferable to use the term **special somatic efferent**. Otherwise, it is of no great importance which terms are used to group the neurons of the cranial nerves; the important thing is to know where the cell bodies of the various cranial nerves are located, the course of their fibers, and their functions.

Most laterally, we find the **somatic afferent nuclei**. This group comprises the **sensory trigeminal nucleus** (5), the **vestibular nuclei** (8), and the **cochlear nuclei** (8). The sensory trigeminal nucleus consists of three functionally different parts (the spinal, the principal, and the mesencephalic nuclei). Together, the three parts form one continuous column, which extends from the upper cervical segments of the cord into the mesencephalon (Fig. 27.2).

The fibers of the vestibulocochlear nerve are often classified as **special somatic afferent** because they originate from special sense organs; the trigeminal fibers are then termed **general somatic afferent**.

Figure 27.4 and the discussion here show that fibers of one kind all come from one of the columns of nuclei only, even though the fibers peripherally may follow several of the cranial nerves. Thus, all (general) somatic afferent fibers end in the sensory trigeminal nucleus, whereas the fibers peripherally follow not only the trigeminal nerve but also the glossopharyngeal and the vagus nerves.

Brain Stem Reflexes

Like the spinal nerves that are links in spinal reflex arcs, the cranial nerves constitute afferent and efferent links of reflex arcs with reflex centers in the brain stem (Fig. 27.4). Some brain stem reflexes are simple, such as the monosynaptic stretch reflex—the **masseter reflex**—that can be elicited of the masticatory muscles. Often, however, the reflex centers are more complex, comprising neurons at several levels of the brain stem (for some, even at the cortical level). Thus, the afferent fibers may enter the brain stem at one level, whereas the efferent fibers leave at another. One example is the **corneal reflex** (touching of the cornea elicits an eye wink), in which the afferent fibers of the trigeminal nerve, entering at the midpontine level, descend in the brain stem and form synapses in the lower medulla (the spinal trigeminal nucleus). From the medulla, the signals travel by interneurons to the facial nucleus on both sides, located in the lower pons. The reflex center is in this case rather extensive; consequently, lesions at various levels of the brain stem may produce a weakened or abolished corneal reflex. Depending on the location of the lesion, however, the change of the corneal reflex will be accompanied by various other symptoms, which helps in determining the exact site of the lesion.

Several other brain stem reflexes are treated below in conjunction with the cranial nerves that mediate them.

The Cranial Nerve Nuclei Are Connected with Central Sensory and Motor Tracts

As mentioned, the cranial nerves and their nuclei are organized in accordance with the same general rules as

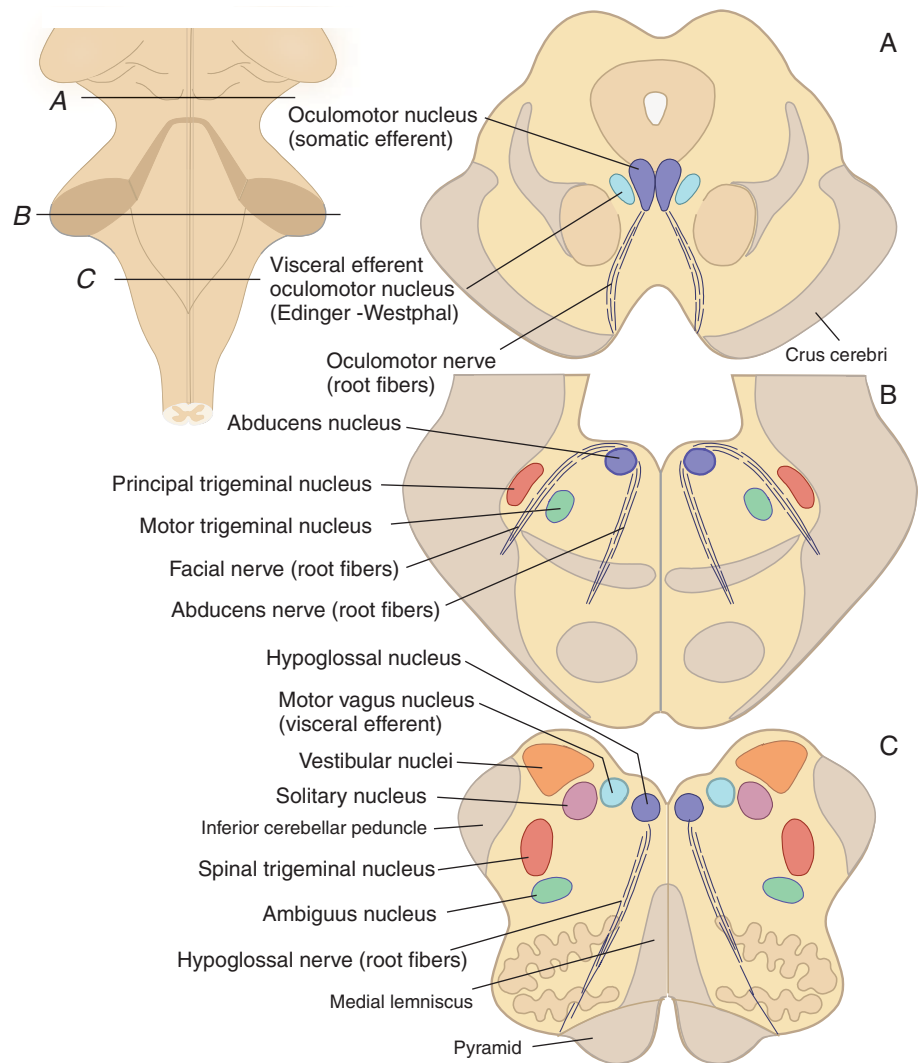


FIGURE 27.5 Position of the cranial nerve nuclei at three levels of the brain stem. A: Mesencephalon. B: Pons. C: Medulla oblongata. Compare with Fig. 27.2.

the spinal nerves (with some exceptions). This means that the cranial nerves are the first links in sensory pathways corresponding to the **dorsal column—medial lemniscus system** and the **spinothalamic pathway** (Fig. 27.4; see also Figs. 14.2 and 14.4). Further, as the nuclei involved in the sensory pathways conducting from the spinal cord, those of the brain stem are subjected to **descending control** of the sensory transmission. This concerns influences from parts of the reticular formation and from the cerebral cortex.

Several of the somatic efferent (motor) cranial nerve nuclei are influenced by the **pyramidal tract**—that is, by fibers forming the **corticobulbar tract** (Fig. 27.4; see also Fig. 22.1). An important difference between the corticospinal and corticobulbar fibers is that several of the cranial nerve nuclei receive both crossed and uncrossed fibers. Unilateral damage to the descending fibers (e.g., in the internal capsule) produces clear-cut pareses only in some of the muscle groups innervated

by the cranial nerves (most marked in the mimetic muscles).²

Examination of the cranial nerves is of great importance in clinical neurology because it can provide exact information about the site of a disease process. A prerequisite is that the examiner has reasonably precise knowledge of where the cranial nerves exit from the brain stem (Fig. 27.1; see also Fig. 6.16) and the position of their nuclei both rostrocaudally and mediolaterally (Figs. 27.2 and 27.5). Further, the functions of the various nerves must be known in sufficient detail as a basis for the necessary tests. On the basis of such knowledge, together with knowledge of the positions of the long motor and sensory tracts passing through the brain

2 The accessory, the hypoglossal, and the part of the facial nucleus supplying the lower part of the face—and often also the motor trigeminal nucleus—receive only crossed fibers, as judged from clinical observations (Monrad-Krohn 1954).

stem, the clinician can make a fairly precise topographical diagnosis in most cases (Fig. 27.6; see also Fig. 14.3).

Figures 6.16–6.18 and 6.20 show the location of the cranial nerve nuclei in cross sections of the brain stem.

Brain Stem Lesions Can Produce Symptoms from Several Cranial Nerves and Long Tracts

Many disease processes can cause brain stem lesions. Often, brain stem symptoms are caused by diseases that produce symptoms also from other parts of the central nervous system (CNS; e.g., multiple sclerosis, amyotrophic lateral sclerosis, metastatic brain disease). Lesions limited to the brain stem are most often due to vascular occlusions of arterial branches causing infarctions (see Figs. 8.3–8.5). Any branch can be affected (although some more often than others), and the resulting symptoms depend on which structures are supplied by the occluded branch. There are, however, large **individual variations** regarding the area supplied by a particular artery, and between the right and left side in the same individual. Thus, symptoms will vary after occlusion of a particular artery. Memorizing detailed lists of symptoms typical for each arterial branch has therefore limited value. A fairly precise localization of the lesion can usually be made on the basis of a thorough clinical examination of the cranial nerves and the long ascending and descending tracts (the dorsal column–medial lemniscus, the spinothalamic tract, and the pyramidal tract).

Figure 27.6 gives an overview of the structures that can be damaged by lesions in the **medulla**. Most common are lateral infarctions in the area supplied by the

posterior inferior cerebellar artery (see Fig. 8.3), even though the occlusion often sits in the vertebral artery. The German neurologist Adolf Wallenberg described the clinical picture of such infarctions in 1895, and it has since been known as **Wallenberg's syndrome**. Typical cases present with dizziness and vertigo (vestibular nuclei), gait ataxia (spinocerebellar and olivocerebellar tracts in the inferior cerebellar peduncle), difficulties with swallowing and hoarseness (ambiguous nucleus and root fibers of the vagus and glossopharyngeal nerves). A characteristic symptom constellation is reduced pain and temperature sensation in the face on the side of the lesion and in the body of the opposite side (face: spinal trigeminal nucleus and descending trigeminal root fibers; body: spinothalamic tract). Most cases also present **Horner's syndrome** (loss of sympathetic innervation of the face on same side as the lesion, see Fig. 28.10) due to interruption of descending fibers to the preganglionic sympathetic neurons in the cord. Individual cases may present only some of these symptoms or other symptoms in addition, depending on the exact site and extension of the lesion.

THE HYPOGLOSSAL NERVE

The **twelfth cranial nerve**, the hypoglossal nerve (Fig. 27.1), is the motor nerve of the tongue. It is composed of only **somatic efferent fibers**. The fibers come from the **hypoglossal nucleus**, which forms a slender, longitudinal column close to the midline in the medulla (Figs. 27.2 and 27.5). The nucleus produces an elongated elevation

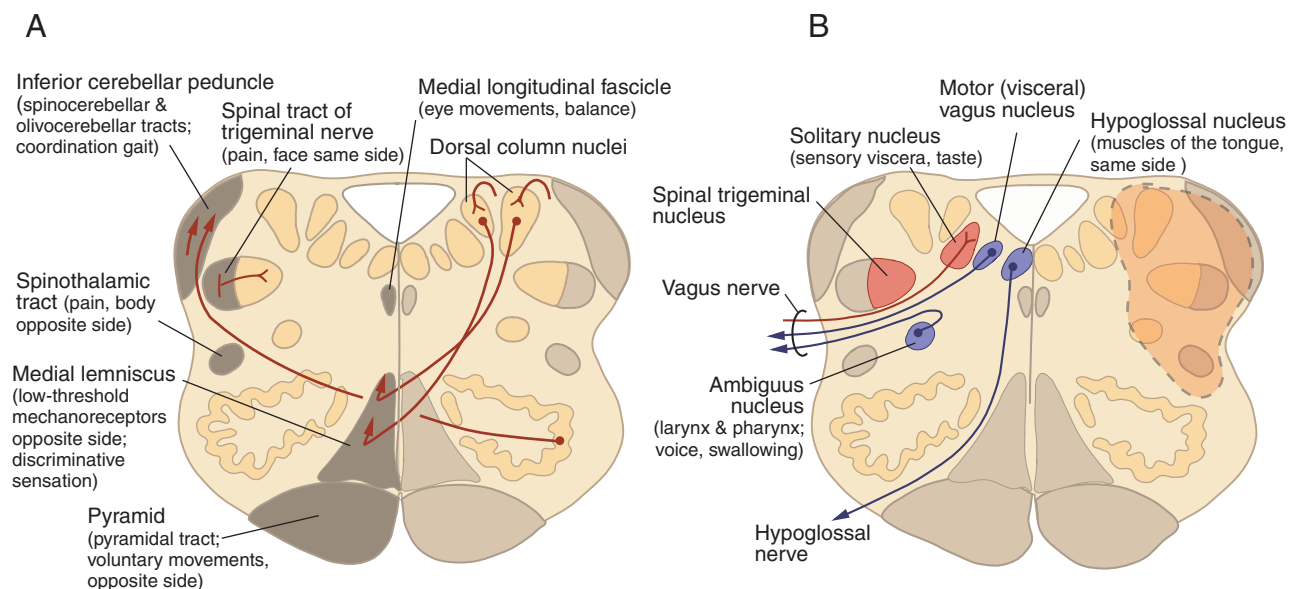


FIGURE 27.6 Position of cranial nerve nuclei, some other nuclei, and the main tracts in the medulla. **A:** Major tracts and typical symptoms produced by their interruption. **B:** Cranial nerve nuclei. The area of

infarction in a case of Wallenberg's syndrome is indicated with a stippled line. The figure helps identify the symptoms that are likely to occur after lesions of lateral or medial parts of the medulla, respectively.

in the floor of the fourth ventricle (the hypoglossal trigone; see Fig. 6.19). The **root fibers** of the nerve pass ventrally and leave the medulla just lateral to the pyramid (Figs. 27.1, 27.5, and 27.6). Several small fiber bundles join to form the nerve, which leaves the skull through the **hypoglossal canal** in the occipital bone. The nerve then forms an arc as it courses downward and forward in the upper neck—external to the carotid artery—to the root of the tongue. The fibers innervate the striated muscle cells of the tongue.

The muscles of the tongue are used **voluntarily** during speech and eating. In such activities, the neurons of the hypoglossal nucleus are influenced by fibers of the pyramidal tract coming from the face region of the motor cortex of the opposite hemisphere. The descending fibers cross in the medulla just above the nucleus. A central motor lesion above the nucleus (e.g., in the internal capsule) can produce pareses of the opposite half of the tongue, together with the more obvious pareses of the extremities. No clear-cut atrophy of the tongue occurs in cases of central motor lesions.

Reflex movements of the tongue occur in swallowing (and vomiting). The hypoglossal motoneurons are then activated from the brain stem reflex centers located in the reticular formation. Various kinds of stimuli can activate the reflex center for swallowing, notably touch- and pressure receptors at the back of the tongue.

A unilateral **lesion** of the hypoglossal nerve or nucleus produces paralysis of the tongue on the same side. As this is a peripheral paresis, a pronounced atrophy of the tongue muscles ensues. This is witnessed by a wrinkled surface of the tongue because the mucous membrane becomes too big for the reduced volume (Fig. 27.7). When stretching out the tongue, it **deviates** to the paretic side, because of paresis of the **genioglossus muscle**. This muscle passes backward and laterally into the tongue from its origin at the inside of the mandible; its normal

action when acting unilaterally is to draw the tongue forward and to the opposite side.

The Ansa Cervicalis

Some of the motor fibers from the first cervical spinal segment join the hypoglossal nerve and follow it for some distance before leaving it and descending in the neck. The descending fibers join other motor fibers from the second and third cervical segments and thereafter form an arc external to the internal jugular vein, called the **ansa cervicalis** (*ansa*, handle). The fibers of the ansa cervicalis innervate the **infrahyoid muscles** and are not related to the hypoglossal nucleus or the muscles of the tongue. Damage to the hypoglossal nucleus or the proximal part of the nerve (before the fibers from C₁ join it) therefore produces no pareses of the infrahyoid muscles.

THE ACCESSORY NERVE

The **eleventh cranial nerve**, the accessory nerve, brings somatic efferent fibers to two muscles in the neck—the **sternocleidomastoid** and the **trapezius**. The accessory **nucleus** is located in a column in the upper part of the cervical cord (Fig. 27.2) and contains neurons that are of the ordinary motoneuron type. The root fibers leave the cord and ascend to enter the posterior fossa through the foramen magnum (Fig. 27.8). The nerve then leaves the skull through the **jugular foramen** together with the vagus and the glossopharyngeal nerves. Outside the skull, the nerve passes internal to the sternocleidomastoid muscle and superficially through the upper part of the lateral triangle of the neck before continuing under the upper part of the trapezius muscle.

Some fibers from the nucleus ambiguus join the accessory nerve for a short distance intracranially. These fibers leave the nerve just outside the jugular foramen and follow the vagus nerve in their further course. They should therefore be considered a part of the vagus with a somewhat aberrant course rather than a part of the accessory nerve.

In **central motor lesions** (of the corticobulbar component of the pyramidal tract), pareses of the contralateral sternocleidomastoid and the trapezius are usually observed. Because of its superficial position, the nerve may be damaged in the **lateral triangle** of the neck, producing a **peripheral paresis**. In such a case only the trapezius is paretic because the fibers innervating the sternocleidomastoid leave the nerve higher up. Further, the upper part of the trapezius muscle is usually most seriously affected, the lower part being supplied also from the cervical plexus. Paresis of the trapezius muscle makes it difficult to elevate the arm, because the trapezius is necessary for the rotation of the shoulder blade around its anteroposterior axis.

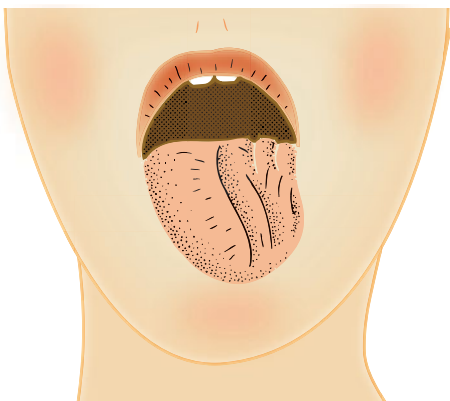


FIGURE 27.7 *Peripheral paralysis of the hypoglossal nerve on the left side.* The tongue deviates to the side of the lesion when the patient tries to stretch it out. Atrophy of the intrinsic muscles of the tongue makes the surface wrinkled.

THE VAGUS NERVE

The vagus nerve is the **tenth cranial nerve** and is characterized by containing fibers of all four kinds described above. Correspondingly, it is connected with four different nuclei in the medulla. The root fibers emerge in a row at the lateral aspect of the medulla (Figs. 27.1 and 27.3) and join to form one nerve, which leaves the skull through the jugular foramen (Fig. 27.8). Embryologically, the vagus nerve belongs to the fourth, fifth, and sixth branchial arches, and this explains the peculiar course of some of its branches. In the **jugular foramen** and immediately below, the nerve has two swellings—the **jugular** and the **nodose ganglia** (Fig. 27.9)—that contain the pseudounipolar cell bodies of the sensory vagus fibers.

As implied by the name (Latin: *vagus*, wandering), the vagus nerve sends branches to widespread regions of the body. After leaving the skull, the nerve passes as a fairly thick cord downward in the neck together with the common carotid artery, further through the thorax, and into the abdomen through the diaphragm. It gives off branches in the neck, in the thorax, and in the abdomen.

Visceral Efferent (Parasympathetic) Vagus Fibers

The visceral efferent neurons belong to the parasympathetic part of the autonomic system. The cell bodies lie

in the (dorsal) **motor nucleus of the vagus** (Figs. 27.2 and 27.6; see Fig. 6.17).³ The fibers do not pass directly to the organs but end on a second set of neurons in parasympathetic ganglia close to or in the walls of the organs (Fig. 27.9), as is the case for all parasympathetic nerves. The neurons leading from the CNS to the ganglia are called **preganglionic**, and those leading from the ganglion to the organ are called **postganglionic** (see Fig. 28.1).

The vagus gives off visceral efferent fibers descending to the **heart** in the neck (Fig. 27.9). The cell bodies of the postganglionic neurons are located in the wall of the heart and around the great vessels near the heart, and the postganglionic fibers end among the muscle cells of the heart, especially those of the **sinus node**, which determines the heart rate.

Other branches from the vagus supply the **esophagus** and the **trachea** and, further down in the thorax, the **bronchi** of the lung. The postganglionic fibers innervate smooth muscles and glands in these structures.

3 Experiments with tracing techniques show that the neurons supplying the **heart** (and probably also the lungs) have their cell bodies in an external, distinct part of the **nucleus ambiguus**. The cell bodies are morphologically like those of the dorsal motor vagus nucleus, but they differ from the larger motoneurons in the main nucleus ambiguus, which innervate striated muscles (of the pharynx and larynx).

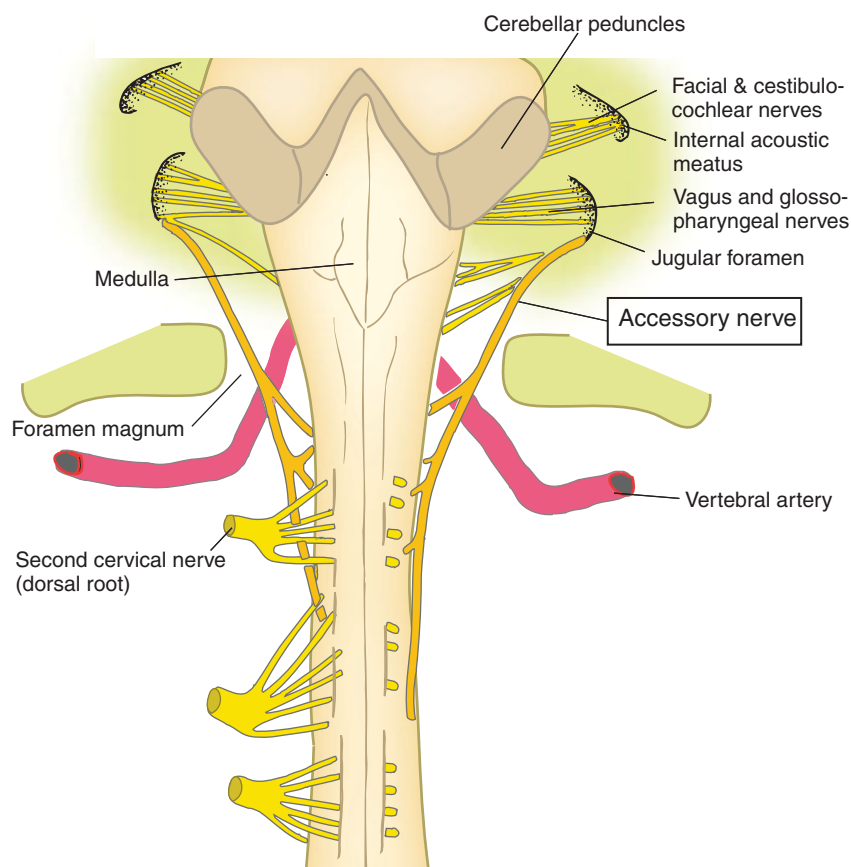


FIGURE 27.8 The accessory nerve. The lower part of the brain stem and the upper cervical cord, as viewed from behind. The cerebellum is removed. The fibers of the accessory nerve proper emerge from the three upper cervical segments and exit the cord between the ventral and dorsal roots (the ventral roots are not visible in the figure). Compare with the emergence of the vagus fibers in Fig. 27.1.

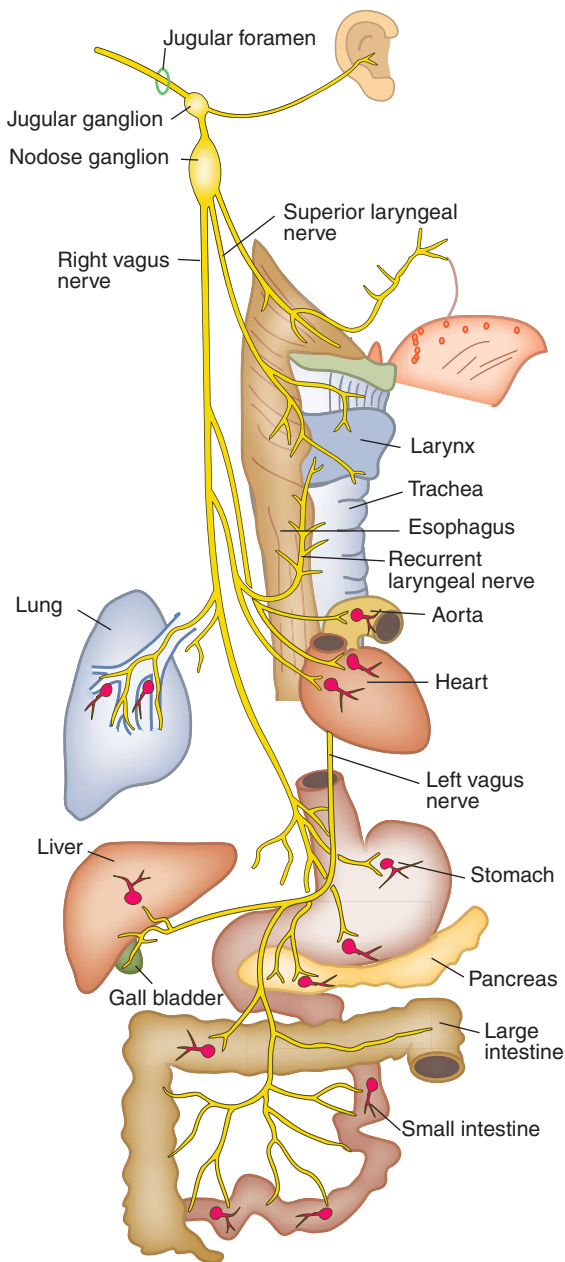


FIGURE 27.9 Course and distribution of the vagus nerve.

In the **abdomen**, the vagus sends fibers to the **stomach**, the **small intestine**, and the first half of the **large intestine**. The vagus also supplies the **liver**, the **gallbladder**, and the **pancreas** with parasympathetic fibers. To reach the various organs, the fibers follow the arteries and form plexuses around them together with sympathetic fibers.

Functionally, signals in the vagus nerve reduce the heart rate, constrict the bronchi, and increase bronchial secretion, whereas the peristaltic movements and secretion are increased in the stomach and intestine. The secretion of the pancreas is also increased.

Visceral Afferent Vagus Fibers

All branches of the vagus with visceral efferent fibers also contain afferent (sensory) fibers. These neurons have their pseudounipolar cell bodies in the large **nodose ganglion** (Fig. 27.9). When the fibers enter the brain stem, they form a small bundle that passes caudally: the **solitary tract** (this is not a true tract, however, but the central course of a nerve). The fibers establish synapses in the **solitary nucleus** (Figs. 27.2 and 27.6; see Fig. 6.17). The efferent fibers from this nucleus pass to the reticular formation and other cranial nerve nuclei in the vicinity and to higher levels such as the thalamus and the hypothalamus (Fig. 27.4).

All of the sensory fibers from the **larynx** and some from the **pharynx** follow the vagus. The sensory fibers from the larynx follow the **superior laryngeal nerve** (from the part above the vocal cords) and the **recurrent laryngeal nerve** (from the lower part of the larynx). On an **embryological** basis, all sensory fibers from the pharynx (and the posterior part of the tongue) should be classified as visceral afferent. There is evidence, however, that such fibers end in the somatic afferent trigeminal nucleus and not in the visceral afferent solitary nucleus. Functionally, they may therefore be considered somatic afferent rather than visceral afferent. Vagus may also carry nociceptive signals from parts of the **heart** (see Chapter 29, under “Central Pathways for Visceral Afferent Signals”).

Visceral afferent signals in the vagus nerve most likely contribute to conscious **feelings** like hunger or satiety. There is also evidence that signals in the vagus nerve contribute to the feelings associated with infections (loss of appetite, tiredness, somnolence, and so forth) and the induction of **sickness behavior**. Together with other visceral afferent fibers, the vagus contributes to our general bodily feeling—also called the “sense of the physiological condition of the body” (Craig 2002). This is discussed further in Chapter 30, under “The Hypothalamus and the Immune System.”

Visceral Reflexes

The visceral afferents in the vagus nerve are links in reflex arcs that control **secretion** and **peristaltic movements** of the gastrointestinal tract (and vomiting). Such reflexes also mediate alterations of **airway secretion** and of the **airway resistance** by changing the tone of the bronchial smooth muscles. The **reflex centers** of all these reflexes are located in the medulla, and the efferent links are visceral efferent fibers coming from the dorsal motor nucleus of the vagus.

Visceral afferents from **baroreceptors** in the wall of the large vessels provide information about the blood pressure in the aorta. Increased blood pressure gives rise to increased firing frequency of the afferent fibers. In turn,

this produces increased firing of visceral efferent vagus fibers, which reduces the heart rate. The **reflex center** must involve connections from the solitary nucleus to the motor nucleus of the vagus (these two nuclei are close neighbors, as can be seen in Fig. 27.2).

The motor nucleus of the vagus can be influenced also by signals other than those coming from the viscera. For example, the sight, the smell, or even the thought of food can produce increased secretion of gastric juice. These are examples of **conditioned responses**, whereas the stimulation of taste receptors produces an unconditioned (true reflex) response.

Other examples of visceral reflexes are discussed in Chapter 29, under “Visceral Reflexes.”

The Vomiting Reflex

Vomiting is usually caused by marked dilatation of the stomach or irritation of its mucosa. The biologic significance of the reflex is presumably to rid the stomach of potentially harmful contents. As we all know, however, vomiting can also be provoked by irritation of the pharynx (putting a finger in the throat) and by foul odors, strong emotions, and travel sickness. Several different afferent links to the reflex center must therefore exist. The emptying of the stomach is caused by coordinated contractions of the smooth muscles in the stomach wall and striated muscles in the diaphragm and the abdominal wall. In addition, laryngeal muscles (closing the airways) and muscles of the pharynx, the soft palate, and the tongue also participate. Thus, the reflex center must activate visceral efferent neurons and α motoneurons at several levels of the brain stem and the spinal cord in a specific sequence. The **reflex center** is actually quite widespread, but usually matters are simplified by restricting it to the medulla, including the **solitary nucleus**. This receives the visceral afferent fibers from the stomach, forming the afferent link when the reflex is elicited from the stomach itself. From the reflex center in the medulla, signals pass to the motor nuclei via synaptic interruption in the reticular formation and reticulospinal fibers. In addition, there are direct spinal projections from the solitary nucleus to the motoneurons of the diaphragm and abdominal wall.

Substances in the bloodstream cause vomiting by direct action at the **area postrema** of the medulla (see Chapter 8, under “Some Parts of the Brain Lack a Blood–Brain Barrier”). Neurons in the area postrema project to the solitary nucleus. **Apomorphine** and other alkaloids are given orally or subcutaneously to provoke vomiting.

Somatic Efferent Vagus Fibers

The somatic efferent vagus fibers come from the **ambiguus nucleus** (Figs. 27.2 and 27.5), which belongs to the

special somatic efferent nuclei. The fibers supply all striated muscles of the **larynx** and parts of the muscles of the **pharynx**. The fibers to the pharynx take off from the vagus as several small branches, whereas most of the fibers to the larynx are collected in the **recurrent laryngeal nerve** (Fig. 27.9). This nerve takes off from the main vagus trunk at the level of the aortic arch on the left side and the subclavian artery on the right. It then arches behind the vessels and ascends in the furrow between the trachea and the esophagus, to reach the larynx. One of the laryngeal muscles located on the outside, the **cricothyroid muscle**, receives motor fibers in the superior laryngeal nerve (which is a predominantly sensory nerve, as mentioned earlier). The vagus also innervates one of the muscles of the soft palate, the **levator veli palatini** muscle.

A **lesion** of the vagus nerve above the exit of the motor branches to the pharynx and the soft palate produces deviation of the uvula and the posterior pharyngeal wall to the normal side (as can be seen, e.g., when a patient is asked to say “aah”; Fig. 27.10). Pareses of the soft palate and the pharynx cause fluid and food to enter the nasal cavity when swallowing (owing to inadequate closure of the nasopharynx). Further, the voice becomes hoarse because the vocal cords cannot be properly adducted. Such a symptom will obviously also occur after a lesion of the recurrent laryngeal nerve anywhere along its course. In case of a unilateral lesion, the voice hoarseness will gradually disappear, because the muscles of the normal side adapt to the changed conditions.

The neurons of the nucleus ambiguus are influenced by, among other sources, the **pyramidal tract** during speech. They can also be activated involuntarily in the **cough reflex** by irritating stimuli of the respiratory tract.

Somatic Afferent Vagus Fibers

This is the smallest contingent of fibers in the vagus nerve. They have their cell bodies in the small **jugular**

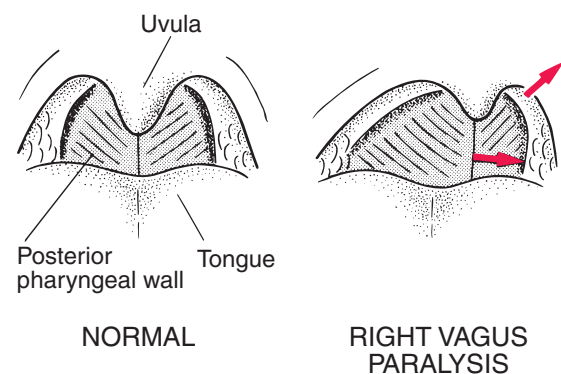


FIGURE 27.10 Paralysis of the right vagus nerve. The uvula and the posterior pharyngeal wall are pulled toward the normal side when the patient says “aah.” (Redrawn from Mumenthaler 1979.)

ganglion and come from a small region of the skin of the external ear—the **auricular ramus** (Fig. 27.9). The fibers terminate in the **trigeminal sensory nucleus**. Touching the innervated area, for example, by an otoscope in the external meatus may evoke a cough reflex and in some individuals even a vomiting reflex. The causes of these phenomena are unknown. They might be due to connections from the trigeminal nucleus to the solitary nucleus or by abnormal signal transmission between sensory fibers of the vagus nerve (**ephaptic transmission**).

THE GLOSSOPHARYNGEAL NERVE

The **ninth cranial nerve**, the glossopharyngeal, resembles the vagus but is smaller and innervates a more restricted region. The root fibers leave the medulla immediately rostral to the vagus fibers (Fig. 27.1). The root fibers fuse to form one trunk that leaves the cranial cavity through the **jugular foramen** (together with the vagus and the accessory nerves). The nerve follows an arched course (ventrally) lateral to the pharynx, which it penetrates to reach the base of the tongue. Close to the jugular foramen, the nerve contains two small **sensory ganglia** with pseudounipolar ganglion cells, the **superior** and **petrous ganglia**.

Of the peripheral branches, some innervate the muscles and the mucous membrane of the **pharynx** (together with the vagus, which appears to be the most important quantitatively); other sensory fibers reach the posterior part of the tongue, to innervate **taste buds**, and the mucous membrane (and also the mucous membrane of the soft palate and the tonsillar region). The glossopharyngeal nerve also contains **visceral efferent** (parasympathetic) fibers to the **parotid gland** and to the salivary glands in the posterior part of the tongue (see Fig. 28.11 for the course of the parasympathetic fibers).

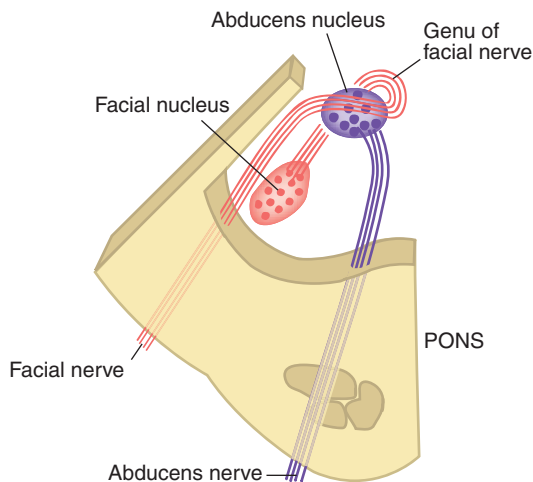


FIGURE 27.11 Course of the facial nerve and its relation to the abducens nucleus. Schematic cross section through the lower pons.

The Sinus Nerve and Baroreceptors

A special contingent of visceral afferent fibers in the glossopharyngeal nerve comes from the wall of the **carotid sinus** (the thin-walled, dilated part of the internal carotid artery). The fibers conduct signals from mechanoreceptors recording the tension of the arterial wall; that is, the receptors monitor the blood pressure and are therefore called **baroreceptors** (cf. the same kind of afferents from the aorta running with the vagus nerve). The afferent fibers end in the **solitary nucleus**, and from there the signals are conveyed to the **motor nucleus of the vagus**. Increased signal frequency of the cardiac vagus fibers reduces the heart rate, and thereby the blood pressure is reduced. When the blood pressure falls, there will be reduced firing of the cardiac vagus fibers, with increased heart rate and blood pressure. This is one of several mechanisms to keep the **blood pressure** within certain limits and that the cerebral blood flow is sufficient at all times.

The Nuclei of the Glossopharyngeal Nerve

The **somatic efferent** fibers to the striated pharynx muscles come from the **ambiguus nucleus**, whereas the **visceral efferent** fibers have their cell bodies in the small **inferior salivatory nucleus** (Fig. 27.2). The signals from this nucleus follow a somewhat complicated course to reach the parotid gland (as is the case for several of the parasympathetic fiber components of the cranial nerves). The preganglionic parasympathetic fibers from the inferior salivatory nucleus end in the small **otic ganglion** just outside the cranial cavity. The postganglionic fibers from the ganglion cells join one of the trigeminal branches—the **auriculotemporal nerve** that passes close to the ganglion—to reach gland (see Fig. 28.11).

The **visceral afferent** fibers carrying signals from the **taste buds** in the posterior third of the tongue end in the **solitary nucleus**. From there, the signals pass to the thalamus and further on to the cerebral cortex. The sensory fibers from the posterior part of the tongue, the tonsils, the soft palate, and the pharynx end in the **sensory trigeminal nucleus**.

THE VESTIBULOCOCHLEAR NERVE

The **eighth cranial nerve**, although consisting of only one trunk, is in reality two functionally different nerves: the **cochlear nerve** (see Chapter 17) and the **vestibular nerve** (described with the sense of equilibrium in Chapter 18). The nerves fuse after leaving the labyrinth (the cochlea and the vestibular apparatus) and follow the **internal acoustic meatus**. At the lower end of the pons, in the **cerebellopontine angle**, the nerve enters the brain stem. Most of the fibers in the vestibular nerve

end in the **vestibular nuclei**; some end in the cerebellum. The cochlear nerve ends in the **cochlear nuclei** (Fig. 27.2). The cell bodies of the primary afferent fibers are located at the bottom of the internal meatus, forming the **vestibular ganglion** and the **spiral ganglion**. The main structural features of the labyrinth are described in Chapters 17 and 18.

THE FACIAL AND INTERMEDIATE NERVES

The **seventh cranial nerve** belongs to the **second branchial (visceral) arch** and innervates structures developed from this. The facial nerve is the motor nerve of the facial (mimetic) muscles, supplying them with **special somatic efferent** fibers. The small **intermediate nerve** follows the facial nerve and is usually considered to belong to it. Thus, parts of the intermediate nerve might be regarded as a sensory root of the facial nerve with its **visceral afferent** fibers. The intermediate nerve also contains **visceral efferent** (parasympathetic) fibers, however.

The facial nerve and the intermediate nerve leave the brain stem together laterally at the lower border of the pons, just ventral to the eighth nerve (Fig. 27.1). They then follow the eighth nerve to the bottom of the internal auditory meatus, where the intermediate nerve leaves the facial nerve. The facial nerve proper (with the somatic efferent fibers) then arches first posterolaterally and thereafter downward in the facial canal. It leaves the skull through the **stylomastoid foramen** immediately medial and anterior to the mastoid process. The nerve then passes forward through the parotid gland and divides into several branches, spreading out like a fan to all the facial muscles.

The **visceral efferent** (parasympathetic preganglionic) fibers in the intermediate nerve bring secretory signals to the **lacrimal gland** and the **submandibular** and the **sublingual salivary glands** (see Fig. 28.11 for the complicated route followed by the fibers). The **visceral afferent** fibers of the intermediate nerve come from **taste buds** in the anterior two-thirds of the tongue.

The Facial Nerve

The motor fibers of the facial nerve have their cell bodies in the **facial nucleus**, located in the lower pons. It belongs to the column of special somatic efferent nuclei (Figs. 27.2 and 27.3). The nucleus consists of several subdivisions, each supplying small groups of muscles. The root fibers of the facial nerve have a peculiar course before they leave the brain stem (Fig. 27.11). First, the fibers pass medially and lie dorsal to the abducens nucleus, forming the **genu of the facial nerve**, before they bend in the lateral and ventral direction. The facial fibers pass just beneath the floor of the fourth ventricle and form a

small elevation; the **facial colliculus** (see Fig. 6.19). Owing to the course of the nerve, symptoms of a peripheral facial paresis may occur in lesions that are located considerably more medial and dorsal than the nucleus itself. Figure 27.11 shows that damage to the abducens nucleus (affecting the lateral rectus muscle of the eye) is also likely to be accompanied by signs of pareses of the facial muscles of the same side.

The Mimetic Muscles

The facial or mimetic muscles originate from the facial skeleton and insert with elastic tendons in the dermis. There are many small muscles, with the majority located around the mouth and the eyes. Of particular practical value are the muscles responsible for blinking and closure of the eye (the **orbicularis oculi muscle**) and the muscles around the mouth (the **orbicularis oris** and several other muscles that move the lips). The **buccinator muscle** prevents the cheeks from being pressed out when the intraoral pressure is increased and, perhaps more importantly, prevents the cheeks from being sucked in between the teeth.

Central and Peripheral Facial Pareses

Signals from the facial nucleus evoke contractions of the mimetic muscles and are therefore responsible for our facial expressions. These muscles function also in conjunction with speech, eating, blinking, and so forth. The **pyramidal tract** (corticobulbar fibers) conveys signals for voluntary movements of the facial muscles. The fibers arise from the face region of the MI in the precentral gyrus (see Fig. 22.5).

The part of the facial nucleus supplying the muscles in the forehead and around the eyes receive both uncrossed and crossed pyramidal tract fibers, whereas the muscles in the lower part of the face receive purely crossed fibers. A **lesion of the pyramidal tract** (e.g., in the internal capsule) therefore produces clear-cut pareses only in the lower part of the face on the opposite side. Most obvious is the sagging corner of the mouth. The patient can still wrinkle the forehead and close the eyes voluntarily. A **peripheral lesion** (of the facial nucleus or the nerve), however, produces pareses of all the facial muscles on the same side as the lesion (Fig. 27.12). For example, the eye cannot be closed, so there is danger of drying and ulceration of the cornea (and permanent loss of vision).

Peripheral lesions of the facial nerve may be caused by hemorrhage, infarctions, or tumors in the pons, by infections of the middle ear (to which the nerve passes in close proximity), or by damage to the branches in the face. Most often, however, the cause of peripheral facial paresis is unknown (**Bell's palsy**). In such cases, the muscle power usually returns after some time.



FIGURE 27.12 *Peripheral facial paralysis (right side).* The patient is asked to close her eyes and to retract the corners of the mouth. (Based on Monrad-Krohn 1954.)

Facial Expressions of Emotion Do Not Depend on the Pyramidal Tract

Whereas signals mediated by the pyramidal tract activate the motoneurons of the facial nucleus in voluntary movements (such as speech and eating), other descending pathways are responsible for facial expressions of emotions, such as sorrow and pleasure. As most of us know from personal experience, a genuine smile cannot be produced on command but arises independent of any conscious will. Indeed, our facial expressions often reveal emotions we would rather have concealed. A voluntary effort is required to suppress spontaneous facial expressions, which most likely are controlled by descending connections from the **hypothalamus** and possibly the **basal ganglia**. Thus, lesions of the pyramidal tract do not abolish spontaneous facial expressions. The patient smiles and laughs when told a good joke but cannot present a polite social smile. In **central pareses** (such as capsular hemiplegia) emotional facial expressions are in fact often exaggerated, and the patient cannot suppress a smile or prevent crying. Diseases of the basal ganglia, such as **Parkinson's disease**, present the opposite picture: the emotional, spontaneous expressions are lacking, whereas a voluntary, social smile is possible.

The Facial Nerve and Reflexes

The facial nucleus is also a link in some important reflex arcs. One is the **corneal** or **blink reflex**. It is elicited by touch or irritation of the cornea, and the sensory signals are conducted centrally in the trigeminal nerve to the

spinal trigeminal nucleus (Fig. 27.13). From there the signals pass via interneurons in the reticular formation to the facial nucleus of both sides, and a contraction of the muscles of the eyelid is produced. The corneal reflex can be weakened or abolished by a lesion anywhere along the course of the afferent and efferent links or in the rather extensive reflex center.

Another reflex mediated by the facial nerve is the **stapedius reflex**, in which the response is contraction of the tiny **stapedius muscle** in the middle ear. The stimulus is an intense sound conducted centrally in the cochlear (acoustic) nerve to the cochlear nuclei. Most likely, interneurons in the reticular formation transfer the signals to the facial nucleus. The stapedius muscle pulls the stapes a little out of the oval window (see Fig. 17.7) and thereby dampens the transmission of sound waves to the cochlear duct. Accordingly, peripheral facial paresis can produce hypersensitivity to sounds, or **hyperacusis**.

Secretion of Tears and Saliva

The preganglionic parasympathetic fibers of the **intermediate nerve**—acting on the **lacrimal**, the **submandibular**, and the **sublingual glands**—have their cell bodies in the small **superior salivatory nucleus**. This nucleus belongs to the column of visceral efferent nuclei (Fig. 27.2). As mentioned, preganglionic parasympathetic fibers acting on the parotid gland have their cell bodies in the inferior salivatory nucleus and leave the brain stem in the glossopharyngeal nerve.

The **secretion of saliva** is brought about primarily by stimulation of the taste receptors but also by signals from higher levels of the brain (such as the thought of tasty food; it is especially effective to imagine that one

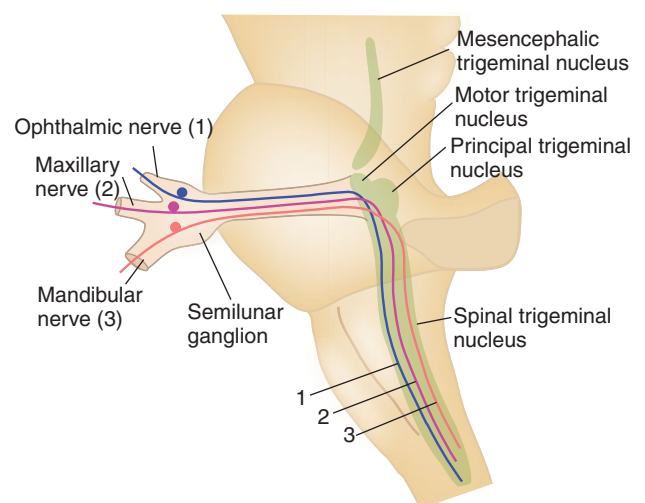


FIGURE 27.13 *The trigeminal nuclei.* In addition, the figure shows the topographic arrangement in the spinal trigeminal nucleus of the fibers from the three main trigeminal branches.

is eating a lemon). Strong emotions, for example, anxiety before a performance, can inhibit the secretion of saliva, as experienced by mouth dryness.

The **secretion of tears**, even more than the salivary secretion, is an example of how visceral functions can be influenced from higher levels of the brain. The continuous secretion of tears is of course primarily a physiological protection of the eyes and increases in response to any irritation of the cornea or the conjunctiva; nevertheless, the most profuse tear production occurs when we express strong emotions by crying. The signals to the superior salivatory nucleus producing the flow of tears when crying are not mediated by the pyramidal tract or other efferent cortical fibers descending in the internal capsule, in correspondence with the fact that tears cannot be produced voluntarily nor can the secretion of tears be suppressed. Most likely, fibers from the **hypothalamus** are responsible for the activation of the visceral efferent neurons during crying. Nevertheless, the hypothalamus is under the influence of higher levels, such as parts of the cerebral cortex and the limbic structures. Thus, the conscious experience of the emotions (such as sorrow or pity) starts the train of neural events leading to tear secretion.

The Intermediate Nerve

The small **geniculate ganglion**, containing the cell bodies of the sensory fibers of the intermediate nerve, is found where the facial nerve bends posteriorly in the temporal bone. Here a branch of the intermediate nerve, the **greater petrosal nerve**, leaves the main trunk of the facial nerve to course anteriorly. It contains **visceral efferent** (parasympathetic) fibers that end in the small parasympathetic **pterygopalatine ganglion** located behind the orbit. From there postganglionic fibers follow trigeminal branches to the **lacrimal gland** and **glands in the nasal cavity** (see Fig. 28.11). The rest of the intermediate nerve fibers leave the facial nerve as it passes downward posterior to the middle ear. This branch is called the **chorda tympani** because it passes through the middle ear (tympanic cavity) on its way forward to join the **lingual nerve** (a trigeminal branch) outside the skull. The chorda tympani contains **visceral afferent** fibers from **taste buds** in the anterior two-thirds of the tongue. These fibers have their cell bodies in the geniculate ganglion. In addition, the chorda tympani carries **visceral efferent** (parasympathetic) fibers that end in the small **submandibular ganglion**. From this ganglion, postganglionic parasympathetic fibers pass to the submandibular and the sublingual (salivary) glands.

THE TRIGEMINAL NERVE

The **fifth cranial nerve** is primarily the sensory nerve of the face, with mainly **somatic afferent fibers**. In addition,

it contains a small portion with **special somatic efferent** fibers to the masticatory muscles. The trigeminal nerve is the nerve of the **first branchial (visceral) arch** and innervates structures that are developed from this arch.

The nerve leaves the brain stem laterally on the pons (Fig. 27.1) with a small (medial) motor root and a large (lateral) sensory root. Shortly after leaving the pons, the nerve expands to form the large **semilunar ganglion**, which contains the cell bodies of the pseudounipolar (sensory) ganglion cells. Three large branches continue anteriorly from the ganglion: the **ophthalmic**, the **maxillary**, and the **mandibular** nerves (Fig. 27.13).

The **ophthalmic nerve** enters the orbit and supplies the eye bulb (including the cornea), the upper eyelid, the back of the nose, and the skin of the forehead with sensory fibers (Fig. 27.14). It also sends fibers to the mucous membranes of the anterior part of the nasal cavity. The **maxillary nerve** runs forward in a sulcus in the bottom of the orbit and sends fibers to the lower eyelid, the skin above the mouth, the upper teeth and the gingiva, and, finally, the hard palate and the posterior (major) part of the nasal cavity. The **mandibular nerve** innervates the lower teeth and gingiva, the tongue, and the skin of the lower jaw and upward, well into the temporal region (Fig. 27.14). The branch of the mandibular nerve supplying the tongue with somatic sensory afferent fibers is called the **lingual nerve**. This nerve receives visceral afferent (taste) fibers from the **chorda tympani**, destined for the anterior two-thirds of the tongue.

The **motor fibers** of the trigeminal nerve follow the mandibular nerve but leave this in several smaller twigs

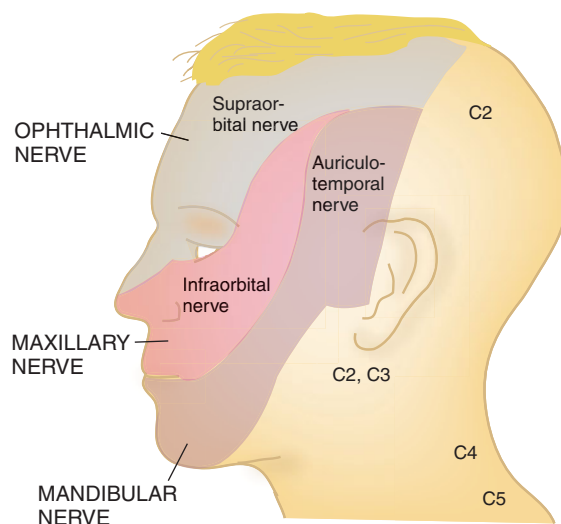


FIGURE 27.14 Distribution in the facial skin of the three main trigeminal branches. The names of some further branches and the segmental origins of sensory fibers to the rest of the head and the neck are indicated.

to the masticatory muscles (and some other muscles with relation to the lower jaw and the soft palate).

The Sensory Trigeminal Nucleus

With regard to function and fiber composition, the sensory part of the trigeminal nerve corresponds to the spinal dorsal roots. The trigeminal nerve, therefore, belongs to the somatosensory system and conducts signals from **low-threshold mechanoreceptors, thermoreceptors, and nociceptors** in the face and in the mucous membranes of the face. As with other spinal nerves, fibers leading from different kinds of receptors are intermingled in the nerve but are arranged by receptor type when entering the CNS. Then the fibers distribute to the three subdivisions of the long sensory trigeminal nucleus (Figs. 27.2 and 27.13). Fibers from proprioceptors (muscle spindles, joint receptors) end in the **mesencephalic nucleus**, fibers from low-threshold mechanoreceptors end in the main or **principal nucleus**, whereas signals from **nociceptors** end in the **spinal trigeminal nucleus**.⁴

Central Transmission of Signals from the Trigeminal Nucleus

Functionally, the **spinal trigeminal nucleus** (especially its caudal part) corresponds to the dorsalmost laminae of the cord, whereas the **principal sensory nucleus** corresponds to the dorsal column nuclei. These similarities are evident also in the central pathways. The secondary sensory fibers from the cells in the spinal nucleus join the **spinothalamic tract** (see Fig. 14.4) and end in the thalamus. Fibers from the main nucleus join the **medial lemniscus** (see Fig. 14.2). The somatotopic pattern within the thalamic terminal region is such that the fibers from the trigeminal nucleus—carrying signals from the face—end most medially, in the VPM (see Fig. 14.6). The ascending fibers from the trigeminal nucleus cross to the opposite side before they join the large sensory tracts. As discussed (under “Brain Stem Lesions Can Produce Symptoms from Several Cranial Nerves and Long Tracts”), a lesion affecting lateral parts of the medulla is likely to interrupt the spinothalamic tract and the spinal trigeminal nucleus. This will usually cause reduced or abolished pain and temperature sensation in the opposite body half but on the same side of the face (Fig. 27.6).⁵ From the thalamus, the

signals are transmitted to the **face region** of the SI in the postcentral gyrus.

More about the Subdivisions of the Sensory Trigeminal Nucleus

The thinnest fibers of the trigeminal nerve (A δ and C fibers)—conducting primarily from nociceptors and thermoreceptors—bend caudally after entering the pons (Fig. 27.13). They continue as a small bundle, the **spinal tract of the trigeminal nerve**, located just beneath the medullary surface. It is joined by somatic afferent fibers that have followed the glossopharyngeal and the vagus nerves peripherally. Just like the solitary tract, the spinal tract, strictly speaking, is not a tract, as it consists of the central process of the pseudounipolar ganglion cells. The spinal tract continues down into the upper cervical segments and corresponds to the zona terminalis (bundle of Lissauer) in the cord (see Fig. 13.16). The fibers enter the **spinal trigeminal nucleus** (nucleus of the spinal trigeminal tract), which corresponds largely to the dorsalmost laminae of the cord. For example, a layer very similar to the substantia gelatinosa is present. The spinal trigeminal nucleus can be further subdivided in a rostrocaudal sequence. The **caudal subnucleus** appears to be especially involved in pain mechanisms and corresponds most closely with the dorsal laminae of the cord. It also receives dorsal root fibers from the upper cervical segments. This may perhaps explain why a certain condition with paroxysms of facial pain of unknown origin—**trigeminal neuralgia**—may sometimes irradiate outside the area innervated by the trigeminal nerve. The spinal trigeminal nucleus furthermore shows a dorsoventral topographic localization. Thus the main trigeminal branches end sequentially, with the ophthalmic nerve ending most ventrally and the mandibular nerve most dorsally (Fig. 27.13).

Thick myelinated fibers (A β) in the trigeminal nerve from the skin end mostly in the principal sensory trigeminal nucleus (Fig. 27.13) with a precise somatotopic pattern.

The **mesencephalic trigeminal nucleus** stretches as a slender column from the upper part of the pons and into the mesencephalon. This is a very unusual nucleus, as its neurons look like pseudounipolar ganglion cells and, indeed, send one process peripherally into the trigeminal nerve. Afferent fibers from the **muscle spindles** of the **masticatory muscles** follow the mandibular nerve, whereas those from the **extraocular muscles** follow the ophthalmic and perhaps the oculomotor nerve. Signals from mechanoreceptors in the root sheaths of the **teeth** end in the mesencephalic nucleus.

Reflexes Involving the Trigeminal Nerve

Like the spinal dorsal root fibers, the trigeminal nerve constitutes the afferent link of several reflex arcs. The trigeminal nucleus, especially the spinal subdivision,

4 The separation of fibers of different types is not quite as sharp as this account may indicate. Many trigeminal fibers divide after entering the brain stem into an ascending and a descending branch (just like the sensory fibers entering the cord). In this manner, single ganglion cells may end in more than one nuclear subdivision.

5 If the lesion is situated high in the medulla or in the lower pons the ascending (secondary sensory) fibers from the spinal trigeminal nucleus have crossed and joined the spinothalamic tract. In such cases, loss of pain and temperature sensation in the face occurs on the opposite side of the lesion.

has numerous connections with the reticular formation. This is partly by means of reflex arcs involving the trigeminal nerve as the afferent link but also as links in ascending pathways to the thalamus with signals from nociceptors.

We described the **corneal reflex** earlier, in conjunction with the facial nerve that constitutes the efferent link of this reflex. Other reflexes are the **sneeze reflex**, elicited from the mucous membrane of the nasal cavity, and the **sucking reflex**, elicited in the newborn from mechanoreceptors of the lips. A brief tap downward on the chin can produce a stretch reflex, stretching the masticatory muscles (among them, the masseter muscle). This **masseter reflex** is monosynaptic and involves the mesencephalic trigeminal nucleus. The peripheral processes of the neurons of this nucleus innervate the muscle spindles, and the central processes reach the motor trigeminal nucleus in the pons. This reflex, with its reflex center in the mesencephalon and pons, is one of those routinely tested in clinical neurology.

The Motor Trigeminal Nucleus

The special somatic efferent fibers in the trigeminal nerve come from the motor trigeminal nucleus, located in the pons at the level of the main sensory nucleus (Fig. 27.13, see also Fig. 27.2). All of the motor fibers follow the mandibular nerve. A peripheral paresis of the motor fibers of the trigeminal nerve (or the nucleus) leads to atrophy of the masticatory muscles and reduced force of biting on the side of the lesion. A **unilateral paresis** of the masticatory muscles is most easily detected by asking the patient to open his mouth widely; the lower jaw then **deviates** toward the side of the paretic muscles (Fig. 27.15). This is caused by paresis of the **lateral pterygoid muscle**, which normally pulls the mandible forward in conjunction with opening of the mouth.

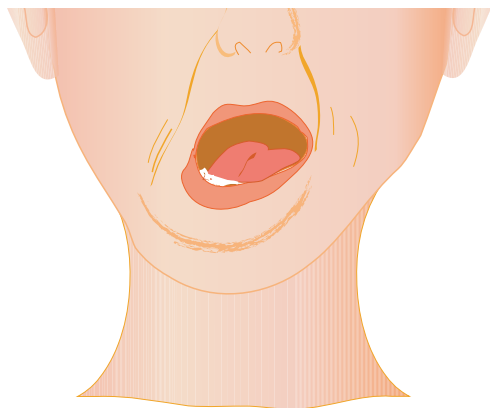


FIGURE 27.15 Deviation of the jaw as a sign of peripheral trigeminal pareses. Paresis of the right trigeminal nerve. Normally, the two lateral pterygoid muscles pull the jaw forward when opening the mouth. When unopposed by the paretic right pterygoid, the left (normal one) moves the jaw to the right when the patient opens her mouth.

Voluntary movements of the jaws during speech and chewing depend on the pyramidal tract (corticobulbar fibers). The fibers from the MI to the motor trigeminal nucleus are often both crossed and uncrossed (bilateral), but in some persons, they appear to be purely crossed. Only in the latter case, will there be clear-cut signs of pareses of the masticatory muscles after a lesion of the internal capsule (capsular hemiplegia). There may also be other signs of a central paresis, such as an increased masseter reflex on the paretic side.

Reflex movements of the masticatory muscles occur during swallowing, sucking, and vomiting. The afferent fibers of such reflexes are sensory trigeminal fibers from the oral cavity and sensory fibers passing in the vagus and the glossopharyngeal nerves.

Lateral Pontine Infarction

Infarction in the lateral part of the pons is usually due to occlusion of the **anterior inferior cerebellar artery**, which leaves the basilar artery in the lower third of the pons (see Fig. 8.3). Besides the pons, the artery often supplies the brachium pontis, the flocculus, and middle parts of the cerebellar hemispheres. The most common symptoms are due to damage of the root fibers and the nuclei of the trigeminal, the facial, and the vestibulocochlear nerves (Fig. 27.1). Cerebellar symptoms and symptoms due to damage to the long ascending and descending tracts are more variable, presumably due to individual variations in the area supplied by the artery. A lateral pontine infarction will usually produce ipsilateral **peripheral facial paresis** (the root fibers leave laterally in the lower pons), ipsilateral **sensory loss in the face** (all sensory qualities due to interruption of trigeminal root fibers), **deafness** and **dizziness** (the vestibulocochlear nerve, vestibular nuclei), and in some **gaze paralysis** to the side (ascending connections from the vestibular nuclei; see Fig. 25.6). In addition, there is usually ipsilateral **ataxia** (the cerebellar hemispheres) and occasionally contralateral **central pareses** and an inverted plantar reflex (pyramidal tract) or contralateral **reduced pain and temperature sensibility** (the spinothalamic tract). **Horner's syndrome** is often present as well (see Chapter 28, under "Interruption of the Sympathetic Innervation of the Head").

THE ABDUCENS, TROCHLEAR, AND OCULOMOTOR NERVES

These three cranial nerves—the **sixth**, **fourth**, and **third**—supply the **extraocular muscles** with **somatic efferent** fibers (Fig. 27.16). In addition, the oculomotor nerve contains **visceral efferent** (parasympathetic) fibers to the smooth, **intrinsic eye muscles**. Simplified, we may say that by moving the eye in the orbit, the extraocular

muscles ensure that the images we look at fall on the central part of the retina (the macula). The intrinsic muscles control the amount of light that reaches the retina and adjust the curvature of the lens so that the retinal image is always in focus. Most of these tasks are performed reflexly, and usually eye position, light access, and lens curvature are controlled simultaneously. In Chapter 25, we discussed the extraocular muscles and the control of eye movements.

Typical **symptoms** resulting from lesions of the extraocular muscles or their nerves are **strabismus** (squint), **double vision**, and **dizziness** (probably due to double vision). In addition, the patient usually keeps his head a little bent or turned in order to reduce the double vision (that is, trying to keep a position in which the paretic muscle(s) is used as little as possible). Figure 25.3 shows a simple scheme for examination of the six extraocular muscles.

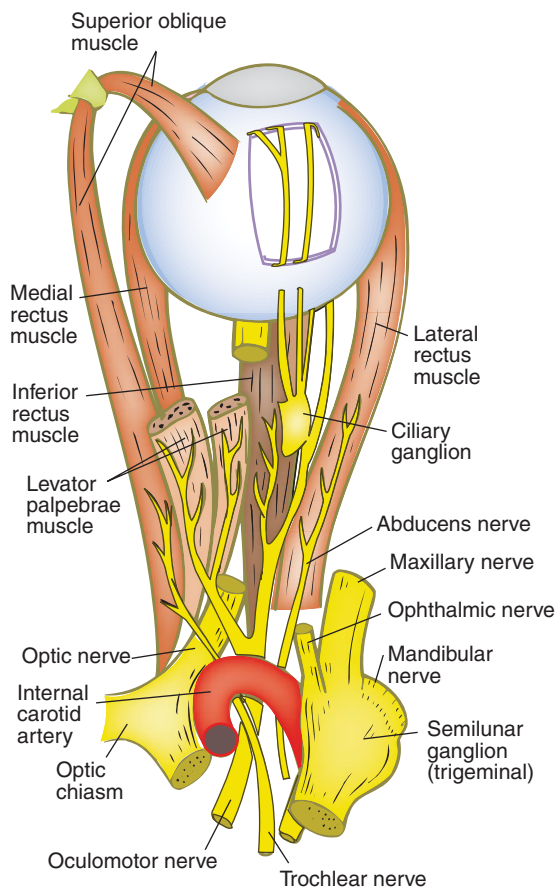


FIGURE 27.16 The right eye with muscles and nerves, as viewed from above. Three cranial nerves innervate the muscles of the eye. Some of the extraocular muscles and the optic nerve have been cut. The external layers of the eye bulb have been partly removed to expose the postganglionic fibers from the ciliary ganglion on their way to the intrinsic eye muscles. Note the position of the trigeminal ganglion and the internal carotid artery.

The Abducens Nerve

The abducens nerve leaves the brain stem close to the midline at the junction of the medulla and the pons (Fig. 27.1; see Fig. 6.15). Figure 27.11 shows the location of the abducens nucleus and its relation to the facial nerve. The abducens nerve runs forward intracranially and passes through the **cavernous sinus** before entering the orbit through the superior orbital fissure (Fig. 27.16). It supplies only one muscle, the **lateral rectus**. The muscle pulls the eye so the cornea faces laterally (abducts the eye; see Fig. 25.2). Even though other muscles can abduct the eye somewhat (the superior and the inferior oblique muscles), the lateral rectus is necessary for more than a slight lateral movement (Fig. 27.17). A person with a unilateral lesion of the abducens nerve usually keeps the head turned somewhat to the side of the lesion to compensate for the loss of lateral motion of the eye.

The Trochlear Nerve

This is the only one of the cranial nerves to leave the brain stem on the dorsal side (just below the inferior colliculus) (Fig. 27.1; see Fig. 6.19).⁶ The **trochlear nucleus** lies a little ventrally to the aqueduct in the mesencephalon (Fig. 27.2). Like the abducens nerve, the trochlear traverses the cavernous sinus and enters the orbit through the **superior orbital fissure**. It innervates the **superior oblique muscle** (Fig. 27.16; see Fig. 25.1), which directs the gaze downward and laterally (see Fig. 25.2).

The Oculomotor Nerve

The nerve emerges from the ventral aspect of the mesencephalon, in the interpeduncular fossa (Fig. 27.1; see Fig. 6.20). This is the largest of the three nerves supplying the extraocular muscles; as mentioned, it contains somatic efferent and visceral efferent fibers. The **somatic efferent** fibers come from the large **oculomotor nucleus** (nucleus of the oculomotor nerve) situated close to the

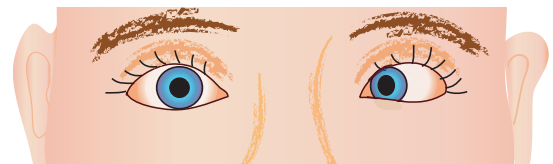


FIGURE 27.17 Paresis of the right abducens nerve. When the patient looks to his right, the right eye does not follow the left, because the right rectus muscle is paralyzed.

⁶ Another peculiarity of the trochlear nerve is that it crosses the midline before leaving the brain stem, so that the left trochlear nucleus innervates the right superior oblique muscle, and vice versa. Some of the fibers of the oculomotor nerve also cross before leaving the brain stem.

midline in the mesencephalon, ventral to the aqueduct (Figs. 27.2 and 27.5). The medial longitudinal fasciculus, with ascending fibers from the vestibular nuclei, lies close to the oculomotor nucleus (and to the abducens and trochlear nuclei as well). The **visceral efferent** (preganglionic parasympathetic) fibers come from the small **nucleus of Edinger-Westphal** located near the oculomotor nucleus. Often the term “oculomotor complex” is used for the somatic efferent and visceral efferent nuclei together.

The **oculomotor nerve** passes forward to the orbit through the **cavernous sinus** together with the other nerves to the eye (entering through the superior orbital fissure). The somatic efferent and parasympathetic fibers part in the orbit (Fig. 27.16). The somatic efferent fibers innervate the following extraocular muscles: the **superior** and **inferior rectus**, the **medial rectus**, and the **inferior oblique**. These muscles can move the eye medially, upward, and downward and rotate it around the sagittal axis (see Fig. 25.2). In addition, the oculomotor nerve supplies the **levator palpebrae** muscle, which serves to lift the upper eyelid.

The **visceral efferent** oculomotor fibers end in the small **ciliary ganglion** situated behind the eye (Fig. 27.16). Here the fibers establish synapses with the postganglionic neurons, which send their axons anteriorly in the wall of the eye to innervate the intrinsic (smooth) muscles of the eye: the **pupillary sphincter** and the **ciliary muscle** (see Fig. 16.2). Contraction of the ciliary muscle increases the lens curvature when looking at near objects (see Chapter 16, under “The Lens and the Far and Near Points of the Eye: Accommodation”). The sphincter constricts the pupil to reduce the amount of light reaching the retina.

A **lesion** of the **oculomotor nerve** produces, among other symptoms, an abnormal position of the eye, which is directed laterally (due to the unopposed pull of the lateral rectus) and downward (due to the superior oblique

muscle). As in lesions of the abducens or the trochlear nerves, the patient will have double vision. In addition, the upper eyelid droops—**ptosis**—because of paralysis of the levator palpebrae (Fig. 27.18). The **interruption** of the **parasympathetic fibers** makes the pupil larger (due to loss of action of the pupillary sphincter), and the light reflex is absent (in an incomplete lesion of the nerve, the pupil may be slightly larger and the reaction to light more sluggish than on the normal side). **Accommodation** of the lens is abolished, making it impossible to see near objects sharply. The intracranial course of the oculomotor nerve makes it especially vulnerable in cases of temporal herniation caused by increased intracranial pressure (see Chapter 3). Thus, examination of the size of the pupils, and their reaction to light, is of great practical value in patients who are unconscious after **head trauma**.

The Light Reflex and the Accommodation Reflex

The oculomotor nerve is the efferent link of both of these reflexes, even though they are quite different in other respects. The **light reflex** is relatively simple, with its reflex center in the brain stem (Fig. 27.19). Increased amount of light hitting the retina elicits a contraction of the pupillary sphincter muscle. Both pupils constrict even when the light hits only one eye. The afferent link consists of fibers of the optic nerve that leave the optic tract before it reaches the lateral geniculate body. The fibers end in the pretectal nuclei on both sides. From these nuclei, the signals pass to the Edinger-Westphal nuclei on both sides, and by means of the ciliary ganglion, the signals reach the sphincter. The bilaterality of the connections explains why a unilateral stimulus produces a bilateral response.

Unilateral **interruption** of the **oculomotor nerve** abolishes the light reflex in the eye on the side of the lesion, but the reflex is present in the other eye. In case

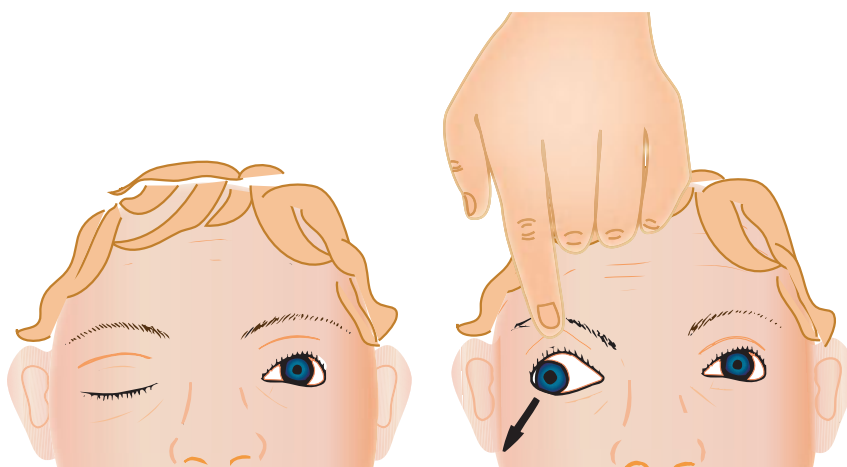


FIGURE 27.18 Paresis of the oculomotor nerve (right side). **Left:** The upper eyelid droops (ptosis) due to paresis of the levator palpebrae muscle. **Right:** The examiner lifts the eyelid and reveals that the right eye is abducted and lowered due to the unopposed actions of the lateral rectus and the superior oblique muscles.

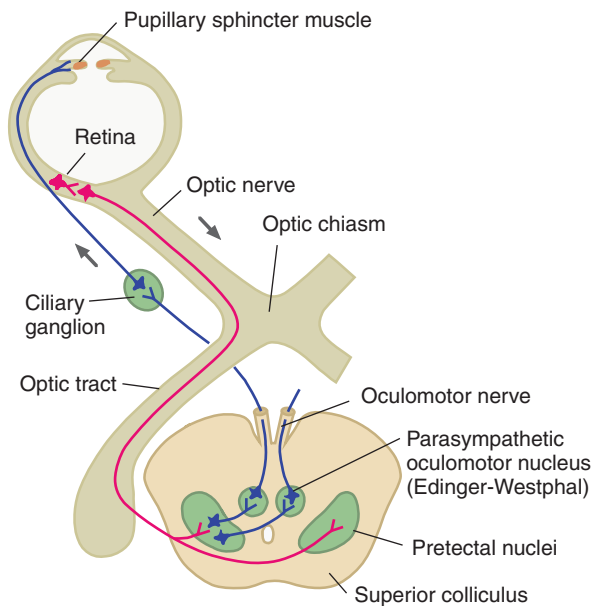


FIGURE 27.19 The reflex arc for the light reflex.

of interruption of the **afferent link** on one side (damage to the retina or the optic nerve), the light reflex is absent in both eyes when light is shone into the eye on the lesioned side but is present in both sides when the other eye is illuminated. Thus, examination of the light reflex can provide valuable information with regard to the site of a lesion.

The **accommodation reflex** is a cortical reflex: the reflex arc passes through the cerebral cortex. The afferent link is fibers passing in the optic nerve from the retina, and the efferent link consists of parasympathetic fibers in the oculomotor nerve to the ciliary ganglion (Fig. 27.20). From there the postganglionic fibers pass to the ciliary muscle. The reflex center is not known in detail, but recent studies indicate that the cortical fibers pass to the superior colliculus (not the pretectal nuclei,

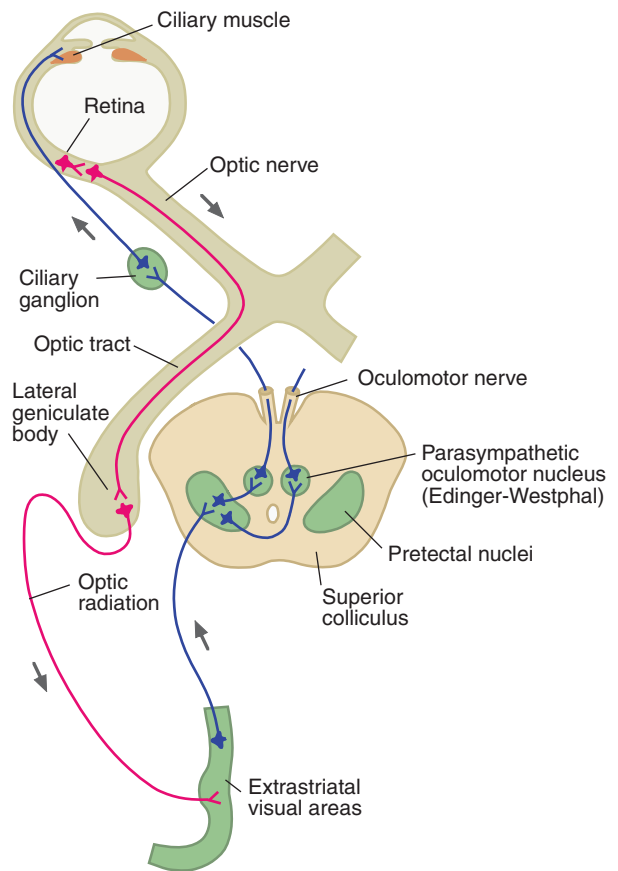


FIGURE 27.20 The reflex arc for the accommodation reflex.

as shown in Fig. 27.18) and from there to the nearby reticular formation. Neurons in the reticular formation mediate the signals to the Edinger-Westphal nucleus. The accommodation reflex is elicited only when we fix the gaze on an object that is moving toward us. Together with the accommodation, a **pupillary constriction** occurs as the object comes closer.

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VI

THE AUTONOMIC NERVOUS SYSTEM

THE autonomic nervous system (or visceral system) is not a term that can be defined precisely, either anatomically or functionally. The old belief that the somatic and the autonomic parts of the nervous system are completely independent is not tenable. The more we have learned about the nervous system, the clearer it has become that simplistic divisions like this are arbitrary. Some authors therefore maintain that the term “autonomic nervous system” should be abandoned and replaced by simply referring to **visceral neurons**—that is, the neurons that innervate visceral organs. For practical reasons, it is nevertheless helpful to use the term “autonomic nervous system” and to **define** it very broadly as the neuronal groups and fiber connections that control the activity of **visceral organs, vessels, and glands** (also the vessels and glands that are not parts of visceral organs). Visceral organs contain smooth-muscle cells and glandular cells. We can therefore also define the autonomic system as the parts of the nervous system that control the activity of **smooth muscles and glands**, regardless of their location in the body (in contrast to the somatic or cerebrospinal system, which controls striated skeletal muscles). Mostly, we are not aware of the processes going on in the organs controlled by the autonomic nervous system, and their activities are not subject to voluntary, conscious control.

The autonomic system can be subdivided in different ways. As with the somatic system, we distinguish **peripheral** and **central** parts. Whereas the peripheral parts of the autonomic and somatic systems can be separated fairly well, the division becomes much less clear within the central nervous system. The peripheral parts of the autonomic system are described in Chapters 28 and 29. Although the nerves to visceral organs contain both efferent (motor) and afferent (sensory) fibers, the anatomic differences between the autonomic and the somatic systems concern primarily the efferent side. Chapter 28 deals with the **visceral efferent neurons**,

while Chapter 29 describes the characteristic features of **visceral afferent neurons**—that is, the sensory innervation of visceral organs. Even though visceral afferent neurons do not differ structurally from somatic afferent neurons, there are nevertheless physiological differences that are important in a clinical context—in particular, in relation to pain arising in the visceral organs. Although Chapter 29 contains discussion of some visceral reflexes and their central control, main topics concerning the central control of the autonomic system, notably the hypothalamus, are discussed in Chapter 30.

The autonomic system can also be divided into a **sympathetic** and a **parasympathetic** part, differing anatomically and functionally (again, the separation is most obvious in the peripheral nervous system). The actions of the two parts of the autonomic system are often antagonistic: where the sympathetic system activates, the parasympathetic reduces the activity, and vice versa. In addition, the **enteric nervous system** is now regarded as a distinct part of the autonomic nervous system. It consists of some million neurons and an extensive network in the wall of the digestive tract. The enteric system is influenced by both the sympathetic and the parasympathetic systems but functions independently as well.

In a superior perspective, a major function of the autonomic system is to contribute to **bodily homeostasis**, that is, the maintenance of a relatively constant internal milieu. The autonomic nervous system is not the only means by which the central nervous system can control homeostasis, however. In addition, the **endocrine system**, which is controlled from the central nervous system via the pituitary gland, serves this purpose. Further, homeostasis requires appropriate somatic motor activity to secure, for example, the supply of water and nutrients. The role of the **hypothalamus** is to organize the autonomic, endocrine, and somatic-motor processes into behavior that is appropriate for the immediate and long-term needs of the organism.

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Visceral Efferent Neurons: The Sympathetic and Parasympathetic Divisions

OVERVIEW

In general, the actions of the sympathetic system suggest that it is of special importance in situations of stress, requiring mobilization of bodily resources. The parasympathetic system, in contrast, contributes primarily to processes of maintenance, such as digestion and reproductive behavior. Such generalized statements are useful only as rules of thumb, however, because the actions of the two systems are much more varied than this. Table 28.1 gives an overview of the main actions of the two systems in various organs.

The sympathetic and parasympathetic system both consist of two consecutive neurons leading from the central nervous system (CNS) to the target organs. The **preganglionic neuron** has its cell body in the cord or in the brain stem and the axon terminates in a **ganglion**. **Postganglionic neurons** have their cell bodies in the ganglia while their axons terminate close to **smooth muscle cells** and **glandular cells**. The **preganglionic sympathetic neurons** are found in the **intermediolateral column** of the cord, stretching from T_1 to L_2 , whereas the **preganglionic parasympathetic cell bodies** reside in the **brain stem** and in the **sacral cord**. All **preganglionic neurons** use **acetylcholine** as transmitter in the ganglia, where they act on **nicotinic receptors**. Most **sympathetic neurons** use **norepinephrine** (and various neuropeptides) as transmitter, while **parasympathetic neurons** use **acetylcholine** binding to **muscarinic receptors**.

Preganglionic sympathetic and parasympathetic fibers leave the cord and the brain stem through ventral roots and motor cranial nerves. Their further course differs, however. The **parasympathetic fibers** pass directly to ganglia that lie close to or in the wall of the target organ. The **sympathetic fibers** leave the spinal nerves just outside the intervertebral foramen and enter the **sympathetic trunk**, which stretches from the base of the skull to the lower end of the cord. The **sympathetic trunk** forms a chain of interconnected sympathetic ganglia. From these ganglia, **sympathetic postganglionic fibers** follow the spinal nerves to the extremities and

somatic structures of the trunk. **Preganglionic sympathetic fibers** destined for visceral organs pass through the ganglia and leave the sympathetic trunk in separate **splanchnic nerves**. These end in **prevertebral ganglia**, and from there **postganglionic fibers** follow arteries to the target organs.

The **enteric system** consists of some million neurons in the wall of the gastrointestinal tract, controlling peristaltic movements, secretion, and absorption. While it operates to a large extent independently of the CNS, this “mini-brain of the gut” integrates local information from the gut with central commands through the sympathetic and parasympathetic systems.

GENERAL ORGANIZATION

Two Succeeding Neurons Constitute the Efferent Pathway

Despite their different actions, the sympathetic and the parasympathetic systems have certain features of peripheral organization in common that should be known before discussing the differences between them. In contrast to the somatic efferent fibers (leading from motoneurons to skeletal muscles), the visceral efferent fibers do not pass nonstop to the organs. The transmission of signals is synaptically interrupted in **autonomic ganglia** (Figs. 28.1 and 28.2). The multipolar neurons in the ganglia send their axons to the effectors (smooth-muscle cells and glands) in the visceral organs. The neurons conducting signals to the ganglia are called **preganglionic**, whereas the ganglion cells and their axons are called **postganglionic**. The cell bodies of the **preganglionic neurons** are located in the spinal cord and the brain stem. The **preganglionic neurons**, both the sympathetic and the parasympathetic ones, use **acetylcholine** as a neurotransmitter.

Another typical feature of the visceral efferent neurons is that, as a rule, the **postganglionic fibers** form extensive **plexuses** around the organs they innervate.

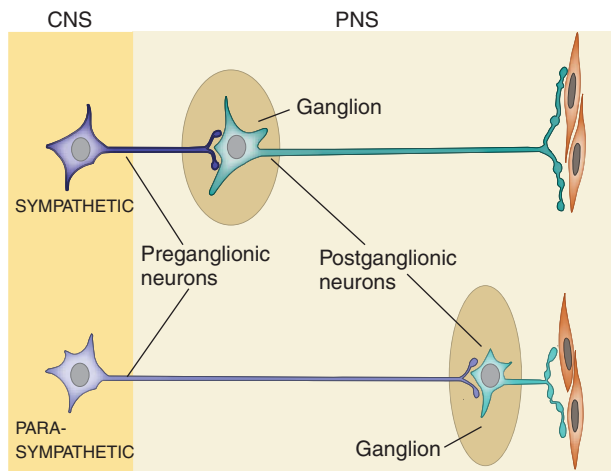


FIGURE 28.1 Basic organization of the peripheral part of the autonomic system. Two consecutive neurons conduct the signals from the central nervous system to the effectors. Note the difference in length between the pre- and postganglionic fibers in the sympathetic and the parasympathetic systems. Effectors may be glandular cells and not only smooth-muscle cells, as shown here.

Postganglionic Fibers Do Not Establish Typical Synapses

The thin, unmyelinated postganglionic fibers do not form typical synapses with the effector cells; in contrast to the somatic efferent fibers innervating skeletal muscle cells (see Fig. 21.5). When the postganglionic fibers reach the vicinity of the effector cells, they branch extensively, and there are small swellings—**varicosities**—along the branches (Fig. 28.1). In each varicosity, there are vesicles with neurotransmitters. The varicosities do not form synapses but can be located fairly close to the effector cells. In some places, the distance is only 50 nm, thus permitting very direct and precise control of the effector cell. Most often, however, the distance between the varicosities and the effectors is so great that the neurotransmitter, after being released, must diffuse over a considerable distance to reach its target. This means that the neurotransmitter acts on several effector cells within a certain distance. When the diffusion distance is great, the transmitter acts slowly, with a long latency and a prolonged action (compared with the fast action at the neuromuscular junction). Often, only a few of the smooth-muscle cells in the wall of a hollow organ (such as a vessel) are close enough to the varicosities to be directly influenced by the transmitter. In such cases, the action potential elicited in some smooth-muscle cells is propagated from cell to cell via **gap junctions** between them (electric coupling). In general, therefore, the actions of the autonomic nervous system are more **diffusely distributed**, both **spatially** and **temporally**, than is the case in the somatic system. The properties of the smooth-muscle cells contribute to increase this difference, because their action potentials last much longer

than those of skeletal muscle cells do. Further, smooth-muscle cells (e.g., in the wall of the gastrointestinal tract) can be made to contract by stimuli other than nervous ones, such as by stretching and by the actions of hormones. The autonomic system therefore contributes to the regulation of the contraction of smooth-muscle cells in the gastrointestinal tract and in the walls of the vessels, but it is not alone in this capacity (again in contrast to the control of skeletal muscle cells by the motoneurons).

Some Organs Are Subject to More Precise Autonomic Control than Others

There are, nevertheless, great differences between different organs with regard to the precision of their autonomic innervation. Some visceral organs require much faster and more accurate control than others. The smooth muscles of the **eye** (the ciliary muscle and the pupillary sphincter, regulating the curvature of the lens and the diameter of the pupil) must be subject to very precise control. The same holds for the muscles of the **ductus deferens**, in which the propulsive contractions must be fast and well coordinated. Such demands are not made of the smooth muscles of the gastrointestinal tract and the vessels. Corresponding to different functional requirements, the pattern of autonomic innervation varies in different organs. The intrinsic eye muscles, for example, receive a large number of nerve fibers so that all muscle cells come in close contact with varicosities of nerve fibers. This is called **multiunit arrangement**, because it is similar to the conditions in the somatic system with many precisely controlled motor units. In places with few nerve fibers to supply a large number of smooth-muscle cells, so that only a few cells are close to nerve varicosities, we use the term **single-unit arrangement**. Thus, numerous muscle cells behave as a unit when one or a few of them are activated.

Autonomic Ganglia and Plexuses

The autonomic ganglia contain **multipolar neurons** of various sizes (Fig. 28.3) with long, branching dendrites. The axons are mostly unmyelinated and very thin. The cell bodies of the ganglion cells are embedded in a meshwork of fibers consisting of afferent fibers, ganglion cell dendrites, and axons of the ganglion cells. At least some of the ganglia receive **sensory fibers** from visceral organs. There are also **interneurons** in the autonomic ganglia. Such features, and other data, indicate that the ganglia are not just simple synaptic interruptions of purely motor (efferent) pathways but can serve as **reflex centers** for some visceral reflexes.

When an organ is innervated by both sympathetic and parasympathetic fibers, these two components intermingle and form **autonomic plexuses** just outside

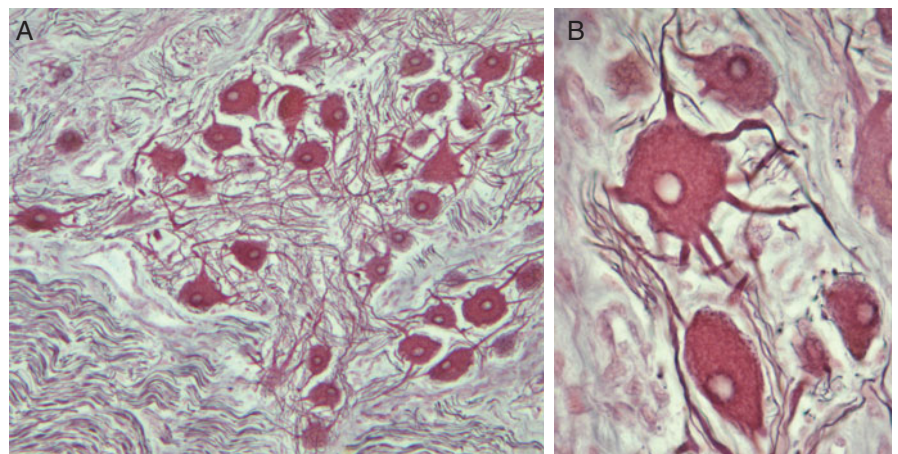
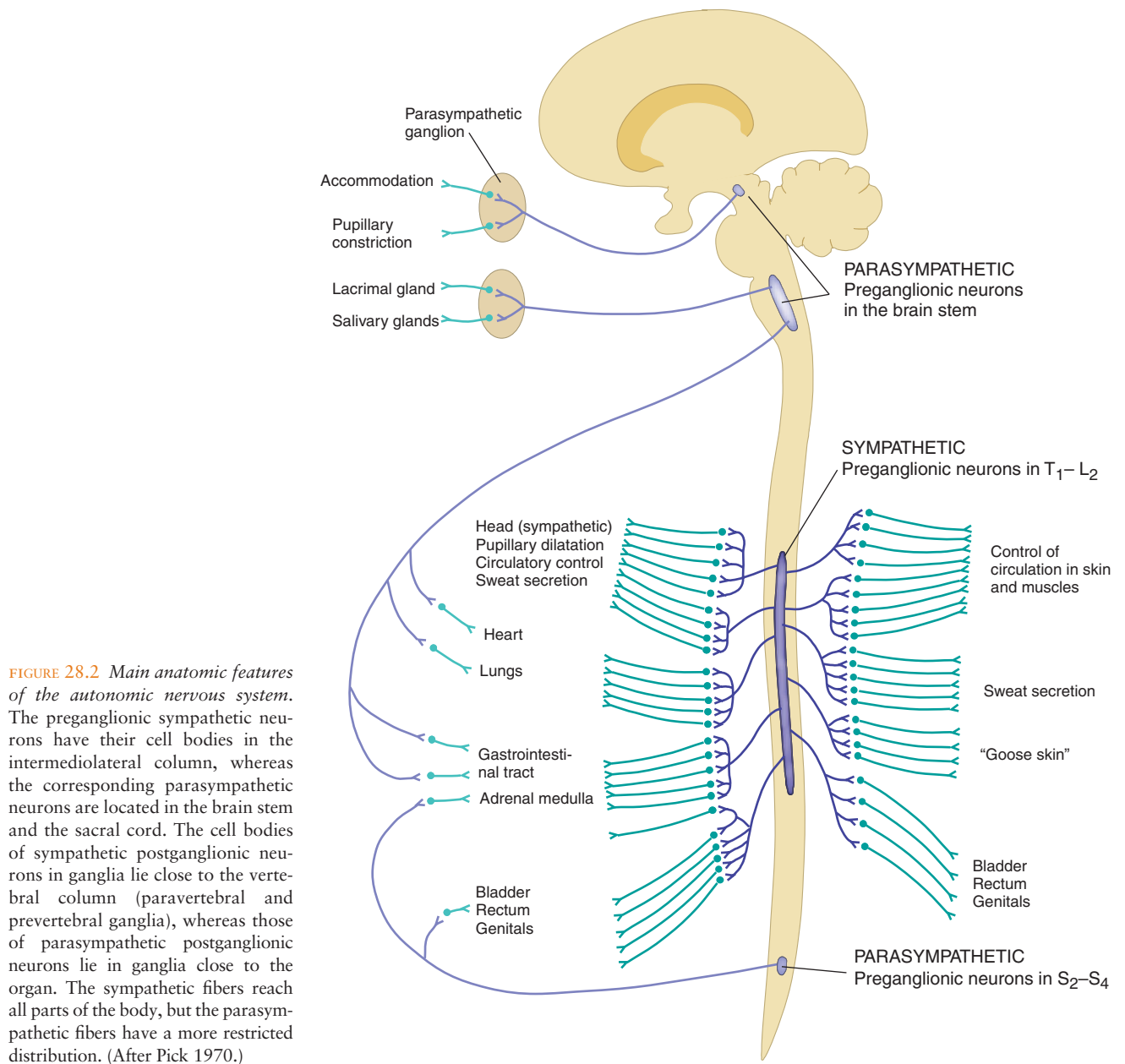


FIGURE 28.3 Autonomic ganglion. Photomicrograph of section through the celiac ganglion impregnated with heavy metals to show nerve fibers (black) and neurons (red). **A:** The multipolar postganglionic neurons lie in a wickerwork of preganglionic and postganglionic nerve fibers. Thus, the autonomic ganglia consist of groups of postganglionic neurons embedded in a plexus. **B:** Higher magnification of postganglionic neurons.

or in the wall of the organs. At many places, cell bodies of postganglionic cells form small clumps intermingled with the fibers of the plexuses. Thus, ganglia and plexuses often coexist, and the ganglia then have the same prefix as the plexuses. For example, the **cardiac plexus** on the outside of the heart contains both kinds of fibers and, in addition, the cell bodies of the postganglionic parasympathetic neurons. Especially well developed plexuses are found around the large vessels in the upper abdomen, where “parasympathetic vagus fibers intermingle with sympathetic fibers. These **prevertebral plexuses** get their names from the arteries they surround and follow peripherally: the **celiac plexus**, the **superior** and **inferior mesenteric plexus**, and the **renal plexus**. The prevertebral plexuses continue into the pelvis as the **hypogastric plexus**.

Differences between the Sympathetic and the Parasympathetic Systems

The two systems differ in several respects. The location of **preganglionic neurons** differs: the sympathetic ones are found in the T₁–L₂ spinal segments, whereas the parasympathetic preganglionic neurons lie in the brain stem and the S₂–S₄ spinal segments (Fig. 28.2). Further, the **ganglia** are located differently: sympathetic ones lie close to the CNS, whereas the parasympathetic ganglia are located close to the target organs. Thus, the sympathetic preganglionic fibers are short, and the parasympathetic ones are long (Figs. 28.1 and 28.2).

Another difference between the two systems is the **neurotransmitters** used by the **postganglionic neurons**: the sympathetic fibers release **norepinephrine**, whereas the parasympathetic fibers release **acetylcholine** from their varicosities.

Finally, the **distribution** of postganglionic fibers is different. Thus, virtually all parts of the body receive sympathetic fibers, whereas the parasympathetic fibers

are mostly restricted to the true visceral organs. The body wall and the extremities (skin, muscles, joints) do not receive parasympathetic fibers.

Table 28.1 gives a broad overview of the functions of the two systems; a more comprehensive discussion is provided later in this chapter.

The Parasympathetic Innervation Is Usually More Precise than the Sympathetic

As a rule (with notable exceptions), the sympathetic system is more diffusely organized than the parasympathetic. This is evident, for example, in the relation between the number of preganglionic and postganglionic fibers. In the parasympathetic **ciliary ganglion** (see Fig. 27.16), two postganglionic fibers leave the ganglion for each preganglionic fiber reaching it—that is, a 2:1 relationship (in the cat). For the sympathetic **superior cervical ganglion** (Fig. 28.7), the relationship is 30:1 in the cat and 60 to 190:1 in humans (this ganglion contains more than 1 million neurons in humans). Further, the **parasympathetic** innervation is in several places arranged with **multiunits**—that is, small “motor units” with the possibility of precise control. This concerns, for example, the innervation of the intrinsic eye muscles. The **sympathetic** innervation is often (but not always) arranged with **single units**—that is, a large number of smooth muscle cells are activated from one postganglionic fiber and behave as a functional unit.

Another difference concerns the **topographic arrangement** of the preganglionic neurons. Whereas the sympathetic neurons show only a fairly rough topography within the intermediolateral column in relation to the location of the target (Fig. 28.7; Table 28.2), the parasympathetic neurons are as a rule collected in distinct nuclei (Figs. 28.2 and 28.11) or subdivisions of a nucleus (like the motor nucleus of the vagus), each related to one target organ.

TABLE 28.1 Summary of Some Main Actions of the Autonomic System

Organ	Sympathetic System	Parasympathetic System
Arteries (arterioles)	Vasoconstriction	Vasodilatation in glands and genital organs
Skin	Vasoconstriction; sweat secretion; goose skin	No innervation
Skeletal muscles	Vasoconstriction	No innervation
Heart	Increased rate and stroke volume	Reduced heart rate
Airways	Relaxation of bronchial smooth muscle (reduced airway resistance)	Contraction of bronchial smooth muscle; secretion from mucous glands
Gastrointestinal tract	Reduced blood flow and peristaltic movements	Increased peristaltic movements; secretion of gastric juice, bile, and pancreatic juice
Urinary bladder	Relaxation of muscle for emptying (detrusor)	Contraction of detrusor; relaxation of urethral sphincter
Rectum	Relaxation of muscle for emptying; closure of internal sphincter (smooth muscle)	Contraction of muscle for emptying; relaxation of internal sphincter
Genital organs	Contraction of the ductus deferens (ejaculation)	Erection (penis, clitoris)
Eye	Pupillary dilatation	Pupillary constriction; accommodation; secretion of tears

see Table 28.2 for further details.

TABLE 28.2 *Principal Features of Autonomic Innervation*

Organ	Sympathetic			Parasympathetic		
	Preganglionic Neuron	Postganglionic Neuron	Action	Preganglionic Neuron	Postganglionic Neuron	Action
Eye	T ₁ –T ₂	Superior cervical ganglion	Pupillary dilatation	Parasympathetic oculomotor nucleus (Edinger Westphal)	Ciliary ganglion	Pupillary constriction accommodation
Lacrimal gland	T ₁ –T ₂	Superior cervical ganglion	?	Superior salivatory nucleus	Pterygopalatine ganglion	Secretion
Submandibular and sublingual glands	T ₁ –T ₂	Superior cervical ganglion	Vasoconstriction	Superior salivatory nucleus	Submandibular ganglion	Secretion, vasodilatation
Parotid gland			Vasoconstriction	Inferior salivatory nucleus	Otic ganglion	Secretion, vasodilatation
Heart	T ₁ –T ₄₍₅₎	Superior, middle, and inferior cervical 1 upper thoracic ganglia	Increased rate and stroke volume	Dorsal motor vagus nucleus	Cardiac plexus	Reduced heart rate
Bronchi, lungs	T ₁ –T _{3–4}	Inferior cervical 1 upper thoracic ganglia	Bronchial dilatation	Dorsal motor vagus nucleus	Pulmonary plexus	Bronchial constriction and secretion
Stomach	T ₆ –T ₁₀ (greater splanchnic nerve)	Celiac ganglion	Inhibition of peristaltic movement and secretion; contraction of pyloric sphincter	Dorsal motor vagus nucleus	Gastric plexus	Peristaltic movement and secretion
Pancreas	T ₆ –T ₁₀ (greater splanchnic nerve)	Celiac ganglion	?	Dorsal motor vagus nucleus	Periarterial plexus	Secretion
Small intestine, ascending and trans-verse large intestine	T ₆ –T ₁₀ (greater splanchnic nerve)	Celiac ganglion 1 superior and inferior mesenteric ganglia	Inhibition of peristaltic movement and secretion	Dorsal motor vagus nucleus	Myenteric and submucous plexus (ganglion)	Peristaltic movement and secretion
Descending large intestine, sigmoid and rectum	T ₁₀ –L ₂	Inferior mesenteric, hypogastric, and pelvic ganglia (plexuses)	Inhibition of peristaltic movement and secretion	Sacral cord S ₃ –S ₄	Myenteric and submucous plexus (ganglion)	Peristaltic movement and secretion
Ureter, bladder	T ₁₀ –L ₂	Hypogastric and pelvic ganglia (plexuses)	Inhibition of detrusor	Sacral cord S ₃ –S ₄	Pelvic plexus	Contraction of detrusor
Head and neck (skin and skeletal muscles)	T ₁ –T _{2–4}	Superior cervical ganglion	Vasoconstriction sweat secretion, and piloerection	—	—	—
Upper extremity	T ₃ –T ₆	Stellate ganglion, upper thoracic	Vasoconstriction sweat secretion, and piloerection	—	—	—
Lower extremity	T ₁₀ –L ₂	Lower lumbar and upper sacral ganglia	Vasoconstriction, sweat secretion, and piloerection	—	—	—

The difference in innervation precision between the sympathetic and the parasympathetic systems mentioned above have several **exceptions**, however. One example is the ductus deferens, in which the rhythmic contractions are elicited by sympathetic fibers. A multi-unit arrangement is required to ensure the necessary precision and speed of the contractile wave moving the sperm during ejaculation. Another example is that the parasympathetic innervation of the gastrointestinal tract is rather diffuse (single-unit arrangement).

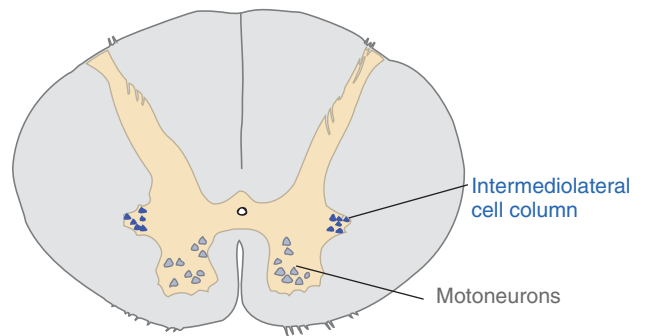


FIGURE 28.4 The intermediolateral cell column contains the cell bodies of the preganglionic sympathetic neurons. Cross section of the thoracic cord.

PERIPHERAL PARTS OF THE SYMPATHETIC SYSTEM

Preganglionic Fibers and the Sympathetic Trunk

The peripheral parts of the sympathetic system consist of both neurons conveying signals to visceral organs and sensory fibers leading in the opposite direction. The efferent, preganglionic sympathetic neurons have their cell bodies in the **intermediolateral column** in the spinal cord (Fig. 28.4). The preganglionic fibers leave the cord (like other efferent fibers) through the ventral roots, but because the intermediolateral column is present

in only the T_1 – L_2 segments, only the ventral roots of these segments contain preganglionic sympathetic fibers (the sympathetic system is also called the thoracolumbar system). The sympathetic fibers follow the somatic ones for a short distance, however. Just after the ventral and the dorsal roots fuse, the sympathetic preganglionic fibers leave the spinal nerve to end in a **sympathetic ganglion** (Fig. 28.5). In early embryonic life, one ganglion is

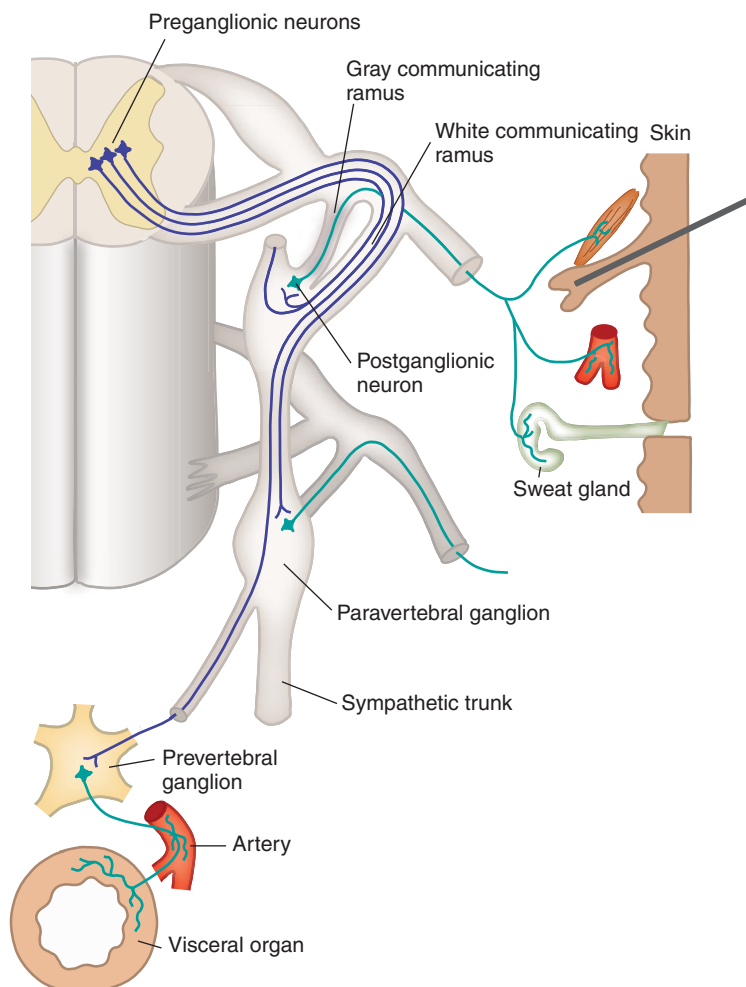


FIGURE 28.5 The sympathetic system. Postganglionic neurons are located in the ganglia of the sympathetic trunk and in the prevertebral ganglia. Sympathetic fibers to the trunk and the extremities follow the spinal nerves, whereas fibers to the visceral organs form separate nerves and follow the main vessels to the organs.

produced on each side for every spinal segment, but during further development, some ganglia fuse, so the final number is smaller than the number of segments. This reduction is most marked in the cervical region.

The ganglia are located just outside the intervertebral foramen, laterally on the vertebral column (Fig. 28.6). The row of such **paravertebral ganglia** extends from the base of the skull to the coccygeal bone in the pelvis minor. Because fiber bundles interconnect the ganglia, a continuous string called the **sympathetic trunk** is formed (Figs. 28.5 and 28.6). The ganglia form small swellings along the trunk. There is usually one ganglion for each pair of spinal nerves, except in the cervical region, where there are only three: the **superior, middle, and inferior cervical ganglia**. The middle ganglion can be missing, and the inferior cervical ganglion is usually fused with the uppermost thoracic ganglion to form the large **stellate ganglion** (Fig. 28.6). Some cross connections are present in the lumbar and sacral parts of the sympathetic trunks.

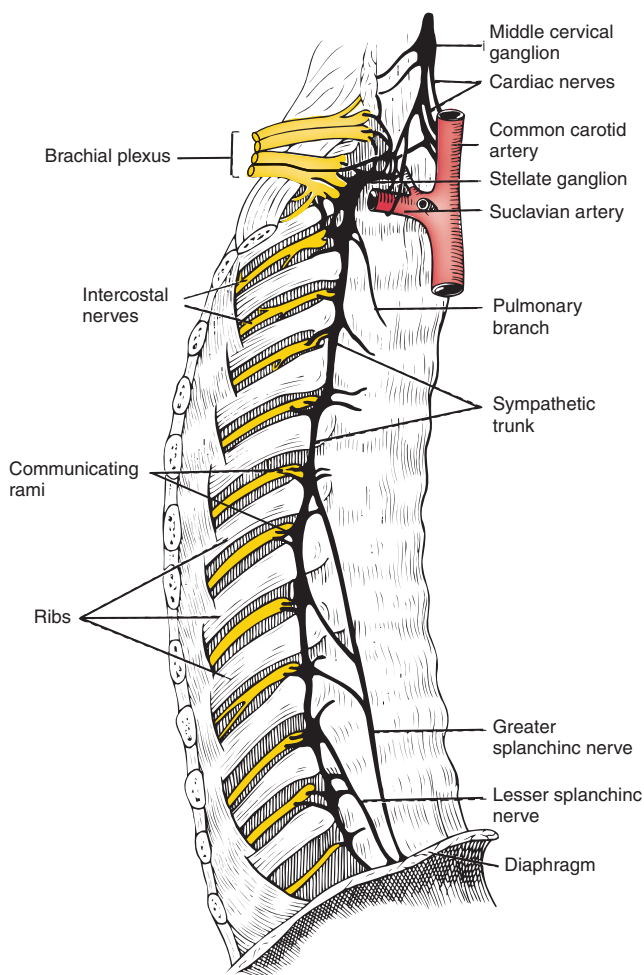


FIGURE 28.6 *The sympathetic trunk.* Part of the thoracic vertebral column and the ribs, as viewed from the right. Note the communicating rami and the splanchnic nerves. (Redrawn from Spalteholz 1933.)

The preganglionic sympathetic fibers leave the spinal nerve as a small bundle called the **white communicating ramus** (branch), which connects the nerve with the sympathetic trunk (Figs. 28.5 and 28.6). The whitish color arises because the preganglionic fibers are myelinated. Because of the restricted extension of the intermediolateral column, only the thoracic and the upper two lumbar spinal nerves give off white communicating rami.

When reaching the ganglia of the sympathetic trunk, some preganglionic fibers establish synapses with **postganglionic neurons** in that ganglion, whereas others continue uninterrupted through the ganglion (Fig. 28.5). In the upper part of the trunk, the fibers continue rostrally, in the lower part caudally, to establish synapses with ganglion cells in ganglia at levels above and below the intermediolateral column (Figs. 28.7, 28.8, and 28.9). This arrangement ensures that preganglionic fibers reach all the ganglia of the sympathetic trunk (the paravertebral ganglia).

Some of the preganglionic fibers pass directly through the ganglia and form separate nerves—**splanchnic nerves**—destined for **prevertebral sympathetic ganglia** (Figs. 28.5 and 28.7). We return to this point later in this chapter.

Postganglionic Sympathetic Fibers

The postganglionic fibers from the ganglia of the sympathetic trunk take different routes. From all of the ganglia, some fibers pass back to the spinal nerve as the thin **gray communicating ramus** (Figs. 28.5 and 28.6). This ramus is more grayish than the white ramus because most of the postganglionic fibers are unmyelinated.¹ All of the spinal nerves receive postganglionic fibers through the communicating rami (Fig. 28.6). The postganglionic fibers follow the spinal nerves out into all their branches (Figs. 28.8 and 28.9). Some postganglionic fibers leave the trunk, pass to larger arteries in the vicinity, and innervate the smooth-muscle cells of the artery. Many of the postganglionic sympathetic fibers that follow the spinal nerves leave them peripherally to innervate small vessels. Other fibers from the spinal nerves innervate the sweat glands and the smooth muscles attached to hair follicles (Fig. 28.5).

Sympathetic Innervation of the Head and Extremities

The head, neck, and upper extremity receive **preganglionic** sympathetic fibers from the **upper thoracic segments** (Table 28.2). The fibers enter the sympathetic trunk

¹ White and gray rami often fuse into one, so that even at the levels T₁–L₂ there may be only one communicating ramus on each side. This contains, as will be understood, both the pre- and postganglionic fibers. In the case of two rami, the color difference between them is not very marked.

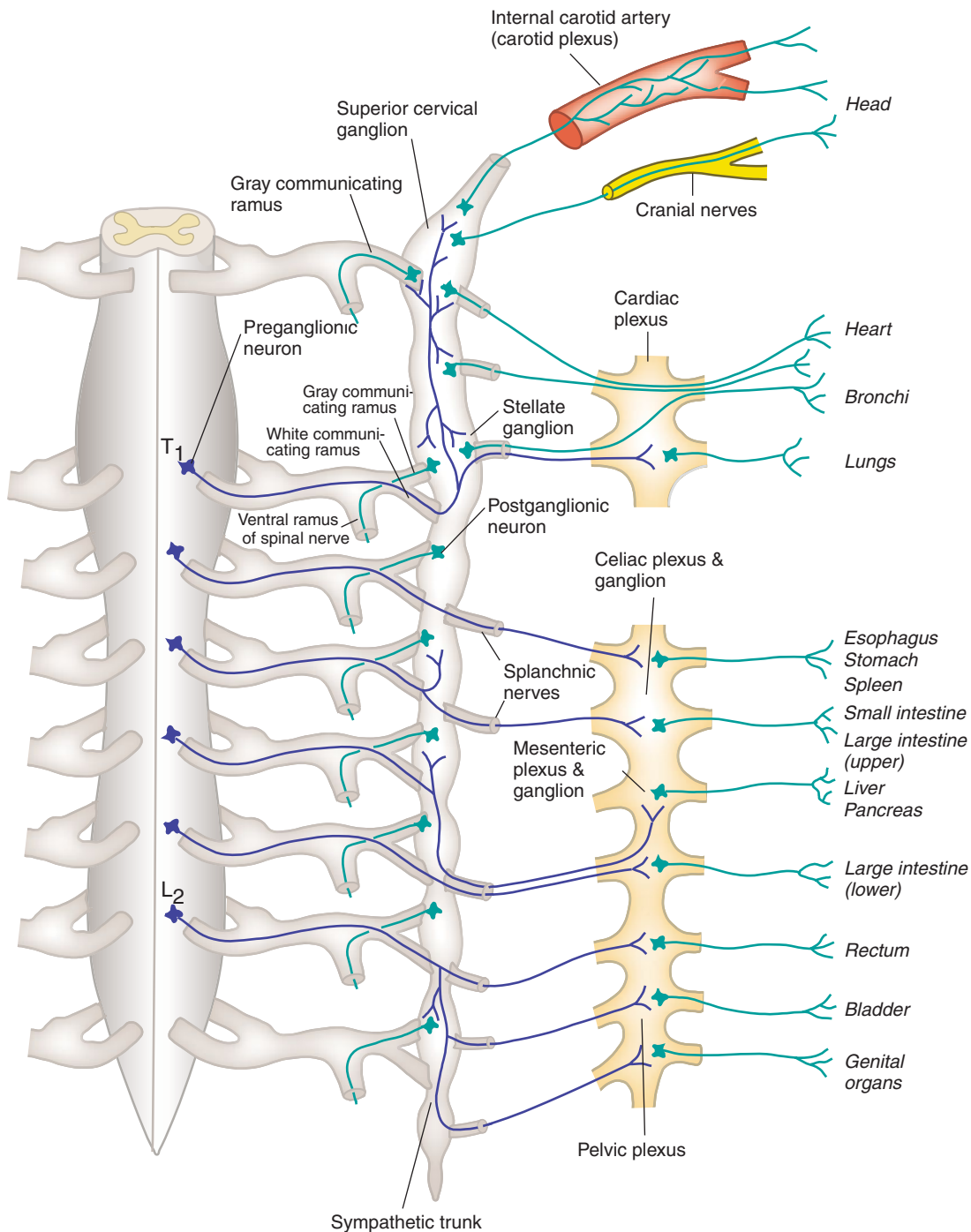


FIGURE 28.7 *The sympathetic system.* The sympathetic trunk with the paravertebral ganglia, and the prevertebral ganglia and plexuses

are shown. The organs innervated from the various ganglia are also specified.

through the communicating rami. Some establish synaptic contacts with postganglionic neurons in the upper thoracic ganglia, whereas other fibers pass through these ganglia to end in the cervical ganglia (Figs. 28.7 and 28.8). From these ganglia, postganglionic fibers enter the spinal nerves to the neck (C₁–C₄) and the upper extremity (C₅–T₁).

The **head** receives **postganglionic** fibers from the **superior cervical ganglion**. From the ganglion, the fibers follow arteries and cranial nerves to the skin, the eye, the lacrimal gland, and the salivary glands (Table 28.2).

With regard to the **lower extremities**, the arrangement corresponds to that described for the upper extremity,

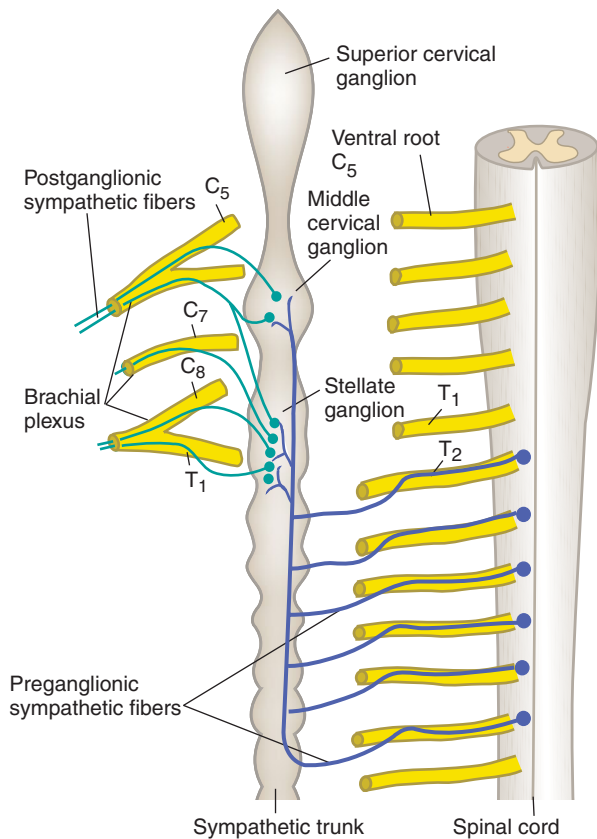


FIGURE 28.8 *Sympathetic innervation of the upper extremity.* The preganglionic fibers come from the upper thoracic segments of the cord and synapse in the ganglia of the sympathetic trunk up to the middle cervical ganglion. The postganglionic fibers follow the spinal nerves to the arm. (Redrawn from Haymaker and Woodhall 1945.)

with postganglionic fibers following the spinal the spinal nerves (Fig. 28.9).

Interruption of the Sympathetic Innervation of the Head: Horner's Syndrome

A lesion of the sympathetic trunk in the neck is the most common cause of sympathetic denervation of the head. Because all preganglionic fibers enter the trunk below the inferior cervical ganglion (Fig. 28.8), a lesion anywhere above the level of the thoracic outlet can interrupt the sympathetic innervation of the head. It may be caused, for example, by a tumor in the apex of the lung, in the thyroid gland, or in any of the numerous lymph nodes in the neck. The ensuing symptoms can be understood based on the effects of sympathetic fibers on the skin and the eye (Table 28.1). In the case of a unilateral lesion, the facial skin on the side of the lesion becomes redder (warmer) and drier than that on the other side (caused by vasodilatation and lack of sweat secretion). The pupil is miotic (smaller) on the side of the lesion (paralysis of the pupillary sphincter), and there is slight ptosis (the eyelid droops) owing to paralysis of the

smooth tarsal muscle. This constellation of symptoms—**red and dry skin, miosis, and ptosis**—affecting half of the face is called **Horner's syndrome** (Fig. 28.10).

Horner's syndrome can also be caused by lesions in the brain stem, interrupting the descending fibers to the intermediolateral column (cf. Chapter 27, under “Brain Stem Lesions Can Produce Symptoms from Several Cranial Nerves and Long Tracts” and “Lateral Pontine Infarctions”).

Sympathetic Innervation of the Viscera

As mentioned, the postganglionic sympathetic fibers to the extremities and the body wall follow mainly the spinal nerves. The visceral organs of the thorax and the abdomen do not receive branches from the spinal nerves, however. Therefore, the sympathetic fibers destined for these organs have to form separate nerves. Such **splanchnic nerves** are thin twigs that leave the sympathetic trunk at various levels.

The sympathetic fibers to the **heart** follow the superior, middle, and inferior **cardiac nerves** leaving the corresponding cervical ganglia. The cell bodies of the postganglionic neurons lie in the cervical ganglia. In addition, some smaller branches to the heart take off from the upper thoracic sympathetic ganglia.

Most sympathetic fibers to the **abdominal viscera** form the **greater and lesser splanchnic nerves** (Fig. 28.6). These nerves consist mainly of preganglionic fibers from spinal segments T₆–T₁₁, which pass uninterrupted through the ganglia of the sympathetic trunk (Figs. 28.5 and 28.7). The nerves penetrate the diaphragm and end in ganglia located on the ventral side of the abdominal aorta in the upper abdomen. As mentioned, these are called **prevertebral ganglia**, to distinguish them from the paravertebral ganglia of the sympathetic trunk.

The largest among the prevertebral ganglia is the **celiac ganglion**, located where the celiac artery emerges from the aorta. Smaller prevertebral ganglia occur where the **superior and inferior mesenteric arteries** emerge. The postganglionic fibers follow the arteries to the various organs.

The greater and lesser splanchnic nerves and the corresponding prevertebral ganglia supply the visceral organs in the **upper and middle part of the abdomen**, such as the stomach, pancreas, gallbladder, small intestine, and the large intestine to the descending part. The **adrenal medulla**, which also receives preganglionic fibers from the splanchnic nerves, is special. The medullary endocrine cells (**chromaffin cells**)—which are transformed postganglionic neurons—release **epinephrine** (and small amounts of norepinephrine) into the bloodstream on sympathetic stimulation.

The visceral organs of the **lower abdomen** and the **pelvis** receive their sympathetic innervation from the intermediolateral column in the lower thoracic and upper two

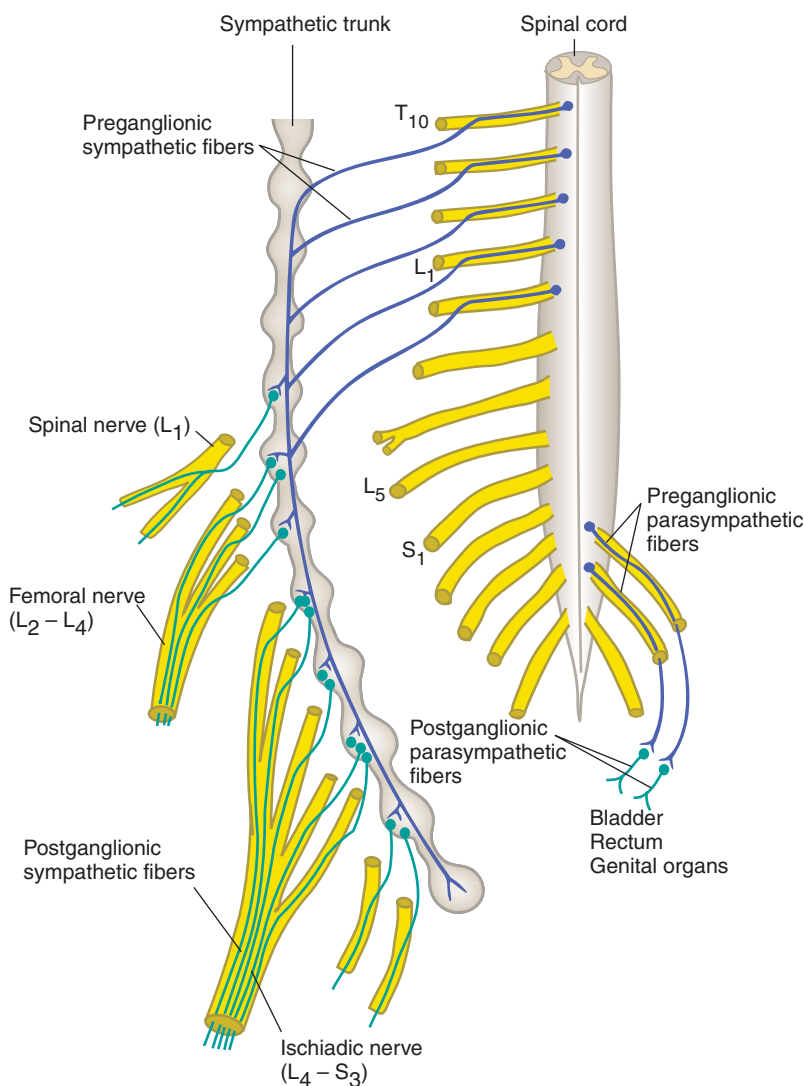


FIGURE 28.9 Sympathetic innervation of the lower extremity. On the right side, pre- and postganglionic parasympathetic fibers from the sacral cord are also shown. (Redrawn from Haymaker and Woodhall 1945.)

lumbar segments (Table 28.2). These fibers also leave the sympathetic trunk as separate nerves (lumbar splanchnic nerves) to reach prevertebral ganglia. The postganglionic fibers follow the arteries to the organs (Fig. 28.6).

The prevertebral ganglia are embedded in a meshwork of fibers, forming **prevertebral plexuses**, with names corresponding to those of the ganglia (Fig. 28.7). The plexuses formed mainly by sympathetic fibers continue

from the lower part of the abdominal aorta into the pelvis minor as the **hypogastric plexus**. In the pelvis, the hypogastric plexus mixes with parasympathetic fibers from the pelvic nerves and forms the **pelvic plexus** around the pelvic organs, as mentioned earlier.

PERIPHERAL PARTS OF THE PARASYMPATHETIC SYSTEM

As mentioned, the **preganglionic parasympathetic** (visceral efferent) neurons have their cell bodies in the brain stem and in the sacral cord. The neurons look like the sympathetic preganglionic neurons of the intermediolateral column.

The Cranial Nerves Contain Preganglionic Parasympathetic Fibers

The preganglionic fibers of the **cranial part** of the parasympathetic system follow the **oculomotor**, the **facial**

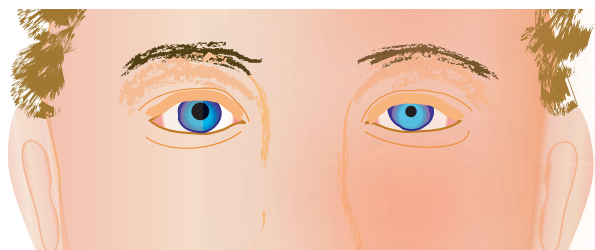


FIGURE 28.10 Horner's syndrome. The left half of the face shows the characteristic symptoms of loss of sympathetic innervation: red and dry skin, constricted pupil (miosis), and drooping of the upper eyelid (ptosis).

(intermediate), the **glossopharyngeal**, and the **vagus** nerves. The fibers come from the visceral efferent column of cranial nerve nuclei (see Figs. 27.2 and 27.3). The preganglionic fibers of the cranial nerves supplying structures in the head end in several parasympathetic **ganglia**, located outside the skull close to large cranial nerve trunks (Fig. 28.11). These are the **ciliary**, the

pterygopalatine, the **otic**, and the **submandibular ganglia**. From these ganglia, the postganglionic fibers pass to the effector organs (the intrinsic muscles of the eye, the lacrimal gland, and the salivary glands).

The preganglionic fibers of the **vagus** nerve do not end in well-defined ganglia but in more diffusely distributed collections of postganglionic neurons in the

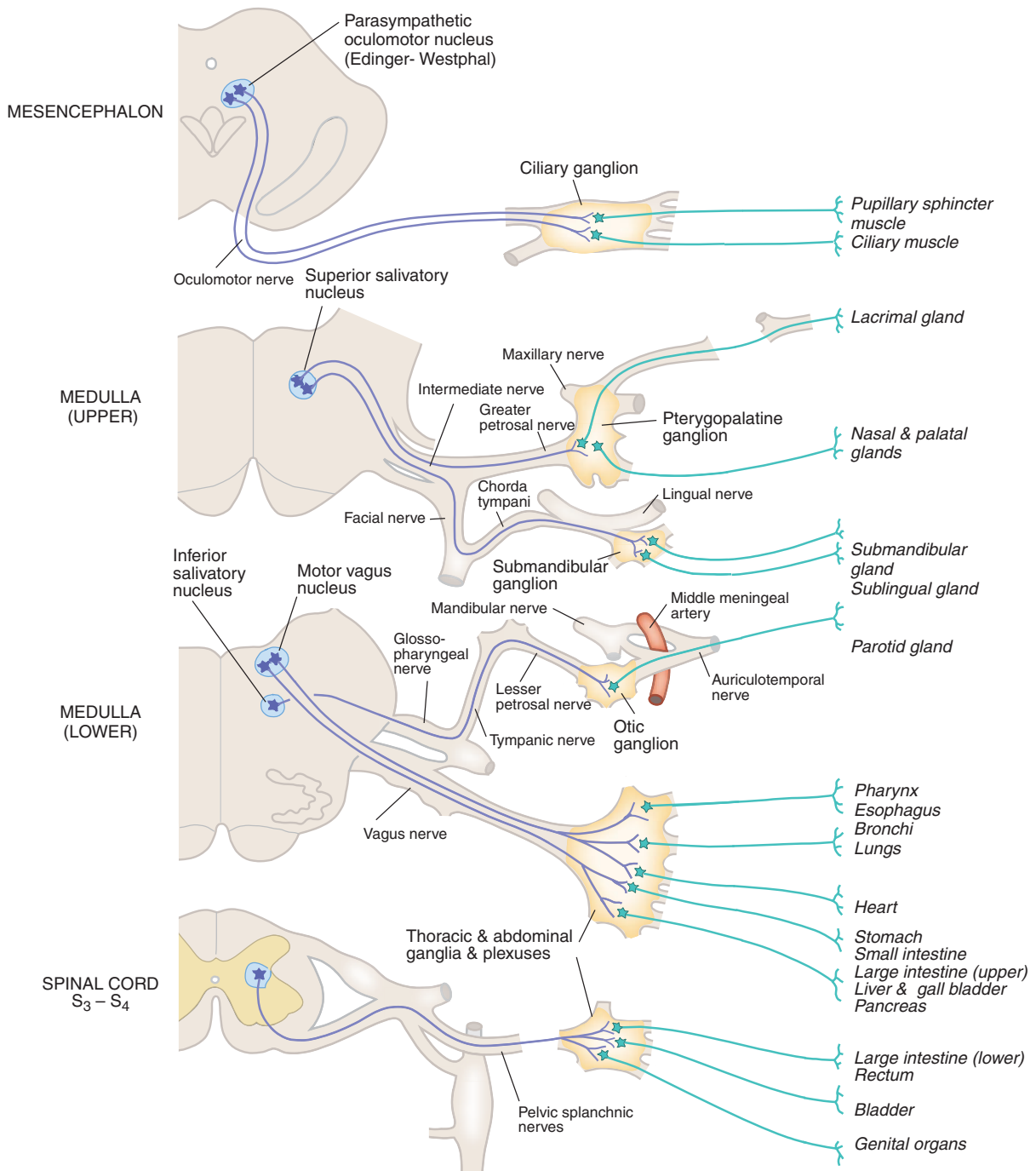


FIGURE 28.11 *The parasympathetic system.* The cell bodies of the preganglionic neurons are located in the brain stem and in the sacral cord. The peripheral course of the parasympathetic fibers is often

quite complicated because they “jump” from one cranial nerve to another on their way to the target.

walls of (or just outside) the thoracic and abdominal organs. These postganglionic neurons have short axons running in the wall of the organ and innervate smooth-muscle cells and glands. As mentioned in Chapter 27, the vagus nerve sends parasympathetic fibers to the heart, the lungs, the gastrointestinal tract down to the descending colon, the gallbladder, the liver, and the pancreas (see Fig. 17.9). With postganglionic sympathetic fibers, the vagus nerve forms the **cardiac plexus**, which lies around the aortic arch and extends down onto the heart. The cardiac plexus also contains scattered groups of postganglionic parasympathetic cell bodies, with the largest group just underneath the aorta. The **sinus node** and the **atrioventricular node** receive the densest innervation of parasympathetic postganglionic (vagus) fibers, whereas the ventricles receive few such fibers (they receive a dense sympathetic postganglionic innervation, however). In the **airways**, the vagus participates in plexuses around the trachea and the bronchi—the **tracheobronchial plexus**—containing scattered postganglionic neurons.

The Sacral Part of the Parasympathetic System Supplies the Genitals, Bladder, and Rectum

The cell bodies of the preganglionic neurons in the sacral cord are located in the (S_2) S_3 – S_4 segments, with a position corresponding to the intermediolateral column of the sympathetic system (Figs. 28.11 and 28.12). The parasympathetic preganglionic fibers leave the cord through the ventral roots and follow the spinal nerves for a short distance. Then they leave the spinal nerves as separate, small **pelvic splanchnic nerves**. In contrast to the sympathetic preganglionic fibers, the parasympathetic ones do not pass to the sympathetic trunk but join postganglionic sympathetic fibers in the **pelvic plexus**, located immediately lateral to the rectum, the bladder, the prostate gland (in the male), and the cervix (in the female). Many of the postganglionic parasympathetic neurons are located in the pelvic plexus (the **pelvic ganglion**).

Postganglionic parasympathetic fibers innervate all the **organs of the pelvis** and, in addition, the corpora cavernosa (erectile tissue) of the **penis** and the **clitoris**. The **descending** and **sigmoid colon** and the **rectum** are innervated from the sacral parasympathetic division. Of particular practical importance is the parasympathetic innervation of the rectum and the bladder, which is responsible for **emptying** these organs (Fig. 28.12). We treat the emptying reflex of the bladder later in this chapter.

THE ENTERIC NERVOUS SYSTEM

The “Mini-Brain of the Gut”

As mentioned, the digestive tract contains millions of neurons embedded in two plexuses in the wall. Largest is the **myenteric plexus**, which is located between the

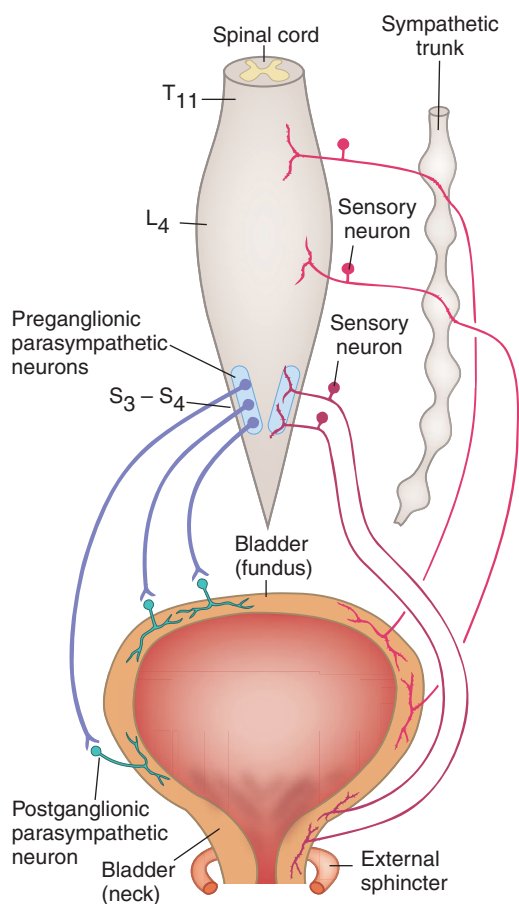


FIGURE 28.12 *Innervation of the bladder.* **Left:** The parasympathetic innervation of the smooth muscles that are responsible for emptying the bladder. The preganglionic neurons are located in the spinal segments S_3 – S_4 , whereas the postganglionic cell bodies are found just outside or in the wall of the bladder. **Right side:** The course of the sensory fibers from the bladder. From the lower part, the sensory fibers follow the efferent parasympathetic fibers, whereas the sensory fibers from the upper part (the fundus) follow the sympathetic fibers (via the sympathetic trunk). The striated, external sphincter muscle is innervated by motoneurons in the sacral segments S_3 – S_4 (not shown).

circular and the longitudinal external smooth-muscle layers; the smaller **submucous plexus** is situated below the mucous membrane. The plexuses are formed by the axons of the enteric neurons, together with postganglionic sympathetic fibers (with their cell bodies in the prevertebral ganglia). The neurons of the enteric system monitor the **tension** in the intestinal wall and the **chemical milieu** of the mucosa, and they control the **peristaltic movements** of the bowel, moving the content toward the anus. In addition, they influence blood flow, the secretion from mucosal glands, and the absorption of nutrients.

It was formerly believed that the enteric neurons were parasympathetic postganglionic. Only a minority contains acetylcholine, the classic transmitter of postganglionic parasympathetic neurons, however. It has become increasingly clear that these plexuses and ganglia represent a part of the peripheral nervous system that is, to a large extent, independent of the rest of the nervous

system. Some even use the term “the mini-brain of the gut.” Thus, the neurons in the myenteric and submucous ganglia are able to function even after the sympathetic and parasympathetic nerves to the gastrointestinal tract have been cut. The number of neurons in the enteric system is large, and only a few of these are under direct control from the CNS (by means of parasympathetic preganglionic fibers).

Broadly speaking, the enteric system integrates information about the local conditions with central commands through sympathetic and parasympathetic postganglionic fibers.

Neuronal Types and Neurotransmitters

The enteric plexuses contain not only **motor neurons**—innervating smooth-muscle cells and glands—but also **interneurons** and most likely **sensory neurons**. Numerous neurotransmitters—in particular, many **neuropeptides**—have been found in the enteric ganglion cells. With few exceptions, however, they are not yet functionally characterized. Different kinds of neurons contain different combinations of neuropeptides and classic transmitters (acetylcholine, GABA, and serotonin). **Excitatory neurons**, sending their axons upward (toward the mouth), ensure constriction of the bowel above a bolus of intestinal contents. Such neurons typically contain **acetylcholine** and **substance P**. **Inhibitory neurons** send their axons downward (toward the anus) and inhibit the circular muscle layer in front of a bolus, thus easing its propulsion. Such neurons may contain **ATP**, **vasoactive intestinal peptide (VIP)**, or **nitric oxide (NO)**. Neurons acting on epithelial cells to increase the transport of **water** and **electrolytes** may contain either **VIP** or **acetylcholine** (with a variety of peptides, such as **galanin**, **neuropeptide Y**, **cholecystokinin [CCK]**, or **calcitonin gene-related peptide [CGRP]**).

A special kind of neuron contains the **gastrin-releasing peptide (GRP)**. A group of such neurons are found in the wall of the stomach and send processes to the gastrin-producing cells in its distal part (the antrum). Stimulation of the vagus causes release of GRP, which then induces release of gastrin to the bloodstream. It affects acid-producing cells (parietal cells) in the upper part of the stomach.

Finally, some enteric neurons are most likely **sensory** and send their central process to prevertebral ganglia. Thus, local reflex arcs not involving the CNS may exist.

FUNCTIONAL ASPECTS OF THE AUTONOMIC NERVOUS SYSTEM

Defense and Maintenance

When the autonomic system is considered as a whole, certain main functional features emerge (Table 28.1).

The **parasympathetic system** controls primarily processes that are necessary for the maintenance of the organism over the long term. The parasympathetic system thus activates the digestive processes, ensures that waste products are expelled by contraction of the bladder and rectum, protects the eye against strong light, ensures focused vision, reduces the activity of the heart, reduces the diameter of the airways, and increases bronchial secretion. The **sympathetic system** as a whole is more concerned with mobilizing the resources of the body when an extra effort is required. In situations of fear and anger, there are usually signs indicating increased sympathetic activity, such as increased blood pressure, increased heart rate, and dilatation of the pupils. At the same time, stored energy is mobilized by epinephrine secreted into the bloodstream from the adrenal medulla (increased blood level of glucose and fatty acids). Epinephrine also activates the heart and dilates the bronchi (relaxation of smooth-muscle cells). The activity of the gastrointestinal tract is inhibited. In general, such responses are adequate in flight-or-fight situations.

A More Nuanced View

The simple dichotomy presented in the preceding text disregards that, in addition, the sympathetic system takes part in daily maintenance and is active not just in stressful situations. For example, the sympathetic system is crucial for the temperature-regulating functions of the skin, and sympathetic activity is required every time we rise from the sitting position to avoid fall in blood pressure and fainting. Further, the sympathetic system cooperates with the parasympathetic system in control of reproductive functions. Moreover, the sympathetic system is not activated in an **all-or-none** fashion: some parts may be active, while others are not. Thus, the sympathetic output to an organ may be regulated up and down independently of sympathetic outputs to other organs. Finally, both systems may send signals at the same time to an organ (the heart, e.g., receives both kinds of signals at the same time). The traditional scheme with reciprocal autonomic innervation—with autonomic “tone” determined by a point on a continuum from sympathetic to parasympathetic—is therefore not tenable.

Actions of Sympathetic Fibers on the Circulatory and Respiratory Organs

The postganglionic sympathetic fibers innervate vessels in all parts of the body. In general, the sympathetic innervation of the circulatory system ensures that the **cardiac output** can be increased, that the **blood pressure** can be maintained, and that the **blood flow** is directed to the organs needing it the most.

The sympathetic system increases the **heart rate** by acting on pacemaker cells in the sinus node. Further, the

stroke volume increases. Cardiac muscle cells with a lower spontaneous firing frequency than the cells of the sinus node are activated by spread of the signal from the sinus node before the spontaneous depolarization has reached the threshold for an action potential. At rest, the heart is under a certain dominance of the parasympathetic system (the vagus nerve), which “restrains” the cardiac activity. Actions on the stroke volume by the autonomic system are mediated by fibers ending near the muscle cells of the ventricles.

Because the vascular smooth-muscle cells are arranged circularly, contraction reduces the diameter and increases the vascular resistance. Such vasoconstriction is most marked in the smallest arteries, the **arterioles**, which are especially concerned with the regulation of blood flow to the organs. Action potentials in the sympathetic fibers ending in the vessel walls produce vasoconstriction and, thus, reduced blood flow. By varying the signal frequency of the sympathetic nerves, the CNS can vary the diameter of the vessels.² When there are no signals in the sympathetic fibers innervating the vessel (and no other substances act to produce contraction), the arterioles are maximally widened by the internal blood pressure. This is called **vasodilatation** (vasodilation).

In many situations, the task of the sympathetic system is to ensure that there is a sufficient blood flow through **high-priority organs**, primarily the brain and the heart. When the blood flow through these organs diminishes, sympathetic neurons to vessels in other parts of the body increase their firing rate. Thus, the blood flow through skeletal muscles and the visceral organs is reduced. Sudden vasodilatation in large parts of the body leads to a fall in blood pressure and fainting, because the cerebral blood flow is reduced.

Sympathetic innervation of the **large veins** is also important for the maintenance of adequate blood pressure. Constriction of such capacity vessels distributes more of the blood volume to the arterial side—that is, the **effective blood volume** increases. This mechanism is important in case blood volume is reduced (on bleeding or dehydration).

Signals in sympathetic fibers to arterioles in **skeletal muscles** produce (as their main effect) vasoconstriction. Whether some sympathetic fibers may have the opposite effect in humans is not settled. **Epinephrine**—released from the adrenal medulla on sympathetic stimulation—may, however, inhibit the vascular smooth-muscle cells and thereby produce vasodilatation.

The vessels of the **lungs** receive sympathetic fibers producing vasodilatation. It is doubtful whether postganglionic sympathetic fibers act on the bronchial smooth

musculature in humans, even though such innervation is present in several animal species. Thus, using histo-fluorescence techniques, which visualize catecholaminergic nerve fibers in tissue sections, studies in humans have shown the presence of such fibers around the vessels but not around the bronchi. **Epinephrine**, however, has a powerful inhibitory effect on the bronchial musculature; that is, it produces bronchial dilation. Most likely, therefore, the sympathetic system acts on the resistance of the airways by its stimulation of the adrenal medulla.

Control of Blood Pressure and Blood-Flow Distribution

Normally, the sympathetic neurons are activated reflexly, and many of them are links in arcs for so-called **vasomotor reflexes**, the reflexes in which the response is a change of vascular diameter (and thus resistance). The reflex arcs go through the cord or higher levels (the reticular formation or the hypothalamus). The superior aim of the control of **blood pressure** is to ensure that the brain (and the heart) always has a sufficient blood flow. **Baroreceptors** in the large arteries in the neck and the aortal arch record the slightest fall in blood pressure and produce an automatic increase of the signal frequency of sympathetic fibers. This is most marked for the skeletal muscles, but, if necessary, the heart rate is also increased. In this manner, vasoconstriction of the skeletal muscle arterioles is produced, thus increasing the vascular resistance and elevating the blood pressure, with the end result that the blood flow to the brain is increased to an adequate level.

Vasomotor reflexes have been studied with the **microneurographic technique**, enabling the recording of the activity of small groups of postganglionic sympathetic fibers in humans. This makes it possible to study the relationship between the sympathetic signal frequency and, for example, blood pressure. The postganglionic sympathetic fibers to muscle arterioles fire in bursts in pace with the pulse. The bursts are evoked by baroreceptor activation during the diastole of the heart. The overall firing frequency changes in association with changes of blood pressure. Considering the enormous blood flow that can pass through working muscles, we must obviously have central control of this part of the vascular system. Only if the heart increases its output and other vascular beds are constricted (e.g., in the abdominal organs) can the sympathetic “throttling” of the muscles be relieved without fall in the blood pressure.

Individual Differences in Sympathetic Activity

There are striking differences among persons with regard to the level of activity of sympathetic fibers under identical circumstances, as shown with the microneurographic technique. Postganglionic fibers to skeletal muscles,

2 The degree of vasoconstriction is influenced not only by the nervous system but also by circulating hormones, in particular, epinephrine. Further, substances produced by the local metabolism in the tissue influence the degree of contraction of the vascular smooth-muscle cells.

which constitute a large fraction of all postganglionic fibers in peripheral nerves, have been studied in particular. As mentioned, the activity of these fibers changes in close correlation with changes of the central blood pressure. Comparison of persons with normal blood pressure shows that the resting activity of sympathetic fibers varies by a factor of 10 from person to person. Thus, each person appears to have his own characteristic pattern, which is unchanged over a long time. From this “baseline” value, the signal frequency is up- or down-regulated in response to alterations in blood pressure caused by, for example, the change of body position from sitting to standing. No clear correlation has been found between the level of activity in sympathetic fibers to muscles and elevated blood pressure (hypertension).

Effects of Sympathetic Fibers in the Skin

The sympathetic innervation of the skin serves first and foremost the control of **body temperature**. The activity of the postganglionic sympathetic fibers to the skin appears not to be clearly related to the blood pressure, in contrast to the activity of fibers supplying vessels in skeletal muscles, but skin sympathetic fiber activity is closely correlated with the ambient temperature.

The **sweat glands** of the skin are innervated by sympathetic fibers, producing sweat secretion. Increased activity of fibers innervating the sweat glands occurs together with reduced activity of fibers to the small vessels. This produces **vasodilatation** and **sweat secretion** with increased loss of heat. A special feature of the innervation of sweat glands is that the neurotransmitter released from the postganglionic fibers is **acetylcholine** (and not norepinephrine).³

Sweat secretion may also occur in **extreme situations** with a large drop in blood pressure or with strong pain. In such situations, the skin vessels are maximally constricted and the skin is, consequently, pale and cold (**cold sweat**).

Signals in sympathetic fibers also activate the small smooth muscles attached to the **hair roots**, which make the hairs stand up. This is called **piloerection** (piloarrection). At the same time, the muscles compress the **sebaceous glands**, so that they empty their product into the hair follicle. In humans, the sympathetic control of hair position is of minor importance, whereas in animals it is of great significance for control of body temperature.

Irritation of Peripheral Nerves Can Produce Changes in the Skin

The effects of sympathetic fibers to the skin—that is, sweat secretion, vasoconstriction, and piloerection—can be reproduced by electrical stimulation of the ventral roots or the peripheral branches of the spinal nerves. The sympathetic fibers can be irritated by infections or by compression or traction of the nerves. In such cases, there is abnormal sweat secretion from pale and cold areas of the skin. Destruction of the sympathetic fibers (by, e.g., prolonged compression) leads to abolished sweat secretion and vasodilatation, resulting in areas of the skin that are abnormally warm and red and at the same time dry. Observations of such local changes of the skin can be helpful in the diagnosis of diseases that affect the peripheral nerves.

Effects of Sympathetic Fibers in the Gastrointestinal Tract, the Genital Organs, and the Eyes

In the **abdominal viscera**, signals in sympathetic fibers produce **vasoconstriction** and reduced contractile activity of the smooth muscles of the walls of hollow organs (i.e., reducing the amplitude and frequency of the peristaltic movements). At the same time, the secretion of the glands of the digestive tract is reduced. In sum, these effects result in a marked reduction of the digestive processes. The sympathetic system also inhibits the emptying of the **rectum**, both by inhibition of the smooth muscles of the wall and by activating the smooth muscles of the internal anal sphincter. The sympathetic system seems to play a minor role in the control of the bladder in humans (see later, “Normal Emptying of the Bladder”).

The sympathetic innervation of the **genital organs** concerns vessels and the smooth musculature. The innervation of the **ductus deferens** is of particular importance because signals in sympathetic fibers are responsible for the rhythmic contractions during ejaculation. The **uterus** receives sympathetic fibers but their functional role is not clear. Thus, even after complete denervation, the uterus may function normally in pregnancy and in parturition.

The sympathetic fibers to the **eye** have their cell bodies in the **superior cervical ganglion**. They produce dilation of the pupil by activating the **pupillary dilator muscle** and by causing contraction of the radially oriented vessels of the iris (the latter effect is probably most important for pupillary dilation). A small smooth muscle attached to the upper eyelid, the **tarsal muscle**, is also innervated by sympathetic fibers. The tonic activity of this muscle helps to keep the eyelid up while we are awake (the paresis of the tarsal muscle is responsible for the slight drooping of the eyelid occurring in Horner’s syndrome).

3 Human hairy skin (but not glabrous skin) receives sympathetic fibers of two kinds: the ordinary noradrenergic vasoconstrictor fibers innervating vessels in all parts of the body, and cholinergic vasodilator fibers. The latter fibers mediate reflex responses to rise in core temperature (whole body heating) when the need to dissipate heat is great. The cholinergic vasodilator fibers co-release neuropeptides (e.g., VIP). In addition, local NO production contributes to vasodilatation.

Orthostatic and Postprandial Hypotension

Some people have poor control of the blood-flow distribution in situations with change of body position from supine to standing (orthostatic hypotension). This is believed to be due to a failure of the sympathetic system. Such persons also often feel limp and uncomfortable after a meal, because of a fall in the blood pressure (postprandial hypotension). Microneurographic studies suggest that this is caused by a lack of sympathetic activity. Normally after a meal, the sympathetic signal activity increases in nerves to the lower extremities, whereas in the patients with postprandial hypotension no such increase occurs. (To maintain the blood pressure, the blood flow to the lower part of the body must be reduced when that to the digestive tract increases.)

Function-Specific Sympathetic Control

The preceding discussion showed that various categories of sympathetic fibers can be controlled independently. There is not a uniform “sympathetic tone” for all parts of the sympathetic system. High activity in some parts must coexist with low activity in others if the sympathetic system is to fulfill its tasks in controlling blood pressure and body temperature, in reproduction, and so forth. Accordingly, recent anatomic studies show that sympathetic neurons innervating different targets are more clearly segregated than formerly believed. For example, double labeling with retrograde tracers show that, in the intermediolateral column, neurons supplying the **superior cervical ganglion** and the **stellate ganglion** are largely segregated (although they are found mainly in the same spinal segments, with 90% in T_1 – T_6). Thus, higher levels of the CNS (such as the hypothalamus) can selectively control subdivisions of the sympathetic system.

Microneurographic observations further support that selective control takes place. Recording During rest, the firing frequency is uniform to different muscles. In this situation, a common signal (probably from the brain stem) commands all sympathetic neurons controlling muscle blood flow. As soon as a muscle starts working, however, the sympathetic signal frequency drops in the nerve to this muscle, while it remains unaltered to the resting muscles. In this situation, the sympathetic system exerts a differential control of the muscles; that is, the signal frequencies depend on their individual needs.

The specificity of sympathetic control of muscle blood flow can presumably arise in two ways. One possibility is lowered **central drive** (from the brain stem) to the preganglionic neurons that control specific muscles. In turn, this may be due to sensory signals from **ergoreceptors** in the working muscle, or that the **motor cortex** informs higher levels of the autonomic system about which muscles are being selected for a motor task (efference

copy). The other possibility is that a purely **spinal reflex** inhibits the preganglionic neurons that control blood flow to the working muscles.

The Effects of Parasympathetic Fibers

As mentioned, the sympathetic and the parasympathetic systems have mostly antagonistic effects (on the organs innervated by both; Tables 28.1 and 28.2). Because the postganglionic fibers usually contain neuropeptides in addition to acetylcholine, however, their actions may be more complex than either stimulation or inhibition of the organ.

The parasympathetic postganglionic fibers produce **glandular secretion** (e.g., from the lacrimal gland, salivary glands, and glands of the respiratory and gastrointestinal tracts). Parasympathetic fibers are furthermore responsible for increased strength and frequency of **peristaltic contractions** in the gastrointestinal tract and the bladder. The parasympathetic innervation is particularly important for the emptying of the **bladder** (Fig. 28.12) and the **rectum** (control of micturition is treated in Chapter 29 in connection with visceral reflexes).

As mentioned, the **heart** receives parasympathetic preganglionic fibers through the vagus nerve. The sinus node and the atrioventricular node receive particularly dense innervation of postganglionic (cholinergic) fibers. Lowering of the heart rate—by affecting the sinus node—is the most marked effect of vagus stimulation. In addition, the vagus exerts more complex effects on the ventricles. Most likely, this happens by presynaptic inhibition of sympathetic postganglionic fibers, thus reducing the contractile force of the heart muscle. Some effects of vagus stimulation are not mediated by acetylcholine but, most likely, by neuropeptides such as **somatostatin** and **VIP**. In the resting situation the heart receives parasympathetic signals with a low frequency (higher in endurance-trained than in untrained persons). During work, the influence of the vagus diminishes with the need for increased cardiac output. The effect of the vagus on the **coronary arteries** in humans is believed to be constrictive. In the **airways**, the vagus produces bronchial constriction and secretion.

Most of the **vessels** of the body do not receive parasympathetic innervation. Exceptions are vessels of glands and of the external genitals, in which parasympathetic signals cause vasodilation (i.e., they inhibit the smooth-muscle cells). Increased activity of the parasympathetic fibers to the penis (and the clitoris) produces **erection** (see Chapter 29, under “Control of the Erection and Ejaculation Reflexes”).

In the **eye**, parasympathetic fibers of the oculomotor nerve reduce the diameter of the pupil and produce accommodation of the lens (see Chapter 27, under “The Light Reflex and the Accommodation Reflex”).

Emotions and the Autonomic Nervous System

The autonomic system is not independent of higher mental processes, even though its processes are not under conscious control or are as a rule not consciously perceived. The activity of sympathetic fibers to the skin, for example, are strongly influenced by emotions, as witnessed by **blushing** when “having made a fool of oneself” and **pallor** when frightened. Further, the effect of emotions on the circulatory system may manifest itself as palpitations, hypertension, or bradycardia and peripheral vasodilation, leading to a fall in blood pressure and perhaps **fainting**. In fainting, there are marked changes of both sympathetic and parasympathetic activity. The control of the **gastrointestinal tract** may be altered by emotions—for example, with increased peristaltic movements and secretions leading to diarrhea. Also, **bladder emptying** is under emotional influence, as witnessed by the frequent urge to void when nervous—for example, before an exam or an athletic contest. In case of very **strong fear**, involuntary emptying of the bladder and rectum may occur. Emotional influence on **erection** is another example: sensory stimuli and simple reflexes alone do not determine the parasympathetic actions on the vessels in the penis and the clitoris.

Pathways and nuclei mediating the effects of emotions on preganglionic autonomic neurons (and thus on visceral organs) involve parts of the cerebral cortex, the hypothalamus, the amygdala, the periaqueductal gray (PAG), and the reticular formation. For example, electrical stimulation of the amygdala can produce emptying of the bladder and rectum in experimental animals. Probably, the pathway goes via the PAG (see also Chapter 31, under “Amygdala and Conditioned Fear” and “Cortical Control of Autonomic Functions and Emotions”).

NEUROTRANSMITTERS IN THE AUTONOMIC NERVOUS SYSTEM

Preganglionic Neurons

The signal transmission between neurons of the autonomic system is mediated by neurotransmitters, as elsewhere in the nervous system. The preganglionic fibers end with typical synapses on the dendrites of the postganglionic neurons. As mentioned, all (or the vast majority of) **preganglionic** neurons use **acetylcholine**; that is, they are cholinergic. The released acetylcholine binds to **nicotinic receptors** in the membrane of the postganglionic neurons in the autonomic ganglia.

Many (perhaps all) preganglionic neurons contain in addition **neuropeptides** (enkephalin, somatostatin, neuropeptide Y, and others), as demonstrated with immunocytochemical techniques. The various neuropeptides appear to be expressed differentially in subgroups of autonomic

ganglion cells. The functional significance of these neuropeptides, which coexist with acetylcholine in preganglionic neurons, is so far not clear, but the two substances are most likely released together.

Postganglionic Neurons

Most **postganglionic parasympathetic** neurons release **acetylcholine**. In the peripheral organs, acetylcholine binds to **muscarinic receptors** in the membrane of cardiac, smooth-muscle, and glandular cells.

Most **postganglionic sympathetic** neurons release **norepinephrine** and are **noradrenergic** (norepinephric). The effects on the effector cells are mediated by two kinds of receptors, α - and the **β -adrenergic receptors**, which are distributed differently and have different effects on the postsynaptic cells. In the heart, norepinephrine produces increased heart rate by its binding to β receptors. By binding to α receptors, norepinephrine produces contraction of smooth-muscle cells in most blood vessels, in the ductus deferens, and in the pupillary dilator muscle of the eye. Binding to β -receptors elicits relaxation of smooth-muscle cells in the wall of the bladder, the uterus, and the airways.

Epinephrine, which is released from the **chromaffin cells** of the **adrenal medulla** by sympathetic stimulation, has largely the same effects as norepinephrine. Thus, epinephrine binds to α - and β -adrenergic receptors of the heart, vessels, and the respiratory tract. In addition, epinephrine stimulates the release of free fatty acids from adipose tissue and the breakdown of glycogen to glucose. These **metabolic effects** are mediated by β receptors in fat and liver cells.

Subgroups of Adrenergic Receptors

Each of the two main kinds— α - and β -adrenergic receptors—has several subtypes with different distributions and actions. When norepinephrine binds to the α_1 **receptor**, it produces opening of **Ca^{2+} channels**, which leads to depolarization and, in turn, elicits contraction or secretion. The action of the α_1 receptor is not directly on the Ca^{2+} channel but indirectly via intracellular second messengers (diacylglycerol and activation of protein-kinase C). The α_2 **receptor** is mostly localized presynaptically and modulates the transmitter release. The β_1 **receptor** is mostly localized postsynaptically in the heart, on adipose cells, and in the CNS. It acts through cyclic AMP as a second messenger. The β_2 **receptor** has a different distribution than the β_1 receptor, being primarily found in smooth-muscle cells of the respiratory tract. Binding of epinephrine (or drugs with similar action) to β_2 receptors relaxes the smooth-muscle cells, notably in the walls of the bronchi. This relaxation produces dilatation of the bronchi and reduces airway resistance.

Noncholinergic and Nonadrenergic Transmission in the Autonomic System

In addition to the classical neurotransmitters acetylcholine and norepinephrine, several other neuroactive substances have been demonstrated in the autonomic nervous system. As mentioned, many preganglionic and postganglionic neurons contain neuropeptides, as well as acetylcholine or norepinephrine. Further, some autonomic neurons—notably in the enteric system—contain neither acetylcholine nor norepinephrine. Such **noncholinergic** and **nonadrenergic** (NANC) autonomic fibers are also found in the respiratory tract, the gastrointestinal tract, the bladder, and the external genitals. Some of them release **ATP** or **NO** as a neurotransmitter; others contain neuropeptides such as **somatostatin**, **substance P**, **VIP**, and **CCK**.

The **coexistence** of norepinephrine and other transmitters was first suggested by the observation that blocking the receptors for norepinephrine did not prevent all effects of sympathetic nerve stimulation. In the **ductus deferens**, which receives a very dense sympathetic innervation, stimulation of the nerves produces, first, a fast contraction caused by release of ATP and, subsequently, a slow contraction produced by norepinephrine. In the **salivary glands**, the parasympathetic postganglionic fibers release both acetylcholine and VIP. The acetylcholine produces secretion from the glandular cells, whereas the VIP produces vasodilatation. Another example concerns the arteries of the **penis** and the **clitoris**, which dilate to cause erection. This vasodilatation is caused by parasympathetic postganglionic fibers that release NO (but not acetylcholine).

Some parasympathetic postganglionic fibers in the **heart** release somatostatin and probably VIP, thus increasing the heart rate. In the **stomach**, vagus stimulation can produce release of VIP in addition to acetylcholine. Stimulation of nerves to the human **airways** can produce bronchial dilatation, although not by release of norepinephrine or acetylcholine. The effect appears to be mediated by release of **VIP** from postganglionic nerve varicosities.

Presynaptic Receptors Modulate the Transmitter Release from Postganglionic Nerve Terminals

Neurotransmitters released from the postganglionic neurons bind not only to postsynaptic receptors in the membrane of smooth-muscle and glandular cells but also to presynaptic receptors in the membrane of the varicosities along the fibers (see Fig. 5.1). Thus, for example, norepinephrine that is released from sympathetic fibers can bind presynaptically and inhibit further release of norepinephrine or bind to parasympathetic cholinergic terminals in the vicinity. In the **heart**, sympathetic fibers inhibit the release of acetylcholine in this manner.

The sympathetic inhibiting effect on the peristaltic contractions of the gastrointestinal tract is mediated, at least partly, by binding of norepinephrine to α receptors on the parasympathetic, cholinergic terminals: that is, the release of acetylcholine is inhibited.

Sensitization

When the postganglionic autonomic fibers to an organ are interrupted, the sensitivity of the organ to the transmitter (which is no longer released) is increased. Epinephrine and norepinephrine in the bloodstream, for example, have a more powerful action after an organ has lost its sympathetic innervation, and the same holds for adrenergic drugs. This phenomenon, called **sensitization**, is not restricted to the autonomic system, however. It occurs, presumably, after denervation of any neuron. For example, skeletal muscle cells have increased sensitivity to acetylcholine after having lost their nerve supply. The underlying mechanism is probably increased postsynaptic density of receptors, as though the neuron attempts to maintain normal synaptic activity.

Drugs with Actions on the Autonomic Nervous System

Several drugs influence the synaptic transmission in the autonomic nervous system. **Atropine** blocks the action of acetylcholine (released from postganglionic parasympathetic fibers) on **muscarinic** receptors. Other drugs have similar **anticholinergic** effects, often as a side effect. This is the case for several psychopharmaceuticals. The peripheral actions of the parasympathetic system are inhibited, causing symptoms such as dilated pupils (mydriasis) and reduced accommodation of the lens (causing difficulties in seeing close objects clearly). The heart rate increases, and the secretory activity is reduced in several glands. The reduced salivary secretion causes dryness of the mouth, a very bothersome side effect of anticholinergic drugs. Atropine, for example, is used to reduce secretion of glands in the respiratory tract during surgical anesthesia. The peristaltic contractions of the bowel are reduced, causing constipation. The bladder contractility is reduced, with danger of incomplete emptying (especially in cases of prostatic enlargement causing increased urethral resistance, the danger of urinary retention should be kept in mind). Because the sweat glands receive a cholinergic innervation, their secretion may also be reduced (most antiperspirants contain substances with an anticholinergic action).

Pilocarpine is an example of a drug with a **parasympathicomimetic** action: that is, a cholinergic drug. Administration of pilocarpine causes increased salivation and tear flow, reduced heart rate, and increased secretion from, and peristaltic movements of, the gastrointestinal

tract. The pupil is small (miotic), causing reduced vision in dim light.

Many drugs activate adrenergic receptors—that is, they have **sympathomimetic** effects. Some act on both α and β receptors; others act preferentially on one or the other receptor type (or on subtypes). **Isoprenaline** (isoproterenol) acts selectively on β receptors and produces increased heart rate and bronchial dilatation. **Metaraminol** acts preferentially on α receptors and causes peripheral vasoconstriction and, thereby, increased blood pressure.

Drugs that **block α receptors** (such as **phentolamine**) produce peripheral vasodilatation and a fall in blood pressure, whereas drugs that **block β receptors** mainly cause reduced heart rate and stroke volume, and bronchial constriction. The development of more selective β blockers, acting selectively on β_1 receptors present in the heart, has made it possible to treat hypertension without unwanted bronchial constriction (β_2 receptors are found primarily in the lungs). In contrast, the development of adrenergic drugs acting selectively on β_2 receptors (and not on β_1 receptors) has made it possible

to treat patients with bronchial obstruction (as asthmatics) without such side effects as increased cardiac activity and hypertension.

Drugs can also influence the signal transmission in the **autonomic ganglia**. As mentioned, acetylcholine is the main transmitter in both sympathetic and parasympathetic ganglia. The nicotinic receptors in the ganglia are nevertheless somewhat different from those present at the neuromuscular junction. This makes it possible to influence one of these targets without affecting the other.

All neurotransmitters present in the peripheral parts of the autonomic system are also found in the CNS, together with adrenergic and cholinergic receptors. Therefore, drugs designed to act on peripheral parts of the autonomic nervous system may produce side effects through actions in the CNS—that is, in case they pass the blood-brain barrier. **Beta (β) blockers**, for example, which are used extensively to treat hypertension, can give central side effects, such as dizziness, disturbed sleep, and depression.

Sensory Visceral Neurons and Visceral Reflexes

OVERVIEW

Afferent fibers conveying sensory information from the viscera to the central nervous system (CNS) go together with the visceral efferent (sympathetic and parasympathetic) fibers. Sensory fibers follow the sympathetic **splanchnic nerves**, leaving the sympathetic trunk at various levels, the **parasympathetic cranial nerves** (the oculomotor, the intermediate, the glossopharyngeal, and the vagus nerves), and, finally, the parasympathetic **pelvic nerves** leaving the sacral spinal nerves to innervate pelvic organs. As a general rule, fibers conducting signals from **visceral nociceptors** follow sympathetic nerves, whereas the parasympathetic nerves contain fibers conducting from other kinds of receptors.

The visceral afferent neurons are structurally indistinguishable from the somatic afferent ones—that is, they have their pseudounipolar cell bodies in ganglia of the spinal and cranial nerves. The peripheral process follows the sympathetic or parasympathetic nerves to the organs, whereas the central process enters the dorsal horn or the brain stem sensory nuclei (the solitary nucleus). The vast majority of the visceral afferent fibers are thin **Aδ** and **C fibers**.

The normal function of the sensory innervation of the internal organs is primarily related to their mediation of **visceral reflexes**, such as coughing, vomiting, swallowing, circulatory and respiratory reflexes, emptying of the rectum and bladder, and so forth. These reflexes have their **reflex centers** in the spinal cord or in the reticular formation. In addition, “**silent nociceptors**” may mediate information about the local milieu of the internal organs, contributing to **homeostasis**. Visceral afferents also mediate information from the **immune system** to the brain, evoking fever and other aspects of sickness behavior during infections and other inflammatory diseases. With some exceptions, such as taste and fairly diffuse sensations of hunger, fullness, and so forth, the sensory signals from the viscera are not consciously perceived.

The control of **micturition** is discussed, including not only the spinal reflex mechanisms involved but also the necessary contributions from the **pontine micturition center** and the cerebral cortex.

Under abnormal conditions, however, sensory signals from the viscera can produce the sensations of intense pain and feeling of sickness. We discuss the special features of **visceral pain** and the phenomenon of **referred pain** in particular.

VISCERAL RECEPTORS AND AFFERENT PATHWAYS

Visceral Receptors

Morphologically, most of the visceral receptors are free nerve endings (see Fig. 13.1).¹ They respond to various kinds of stimuli, even though they are structurally identical. A large group consists of **mechanoreceptors** recording the tension of the tissue in which they are located. Such receptors are found, for example, in the heart, lungs, and the walls of hollow abdominal organs. They provide information about the degree of filling of hollow organs and can elicit reflex contractions aimed at emptying the organ or moving the content. Some stretch receptors may give rise to sensations of pain by responding to strong **dilatation** of a hollow organ and to forceful **contractions** of the smooth musculature, but most visceral **nociceptors** are believed to be **chemoreceptors** sensitive to substances in the tissue produced by **inflammation** or **ischemia**.

Other chemoreceptors, strategically placed in the vascular system (at the aortic arch and at the bifurcation of the common carotid artery), react to the **carbon dioxide** and **oxygen** concentration in the blood.

Typically, visceral receptors have large **receptive fields**, with a low density of nerve endings. These features fit with the observations that visceral receptors exhibit marked **spatial summation**—that is, the threshold for eliciting action potentials becomes lower as the area of stimulation increases. This may explain why strong but spatially very restricted stimuli (such as knife cuts) usually do not evoke pain from visceral organs.

¹ There are some encapsulated nerve endings in the visceral organs. Thus, Pacinian corpuscles are present in the pancreas, the mesenteries, and the vessel walls. Their functional role at these sites is unknown.

Peripheral Routes for Nociceptive Signals

As mentioned, most visceral afferents leading from nociceptors follow the sympathetic nerves (Fig. 29.1). Exceptions to this rule are nociceptors in the neck of the **bladder**, the **prostate**, the **cervix uteri**, and the **rectum**. “Pain fibers” from these organs follow the **parasympathetic pelvic nerves** to the sacral spinal nerves (see Fig. 28.12). Signals from nociceptors in the fundus of the uterus and of the bladder, however, follow the sympathetic nerves of the hypogastric plexus, ending in segments T₁₁–L₂. Although nociceptive signals from the **heart** were assumed to follow exclusively the sympathetic nerves, some observations indicate that this may hold only for signals from the anterior wall of the heart, while signals from the posterior wall and the lower surface may be conducted in the vagus nerve.²

Fibers carrying nociceptive signals from the **arteries** in the extremities and the body wall follow the spinal nerves, together with the sympathetic efferent fibers.

Central Pathways for Visceral Afferent Signals

The sensory signals from the visceral organs pass through the dorsal roots and end primarily in the dorsalmost parts of the dorsal horn—that is, **lamina I** (Fig. 29.1; see Fig. 13.16). Notably, the visceral afferents avoid the substantia gelatinosa (lamina II, which receives numerous C fibers from somatic structures). The sensory fibers following the cranial nerves end in the **solitary nucleus** in the medulla (see Fig. 27.2). From these receiving cell groups in the spinal cord and in the brain stem, the signals are transmitted to motor nuclei—especially those consisting of **preganglionic autonomic neurons**—to the **reticular formation**, and to the **hypothalamus** and the **thalamus**. For the most part, the visceral afferent signals that arrive through the dorsal roots are transmitted centrally through the **spinothalamic** and **spinoreticular tracts** (see Figs. 14.4 and 26.10). Details are not known, however, with regard to the central transmission of visceral afferent signals.

² Observations after surgical interventions to relieve pains of **angina** were taken as evidence that all nociceptive fibers from the heart follow the sympathetic nerves. These are the middle and lower cardiac nerves arising from the cervical sympathetic trunk, and some smaller twigs leaving the sympathetic trunk below the stellate ganglion (4–5 upper thoracic ganglia). Although most patients reported complete relief of angina pains after cutting of the dorsal roots in the upper thoracic and lower cervical segments (or removing the corresponding ganglia), some patients were not helped. However, some patients reported pain relief after cutting of the vagus nerve, and electric stimulation of the vagus nerve has been reported to evoke burning pain deep in the chest. Perhaps these findings may be explained if the nociceptive signals follow different nerves from the anterior and posterior cardiac walls. Indeed, animal experiments indicate that sympathectomy does not relieve pain arising in the posterior cardiac wall.

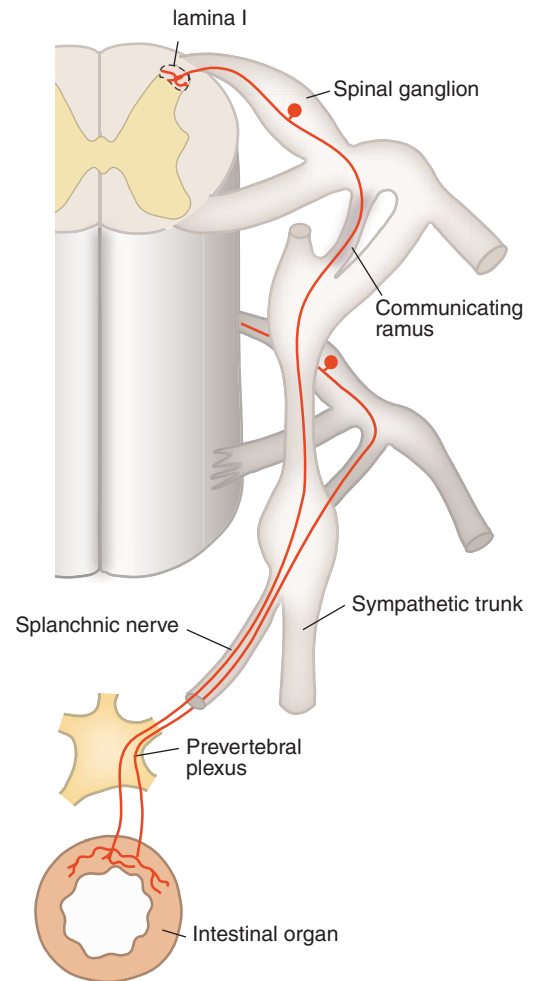


FIGURE 29.1 Sensory fibers follow sympathetic nerves to the visceral organs. The fibers pass through the prevertebral plexuses and ganglia before they enter the spinal nerves via the communicating rami.

The Dorsal Columns Carry Signals from Visceral Nociceptors

Surprisingly, in addition to constituting a major pathway for signals from somatic low-threshold mechanoreceptors, the dorsal columns have also been shown to carry sensory signals from visceral organs. This concerns **postsynaptic** dorsal column sensory units, many of which are activated by noxious stimuli. In the **gracile nucleus**, some neurons receive converging inputs from cutaneous low-threshold mechanoreceptors and high-threshold receptors in the pelvic viscera. Corresponding convergence occurs in the next link, in the VPL of the thalamus. Signals from cutaneous low-threshold mechanoreceptors on the arms and chest and from nociceptors in the heart converge on neurons in the cuneate nucleus and the thalamus. Thus, the dorsal column–medial lemniscus pathway can obviously contribute to signaling from visceral nociceptors, although this is not usually apparent from studies of its functions or the

effects of lesions. Nevertheless, there is evidence that signals in the dorsal columns can contribute to the experience of pain in humans. For example, a patient with intense pain due to cancer of the large bowel was made pain free by bilateral sectioning of the gracile fascicles at the T₁₀ level (the effect lasted until his death three months later). Animal experiments confirm that nociceptive signals from the lower abdomen and the pelvis cease to activate the cerebral cortex after transection of the gracile fascicle. Finally, the convergence mediated by the dorsal columns may also contribute to **referred pain** (discussed later).

VISCERAL REFLEXES

Many of the visceral reflexes elicited by signals from visceral receptors and receptors in the walls of vessels have their **reflex centers** in the spinal cord. The more complex reflexes, however, requiring coordination of activity in several parts of the body, have reflex centers in the brain stem or in the hypothalamus. We return to this in Chapter 30. **Vasomotor reflexes** were discussed earlier in this chapter. Other important visceral reflexes are produced by stimulation of receptors in the **lungs** and the **airways**, such as coughing and respiratory adjustments (see later). The **vomiting reflex** can be elicited by irritation of the mucosa of the stomach but also in various other ways (see Chapter 27, under “The Vomiting Reflex”). The **emptying reflexes** of the rectum and the bladder are elicited by stimulation of stretch receptors in their walls and have reflex centers partly in spinal segments (S₂) S₃–S₄ and partly in the brain stem. These visceral reflexes are unusual because they can be suppressed voluntarily. The emptying reflex of the bladder is discussed further later.

Reflexes Elicited from Receptors of the Lungs

Signals from **stretch receptors** in the bronchial walls contribute to inhibition of inspiratory movements when the lungs have been inflated to a certain extent (the Hering-Breuer reflex). Receptors producing **coughing** are probably free endings between the epithelial cells of the airways, in part located very close to the epithelial surface (irritation receptors). Such free nerve endings contain **substance P** (as do many other sensory neurons), which is released by exposure to irritant gases.

A special kind of receptor—the **J**, or **juxtapulmonary, receptor**—is located close to the lung alveoli. It responds to increased pulmonary capillary pressure. Increased pressure in the left atrium (which receives the blood from the lungs) immediately leads to increased pulmonary capillary pressure, with the danger of developing lung edema. Thus, it seems reasonable that the capillary pressure must be monitored closely. Stimulation of the

J receptors produces rapid and shallow breathing but may probably also cause bronchial constriction (this is known to occur in patients with heart failure and increased pulmonary capillary pressure). It is furthermore believed that signals from the J receptors can reach consciousness and cause a feeling of shortness of breath, or **dyspnea**. J receptors are also believed to elicit the dry cough typical of lunge edema, as occurring in patients with congestive heart failure or persons suffering from altitude sickness.

The Emptying Reflex of the Bladder

As mentioned, the bladder emptying reflex is evoked by stimulation of **stretch receptors** in the wall of the bladder, which record filling. The signals are conducted in myelinated afferent fibers to the lumbosacral cord (Fig. 29.2; see Fig. 28.12). Many sensory units that lead from the bladder are slowly adapting with dynamic sensitivity; that is, they respond more upon rapid than upon slow distension of the bladder. No signals are sent when the bladder is empty, but when urine starts to accumulate the sensory units start firing. Normal adult **bladder capacity** is about 500 mL. The pressure in the human bladder during filling is typically between 5 and 15 mm Hg, whereas emptying is normally elicited at 25 to 30 mm Hg. At night (during sleep), the bladder fills to about the double of daytime volume before evoking an urge to void. This is a prerequisite for 8 hours uninterrupted sleep.³ Urine does not leak out in the filling phase because the **intraurethral pressure** is kept higher than the **intra-vesical pressure**. The intraurethral pressure is maintained by several factors, among them smooth muscles and elastic tissue in the urethral wall. In the filling phase, the smooth muscle of the bladder wall—the **detrusor muscle**—is relaxed, while the striated external sphincter in the pelvic floor is tonically active. When the intra-vesical pressure reaches the critical level, brisk activity of parasympathetic neurons makes the detrusor muscle contract. In addition, the striated sphincter muscle and other muscles in the pelvic floor must relax. Normal emptying of the bladder thus requires **coordinated control** of parasympathetic preganglionic neurons in the S₃–S₄ segments and of α motoneurons in the S₁–S₃ segments.

The **parasympathetic** control of the detrusor muscle is mediated by **acetylcholine**. In addition, **vasoactive intestinal peptide (VIP)** released from parasympathetic fibers might contribute to inhibition of the smooth muscles surrounding the urethra.

3 One or more of the centers of the micturition reflex must therefore be inhibited during sleep. This inhibition develops between the ages of three and five. In children with **enuresis** (bedwetting), this inhibition seems to be lacking, as about the same bladder filling volume elicits emptying during night and during daytime.

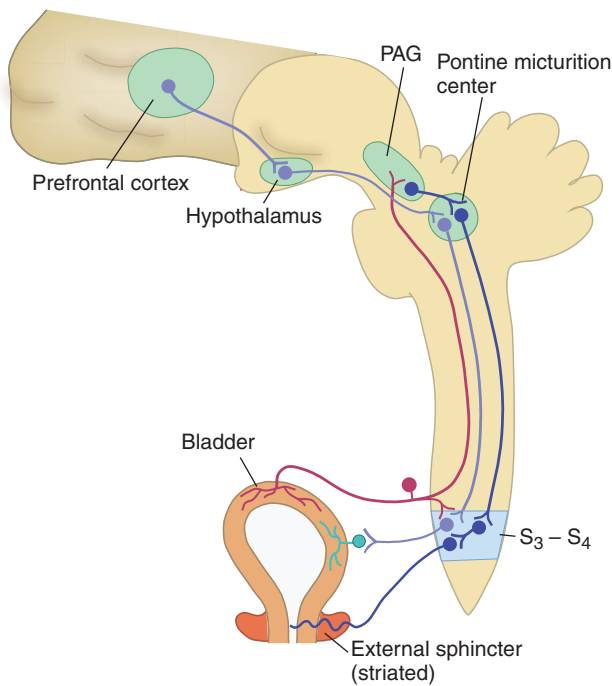


FIGURE 29.2 Pathways and nuclei involved in control of micturition. The reflex center for the emptying reflex is located in the sacral cord but is controlled by descending connections from the pontine micturition center. This is under control of higher centers, like the PAG, the hypothalamus, and the cerebral cortex.

The Role of the Sympathetic System in Bladder Control Is Not Clear

Evidence in experimental animals (cats and dogs) indicates that sympathetic activity increases during the filling phase, inhibiting the detrusor and activating smooth-muscle cells around the urethra in the bladder neck (internal sphincter). Few noradrenergic fibers are found in the human detrusor region (fundus), however, and sympathetic stimulation does not seem to inhibit the detrusor in the filling phase. (Because many noradrenergic fibers are found in the pelvic plexus and associated small parasympathetic ganglia, sympathetic inhibition of the detrusor muscle might theoretically occur by inhibition of the postganglionic parasympathetic neurons.) There is a richer noradrenergic innervation acting on α_1 -adrenergic receptors to produce contraction of smooth-muscle cells in the bladder neck and trigonal area. These smooth muscles function as a “genital sphincter” that prevents back-flow of semen to the bladder during ejaculation. The role of sympathetic innervation of the bladder neck for maintaining the urethral pressure during the bladder-filling phase is not equally clear, however. Nevertheless, drugs acting on α adrenergic receptors affect bladder emptying. Thus, α -adrenergic agonists may worsen urinary retention in patients with an enlarged prostatic gland and,

conversely, selective α antagonists are used to improve bladder outflow in such patients. Alpha (α)-adrenergic receptors are present at many levels of the reflex pathway—centrally in the cord and at higher levels, in the autonomic ganglia, and in the smooth-muscle cell membrane; further, they are located both pre- and postsynaptically and are activated by both norepinephrine released from postganglionic sympathetic fibers and by circulating epinephrine. Thus, deciding the site of action of a certain adrenergic drug is not straightforward.

Central Control of Micturition

Normal emptying of the bladder requires more than intact spinal reflex arches. This is evident in patients with **transection of the cord** above the sacral level. If the lesion is complete, all descending connections acting on the sacral reflex centers are interrupted. The patients experience great difficulties with emptying the bladder because, among other problems, the activities of the detrusor muscle and the internal sphincter are not coordinated. This is called **dyssynergia**. (In addition, the bladder often becomes hyperactive—i.e., the emptying reflex is elicited at a lower pressure than before.) Patients with lesions of the brain stem above the pons do not have dyssynergia. Further observations show that the **dorsolateral pons** (near the locus coeruleus) contains a neuronal network that coordinate the spinal reflexes involved in micturition—the **pontine micturition center** (Fig. 29.2). Ascending spinoreticular fibers, which inform about the filling pressure, join the spinothalamic tract ventrolaterally in the cord. The fibers end in the **PAG**, which is assumed to send signals to the micturition center. The marked **emotional** influence on micturition, such as frequent urination in association with nervousness and fear, is probably mediated by connections from the **amygdala** to the PAG (see Fig. 31.6). Animal experiments indicate that the descending (reticulospinal) fibers from the pontine micturition center lie dispersed in the lateral funiculus, with the majority in its dorsal part. Clinical observations, however, indicate that in humans the fibers might lie more ventrally, close to the pyramidal tract in the lateral funicle. Thus, damage of the **pyramidal tract** in the spinal cord is often associated with **urge incontinence** (inability to inhibit the emptying reflex) as a sign of impaired central bladder control.

Although lesions above the pons do not disturb the normal emptying of the bladder, patients with such lesions may have problems with controlling of the initiation of micturition (inhibition in particular). The cell groups responsible for the voluntary control of micturition are not known in detail, although medial parts of the **frontal lobe** and the **hypothalamus** appear to be involved (Fig. 29.2). Clinical observations indicate that damage to the frontal lobes may cause **urge incontinence**.

Cortical Activity and Bladder Control

Positron emission tomography (PET) and functional magnetic resonance imaging (fMRI) studies indicate that neurons in the dorsolateral pons and the prefrontal cortex increase their activity during **micturition**. The activity is largest on the right side in both men and women. In persons trying to void without succeeding, the activity increases in the frontal lobe but not in the dorsolateral pons. There is sign of increased activity, however, in a slightly more ventral pontine region; in animals, this region controls the pelvic muscles (presumably including the external urethral sphincter) by descending reticulospinal fibers. When trying to **inhibit voiding** when the bladder was full, activity increases in the anterior **cingulate gyrus** and the **insula**. This fits with data showing that the insula receives visceral sensory inputs (see Chapter 34, under “The Insula”). Both the insula and the anterior cingulate gyrus are furthermore known to influence the autonomic system. In addition, the anterior cingulate gyrus is active during focused attention and when selecting appropriate behavior. The insula may be responsible for sending descending commands—eventually inhibiting the detrusor—when micturition should be postponed.

The Erection and Ejaculation Reflexes

As mentioned, erection depends on parasympathetic signals to the vessels of the penis or the clitoris, whereas ejaculation is due mainly to sympathetic signals to the smooth muscles of the ductus deferens and the prostate. The preganglionic neurons of these functions have their cell bodies in the lumbosacral cord (see Figs. 28.7 and 28.11). In addition, successful expulsion of the semen requires coordination of smooth muscles closing the upper urethra (the genital sphincter, mentioned above) and striated **muscles of the pelvic floor**—that is, sphincter muscles and muscles around the penile cavernous bodies, especially the **bulbospongiosus** (bulbocavernosus) **muscle**. The α motoneurons of the latter muscles are found in the sacral cord. In male rats, these groups of motoneurons are three times larger than in female rats. This **sexual dimorphism** arises early in the prenatal development and depends on the presence of minute amounts of androgens.

Evidently, erection and ejaculation require precise temporal coordination of several muscles. The coordination is carried out in the spinal cord by a complex **premotor network**. For example, the activity of sympathetic preganglionic neurons in the lower thoracic–upper lumbar cord must be coordinated with the activity of parasympathetic preganglionic neurons and α motoneurons in the sacral cord. Cell groups in the **reticular formation** of the **pons**, close to those controlling micturition, and in the **hypothalamus** control the spinal

reflex centers for erection and ejaculation. They are not necessary for erection and ejaculation, however. Thus, provided the lesion is above the midthoracic level, both erection and expulsion of the semen can occur after transection of the cord. In such patients, erection is evoked by sensory stimuli from the penis (the patient has no conscious sensations, however, because the ascending sensory tracts are interrupted). The hypothalamic cell groups are influenced, among other areas, from the cerebral cortex and the amygdala. Normally, of course, descending connections from these higher levels are involved in the induction of erection, more or less independently of the sensory signals that act through the spinal reflex arc. The spinal reflex centers of erection and ejaculation are **inhibited** from an area in the ventral part of the upper medulla by way of descending connections that act on parasympathetic and sympathetic preganglionic neurons and α motoneurons. The lack of such inhibitory connections may explain why some patients with transection of the cord above the sacral level suffer from **priapism**—that is, a hyperactive erection reflex.

VISCERAL PAIN

Salient Features

Visceral pain differs in many respects from pain originating in somatic structures; it is usually more diffuse, less precisely graded, and of a different quality than pain arising in the skin. Visceral disease processes, such as **appendicitis**, initially cause a vague discomfort, with nausea rather than pain. Also, a feeling of fainting may occur, probably due to lowered blood pressure. The initial discomfort or pain is typically felt deep in the midline in the thorax or the abdomen, regardless of the organ involved. This first phase usually lasts for minutes to some hours. Thereafter, the pain moves to sites more specific to the diseased organ, but not necessarily where the organ is located. The pain is typically felt in the body wall or the extremities innervated from the same spinal segments as the diseased organ. This **referred pain**—described in more detail later—is easier to localize and describe than the initial discomfort or pain. Skin areas with referred pain may become **hyperesthetic** (increased sensitivity for touch) or **hyperalgesic** (increased pain upon nociceptor activation). The hyperalgesia is usually most marked in muscles and can be accompanied by tonic contractions. Appendicitis, for example, usually leads to a bandlike hyperesthetic region of the skin corresponding to the dermatomes T_1 – T_{11} (see Fig. 13.14) on the right side. In addition, **circulatory changes** may occur in the same area, presumably caused by reflex actions on preganglionic sympathetic neurons. Thus, looking for such sensory and

circulatory changes may provide valuable diagnostic clues when suspecting diseases of visceral organs.

Typically, visceral pain is accompanied by **slow pulse**, **lowered blood pressure**, and **cold sweat**—that is, effects mediated by the autonomic nervous system. **Nausea** and vomiting, and a tendency to keep still are also frequent in patients with visceral pains.⁴ Many of these effects are probably orchestrated by the **periaqueductal gray (PAG)**. The PAG receives nociceptive signals and coordinates a varied specter of autonomic and somatic reactions to stressful stimuli (see Chapter 15, under “The Functional Role of the Pain-Suppressing Systems”). Several factors might contribute to the **diffuseness** of visceral pain. As mentioned, the density of sensory innervation is generally low in visceral organs with, consequently, very large receptive fields of the sensory units. In addition, the central branches of the visceral spinal ganglion cells run for a long distance, both rostrally and caudally in the cord. Thus, signals entering in one dorsal root can activate spinothalamic cells at many segmental levels.

Stimuli that Activate Visceral Nociceptors

The adequate stimuli for visceral nociceptors differ from those of somatic nociceptors. This may be so because visceral organs are not normally subject to stimuli that are potentially tissue damaging and thus require a change of behavior. In fact, stimuli that harm the visceral tissue may not provoke pain, while apparently innocuous stimuli may cause intense pain. Surgery of visceral organs (e.g., of the abdomen) can be performed without the patient feeling pain as long as the abdominal wall is anesthetized. Pain is felt only when the mesenteries or the peritoneum is pulled. Cutting with a sharp knife, for example, is not painful, and neither is tearing or pinching. Indeed, cancer of the lungs, kidneys, the liver, and the gut may cause considerable tissue destruction without evoking pain. Still, abnormal **distension** of a hollow organ or forceful **contraction of smooth muscles** in its wall produces intense pain. This occurs when a stone blocks the ureter or the gall ducts, or when something obstructs the intestine. The wall of the organ is distended above the obstruction, and forceful contractions of the smooth musculature are produced in an attempt to overcome the obstruction. Such spasmodic contractions usually occur with regular intervals, explaining the typical bouts of pain experienced in such cases. Another cause of visceral pain is **ischemia**, which can occur when an artery is narrowed by thrombosis or embolism. The most common example is the pain felt when a coronary artery is narrowed

(angina pectoris) or occluded (heart infarction). Pain may also be felt when viscera are **inflamed** and when **irritating substances** come in contact with the peritoneum, as in cases of perforation of a gastric ulcer or an inflamed gallbladder. The spread of bile, gastric content, or blood to the peritoneal cavity causes extreme pain and shock. Sensory units leading from the bladder change their behavior when the mucosa is inflamed. First, mechanoreceptors recording stretching of the wall become more sensitive (lowered thresholds); second, many “silent nociceptors” (see Chapter 13, under “Not All Nociceptors May Signal Tissue Damage”) start firing. Under normal conditions, even marked stretching of the wall does not activate such receptors, whereas in inflammation they respond to even moderate distension of the bladder. This may explain the frequent voiding in patients with cystitis and the urge to void even when the bladder is almost empty.

The **arteries** are also sensitive to painful stimuli. An arterial puncture (e.g., to draw a blood sample) is painful. Further, spasmodic contractions or strong dilatation of arteries are also painful. For example, the pain of **migraine** is most likely due to dilatation of extracranial arteries and subsequent inflammation.

Sympathectomy

Knowledge of the special features of visceral pain and of the segmental innervation of visceral organs (see Table 28.2) is of importance for surgical interventions of the autonomic nervous system. Such operations are performed most often on the sympathetic system (sympathectomy). Indications are chronic pain conditions, especially **complex regional pain syndromes (CRPS)**; see Chapter 15, under “Complex Regional Pain Syndromes (CRPS)—Special Type of Pathologic Pain”) but sympathectomies are also performed to increase the **blood flow** of the extremities (to interrupt vasoconstrictor fibers) and to reduce excessive sweating (**hyperhidrosis**). For example, interruption of the lumbar sympathetic trunk interrupts the sympathetic outflow to the lower extremities (see Fig. 28.9). The use of sympathectomies for these conditions is controversial, due to unwanted side effects and lack of convincing documentation of effects.

Sympathectomy or sympathetic block is also performed occasionally to interrupt the **pathways for pain** signals from visceral organs. For example, in some patients with intractable pain arising in the upper abdomen (usually because of terminal cancer of the pancreas or the stomach), the transmission of sensory signals can be blocked by alcohol injections into the **celiac plexus**. Because this procedure also blocks the efferent sympathetic signals, the patients experience transitory orthostatic hypotension due to vasodilatation in the abdominal organs.

4 The bouts of pain due to obstruction of a hollow visceral organ (the ureter or the intestine) are usually accompanied by an intense urge to move.

Referred Pain

As mentioned, pain of visceral origin is often not felt where the organ is located but in some other place, often in the body wall or the extremities (Fig. 29.3). This phenomenon is called **referred pain**. The referred pain—for example, in the left arm in the case of angina, and under the right scapula in the case of a gallstone—can be localized fairly precisely by the patient. At the site of the diseased organ, however, there is usually only a diffuse pain, difficult both to localize and to describe. As mentioned, it typically takes some time for referred pain to develop.

The most widely accepted explanation of referred pain is based on the observed **convergence** of signals from somatic structures (especially the skin) and viscera.⁵ Such convergence occurs in the dorsal horn, in the dorsal column nuclei, and in the thalamus, as discussed above and in Chapter 14 (under “Spinothalamic Neurons Receive Signals from Both Somatic and Visceral Structures”). As a rule, the diseased organ and the site of referred pain receive sensory innervation from the same **spinal segments**. Conceivably, the signals coming from the visceral organs are interpreted as arising in the skin and not in the visceral organ because signals from the visceral organs are never consciously perceived under normal circumstances (e.g., nociceptors of the heart are not normally stimulated). The referred pain usually takes some time to develop, and the occurrence of (referred) hyperesthesia and hyperalgesia is presumably explained by **sensitization** of central neurons receiving convergent inputs, leading to hyperexcitability and even spontaneous firing (i.e., independent of signals from the periphery). The importance of central sensitization is supported by the observation that the referred pain is felt even after blocking the sensory nerve fibers that lead from the area of referred pain. Further, the expansion of the painful area after some time is best explained by central sensitization of propriospinal neurons—transmitting signals to segments above and below those innervating the diseased organ. Indeed, animal experiments show that the receptive fields of dorsal horn neurons expand dramatically when the tissue from which they receive sensory signals is chronically inflamed.

Another peculiarity is that referred pain shows a tendency to localize to parts of the body that previously have been the site of a painful process. For example, a patient felt the pain of angina in a part of the spine that had been fractured many years ago. It seems as if a pain can be “remembered” as a persistent central sensitization (plasticity).

The peripheral axon of some **spinal ganglion cells** divides, with one branch going to the skin and another

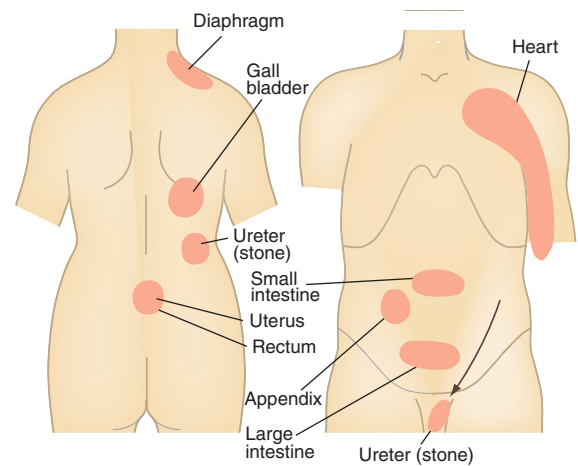


FIGURE 29.3 *Referred pain.* Examples of typical sites of pain in diseases affecting various visceral organs. (Based on Cope 1968.)

going to a visceral organ or a muscle. For example, one study found that about one-fifth of the fibers in the splanchnic nerve could be activated by electric stimulation of a somatic nerve. This may also contribute to the occurrence of referred pain. In such instances, an **axon reflex** (see later) may cause cutaneous hyperesthesia and vascular changes, as sensory signals from the visceral organ are conducted not only into the cord but also peripherally in the branch into the skin. There sensory fibers may release **neuropeptides** such as substance P and VIP, which cause changes of blood flow and sensitize sensory nerve endings (see Chapter 13, under “Primary Sensory Fibers and Neurotransmitters”).

Examples of Referred Pain

The pain arising in the **heart** is usually referred to the ulnar aspect of the left arm or the upper part of the chest (Fig. 29.3). These regions of the skin and the heart both receive sensory innervation from the upper thoracic segments of the cord. The **gallbladder** and the skin in the region of the lower end of the scapula are both innervated from the eighth thoracic segment. The shoulder pain on irritation of the **diaphragm** is explained by the common innervation from the fourth and fifth cervical segments. Pain from the urinary **bladder** can be referred to two areas of the skin: one innervated by the spinal segments S_2 – S_3 and one higher up on the back innervated by the lower thoracic and upper lumbar segments (see Fig. 13.15). This fits with the segments of the cord innervating the bladder (see Fig. 28.12).

Antidromic Signals and the Axonal Reflex

Electrical stimulation of dorsal roots can produce vasodilatation in the dermatome of the root concerned. This is caused by signals conducted in the peripheral direction by the sensory fibers. Signals conducted in the

⁵ That local anesthesia of the diseased organ prevents the development of referred pain proves that it depends on the nervous system.

direction opposite the normal direction are called **antidromic**. Of course, the action potentials are exactly the same as those conducted in the normal—**orthodromic**—direction. Antidromic signals in **C fibers** appear to release **substance P** from the peripheral branches. Substance P probably causes the release of **histamine** (presumably from mast cells). Histamine causes vasodilatation, especially of the capillaries, and at the same time the capillaries become leaky. Thus, a local edema is produced.

Various phenomena can probably be explained by this phenomenon. When the skin is stroked with a fairly sharp object, it reddens (vasodilatation) after a few seconds on both sides of the stripe. This can be explained as follows: the stroking of the skin stimulates C fiber nociceptors, and action potentials are conducted to the CNS (and we experience a sharp pain). At the same time, however, the action potential is also conducted peripherally in the branches of the C fiber that do not

innervate the stimulated skin stripe. These branches end in the skin outside the stripe, where they liberate substance P and cause vasodilatation. The process is called a reflex, and because it uses only the peripheral process of a pseudounipolar ganglion cell, it is called an **axon reflex**. The reflex cannot be elicited in an area of the skin that has been deprived of its sensory innervation, proving that the phenomenon is mediated by nerve fibers.

Under normal conditions, the antidromic signals in sensory fibers hardly play any role, but they may help explain certain pathological phenomena. An example from the **airways** can be mentioned. In disposed individuals, irritating gases can produce a marked edema of the mucous membranes. The edema is caused apparently by histamine release, which, in turn, is caused by substance P released by an axon reflex from the sensory fibers that innervate the mucous membrane (partly coming close to the surface of the epithelium).

The Central Autonomic System: The Hypothalamus

OVERVIEW

Sympathetic and parasympathetic parts cannot be clearly separated in the higher centers that control autonomic functions. This is not unexpected, as there is a constant need for coordination of the activity of preganglionic sympathetic and parasympathetic neurons. As discussed in Chapter 28, it is not so that *either* the sympathetic *or* the parasympathetic system is active. Further, higher levels can select specific subgroups of preganglionic neurons for a certain task. We have also mentioned regions in the brain stem—in the reticular formation in particular—that initiate various automatic behaviors, such as stereotyped, purposeful movement patterns and effects mediated by the autonomic nervous system. Such brain stem “centers” do not operate independently, however; in most instances their activity is modulated and controlled from higher levels of the brain, among which the hypothalamus has a prominent role. In addition, the hypothalamus plays a unique role—by means of the pituitary gland—in the superior control of the **endocrine system**.

Even though the hypothalamus is the part of the brain most closely linked with control of the autonomic nervous system, its functional role is wider. Thus, a central task of the hypothalamus is to **coordinate** autonomic, endocrine, and somatic motor responses to behavior that is appropriate for the immediate needs of the body, such as feeding, drinking, and reproduction. The overall aim is to maintain bodily **homeostasis** in a wide sense. The hypothalamus is also connected with higher parts of the central nervous system (CNS), however, including the cerebral cortex and the amygdala. These connections are important links in the interactions between **bodily** and **mental processes** (psychosomatic interrelations), which play a role in both health and disease. Especially, interactions between the nervous system and the **immune system** may help explain how mental states influence health and disease, and how bodily processes influence our mental states.

The **afferent connections** of the hypothalamus are multifarious, reflecting its integrative role. The hypothalamus receives information about olfactory and taste stimuli; the conditions in the gastrointestinal tract; the

blood pressure, noxious stimuli, and skin temperature; and the intensity of ambient light. Afferents from limbic structures inform about emotional and motivational aspects. Hypothalamic **efferent connections** are also widespread and have three main targets: the pituitary, autonomic preganglionic neurons, somatic-efferent cell groups, and higher levels of the brain (limbic structures and the cerebral cortex).

The role of the amygdala (and some other nuclei) and the cerebral cortex in the control of autonomic functions is discussed in Chapter 31.

CENTERS IN THE BRAIN STEM FOR COORDINATION OF BEHAVIOR

Physiological experiments have localized several regions of the brain stem involved in the control of autonomic processes (see Chapter 26, under “Control of Circulation and Respiration,” and Chapter 29, under “Visceral Reflexes”). Electrical stimulation and lesions of restricted parts of the reticular formation of the pons and medulla produce changes in blood pressure, cardiac activity, respiration, sweat secretion, gastrointestinal activity, and other processes. Such higher-level centers exert control of the lower autonomic reflex centers of the spinal cord and coordinate their activities. The anatomic substrates of the brain stem centers are groups of neurons in the medullary and pontine reticular formation; as a rule, the centers are not well-defined nuclei but are formed by extensive **networks** of interconnected neurons. Visceral afferent fibers ending in the **solitary nucleus** inform the networks of the state of the body. From there, signals pass to various premotor groups in the reticular formation, which, in turn, send efferents to motor nuclei (autonomic and somatic) in the brain stem and the cord. The brain stem centers thus ensure, for example, that the segmental vasomotor reflexes operate together to serve the needs of the whole organism, not only its individual parts. We discussed in Chapter 28 how the blood supply of other organs is subordinate to the needs of the brain and how the micturition center in the pons coordinates the activity of several smooth and striated muscles. Further, cell groups in the reticular

formation initiate movements and postural adjustments as parts of complex behaviors in response to thirst, hunger, external dangers, sexual arousal, and so forth. Several such behaviors are coordinated from the **superior colliculus** and the **periaqueductal gray (PAG)** in the mesencephalon by way of their efferent connections to the reticular formation. Connections from the PAG initiate coordinated alterations of circulation and respiration, pain perception, and automatic movements in response to threatening or novel stimuli. Pain, for example, might be understood not merely as signal of tissue damage but as signaling the need to change behavior.

A perfect coordination cannot be performed by the brain stem autonomic centers, however. This is witnessed by the poor control of autonomic functions such as blood pressure and body temperature in **decerebrate animals**. Optimal autonomic control, and coordination of autonomic with endocrine and somatic processes, requires that the brain stem and spinal centers be supplied with afferent fibers from higher centers, especially in the hypothalamus but also in the amygdala (which acts on the brain stem partly through the hypothalamus).

PAG Coordinates Behavior in Response to Threatening Stimuli

The PAG is not a unit, either anatomically or functionally. It consists of several columnar groups of neurons, each differing with regard to connections. Stimulation of a **dorsolateral column** elicits arousal, tachycardia, rise in blood pressure, and increased respiration, and it facilitates orienting responses. At the same time, a **non-opioid** analgesia is produced. The actions are mediated by, among others, connections to the locus coeruleus, the ambiguous nucleus, the solitary nucleus, and parts of the reticular formation but also by ascending connections to the anterior hypothalamus and the intralaminar thalamic nuclei. Together, this part of the PAG initiates **defensive behavior** in response to strong emotions or aversive external stimuli. Stimulation of a **ventrolateral column** in the PAG elicits inhibition of movements, like the “**freezing**” response to sudden fear (e.g., when a rat sees a cat). In addition, the PAG probably mediates the immobility typical of conditions with strong visceral pain. The immobility is accompanied by fall in blood pressure, bradycardia, and **opioid-dependent** analgesia. These effects are mediated by efferent connections to the nucleus raphe magnus (NRM) and the reticular formation in the ventrolateral medulla, and to the lateral hypothalamus and the basal forebrain.

How the brain chooses between the two kinds of behavior that can be elicited from the PAG is not quite clear. Presumably, descending commands from higher levels (probably routed through the hypothalamus and the amygdala) differ, depending on the nature of the stimuli and their context.

STRUCTURE AND CONNECTIONS OF THE HYPOTHALAMUS

This small part of the brain, weighing only 4 to 5 g in humans, is a mosaic of minor nuclei that can be distinguished on the basis of their cytoarchitectonics, connections, cytochemistry, and physiological properties. Only the main features are discussed here.

Hypothalamic Nuclei

In the wall of the third ventricle, below the hypothalamic sulcus, some identifiable nuclei are embedded in a more diffuse mass of neurons (Figs. 30.1–30.3); together these constitute the hypothalamus. The borders between the hypothalamic nuclei and the neighboring regions are not sharply demarcated. This explains why different authors have drawn the borders differently. Figure 30.3 shows the classic subdivisions of the British neuroanatomist Le Gros Clark. It is now common to distinguish a **medial part** of the hypothalamus, containing several discernible nuclei, and a **lateral hypothalamic area** (or nucleus) with a diffuse structure. Numerous longitudinally running fibers traverse the lateral hypothalamic area. These fibers are often collectively termed the **medial forebrain bundle**, though this is not a single tract anatomically or functionally and has no sharp borders (see Chapter 31, under “The Medial Forebrain Bundle”).

Within the medial part, one can distinguish **anterior**, **middle** (tuberal), and **posterior** (mammillary) nuclear groups. Particularly well defined are two anterior large nuclei—the **paraventricular nucleus** and the **supraoptic nucleus** (Fig. 30.3). The first is located close to the wall of the third ventricle; the latter is just above the optic chiasm (Fig. 30.1). In the same region lies the **suprachiasmatic nucleus** (SCN) that functions as a biologic clock (see later, “Circadian Rhythms”).¹ In the middle, or tuberal, nuclear group we find the **ventromedial**, the **dorsomedial**, and the **arcuate** (infundibular) nuclei. The latter nucleus is located in the bottom of the third ventricle (Figs. 30.1 and 30.3; see Fig. 6.24), below the ventromedial nucleus. In the posterior part of the hypothalamus we find the **posterior nucleus** close to the ventricular wall, whereas the characteristic **mammillary nucleus** (consisting of several subnuclei) is located in the bottom of the ventricle (Fig. 30.3; see also Fig. 33.8). The term **mammillary body** is used for the macroscopically visible part of the mammillary nuclei (see Fig. 6.13). In general, short intrahypothalamic fibers mediate a high degree of cooperation among the hypothalamic nuclei. Notably, there are numerous reciprocal connections

¹ The suprachiasmatic nucleus cannot be identified in ordinary thionine-stained sections in contrast to the supraoptic and paraventricular nuclei. It is clearly visible, however, by use of immunocytochemical identification of several neuropeptides (vasopressin, VIP, neuropeptide Y, and neurotensin).

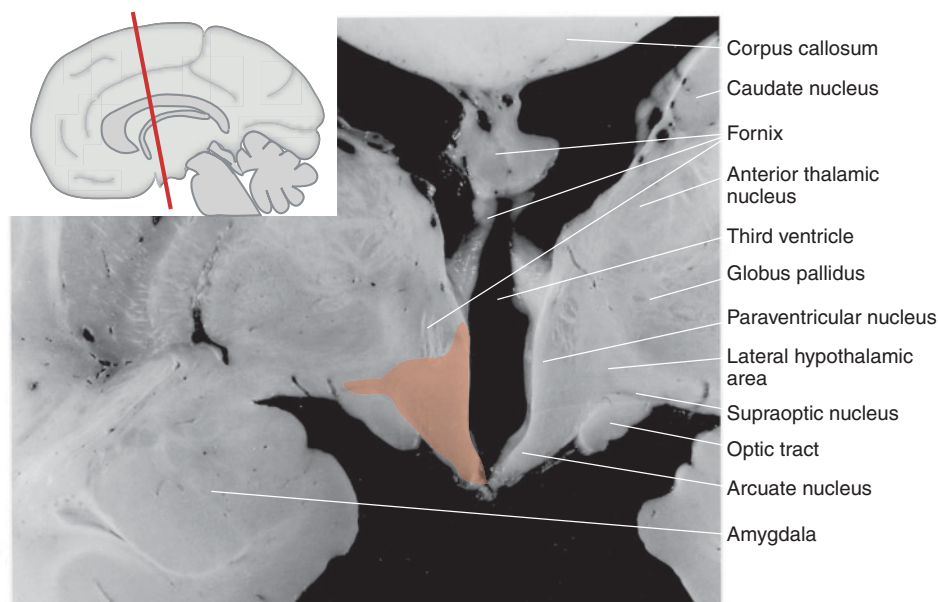


FIGURE 30.1 The hypothalamus. Frontal section through the hemisphere.

between the medial nuclei and the lateral area (nucleus) of the hypothalamus.

The Hypothalamus Contains Many Neurotransmitters

Numerous neurotransmitters are present in the hypothalamus, as demonstrated with immunocytochemical and biochemical methods. **Acetylcholine, norepinephrine, dopamine, serotonin, histamine,** and many **neuropeptides** occur with a differential distribution among the hypothalamic nuclei. Norepinephrine is among the neurotransmitters found in highest concentration (some of the norepinephrine is related to terminals of fibers from the locus coeruleus). Some of the neuropeptides are involved in the hypothalamic control of the pituitary gland or are released to the bloodstream in the pituitary. We return to some of the neuropeptides and

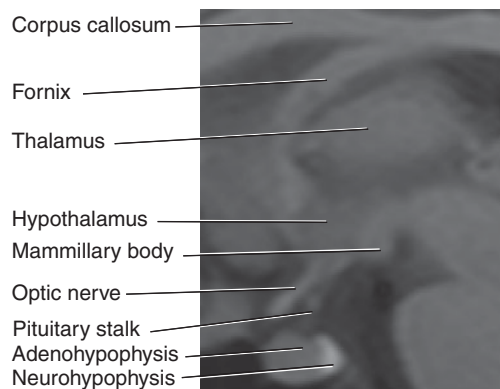
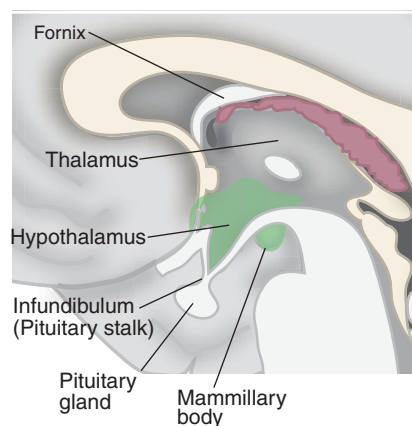


FIGURE 30.2 The hypothalamus and the pituitary as seen in a midsagittal MRI scan. (Courtesy of Dr. S.J. Bakke, Rikshospitalet University Hospital, Oslo, Norway.)

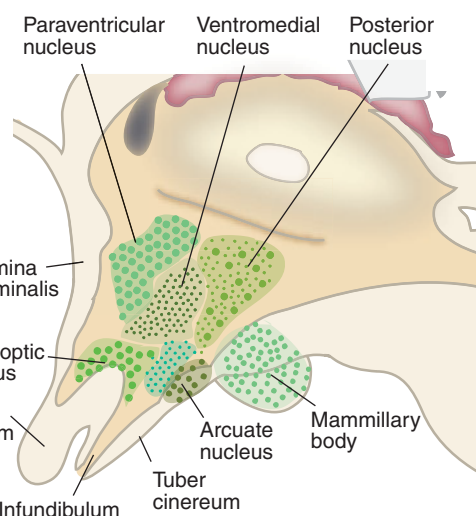


FIGURE 30.3 The hypothalamus. Median section through the third ventricle. Some of the major hypothalamic nuclei are shown with colored dots. The size of the dots indicates the relative size of the neurons of the various nuclei. (Redrawn after Le Gros Clark et al. 1936.)

their possible functions later in this chapter. Here we note that each neuropeptide takes part in different functional tasks, even though only one is mentioned in this discussion. The functional role of the various neurotransmitters in the hypothalamus is still incompletely known.

Afferent Connections and Other Kinds of Input to the Hypothalamus

Figure 30.4 shows diagrammatically the main afferent connections of the hypothalamus. It is immediately clear that many—perhaps most—parts of the brain are able to influence the hypothalamus! (The mammillary nucleus is in several aspects different from the other hypothalamic nuclei and is treated separately later in this chapter.) In addition, the hypothalamus is special because it receives information by **hormones** acting on specific receptors, and that it contains special **sensory neurons** that record blood temperature (**thermoreceptors**) and salt concentration (**osmoreceptors**). In addition, information about temperature and water balance is brought to the hypothalamus from peripheral receptors in the body. These features reflect the functions of the hypothalamus in homeostatic control.

Hypothalamic afferent nerve fibers bring signals from most kinds of **sense organ** and from higher levels of the brain, such as the **cerebral cortex** and the **limbic structures**. Thus, the hypothalamus receives information about **olfactory** and **taste** stimuli; the conditions in the **gastrointestinal tract**; the **blood pressure**, **noxious stimuli**, and **skin temperature**; and the intensity of **ambient light**. The afferent fibers from limbic structures, such as the amygdala, inform about **emotional** and **motivational** aspects.

The many groups of afferents end in at least partially different parts of the hypothalamus, which fits with physiological evidence that the hypothalamus consists of many functionally different parts (we discuss some of these later in this chapter). Nevertheless, both the many intrinsic connections and the properties of single cells show that the various afferent signals are considerably processed and integrated before commands are sent out to various targets.

The Connections and Functions of the Hypothalamic Nuclei Have Been Difficult to Clarify

Determining the exact connections and functional roles of the various nuclei has proved more difficult in the hypothalamus than in most other parts of the brain. This is partly because the nuclei are so small and are located in a part of the brain that is difficult to reach with experimental manipulations and partly because most of the afferent and efferent fibers are unmyelinated and mixed with fibers destined for other parts of the brain. In particular, lesions or stimulations of the lateral hypothalamic area are bound to affect the medial forebrain bundle and, thereby, fibers destined for other regions than the hypothalamus. Further, the rich network of intrahypothalamic connections means that a lesion of one nucleus will interfere with the functioning of several others as well. Modern methods using toxic agents that destroy the cell bodies without affecting fibers of passage have helped to settle some controversies, however. Finally, there are notable differences in both connections and neurotransmitters in various species, and most experimental data have been obtained in rats or cats.

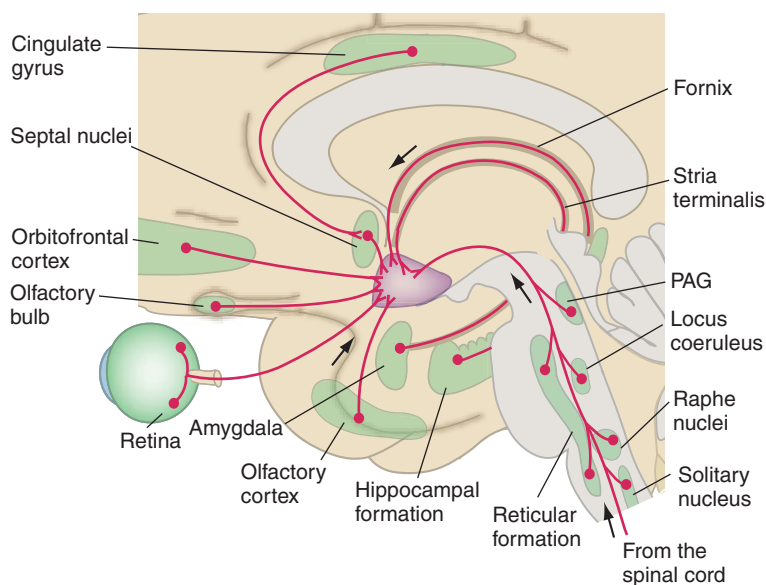


FIGURE 30.4 Main afferent connections of the hypothalamus. Arrows indicate direction of impulse conduction.

Efferent Connections

The efferent connections of the hypothalamus are widespread (Fig. 30.5), enabling this tiny part of the brain to influence the **pituitary**, **autonomic preganglionic neurons**, **somatic-efferent** cell groups, and **higher levels** of the brain. The hypothalamus sends fibers to most of the cell groups from which it receives afferents, such as the amygdala, the septal nuclei, nuclei of the reticular formation, the PAG, some of the cranial nerve nuclei, and the spinal cord. The hypothalamus can also influence the prefrontal cortex.

Hypothalamic influences on **autonomic processes** such as blood pressure, heart rate, temperature regulation, and digestion are mediated by direct and indirect descending connections to the preganglionic sympathetic and parasympathetic neurons. Many hypothalamic neurons send axons to parasympathetic cell groups of the brain stem (such as the motor nucleus of the vagus) and to the sympathetic neurons of the intermediolateral column of the cord. In addition, the reticular formation receives many afferents from the hypothalamus and mediates effects on brain stem nuclei.

Hypothalamic effects on the **cerebral cortex** occur through direct projections and indirectly via the thalamus (the latter route concerns especially the mammillary nucleus; see later). Although the mediodorsal thalamic nucleus (MD) is unlikely to receive direct hypothalamic afferents, several nuclei mentioned above that receive fibers from the hypothalamus project to the MD (such as the amygdala and the septal nuclei). The MD projects to the **prefrontal cortex** (see Fig. 33.7). Direct connections from the hypothalamus pass to the **orbitofrontal cortex**

and **temporal association areas**. Some of these direct hypothalamocortical fibers are **GABAergic**. In addition, a diffusely organized **histaminergic** projection to large parts of the cortex arises in the small **tuberomammillary nucleus**. This projection may be involved with other ascending connections in cortical activation (see Chapter 26).

More about Hypothalamic Afferent Connections

A **retinohypothalamic tract** supplies the hypothalamus with visual information about the **total amount of light** (not patterned vision). This information helps control bodily functions with daily variations (see later, “The Hypothalamus and Circadian Rhythms”). In addition to direct projections from the olfactory bulb, **olfactory** information reaches the hypothalamus indirectly from regions receiving fibers from the olfactory bulb, such as the amygdala and the olfactory cortex (Fig. 30.4). This information is important for **sexual behavior** (shown in rats) and for autonomic control of **digestion**. Physiological studies show that **auditory** information can reach the hypothalamus, and the same holds for signals from **cutaneous receptors**. Anatomically, direct fibers have been traced from the **spinal cord** to the hypothalamus. Many such direct fibers arise in **lamina I** and convey nociceptive and other kinds of information needed for homeostatic control (see Chapter 29, under “Central Pathways for Visceral Afferent Signals”). They may also contribute to autonomic and endocrine responses to noxious stimuli. Most information from the spinal cord reaches the hypothalamus indirectly via the **reticular formation**, however. In that case, the sensory information is presumably highly integrated—that is, not providing specific information about stimulus features. Sensory signals from visceral organs reach the hypothalamus after synaptic interruption in the **solitary nucleus** and, most likely, a further synaptic relay in the reticular formation. Such signals, for example, may inform about the filling of the stomach, which influences the feeling of hunger or satiety.

As mentioned, fibers from the **amygdala** may convey olfactory information, but they also provide signals related to emotions and motivation. The **hippocampal formation** in the medial part of the temporal lobe (another of the limbic structures) sends many fibers in the **fornix** to end in the hypothalamus. We do not know what information these fibers transmit. Presumably, it is a synthesis of several sensory modalities. The **ventral striatum** sends fibers to the lateral hypothalamus, and the **cingulate gyrus** can act via the **septal nuclei**. Part of the **orbitofrontal cortex** has direct connections to the hypothalamus, presumably mediating processed information about taste and smell; that is, information that has been evaluated regarding its significance. Sensory signals linked to **motivation** may also be conveyed to the hypothalamus from the orbitofrontal cortex.

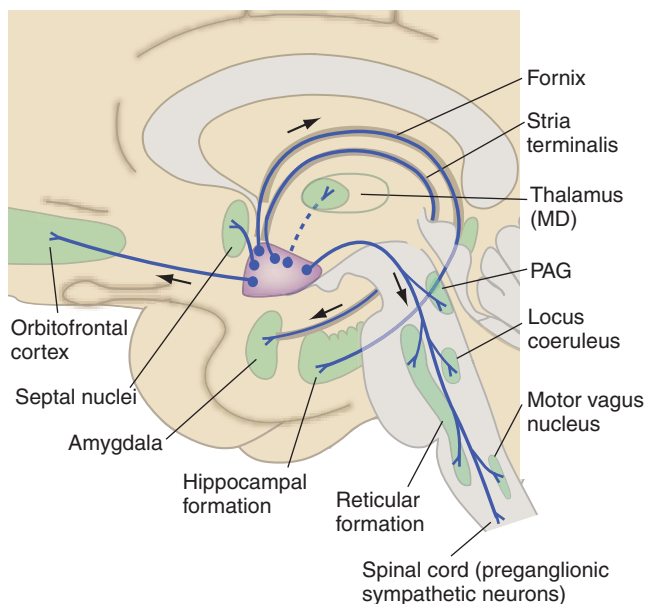


FIGURE 30.5 Main efferent connections of the hypothalamus. The connections to the pituitary (Fig. 30.7) gland are not included, nor are the connections of the mammillary nucleus (see Fig. 30.6).

The Mammillary Nucleus

Situated most posteriorly in the hypothalamus, the mammillary nucleus differs in several respects from the other nuclei. It consists of several subnuclei and forms the characteristically shaped mammillary bodies (one on each side) at the ventral aspect of the diencephalon (Fig. 30.3; see Figs. 6.13 and 6.15). The subnuclei differ to some extent with regard to connections, although we will disregard such differences here. The mammillary nucleus appears to be an important link in the signal transmission from structures in the temporal lobe to the cingulate gyrus. These connections, and thus the mammillary nucleus, are most likely concerned with **spatial memory** and **orientation** in space (see Chapter 32, under “Amnesia Caused by Lesions outside the Medial Temporal Lobe”).

Among the **efferent** mammillary connections (Fig. 30.6), the **mammillothalamic tract** (the bundle of Vicq d’Azyr) is most conspicuous. These fibers, forming a thick bundle, run anteriorly and upward in the wall of the third ventricle to end in the **anterior thalamic nucleus** (see Figs. 6.24 and 33.7), and this nucleus projects to the **cingulate gyrus** (Fig. 30.6). Thus, a major target of the mammillary nucleus is the cerebral cortex. Another large efferent fiber tract descends in the brain stem as the **mammilotegmental tract**. These fibers end primarily in certain nuclei of the mesencephalic and pontine reticular formation. Fibers also pass to the pontine nuclei, enabling signals from the mammillary nucleus to reach the cerebellum.

Most of the **afferent** fibers to the mammillary nucleus originate in the **hippocampal formation** and pass in the **fornix** (Fig. 30.6; see Figs. 6.23 and 32.2). In humans, the fornix contains more than 1 million axons. In addition, the mammillary nucleus receives afferents from the septal nuclei and some brain stem nuclei.

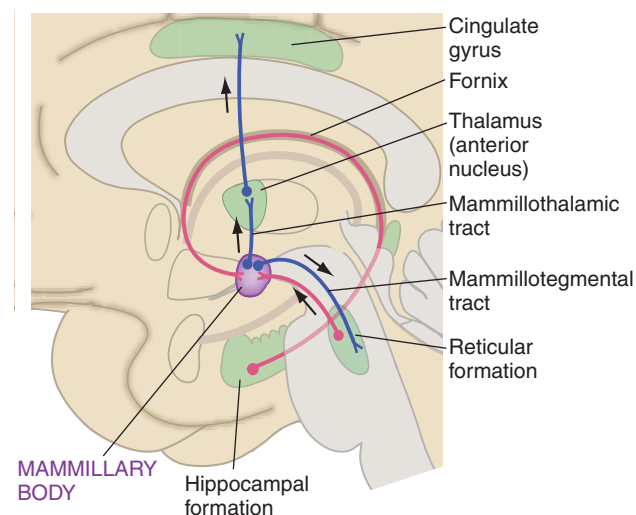


FIGURE 30.6 Main connections of the mammillary nucleus.

Hormones that Act on Hypothalamic Neurons

In addition to receptors for the endogenous neurotransmitters, many hypothalamic cells contain receptors for various hormones, such as for the steroid **sex hormones**, **thyroid hormones**, hormones of the **adrenal cortex**, and hormones released from the **anterior pituitary**. Often these hormones act on the brain as part of feedback loops to regulate the production of hormones from the pituitary. Hormones also influence other functions of the brain, however.

The **lipid-soluble** steroid hormones (sex hormones and the hormones of the adrenal cortex) and thyroid hormones pass the blood–brain barrier easily. The steroid hormones bind to intracellular receptors and thereby influence the activity of hypothalamic cells (and cells in other parts of the brain, particularly in certain limbic structures). The binding of **sex hormones** in the brain influences reproductive behavior, as shown in animal experiments. **Glucocorticoids** (from the adrenal cortex) bind to specific receptors in the paraventricular nucleus, among other places. Binding of glucocorticoids elicits behavior with the aim of increasing carbohydrate intake in experimental animals.

Some hormones that do not pass the blood–brain barrier can nevertheless act on the hypothalamus and certain other brain regions. This is possible because certain regions of the brain close to the ventricles lack a blood–brain barrier (see Chapter 7, under “Some Parts of the Brain Lack a Blood–Brain Barrier”), and the neurons in such regions send axons to other parts of the brain. One example is the **subfornical organ**, located below the fornix close to the wall of the third ventricle. The hormone **angiotensin II**, which is produced in response to a blood-volume reduction, binds to specific receptors in the subfornical organ. The latter sends axons to the **supraoptic** and **paraventricular** hypothalamic nuclei (which produce the antidiuretic hormone). This exemplifies that hormones acting on the hypothalamus (directly or indirectly) provide feedback information of importance for the hypothalamic control of the pituitary gland. We mention other hormonal actions on the hypothalamus and other parts of the brain later in this chapter.

THE HYPOTHALAMUS AND THE ENDOCRINE SYSTEM

Direct and indirect connections to the preganglionic sympathetic and parasympathetic neurons explain the effects of the hypothalamus on organs innervated by the autonomic nervous system. In addition, the hypothalamus acts on the endocrine organs by controlling the superior endocrine organ, the **pituitary**. Accordingly, diseases or lesions of the hypothalamus can disturb functions controlled by the endocrine system, such as sexual functions,

growth, and metabolism. The pituitary gland (the hypophysis) consists of an **anterior lobe**, the **adenohypophysis**, which develops from the epithelium of the primitive foregut and consists of clusters of epithelial cells with a rich supply of wide capillaries (sinusoids). The **posterior lobe** of the pituitary, the **neurohypophysis**, develops from the neural tube and consists of nerve terminals of fibers from the hypothalamus and a special kind of glial cell, the **pituitocytes**.

Hypothalamic control concerns both the anterior and posterior parts of the pituitary (Fig. 30.7). Two different pathways exert the hypothalamo–pituitary interactions. The posterior pituitary receives a direct neural tract—often referred to as the **supraopticohypophyseal** or the **hypothalamohypophyseal tract**—whereas the so-called **tuberoinfundibular tract** and a special **portal vascular system** reach the anterior pituitary (Fig. 30.6C). We return to this later.

The Anterior Pituitary Produces Several Hormones

The epithelial cells of the adenohypophysis produce and secrete the following hormones:

1. **Growth hormone (GH)** or **somatotropic hormone**, which stimulates body growth, particularly growth of long bones
2. **Thyroid-stimulating hormone (TSH)**
3. **Adrenocorticotrophic hormone (ACTH)**, which stimulates the production of steroid hormones, such as cortisol, in the adrenal cortex²
4. Two **gonadotropic hormones**—one **follicle-stimulating hormone (FSH)** that promotes the growth of the oocyte and its surrounding follicle cells and one **luteinizing hormone (LH)** that is necessary for the ovulation and formation of the corpus luteum from the follicular cells
5. **Prolactin** (or lactogenic hormone), which stimulates growth of the mammary gland during pregnancy and maintains the milk production during the nursing period

Various observations indicate that, as a rule, a specific cell type produces each hormone. The cells are named for the hormone they produce, and are called **somatotrophs** (GH), **thyrotrophs** (TSH), **mammotrophs** (prolactin), and so forth. In routine histological sections, however, only three kinds of epithelial cell can be recognized in the anterior pituitary: **acidophils**, **basophils**,

and **chromophobes**. The acidophils produce GH and prolactin, whereas the basophils probably produce the rest. The chromophobes may represent precursors to the acidophils and basophils. It follows that both the basophils and the acidophils are heterogeneous groups, as indeed has been shown with immunocytochemical techniques with antibodies raised against the various hormones.

Relationship between the Hypothalamus and the Posterior Pituitary

Two peptide hormones are released to the bloodstream in the posterior pituitary: **vasopressin** or **antidiuretic hormone (ADH)** and **oxytocin**. Both hormones consist of nine amino acids, are synthesized in the hypothalamus, and are brought to the pituitary by axonal transport. The thin, unmyelinated axons reaching the posterior lobe (in humans, about 100,000) come from two nuclei in the anterior part of the hypothalamus: the **supraoptic nucleus** and the **paraventricular nucleus** (Figs. 30.1, 30.3, and 30.7A). Most of the hormone-producing cells are large, with large vesicles in their cytoplasm that contain precursor molecules of the final hormones. The neurons are collectively termed the **magnocellular neuroendocrine system** (to distinguish them from the parvocellular neuroendocrine system that is discussed later). Even though vasopressin and oxytocin are produced in both cell groups, vasopressin is produced predominantly in the supraoptic nucleus and oxytocin mainly in the paraventricular nucleus. The hormones can also be demonstrated within the axons, which end with large nerve terminals in close contact with the fenestrated capillaries of the posterior lobe (Fig. 30.7B). Action potentials invading the nerve terminals constitute the signal for release of the hormone (just as for the release of neurotransmitter in ordinary nerve cells). The cells of the supraoptic and the paraventricular nucleus are called **neurosecretory** because they have all the characteristic features of neurons but, at the same time, release their product to the bloodstream.

Vasopressin: Control of Osmolarity

Vasopressin (ADH) was extracted from the posterior lobe quite early, before the neural connection between the hypothalamus and the posterior lobe had been ascertained. The hormone acts by increasing the water reabsorption in the kidneys by acting on **aquaporins** (water channels)—that is, it reduces the urine secretion (the hormone also elicits contraction of vascular smooth-muscle cells, which explains why it is also called vasopressin). It was known that destruction of the posterior lobe leads to a condition called **diabetes insipidus**, which is characterized by daily urine volume of 10 to 15 liters; diabetes insipidus also occurs as an inherited disease.

2 The hormone ACTH is synthesized from a large precursor protein called **proopiomelanocortin** or pro-ACTH/endorphin. This precursor molecule is cleaved into other peptides, notably β **lipotropin** (β -LPH) with a yet-unsettled function, and β **endorphin**, which is a potent opioid peptide with inhibitory actions on pain transmission. The functional role of the β endorphin secreted from the pituitary is not clear, however. Beta (β) endorphin is also found in a hypothalamic nucleus (the arcuate nucleus) with projections to brain stem nuclei of importance for pain transmission (see Chapter 15, under “Opiates and Endorphins”).

Patients with this disease have pronounced cell loss in the supraoptic and paraventricular nuclei. Later it was discovered that the disease could also be produced by cutting the pituitary stalk. This was the beginning of a full understanding of the nature of the relationship between the hypothalamus and the pituitary. The production of ADH varies in accordance with the **osmolarity** of the blood.³ Most likely, the cells of the supraoptic nucleus (and in some other nuclei) function as **osmoreceptors**. When the osmotic pressure of the blood increases because of extraordinary loss or reduced intake of fluid (e.g., by heavy sweating, diarrhea, or vomiting), the cells of the supraoptic nucleus are excited and increase the frequency of their action potentials, thus releasing more ADH into the bloodstream. This results in reduced urine volume (the urine becomes more concentrated). At the same time, the synthesis of the hormone is increased. Even a salty meal is enough to stimulate the osmoreceptors. Therefore, the hypothalamus is a control center for the body's "housekeeping" of water.

Oxytocin: Parturition and Milk Ejection

Oxytocin elicits contraction of the smooth-muscle cells in the wall of the **uterus** and thus has a role during parturition. It also produces contraction of the smooth-muscle cells (myoepithelial cells) of the mammary gland, thereby assisting in emptying the breast of milk. When the infant suckles, sensory impulses travel from the nipple (through the spinal nerves) to the cord and further to the hypothalamus, where the neurons of the paraventricular nucleus are influenced. Increased firing frequency leads to increased secretion of oxytocin to the bloodstream, and the hormone reaches the mammary gland in seconds. This is called the **milk ejection reflex**. It is special in that only the afferent link is neural; the efferent link is humoral. Although oxytocin is present in males, its function is so far unknown.

Vasopressin and Oxytocin Act in the Brain as Well as Peripherally

Although the peripheral actions of vasopressin and oxytocin have received most attention, both peptides exert effects on central neurons, influencing many aspects of behavior and cognition. For example, central infusion of oxytocin induces maternal behavior in rats, and oxytocin furthermore facilitates pair bonding (preference for a particular mate) in monogamous species. Central actions of vasopressin include sexual behavior, social

interactions, and anxiety reduction (and several other effects). These effects are produced by binding of vasopressin and oxytocin to specific receptors, modulating the activity of task-specific networks in several parts of the brain.

How the peptides reach their many central targets is less clear, however, because both peptides are mainly synthesized in the magnocellular neurons of the supraoptic and paraventricular nuclei that send their axons to the posterior lobe of the pituitary. However, some small (parvocellular) neurons in the paraventricular nucleus produce the peptides and send their axons to various central nuclei. Further, at least in some species, neurons in the suprachiasmatic nucleus, the amygdala, and in the basal forebrain also express vasopressin and oxytocin. Finally, the neurons of the magnocellular nuclei release peptides from the soma and the dendrites (Fig. 30.7B). Dendritic release probably enables vasopressin and oxytocin to act by volume transmission on neurons expressing the appropriate receptors near the paraventricular and supraoptic nuclei. While both peptides enter the cerebrospinal fluid (where their concentrations are usually higher than in the blood) it is not known whether this plays a functional role.

Influence of the Hypothalamus on the Anterior Pituitary: The Hypophyseal Portal System

It has long been known that altered growth, metabolism, and sexual functions—processes that are controlled by hormones produced in the anterior pituitary—can accompany diseases affecting the hypothalamus. There are no axonal connections from the thalamus to the anterior pituitary, however, and mechanisms other than those concerning the posterior lobe must be responsible for the influence of the hypothalamus on the anterior lobe. The discovery of a special vascular arrangement in the infundibulum (stalk) of the pituitary—the **hypophyseal portal system**—was a breakthrough in this respect (Fig. 30.7C). Most of the arteries reaching the anterior pituitary do not branch into capillaries among the epithelial cells but continue upward into the stalk (some arteries enter the stalk directly). In the upper part of the stalk, the vessels form wide capillaries (sinusoids) that finally collect into large veins. These hypophyseal **portal veins** course back to the anterior lobe, where they form a new set of sinusoids among the epithelial cells. From these sinusoids, the blood collects in veins that leave the pituitary. Because the blood in the sinusoids of the anterior lobe has first been through a capillary net in the stalk, substances can be transported from there to the anterior lobe. Numerous thin axons from the hypothalamus end in the uppermost part of the hypophyseal stalk, forming the **tuberoinfundibular tract**. This region is called the **median eminence** (Fig. 30.7C). The axons that end in contact with the capillaries in the median

³ The ADH secretion is also influenced by other factors, although in primates these appear to be much less potent than changes of osmolarity. For example, reduced blood volume leads to concentration of **angiotensin II** in the bloodstream, which, in turn, affects the supraoptic and paraventricular nuclei by means of the subfornical organ. **Ethanol** reduces the secretion of ADH, thereby increasing urine production.

eminence transport **peptides** by axonal transport and release them into the hypophyseal portal system. From the capillaries in the median eminence, the blood brings the peptides to the second capillary network among the epithelial cells of the anterior lobe.

The neurons that send axons to the median eminence lie mainly close to the wall of the third ventricle (periventricularly). Because the neurons are relatively small, they are collectively termed the **parvocellular neuroendocrine system** (to distinguish them from the magnocellular system). Many of the neurons are found in the **arcuate nucleus** (Figs. 30.1 and 30.2).

Most of the peptides transported by the portal system cause hormonal secretion from the epithelial cells of the anterior lobe and are therefore called **releasing hormones** or factors. A single releasing hormone acts preferentially (but not only) on a specific kind of cells in the anterior pituitary and thus produces secretion of mainly one hormone. Some peptides transported in the tuberoinfundibular tract have an inhibitory effect on the secretion of anterior lobe hormones and are called **inhibitory hormones** or factors. Most of the neurons of the parvocellular system release more than one peptide, however. The neurons may therefore have several different actions on the anterior pituitary. Which releasing (peptide) hormone that is preferentially released from a parvocellular neuron depends both on the level of the anterior pituitary hormone in the bloodstream (feedback) and on afferent connections from other parts of

the brain. Thus, the interplay between the hypothalamus and the endocrine system is complex.

Releasing hormones are present also in the hypothalamic neurons that send axons to other parts of the brain, where they act at conventional synapses. Examples are **somatostatin** (inhibits the release of GH), **thyrotropin-releasing hormone** (TRH), and **corticotropin-releasing hormone** (CRH). Thus, these peptides function as releasing hormones at one site and as neurotransmitters at another. This emphasizes the futility of assigning one function to each peptide. We return to the central actions of some of these peptides in Chapter 31.

More about Some Releasing Hormones

In humans, neurons producing **CRH** (corticotrophin-releasing hormone or factor, CRF) are mainly found in a particular small-celled subdivision of the **paraventricular nucleus**. ACTH stimulates the secretion of glucocorticoids from the adrenal cortex, and such secretion is an important response to stress (see below). Besides multifarious metabolic effects, the glucocorticoids act on the anterior pituitary and the hypothalamus to inhibit the synthesis and release of ACTH and CRH (negative feedback). CRH is also involved in emotional responses (see Chapter 31, under “The Amygdala, Anxiety, and Neurotransmitters”). The neurons containing CRH also express several other peptides (among them vasopressin and enkephalin).

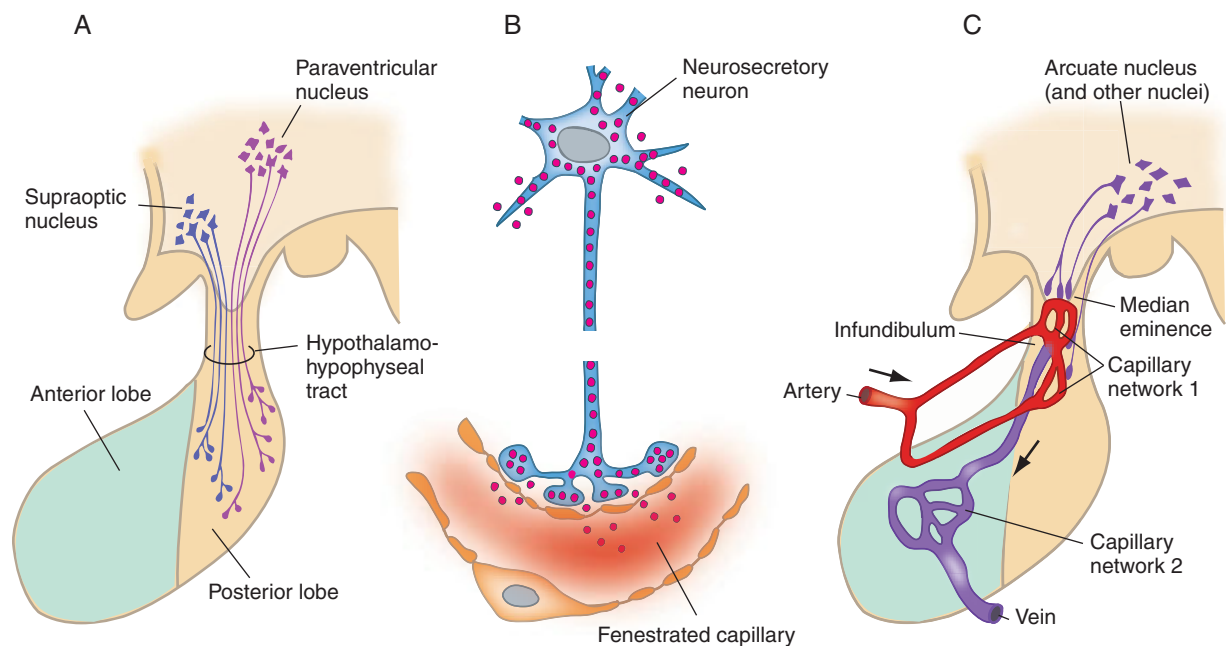


FIGURE 30.7 Relationship between the hypothalamus and the pituitary gland. **A:** Connections from the hypothalamus to the posterior lobe. **B:** Axonal transport of peptide hormones (neuropeptides) from the hypothalamus to the pituitary. **C:** The portal vessels of the

pituitary stalk ensure that releasing hormones (factors) are transported from the median eminence in the upper part of the stalk to the epithelial cells of the anterior lobe.

GRH (Growth hormone–releasing hormone) is found mainly in the **arcuate nucleus**, which, as mentioned, sends its efferents to the median eminence. The peptide **somatostatin**, which inhibits the release of the growth hormone, is present in neurons of the arcuate nucleus but also appears in many other cell groups that do not project to the median eminence. **Prolactin** is also controlled by two hypothalamic peptides, the stimulating **prolactin-releasing hormone** (PRH) and **dopamine** (also called prolactin release–inhibiting hormone, PIH). Hypothalamic neurons containing dopamine are mainly found in two small paraventricular cell groups and in the supraoptic nucleus.

We do not know why some pituitary hormones are controlled by two hypothalamic peptides with opposite effects, while others are controlled by only one.

FUNCTIONAL ASPECTS

The preceding account of its connections suggests that the hypothalamus functions as a coordinator of information of many different kinds, particularly pertaining to homeostasis and “bodily maintenance.” The hypothalamus contains networks of neurons that serve as control centers for several functions, such as blood pressure, body temperature, water balance, metabolism, digestion, reproduction, and defensive responses. Further, the hypothalamus governs the rhythmic variations of several bodily processes. For each of the mentioned broad functional categories, the hypothalamus serves to coordinate endocrine, autonomic, and somatic motor responses into appropriate behavior.

Early lesion experiments indicate that the anterior parts of the hypothalamus are concerned with predominantly parasympathetic effects, whereas the posterior parts are more involved in functions carried out by the sympathetic system. This should serve only as a rough rule of thumb, because further studies show that, as a rule, it is not possible to localize functions to specific hypothalamic nuclei.

Hypothalamic “Centers”

Hypothalamic “centers” are in fact extensive networks of mutually interconnected hypothalamic cell groups. This agrees with the observation that many of the autonomic processes influenced by the hypothalamus are not controlled independently. Some factors, such as the diameter of the vessels, are of importance in several different processes, including temperature regulation and control of blood pressure. Complex behavior, such as that necessary for feeding and drinking, depends on the integrity of both medial and lateral hypothalamic areas, even though subregions play different roles. Early studies with crude methods—often destroying passing fibers

from other nuclei, in addition to parts of the hypothalamus—lead to incorrect identification of hypothalamic centers and misconceptions about the independence of hypothalamic control functions. The altered view on hypothalamic “centers” parallels the change of view during the last 30 to 40 years on the existence of functional centers of the brain in general. Rather than focusing on individual cell groups as responsible (centers) for complex functions, we now emphasize how many cell groups contribute to execution of most tasks. When we nevertheless sometimes use the term “center” for convenience, the reader should realize its limited explanatory value. For example, we have not explained how the CNS controls micturition by placing a “micturition center” in the pons.

A rough functional subdivision of the hypothalamus emerged from the pioneer lesion experiments: The **anterior parts** of the hypothalamus are particularly concerned with effects mediated by the **parasympathetic** system while the **posterior** part controls functions in which the **sympathetic** system has a dominant role.

Control of Body Temperature

The hypothalamus receives information about ambient temperature from thermoreceptors in the skin. It initiates peripheral responses to increase heat production or heat loss. Information from the skin is not sufficient, however, because the superior goal is to keep the **core temperature** constant (i.e., in deep parts of the body)—not the skin temperature that may vary considerably, as we know from everyday experience. To be able to control the core temperature, the hypothalamus must also be informed about the blood temperature in central parts of the body. As mentioned, some neurons in the **anterior hypothalamus** are temperature-sensitive. Accordingly, when the blood flowing through the anterior part of the hypothalamus is warmed, an experimental animal shows signs of increased heat loss. Cats, for example, pant and sweat through the paws. In humans, skin vasodilatation (redness and warmth) and sweating occur. If these parts of the hypothalamus are destroyed (on both sides), the animal no longer reacts to a rise in ambient temperature. The body temperature therefore rises. In contrast, when the posterior parts of the hypothalamus are destroyed, the animal no longer reacts with shivering and vasoconstriction to a fall in ambient temperature. Therefore, the body temperature drops (the vasoconstriction reduces heat loss, and the shivering increases heat production). This exemplifies how the hypothalamus coordinates autonomic (vasoconstriction) and somatic (shivering) responses. These and other observations suggest that neuronal groups in the anterior part of the hypothalamus are necessary for the coordination of the processes ensuring that the body temperature does not rise above normal levels (increasing heat loss), whereas

the posterior parts contain neurons necessary for heat conservation. Together, these parts function as a **thermostat** that tries to keep the body temperature as close as possible to a **set point** of 37°C. If the temperature drops below the set point, the “heater” is turned on—that is, measures are initiated to conserve heat (skin vasoconstriction or putting on more clothes) and, if necessary, to increase heat production (shivering or voluntary muscular activity).

In humans, abnormal rise in the body temperature, **hyperthermia**, can occur in diseases that affect the hypothalamus or by an inadvertent lesion during surgery. The rise can also occur as a side effect of certain **drugs** acting on the brain (e.g., antipsychotics). Occasionally, hyperthermia occurs during **general anesthesia**.

Fever

Fever is part of the so-called **acute-phase response** elicited by immune system activation (see below, “Effects of the Immune System on the Nervous System”). Fever arises when the set point of the hypothalamic thermostat is altered. Especially neurons in the **preoptic area** (POA) in the anterior hypothalamus appear to be crucial, as judged from microinjections of fever-producing substances. If the set point is changed to, for example, 39°C, the normal body temperature is judged to be too low and measures to conserve and produce heat are started (skin vasoconstriction, shivering, curling up, putting on more clothes). The person feels cold while the temperature is below the new set point. Fever has most likely evolved as a defensive response, improving survival from infections (although fever is sometimes harmful).

Fever can be caused (indirectly) by lipopolysaccharides (called **pyrogens**) from bacterial membranes or by simple tissue injury that leads to inflammation (e.g., fever after severe sunburn). The pyrogens or other tissue changes make leukocytes release **cytokines**. The cytokines **interleukin 1 β** (IL-1 β), and **tumor-necrotic factor** (TNF) have been detected in the tissue fluid from the anterior hypothalamus in experimental animals with fever. IL-1 β , which is the most potent fever-inducing substance, appears to act **via interleukin 6** (IL-6). Thus, IL-1 β does not induce fever in knockout mice that lack the IL-6 gene. It is not quite clear, however, how the cytokines accumulate in the hypothalamus. Blood borne cytokines can bind specifically at sites without the blood–brain barrier—such as the **subfornical organ**—where the neurons express IL-1 and IL-6 receptors. In turn, these might induce synthesis of cytokines in the hypothalamus. It is also possible that cytokines are brought through the blood–brain barrier by specific transporters. Finally, there is evidence that sensory impulses in the **vagus nerve** may induce cytokine production in the hypothalamus. Thus, injections of

IL-1 β in the peritoneal cavity (mimicking a local inflammation) do not induce fever after cutting of the vagus nerve.

Whatever the cause of their presence, cytokines in the hypothalamus induce the synthesis of **prostaglandins** (among other substances), which increase the activity of cold-sensitive neurons and reduce the activity of heat-sensitive ones. That antipyretic (reduce fever) drugs, such as **aspirin**, inhibit the synthesis of prostaglandins supports a crucial role of prostaglandins in fever production.

Local adaptations in the hypothalamus ensure that the fever does not reach dangerous levels (**antipyresis**). Cytokines activate neurons in the paraventricular nucleus producing CRH and vasopressin. As discussed, CRH release leads to increased blood levels of cortisol. This would reduce the peripheral production of cytokines and, by that, reduce the fever. Further, fibers from the paraventricular nucleus release vasopressin in the **septal nuclei**. Experimental infusion of vasopressin in the septal nuclei suppresses almost completely the febrile response to intravenous injection of pyrogens. It appears most likely that the septal nuclei act back via septohypothalamic fibers to cancel the effect of cytokines on the thermostat.

The Hypothalamus, Sleep, and Hypocretins

Sleep disturbances often accompany lesions of the hypothalamus in experimental animals and in humans, sometimes as an abnormal amount of sleep and sometimes as insomnia or disturbed sleep rhythm. Animal experiments in the 1940s demonstrated that lesions in the **preoptic area** in the anterior hypothalamus caused insomnia, and later studies identified single neurons with their maximal activity during sleep in the same region. Later studies have identified two neuronal groups in the posterior hypothalamus of importance for wakefulness.⁴

The first is the **tuberomammillary nucleus** that contains histaminergic neurons with widespread axonal ramifications. Histamine binds to histamine receptors (H₁–H₃) in many brain regions. With regard to the histaminergic effect on wakefulness, the projections to the cerebral cortex, the locus coeruleus, and the raphe nuclei are probably most important (histamine has several additional actions, for example on learning and memory). GABAergic neurons in the preoptic area project to the (histaminergic) tuberomammillary nucleus, and may therefore contribute to sleep by inhibiting wake-active neurons.

The second group consists of diffusely spread neurons (about 7000 in humans) in the posterior and lateral

4 The epidemic in the early twentieth century of a presumed viral disease causing (among other complications) extreme somnolence—**encephalitis lethargica**—pointed to the posterior hypothalamus as important for waking.

hypothalamus that release two varieties of the neuropeptide **hypocretin (orexin)**. The hypocretins bind to specific receptors and exert a wake-promoting effect when infused in animals. Further support for the importance of hypocretins came from the observation that patients with **narcolepsy** have reduced levels of hypocretins in the cerebrospinal fluid and loss of hypocretinergic neurons (see Chapter 26, under “Narcolepsy”). The hypocretins exert effects several places but projections to the tuberomammillary nucleus may be of particular importance. Thus, hypocretins increase release of histamine and wakefulness but not in knockout mice lacking expression of the H_1 receptor. In addition, hypocretinergic fibers pass to the locus coeruleus and the raphe nuclei, both related to control of arousal (see Chapter 26, under “Multiple Pathways and Transmitters Are Responsible for Cortical Activation”). Interestingly, the **suprachiasmatic nucleus**, which is responsible for the control of circadian rhythms, projects to regions with hypocretin-expressing neurons.⁵

The hypocretins are yet another example that each of the brain’s many signal substances influences several different processes. Indeed, the hypocretins were discovered because of their ability to induce feeding behavior in experimental animals. Later, their importance for wakefulness was revealed, and now they are implicated in a number of other functions as well.

It should be emphasized that many more neuronal groups and neurotransmitters than those mentioned here are implicated in the control of wakefulness and sleep (see Chapter 26).

The Hypothalamus and Circadian Rhythms

The hypothalamus plays a role as a pacemaker for several functions showing a cyclic, diurnal variation. Such **circadian rhythms** are governed from the small **suprachiasmatic nucleus (SCN)**, situated just above the optic chiasm. γ -Aminobutyric acid (**GABA**) is the transmitter for most of the neurons in the SCN, colocalized with **neuropeptides** (either vasopressin or VIP). **Lesions** of the SCN disturb (but do not necessarily abolish) the cyclic variations of bodily functions. Therefore, SCN is not alone in determining the circadian rhythms, even if it plays the leading part.

Several physiologic parameters—such as hormonal levels, blood pressure, body temperature, wakefulness, and sleep—are subject to circadian alterations. The same holds for several mental functions, such as reaction time and mood. The biologic significance of the cyclic variations of bodily functions is presumably to adapt the level of activity to the most stable and predictable variations

in the environment. For example, in humans it seems appropriate that most bodily functions are at their lowest level during the night, whereas animals hunting at night would need a different activity profile. Persons staying in environments without any information about the time of the day develop a rhythm with cycles slightly longer than 24 hours. The capacity to produce circadian rhythms is inborn, and cyclic activity appears in the hypothalamus (rat) late in intrauterine life. The SCN functions as a **pacemaker** or **biologic clock**, producing a certain rhythm even in the absence of external inputs. Nevertheless, the exact rhythm—that is, at what time of the day the highest and lowest values occur—is modulated by sensory information. **Light stimuli** from the retina (Fig. 30.4), varying with length of the light/dark cycle, appear to be particularly important for the set of such circadian rhythms. Other factors also seem to contribute, however, such as activity level.

The **retinohypothalamic** fibers, which arise from retinal ganglion cells, end in the SCN. They inform only about the amount of light but do not provide specific information about patterns, movements, and so forth. In addition, the SCN receives afferents from a small part of the lateral geniculate nucleus and from the **raphe nuclei** (serotonin).

The SCN has **efferent** connections with several other parts of the hypothalamus, which enable it to synchronize the various functions mentioned above (and other functions as well). This concerns, for example, the **CRH**-containing neurons in the **paraventricular nucleus** that control the blood level of corticosteroids. Other connections reach the **thalamus** and cell groups in the **basal forebrain** (see Chapter 31, under “The Basal Forebrain”). Such connections might influence diurnal variations in motivation and memory. With regard to hypothalamic control of the **sleep–wake cycle**, newly discovered (polysynaptic) connections from the SCN to the **locus coeruleus** (norepinephrine) are of particular interest (see “The Hypothalamus, Sleep, and Hypocretins”). A special polysynaptic efferent pathway from the SCN to the **pineal gland** controls secretion of the hormone **melatonin**. Melatonin binds to high-affinity receptors in parts of the hypothalamus, among them the SCN, and modulates circadian rhythms. Light can influence the SCN pacemaker via variations in the melatonin level (in addition to by the retinohypothalamic fibers). Melatonin is secreted in response to low levels of light and can therefore be said to **signal darkness**.

Melatonin

The **pineal body** or gland (see Chapter 6, under “The Epithalamus and the Pineal Body”) produces a number of neuroactive substances, among them several **neuropeptides** and large amounts of **serotonin**. The hormone **melatonin** has attracted most interest, however.

5 Hypocretinergic connections to dopaminergic neurons in the mesencephalon (in the substantia nigra and the VTA) may explain why hypocretins also influence motivation and “reward behavior.”

Melatonin (*N*-acetylmethoxytryptamine) is synthesized from serotonin in two enzymatic steps. No nerve fibers leave the pineal body, which therefore is assumed to function only as an endocrine gland. The only known **afferent** nerve fibers arise in the **upper cervical sympathetic ganglion** (postganglionic fibers). Ambient **light** influences the secretion of melatonin; increased light (longer days) inhibits secretion. In humans, as in experimental animals, the level of melatonin varies with the light–darkness cycle, with the highest level occurring at night. The influence of light is mediated through a tortuous route: as mentioned, signals from the retina reach the SCN directly; neurons there forward signals directly or indirectly to the preganglionic sympathetic neurons in the upper thoracic cord, and their axons reach the upper cervical ganglion by means of the sympathetic trunk (see Fig. 28.7).

Melatonin acts by binding to G protein–coupled receptors, which occur with the highest concentrations in the SCN, the paraventricular nucleus, and parts of the anterior pituitary. The dominating effect of receptor activation seems to be reduced synthesis of cyclic AMP. Melatonin appears to inhibit the secretion of **gonadotropic hormones** (the effect seems to depend on the time of the day, however). In experimental animals, melatonin also have a number of other actions—for example, influencing **sleep**, **arousal**, body **temperature**, and blood level of **cortisol**. These effects are believed to be mediated mainly by the SCN.

Intake of melatonin in humans alters the phase of such rhythms, probably by setting the “circadian clock” in the SCN. Indeed, melatonin is now widely used to alleviate the unpleasant psychological and physiological effects of **jet lag** (even though its effect in this respect has not yet been convincingly documented). It has also been used to normalize sleep–wake patterns in blind people. The production of melatonin declines with age, and this has been taken to explain sleep disturbances in elderly people. Because melatonin counteracts the accumulation of cytotoxic free radicals, it has been suggested as a drug to halt the progression of degenerative brain diseases (Parkinson’s disease, Alzheimer’s disease, and others). The many reported beneficial effects of melatonin in humans lack documentation, however. It is fair to say that its normal role in humans is not yet clear, especially because most of what we know comes from experiments on rodents.

Control of Digestion and Feeding

The hypothalamus controls various metabolic processes. For example, lesions of certain parts can produce abnormal fat deposition, both in experimental animals and in humans. The gastrointestinal tract is influenced by the hypothalamus. Lesions can produce ulcers of the mucous membranes and bleeding in the stomach and the small

intestine. Stimulation of the anterior hypothalamus, in particular, can elicit increased secretion of gastric juice and strengthening of peristaltic movements.

Not only the digestive processes but also **feeding behavior** is controlled from the hypothalamus. Early stimulation and lesion experiments indicated that the **lateral hypothalamic area** could induce increased eating (or behavior directed at acquiring food), whereas the **medial parts** (especially the ventromedial nucleus) reduce eating (induce satiety). Further studies have shown that, although essentially correct, this scheme is too simple. A complex network within the hypothalamus and between the hypothalamus and other parts of the brain controls food intake and body weight.

The inputs to the hypothalamic “feeding centers” are multifarious, as witnessed by data from animal experiments and by everyday experiences of the numerous factors that influence human feeding behavior. It is nevertheless remarkable how most people maintain their body weight over many years, although even the smallest daily surplus of food would cause a steady weight gain (a surplus of 10 calories a day would increase body weight by about 1 kg in a year). The hypothalamic centers governing food intake are thought to operate as a **homeostat** (a system maintaining a steady state by internal processes.) We also use the term **lipostat**, because the control of body fat is so central to maintaining constant body weight. **Feedback loops** ensure that the food intake oscillates around a set point. Reduced consumption automatically follows overeating.

The protein **leptin**, produced in fat cells, appears to be the single most important factor ensuring negative feedback from adipose tissue to the hypothalamus (**insulin** is another factor). The function of leptin is probably to inform about whether the energy reserves (fat) are sufficient. The so-called **ob gene** (ob for obesity) codes for leptin, while the **db gene** codes for the leptin receptor. Mice lacking either the ob or the db gene develop extreme obesity due to increased food intake and reduced metabolism (in addition, they develop diabetes and increased cortisol levels). Although leptin binds several places in the hypothalamus, a major effect is to inhibit the release of **neuropeptide Y** (NPY).⁶ NPY reduces food intake, as shown by injections in certain parts of the hypothalamus in experimental animals.

Loss of Feeding Control

Of course, psychological factors—or, occasionally, hypothalamic disease—may disturb this finely tuned

⁶ NPY is involved in other hypothalamic tasks, such as control of circadian rhythms, and secretion of releasing hormones. It is often colocalized with catecholamines, GABA, or somatostatin. Finally, NPY is also found in the amygdala where it is involved in stress reactions (having an anxiolytic effect). Obviously, each of the numerous transmitters present in the hypothalamus may be expected to participate in quite diverse functions.

control mechanism. Studies of **obese people** indicate that they have normal leptin production.⁷ A hypothalamic leptin resistance has been proposed, however, to account for the development of obesity (analogous to insulin resistance in some cases of diabetes). If so, the question remains how the resistance arises: is it due to a genetic abnormality, or is the lipostat altered by psychological and other factors? Only in about 35% of overweight people is there evidence of a genetic disposition.

Focusing on single actors like leptin, NPY, and hypothalamic nuclei should not make us forget that the hypothalamus does not operate in isolation. The many connections between the hypothalamus and the cerebral cortex, the amygdala, and other limbic structures tell us that the hypothalamus may be “overruled” by the higher parts of the brain engaged in cognitive and emotional processing. Thus, while the hypothalamus undoubtedly is the organizer for motivated behavior related to digestion and feeding, other regions than the hypothalamus determine whether such behavior occurs. Although we do not fully understand the causes of eating disorders such as **anorexia** and **bulimia**, it is unlikely that the primary cause is to be found in the hypothalamus.⁸

The Arcuate Nucleus, Neuropeptides, and Energy Metabolism

Several transmitters (especially many neuropeptides) are involved in the hypothalamic control of feeding behavior, although much remains before their mutual roles are clarified. Special interest attaches to **NPY**, which is found in high concentration in nerve terminals in the **paraventricular nucleus**. Most of the NPY-containing terminals come from the **arcuate nucleus**. Two groups of neurons in the arcuate nucleus have opposite effects: **NPY neurons** increase food intake while **POMC neurons** (pro-opiomelanocortin) reduce it. For example, injection of NPY near the paraventricular nucleus increases food intake in experimental animals. Further, it decreases the metabolism in **brown fat** and increases fat storage in ordinary (white) fat tissue. This leads to a larger weight gain than expected from the food intake. The NPY-containing connections to the paraventricular nucleus might therefore regulate the energy balance of the body by controlling both appetite and metabolism. How NPY release in the hypothalamus produces this is not known, although some effects may be mediated by

projections from the paraventricular nucleus to sympathetic preganglionic cell groups. In addition to **leptin**, the hormone **insulin** act as one of several **feedback signals** to the arcuate nucleus from the periphery, informing about the energy status of the body. Thus, hypothalamic injection of insulin reduces the food intake in normal rats (but not in rats with genetically determined obesity).

Several other factors contribute in the regulation of food intake. For example, **glucose-sensitive** hypothalamic neurons are likely to be involved in the control of appetite (for instance, the feeling of hunger when the blood glucose is low). The hormone **ghrelin**—produced in the stomach—may also contribute to regulation of hunger and food intake by acting on neurons in the arcuate nucleus. Indeed, ghrelin activates NPY neurons and inhibit POMC neurons, as one might expect from a signal informing about empty energy stores. (As other neuropeptides, ghrelin has diverse actions; it increases, for example, plasticity in the hippocampus.) Gut hormones, such as **cholecystikinin**, do not cross the blood-brain barrier, but can act on neurons in the **solitary nucleus** via the area postrema (lacking a blood-brain barrier). The **vagus nerve** provides the solitary nucleus with information about the filling of the stomach and probably about the levels of glucose and lipids in the liver. From the solitary nucleus, signals travel to the hypothalamus and notably the arcuate nucleus.

Hypothalamus, Sexual Functions, and Sex Differences

The efferent connections of the hypothalamus also enable it to influence sexual behavior—that is, to start and coordinate the autonomic, endocrine, and somatic motor components. This is verified by many animal experiments. For example, injection of the female sex hormone **estradiol** in the **ventromedial hypothalamic nucleus** in castrated male rats elicits copulatory behavior. Lesions of the ventromedial hypothalamic nucleus in female rats reduce their sexual activity. We have little precise knowledge of how the hypothalamus controls sexual functions, however. Neither do we understand fully the structural basis in the hypothalamus for behavioral sex differences, in spite of numerous findings of sex differences among hypothalamic nuclei.

In rodents a nucleus in the anterior hypothalamus—the **sexually dimorphic nucleus of the preoptic area**, **SDN-PO**—is three to eight times larger in males than in females. While several studies in humans have described sex differences in the corresponding region, it is remarkable that they do not agree on which one among three small nuclei that show such sexual dimorphism (neither is there agreement with regard to hypothalamic differences between heterosexual and homosexual men).

Several hypothalamic pathways are likely to be sexually dimorphic. One among many relevant pathways passes from the **paraventricular nucleus** directly to the

7 A small minority of persons with extreme obesity have mutations of the *ob* or *db* genes. Such persons develop overweight very early. They also suffer from low levels of growth hormone and thyroid hormone, and they lack normal pubertal development.

8 A review of 54 patients with eating disorders occurring in conjunction with a brain lesion showed that the majority had lesions in the right fronto-temporal region. Lesions of the hypothalamus were associated with reduced or increased appetite but not with more complex eating disorders (Uher and Treasure 2005).

spinal motoneurons that innervate the **bulbospongiosus** (bulbocavernosus) **muscle**. This muscle, which is important during ejaculation, is large in male rats but tiny in females. Correspondingly, the descending pathway mentioned above is absent in female rats. The motoneuron group in the sacral cord that innervates the bulbospongiosus muscle is three times larger in male than in female rats. The early presence of testosterone prevents programmed cell death among the motoneurons, thereby establishing the sex difference.

As discussed, the hypothalamus controls sexual functions also by its actions on the **pituitary** and its secretion of **gonadotropic hormones**. The sex hormones act on the hypothalamus and other parts of the brain and can therefore influence sexual behavior. As shown in rats and monkeys, transient exposure to small amounts of sex hormones during early development determines later sexual identity and behavior. Structural differences that depend on circulating sex hormones have been described in the ventrolateral hypothalamus and in the amygdala of adult animals. The presence of **testosterone** seems to be crucial: it initiates structural sex difference in the brain, promotes masculine behavior, and suppresses feminine behavior. (See Chapter 34, under “Cognitive Sex Differences and Lateralization” and “Psychological Sex Differences and Biology”.)

THE HYPOTHALAMUS AND THE IMMUNE SYSTEM

Effects of the Nervous System on the Immune System

The relationship between the nervous system and the immune system has attracted increasing attention recently. This topic is important in several respects—for example, regarding the interactions between mental and bodily processes. Besides many anecdotal reports of somatic diseases that start at times of mental stress and imbalance, there is growing experimental evidence of hypothalamic influence on the immune system. This concerns, for example, the activity of **natural-killer cells** (NK cells) and **lymphocytes**. Further, antigen stimulation appears to increase activity among certain hypothalamic cell groups. This also suggests that the hypothalamus is in some way involved in the defense against infections. Other studies show altered T-lymphocyte numbers in deeply **depressed patients**. How these interactions come about is not fully understood, and some findings in this field are contradictory.

The nervous system can influence the immune system in two ways: one by the **endocrine system**, the other by **autonomic innervation** of the lymphoid organs (Fig. 30.8). ACTH and secretion of glucocorticoids from the adrenal cortex primarily mediate the endocrine influence. The inhibiting effects of the glucocorticoids on inflammation and resistance to infections are well known. Long-lasting

stress with increased ACTH secretion might cause increased vulnerability to infections, allergy, autoimmunity, and other diseases.

The bone marrow, thymus, spleen, lymph nodes, and gut-associated lymphoid tissue all receive innervation by **postganglionic sympathetic fibers**. The innervation density is highest in T-lymphocyte areas. Lymphocytes express **β -adrenergic receptors**, and varicosities of sympathetic fibers are found close to lymphocytes. Chemical denervation of lymphoid organs causes reduced T-lymphocyte activity, among other effects (B lymphocytes appear also to be affected). Modulation of immune responses by efferent **parasympathetic fibers** in the **vagus nerve** was recently demonstrated. This seems to happen primarily by acetylcholine binding to peripheral nicotinic receptors (in contrast to other parasympathetic effects that are mediated by muscarinic receptors). One effect of vagus stimulation is inhibition of **cytokine** release from immune cells.

In **conclusion**, the autonomic system is able to modulate the properties of the immune system by endocrine and autonomic pathways, although we have little precise knowledge of the functional significance of such modulation.

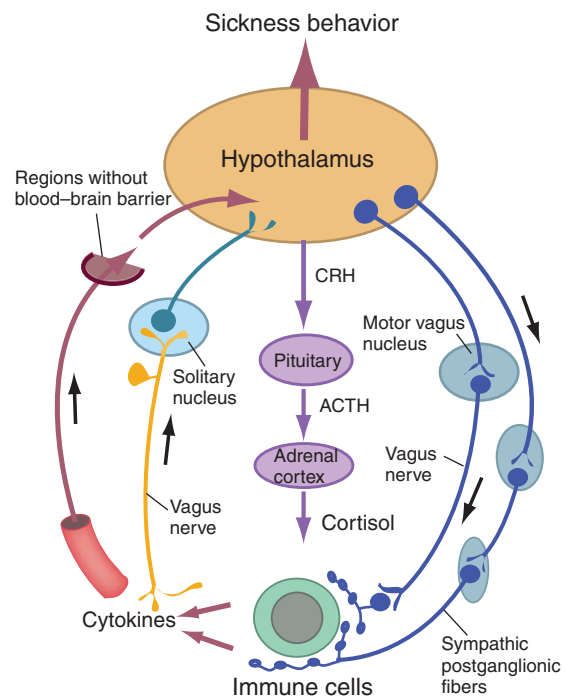


FIGURE 30.8 Interactions between the nervous system and the immune system. Schematic. Cytokines released during immune responses can influence hypothalamic neurons in two ways (left part of the figure). The middle part shows how the hypothalamus can influence the immune system via the pituitary and the adrenal gland. The right part shows how sympathetic and parasympathetic nerves can mediate hypothalamic effects on the cells of the immune system.

Effects of the Immune System on the Nervous System

We deal with immune reactions in the CNS in Chapter 1, as part of defense mechanisms (under “The Reaction of Nervous Tissue to Injury and Inflammation”). However, the immune system can also influence normal nervous processes (Fig. 30.8). One example is that **cytokines** from leukocytes cause fever as one component of the acute-phase response, as discussed above. Further, it seems highly likely that typical symptoms in infectious diseases like drowsiness, loss of appetite (anorexia), social withdrawal, listlessness, and aversion to certain tastes are caused by action of cytokines in the brain. As mentioned, effects are believed to be mediated by binding of cytokines to neurons in regions lacking a blood–brain barrier and perhaps by direct entrance into the hypothalamus. In addition, the **vagus nerve** (and perhaps other nerves) mediates similar effects on the hypothalamus from peripheral tissues with active immune responses (cutting the vagus nerve reduces sickness behavior in animals with abdominal infections). Lymphatic dendritic cells and macrophages release **IL-1 β** that binds to peripheral branches of the vagus. Afferent vagus fibers end in the solitary tract, most likely releasing glutamate. The solitary nucleus projects to several parts of the hypothalamus that may mediate the sickness behavior. In addition, the ACTH secretion from the **pituitary** (and thus the blood cortisol level) can be influenced by signals in the vagus nerve. Thus, fibers from the solitary nucleus end in the paraventricular nucleus and modulate the release of corticotropin-releasing hormone (CRH). Signals in the vagus nerve to the brain stem and the hypothalamus may also participate when **hyperalgesia** develops in patients with infections.

THE HYPOTHALAMUS AND MENTAL FUNCTIONS

Psychosomatic Interrelations

The relationship between the hypothalamus and the pituitary discussed above helps explain why diseases affecting the hypothalamus can produce alterations of hormonal secretion from the pituitary itself, the thyroid, the adrenal cortex, and the gonads. However, it also helps in explaining the relationship between mental and bodily processes—called **psychosomatic interrelations**. Thus, the hypothalamus receives afferents from many parts of the brain, among them the cerebral cortex and the limbic structures, which are closely related to what we term “mental” or “psychic” functions. In short, our mental state—by acting on the hypothalamus—can produce alterations of endocrine organs, of organs innervated by the autonomic system, and of skeletal muscles. Not all such effects are mediated via the hypothalamus, however. Thus, fibers from the cerebral cortex and limbic structures access the autonomic and somatic cell

groups of the reticular formation (see Chapter 31, under “The Amygdala and Emotions”).

We describe in this section just a few examples of psychosomatic interrelations. Women may lose their **menstruation** for some time after psychic stress (loss of a close person, dissatisfaction, depression, and so forth). The mechanism is apparently reduced secretion of gonadotropic hormones from the pituitary, caused by reduced production of the relevant releasing hormone. When subjected to bodily stress (such as infections, trauma, intoxications, and major surgery), the organism responds by, among other things, increasing the secretion of corticosteroids. **Mental stress** can produce the same response by causing increased secretion of releasing hormones from the hypothalamus, which, in turn, increases the secretion of ACTH from the anterior pituitary (see Fig. 31.6). In experimental animals, transection of the stalk of the pituitary prevents the hormonal response to mental stress. Our mental state influences functions controlled by the posterior pituitary (such as urinary volume). A particularly striking example of a psychosomatic interrelation is that of a woman **breast-feeding** her baby, who can cause the milk to trickle from the nipples just by thinking of the child. The inner image of the child in some way influences the hypothalamus, with an increase of oxytocin secretion as the result. This reaches the mammary glands and makes the smooth muscles contract (milk ejection reflex).

Expectation and Health

The relation between **expectation** and disease has attracted much interest. For example, several studies address how **optimism** as a **personality trait** influences psychological adaptation, handling of stress (like major surgery), and self-reported health. Altogether, there are clear positive correlations; that is, an optimistic attitude is associated with better self-reported health, fewer psychiatric symptoms, lower blood pressure, and so forth. While a causal relationship remains to be established, there is solid evidence that our mental state (via the hypothalamus) influences several physiological processes in the body. Further, studies of the **placebo** and **nocebo** phenomena strongly support the importance of expectations for handling of stressful situations (see Chapter 15).

There are exceptions, however, from a positive correlation between optimism and health (or indirect measures, like immunity). Thus, some studies find a negative correlation between optimism and immunity when stressors are complex, long lasting, and unpredictable (Segerstrom 2005). Perhaps, in such situations inevitable frustrations and disappointments might be less adaptive than resignation (as would be chosen by a less optimistic person).

Another question is why people differ so much with regard to how they cope with stressful situations.

Presumably, genetically determined vulnerability factors—both psychological and molecular—vary considerably among people. If vulnerable persons are exposed to many stressful life events in early childhood, they seem at risk to express dysfunctional responses to stress in later life.

Other aspects of the interaction between the mind and the body are discussed in Chapter 31, under “Amygdala and Emotions.”

Actions of Hormones on Psychic Functions

Bodily processes may influence psychic ones, as discussed earlier in this chapter. Such interactions are especially clear with regard to the hormonal effects on the brain. For example, increased production of **thyroid hormones** changes the mental state toward excitement, increased initiative, and lively associations, whereas reduced production leads to apathy, fatigue, and increased need for sleep. Many patients with a thyroid disease have been misdiagnosed as suffering from a mental disorder. Treatment with **corticosteroids** (as used, e.g., with certain kinds of cancer) often elevates the mood (euphoria), making the patient appear unduly cheerful and unconcerned. Effects of female sex hormones on the brain probably cause the changes of mood often associated with the menstrual cycle in women (premenstrual syndrome). Thus, a high density of receptors for **sex hormones** has been demonstrated in various parts of the brain. This concerns, in particular, certain cell groups in the hypothalamus and the amygdala; neurons in other parts, such as the cerebral cortex, can also bind sex hormones. It seems likely, therefore, that circulating hormones influence the excitability of many neuronal groups (steroid hormones cross the blood–brain barrier easily). In cats, implantation of female sex hormones in the posterior part of the hypothalamus produces heat behavior. Many other experiments give further evidence that circulating hormones, presumably by their actions on the brain, can be influenced animal behavior. The male sex hormone **testosterone** can produce male sexual identification and behavior when given to monkeys at an early stage of their development, regardless of the sex of the monkey. Thus, a female monkey may later behave like a male if given a small amount of testosterone for a period shortly after birth.

Psychosomatic Disorders

A **psychosomatic disorder** (syndrome) is usually defined a condition in which psychological processes play a substantial role for the production of the somatic symptoms. The term **somatoform disorder** often implies absence of physical signs that can explain the somatic symptoms. However, the definitions of these terms differ among authors, presumably because the terms embrace a

heterogeneous and poorly understood group of disorders. There is probably no sharp border between conditions in which purely psychic factors cause or contribute to somatic disease (e.g., coronary disease) and those in which clear-cut signs of somatic disease cannot be found (e.g., fibromyalgia). As discussed earlier and in relation to the placebo phenomenon (Chapter 15), all disease processes are more or less influenced by the mental attitude and expectations of the patient, while any somatic disease will lead to psychological reactions. In some conditions—such as irritable bowel syndrome, fibromyalgia, chronic fatigue syndrome, dyspepsia without an ulcer, and noncardiac chest pain—the influences of psychic factors are thought to be important and sometimes decisive, although we cannot explain the causative relationships. In other diseases—like gastric ulcers, coronary disease, asthma, and diabetes—there is good evidence that psychic stress can increase the **vulnerability** (i.e., the risk of developing the disease), without being a main causative factor. Further, in such diseases, which are typically due to the interaction of many risk factors, individuals probably vary as to the importance of each risk factor (see Chapter 31, under “Amygdala, Anxiety, and Transmitters”).

Stress

Used in the present context, stress can be physical or psychological. A common **definition** is something perceived by the individual as a threat to the homeostasis of the organism. Except for cases of extreme physical stress, the crucial point in stress is how the challenge or extra demand is perceived, not the stress by itself. Although the word “stress” in common language denotes something that is bad for the health, extra demands on our physical or mental performance are inevitable and usually not harmful. On the contrary, without challenges to the status quo, there would be no development, no learning, and no improvement of adaptive behavior.

The coordinated sum of endocrine, autonomic, and somatic responses to stress is termed a **stress reaction**. The function of the stress reaction is to maintain homeostasis in a wide sense, and the stressful event itself or the expectation of it can initiate the reaction. The hypothalamus obviously plays a central role in our ability to cope with stressful events; that is, the stress reactions are expressed through the hypothalamic influences on the endocrine, autonomic, and somatic effector systems.

Stress reactions are commonly assessed by measuring the blood level of **epinephrine** (activation of the sympathetic system) and of **cortisol** (activation of the hypothalamic–pituitary–adrenal axis). The bodily response to stress is of course much wider than just these hormonal changes, however, as discussed earlier in this chapter. Indeed, the stress reaction should not be seen as an isolated phenomenon but, rather, as a part of

bodily expressions of **arousal** (arousal was discussed in Chapter 26). In this perspective, stress is a stimulus that increases arousal (with changes of the EEG, attention, muscle tone, respiration, selective autonomic activation, and activation of the CRH–ACTH–glucocorticoid reaction chain).

The purposefulness of the stress reaction is most obvious with physical stress, such as situations demanding extreme endurance or force, or with injuries causing rapid blood loss. However, stress reactions are also adaptive and beneficial before and during extra mental demands such as examinations, performances, or novel situations. Overall, stress reactions are adaptive when they improve our ability to cope with the extra demands. Lack of corticosteroid hormones due to disease of the adrenals (Addison's disease), for example, greatly reduces the ability to tackle even minor challenges to homeostasis. The stress reaction becomes potentially disease provoking only when it by far outlasts the actual stressful situation (with, among other things, a persistently elevated level of “stress hormones”).

Stress and Disease

When talking of stress causing disease, we usually mean psychic stress, and such that lasts for months or years. Whereas serious physical stress produces a stress reaction that is uniform among individuals, the individual variations are larger to psychic stress. Thus, the decisive factor is the subjective perception and interpretation of the situation. This depends on many conditions, such as prior experiences and how they were tackled (especially during early childhood). Among psychological stress factors (stressors), the feeling of **loss of control** is probably the most important. Closely related to this is **unpredictability** of a situation. Animal experiments show that stress reactions to painful stimuli are reduced if the stimulus is preceded by a warning signal. Presumably, this is because not knowing when the stimulus occurs requires constant arousal, whereas the warning enables the animal to relax (reduce arousal) between the stressful events. The importance of the feeling of control is exemplified by the fact that humans with chronic pain reduce their consumption of analgesic drugs if they can decide themselves when to take the drug, rather than having to ask another person. Another important factor determining the level of psychic stress is the possibility of obtaining an **outlet for frustration**—for example, with physical exercise. Finally, animal experiments support that **social attachments** reduce psychic stress.

The Hypothalamus and Emotions

The hypothalamus is among the parts of the brain most directly involved in the expression of emotions or **emotional reactions** (in an experimental context, emotional

reactions can be defined as behavior in response to stimuli producing sensations with an emotional coloring). This agrees well with the fact that the hypothalamus functions as a superior center for control of autonomic processes. Emotions are expressed, as we know from everyday experience, to a large extent through changes of the functions of autonomically innervated organs, such as palpitations, dryness of the mouth, fainting, blushing, paling, alterations of the digestive tract, sweating, frequent micturition, and so forth. In addition, automatic movements, such as rapid, superficial breathing, facial expressions, and postures, witness emotions. Such movements—organized by brain stem premotor networks—are probably influenced from the hypothalamus (see the discussion of emotional smile and facial palsy in Chapter 27, under “Facial Expressions of Emotion Do Not Depend on the Integrity of the Pyramidal Tract”).

The role of the hypothalamus in emotional reactions has been studied in cats and dogs in which the whole cerebral cortex, the basal ganglia, and large parts of the thalamus have been removed. So-called **sham rage** can be provoked in such animals. Because only the hypothalamus is connected with the brain stem in such animals, their expression of rage must depend on the hypothalamus. Such animals react much like normal animals to painful stimuli, with biting, scratching, snarling, and increased ventilation. Because the whole cortex is removed, it is unlikely that true emotions are experienced, however. It seems reasonable to conclude that the hypothalamus contains cell groups that coordinate and put into action the behavior expressing the rage. In contrast to normal animals, the rage of such “hypothalamic” animals is not directed toward anything in particular; they lack the ability to know the nature and the location of the stimulus provoking the pain (as one might expect in an animal lacking the cerebral cortex and most of the thalamus). Further, the expression of the rage dies out very quickly after the stimulus is over, whereas the reactions continue for a while in normal animals (as in humans). This observation suggests that other parts of the brain normally act on the hypothalamus to produce emotional reactions, as we discuss in Chapter 31, under “The Amygdala and Emotions.”

The preceding account indicates that to regard the hypothalamus as the locus of the emotions would be an impermissible oversimplification. Rather, the hypothalamus is a superior center for the **coordination of emotional reactions**. Observations in humans during brain surgery with local anesthesia support this conclusion. Pressure on or traction of the hypothalamic region can elicit reactions of panic, crying, laughter, or profuse talking. The patients sometimes report a change of mood, such as depression or euphoria. It thus appears that the activity of the hypothalamus is significant not only for emotional reactions but also for the emotions

themselves. Conceivably, the emotions are evoked by feedback connections from the hypothalamus to the limbic structures and the cerebral cortex—structures that are necessary for the experience of emotions (as distinct from emotional reactions). The higher regions presumably interpret the hypothalamic activity as evidence of external or internal stimuli that normally evoke strong emotions.

Emotions and Emotional Reactions

When discussing the relations between the hypothalamus and emotions, one must distinguish the emotions themselves from the emotional reactions—that is, the behavior expressing our emotions. We can experience the feelings or emotions only subjectively. Of course, we may learn that certain external stimuli or situations usually produce certain emotions in other people, but such correlations can only be tentative because so many psychological individual variations play a role. We cannot obtain information from animals about their emotions,

but emotional reactions can be directly observed and are often more reliable in animals than in humans. Factors such as upbringing, social conventions, and conscious considerations determine to a large extent the emotional reactions in humans. That emotions in animals can only be inferred indirectly from their behavior explains why it is not quite clear how many basic emotions animals have. Commonly, however, only three basic emotions are identified (in cats, dogs, and monkeys): **rage**, **fear**, and **pleasure** (love). Even though the emotions of animals certainly are less schematic than this, there is no doubt that the emotions of humans have much more variation and nuances. This should be kept in mind when drawing conclusions with regard to emotions and psychosomatic interactions in humans on the basis of animal experiments. The anthropologist Paul Ekman (1984) identifies seven basic emotions in humans, based on their relation to culture-independent facial expressions: **happiness**, **sadness**, **anger**, **fear**, **disgust**, **surprise**, and **contempt**.

VII

LIMBIC STRUCTURES

THIS part of the book deals with parts of the fore-brain that are closely associated with the cerebral cortex with regard to development and connections. The cerebral cortex is divided into two parts (without definite delimitations) on the basis of phylogenetic development: the **neocortex**, which is the most recent part and comprises most of the cortex in higher mammals, and the **allocortex**, which is the oldest part. The neocortex is treated in Chapters 33 and 34, while this chapter deals with the allocortex and closely related subcortical nuclei. The nomenclature used for the oldest part of the cortex varies, but usually the term allocortex is used for the parts of the cortical mantle with a simple, often only three-layered structure instead of the six layers that are typical of the neocortex. In reptiles, the allocortex constitutes what little cortex there is. The allocortex receives afferents from adjacent subcortical

nuclei (in contrast to the neocortex that is closely connected with the thalamus). Together, these subcortical nuclei and the allocortex are commonly said to comprise the so-called **limbic system** (gyrus limbicus is another name for the cingulate gyrus). The cell groups comprising the “limbic system” coincide in part with what was formerly called the **rhinencephalon**, although this term, strictly speaking, comprises only the parts of the brain that receive olfactory fibers. There are wide variations among authors, however, with regard to which neuronal groups are included in the “limbic system,” indicating that the term lacks reasonable precision. Indeed, the term “system” becomes misleading when used to lump—rather arbitrarily—neuronal groups with major functional differences. In this book we therefore use the neutral term **limbic structures** to avoid giving a misleading impression of functional unity.

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31 | The Amygdala, the Basal Forebrain, and Emotions

OVERVIEW

This chapter deals primarily with structures that are of special importance for emotions, motivation, and affective behavior. We include in the discussion the **cingulate gyrus**, the **septal nuclei**, the **amygdala**, and neuronal groups in the **basal forebrain**. In addition, we discuss neocortical regions involved in emotional processing, notably parts of the **prefrontal cortex** and the **insula**.

The **amygdala** is located in the tip of the temporal lobe and consists of several distinct nuclei with different connections. It plays an important role in social behavior and in emotional learning and memory. In particular, a central task of the amygdala is the establishment of links between stimuli and their emotional value. The amygdala has connections with brain stem nuclei (e.g., the periaqueductal gray [PAG]) and influences somatic and autonomic responses to strong emotions (e.g., mediating conditioned fear responses). It also can influence the endocrine system. Amygdaloid connections with the cingulate gyrus, the prefrontal cortex, and the hippocampus mediate influences on cognitive processes, affective behavior, and memory.

Wide areas of the cerebral cortex influence the activity of the **autonomic nervous system**. This influence is exerted mainly via the amygdala, the hippocampal formation, and the septal nuclei, which in turn, influence the hypothalamus and brain stem nuclei. In addition, there are some direct connections from the **insula** and the **orbitofrontal cortex** to the hypothalamus. Further, neocortical areas in the frontal and temporal lobes project to the amygdala and can therefore influence the hypothalamus indirectly.

Extensive cortical areas show altered activity in relation to **emotions**. On the other hand, no area is concerned solely with emotional processing. The **cingulate gyrus** is, among other tasks, important for the choice of behavior in response to conflicting stimuli. The orbitofrontal and medial parts of the frontal lobes are also closely linked with processing of emotions and their integration with cognitive processes.

The term the **basal forebrain** (*substantia innominata*) is used with reference to diffuse neuronal populations below and medial to the well-defined basal ganglia. It contains

many **cholinergic neurons**, and is involved in regulation of in **attention, motivation, and memory**.

WHAT IS THE “LIMBIC SYSTEM”?

The structures included in the term the “limbic system” are often regarded as the substrate of emotions and subconscious processes, in contrast with the cognitive, conscious processes that are assumed to be located in the neocortex. Sometimes terms such as “the emotional brain” or even “the reptile brain” have been used to describe these regions. Each of the cell groups of the “limbic system” participates in various functional domains, however, and confining them to a single conceptual framework only hampers understanding. To avoid the propagation of simplistic concepts is even more important because so much remains before we have a satisfactory understanding of the functional roles of the pertinent structures. The decision as to which structures should be included in the term the “limbic system” has become more and more arbitrary, as new methods have shown that many more parts of the brain participate in emotional processing than believed when the term was coined.¹ For example, parts of the neocortex (not only the allocortex) are strongly implicated in processing of emotions—the **prefrontal cortex** and the **insula**, in particular. Further, cell groups in the basal parts of the hemispheres—the so-called **basal forebrain** (including the nucleus accumbens, discussed in Chapter 23)—are densely interconnected with both limbic structures and the neocortex, and these cell groups participate in both emotional and cognitive processing. We therefore include a description of the basal forebrain in this chapter, while the prefrontal cortex is discussed mainly in Chapters 33 and 34. Our present knowledge indicates that all complex behavioral reactions engage the amygdala, the septal nuclei, the hippocampal region, the reticular formation, the hypothalamus, and large parts of the cerebral cortex.

1 Some authors include in the “limbic system” the mammillary body, the anterior thalamic nucleus, the ventral striatum, and the hypothalamus. Some even include brain stem cell groups such as the ventral tegmental area (VTA), the raphe nuclei, the PAG, and certain cholinergic neuronal groups.

As witnessed by their numerous interconnections, all of these regions cooperate to exert an integrated influence on the peripheral somatic and autonomic effectors. What the American psychologist S.P. Grossman (1976, p. 361) said about the septal nuclei probably holds for the rest of the limbic structures, too: “Just about every behavior and/or psychological function which has been investigated to date has been shown to be affected in some way by septal lesions.”

In **conclusion**, the “limbic system” does not represent a unity that can be defined with a reasonable degree of precision. As stated by the American neurologist Antonio Damasio (1995, p. 20), “the bizarre distinction between cognition and emotion, as if somehow one could have thoughts without emotion, a mind without affect . . . The rift between emotion and cognition acquired a neuroanatomical counterpart in the duality between limbic system and neocortex.”

The Circuit of Papez

In 1937, Papez described what he considered a closed circuit of connections starting and ending in the hippocampus. From the hippocampus, the flow of signals was postulated to pass to the mammillary nucleus, from this nucleus to the anterior thalamic nucleus, from there to the cingulate gyrus, and then finally back to the hippocampus. This circuit of interconnected cell groups was hypothesized to form the anatomic basis of emotional reactions and expressions. These suggestions formed the basis for the concept of “the limbic system,” which was introduced in the early 1950s by Paul MacLean.

THE AMYGDALA

Main Anatomic and Functional Subdivisions

The amygdala (the amygdaloid nucleus) is located in the temporal lobe, underneath the uncus (Figs. 31.1, 31.2, and 31.3; see Fig. 6.29). In humans, the amygdala is a complex of subnuclei, each with a distinctive internal structure, neurotransmitters, and connections.² Here we restrict ourselves to distinguishing between a small **corticomедial** (including a **central nucleus**) and a large **basolateral** nuclear group (including the **lateral nucleus**). The basolateral group increases in size from lower to higher mammals and is particularly well developed in humans. The corticomедial nuclear group lies close to

the olfactory cortex (see Fig. 19.3). To **simplify**, we may say that the corticomедial nuclei are connected primarily with the olfactory bulb, the hypothalamus, and the visceral nuclei of the brain stem, whereas the basolateral nuclei are mainly connected with the thalamus and prefrontal cortex. In addition, the basolateral nuclei send fibers to the ventral striatum and the basal nucleus. This would suggest that the corticomедial part of the amygdala is concerned primarily with autonomic functions, whereas the basolateral parts are more involved in conscious processes related to the frontal and temporal lobes. The many intrinsic connections among the various nuclei show that they must cooperate extensively, however.

The amygdala (or its many components) participates in several higher mental functions, each of which is highly complex. Its “functions” are correspondingly complex and hard to define. Nevertheless, some salient features are clear. Thus, a central task of the amygdala is the establishment of links between **stimuli** and their **emotional value** (put very simply, whether something is good or bad). Most of our memories have some—often quite strong—emotional coloring, which is crucial for our ability to react appropriately to a stimulus. Think of the importance of being able to judge the facial expressions of other people, the emotional aspects of their speech, and so forth. As we discuss in this chapter, damage to the amygdala in monkeys leads to, among other things, difficulties in **social interactions**.

Afferent Connections of the Amygdala

The **corticomедial** nuclei receive afferents from the **olfactory bulb**, the **hypothalamus**, the **intralaminar thalamic nuclei**, and the **septal nuclei** (Fig. 31.4). They also receive **dopaminergic** fibers from the ventral tegmental area in the mesencephalon, as well as fibers from

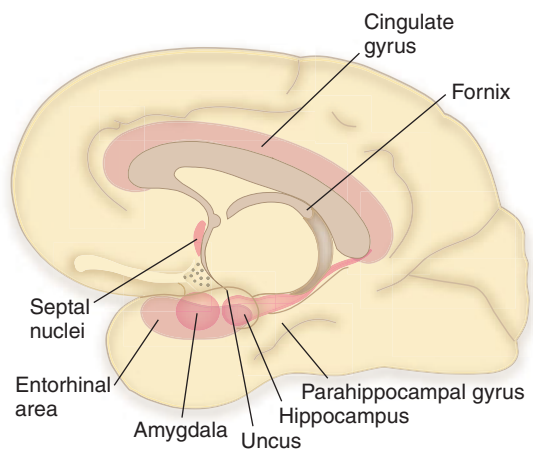


FIGURE 31.1 *The limbic structures.* The right hemisphere, as viewed from the medial aspect. The regions and cell groups indicated in red are usually included in the term “limbic system.”

² The amygdala as we describe it here is structurally and functionally heterogeneous, and we lump the various nuclei under one name purely for convenience. Indeed, in a critical review Larry Swanson and Gorica Petrovich (1998) concluded “...it is necessary to ask whether the concept of a structurally and functionally defined amygdala is indeed valid, or whether the concept is hindering attempts to understand general principles of telencephalic architecture by imposing an arbitrary classification on heterogeneous structures that belong to different functional systems.”

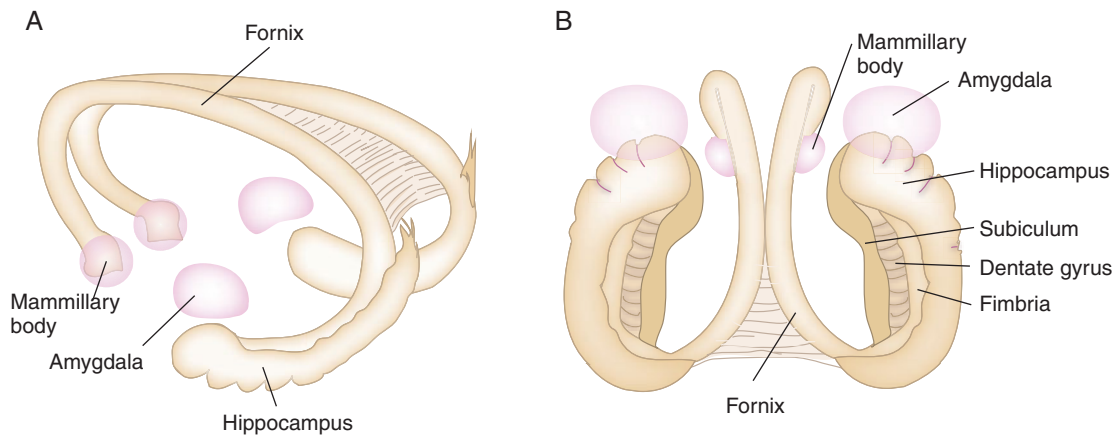


FIGURE 31.2 The hippocampus, fornix, mammillary nucleus, and the amygdala. A: Viewed obliquely from behind. B: Viewed from above.

the **parabrachial area** (in the dorsolateral pons). The latter projection may convey information about taste and, in addition, about painful stimuli. Thus, ascending fibers of spinal **lamina I** nociceptive neurons end in parts of the parabrachial area that project to the central amygdaloid nucleus (among other targets). The sensory units of this pathway have very large receptive fields and receive convergent inputs from the skin and viscera. It seems likely that this lamina I–parabrachial–amygdaloid pathway contributes to the **emotional aspects of pain**.

The **basolateral nuclei** receive fibers from several **thalamic nuclei**, the **prefrontal cortex**, parts of the **temporal**

lobe, and the **cingulate gyrus** (Fig. 31.4). Together, the basolateral nuclei—the **lateral nucleus** in particular—receive all kinds of sensory information. This may be emotionally neutral information from cortical association areas and emotionally laden information about unpleasant and threatening stimuli from the reticular formation, the intralaminar thalamic nuclei, and perhaps parts of the cortex (the insula). Thus, the amygdala receives, for example, information about **fear-provoking stimuli** and their **context**. Efferents from the lateral nucleus reach other amygdaloid nuclei that may influence the cortex (conscious experience of emotions) and

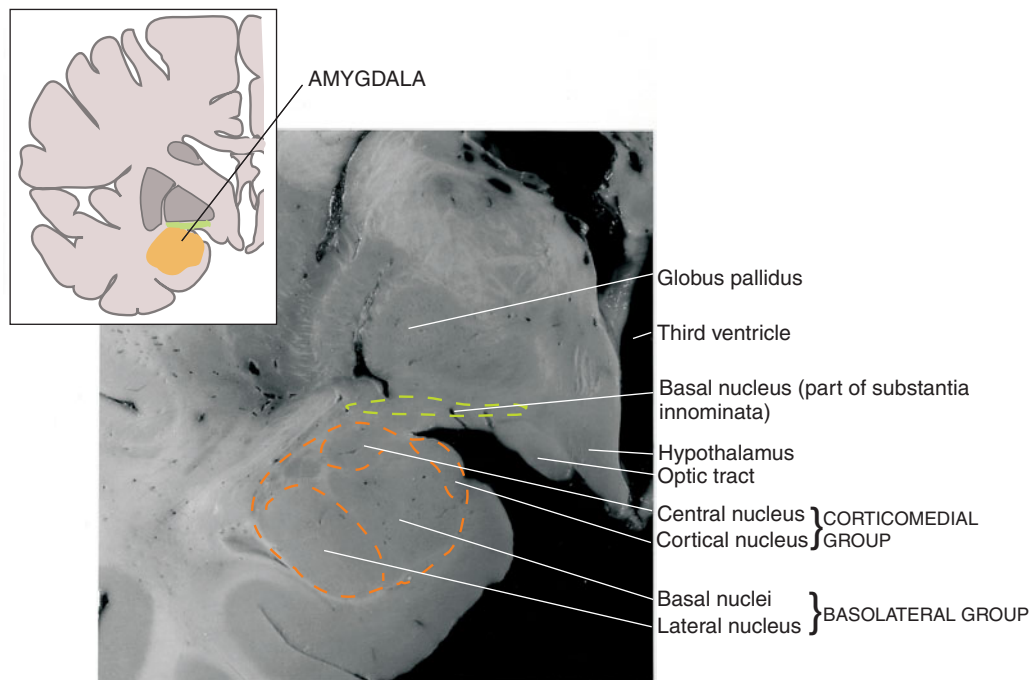


FIGURE 31.3 The amygdala. Frontal section through the left hemisphere (cf. Fig. 31.1). Some of the amygdaloid nuclei are marked with orange stippled lines. The basal nucleus is indicated with green

stippled line. The amygdala and the cerebral cortex of the temporal lobe are closely connected. The section is placed more posteriorly than the one in Fig. 31.9.

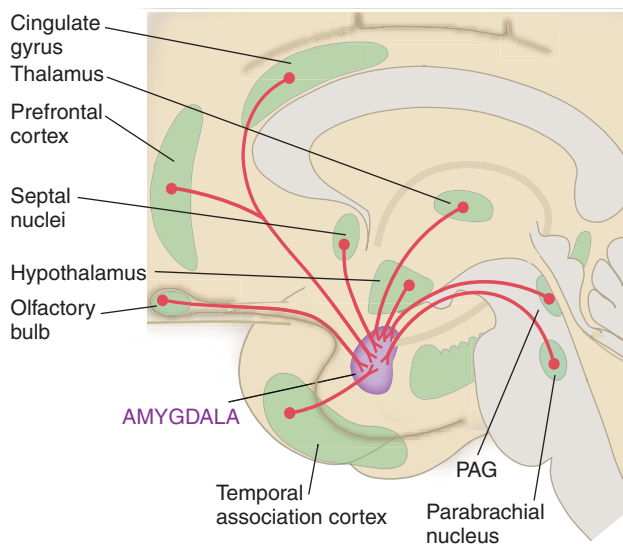


FIGURE 31.4 *Afferent connections of the amygdala.* Note the connections with the hypothalamus, cortical areas in the temporal and the frontal lobes, and brain stem nuclei. Not all known connections are shown.

brain stem cell groups (behavioral reactions, including autonomic responses associated with emotions).

Efferent Connections of the Amygdala

The efferent connections of the amygdala are mostly reciprocal to the afferent ones. One major efferent pathway goes to the **hypothalamus**. Most of these fibers are collected in the macroscopically visible **stria terminalis**, which arches over the thalamus (Fig. 31.5). The fibers end primarily in the ventromedial hypothalamic nucleus (compare with the efferents from the hippocampal formation, which run in the fornix and end in the mammillary nucleus). Other efferents pass to the **thalamus** (especially to the mediodorsal nucleus [MD]), enabling signals from the amygdala to reach the **prefrontal cortex** (Fig. 31.5). As mentioned, these connections may be important for the conscious experience of emotions, such as fear and anxiety. In particular, the amygdala–prefrontal connections might ensure that we spend our limited attentional resources on the most important stimuli (those with emotional coloring), and further, that emotional and cognitive information is integrated prior to decisions and actions.

Parts of the allocortex receive fibers from the amygdala, especially the **hippocampal formation** (the entorhinal area and the subiculum) and the **septal nuclei**. Fibers to the **ventral striatum** (including the nucleus accumbens; see Fig. 23.15) and the basal nucleus (Figs. 31.3 and 31.8; see Fig. 10.1) arise in the basolateral nuclei of the amygdala. The projections to the basal nucleus can induce **activation of the EEG** (arousal and increased attention). There is experimental evidence that synaptic

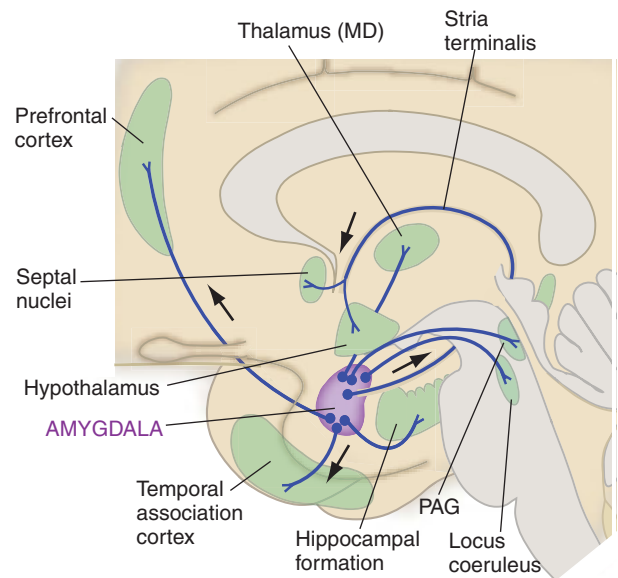


FIGURE 31.5 *Efferent connections of the amygdala.* Note connections to the hypothalamus, the PAG, the hippocampus, and cortical areas in the temporal and frontal lobes. See also Fig. 31.6.

learning effects on neurons in the auditory cortex depend on inputs from the amygdala (and the basal nucleus) besides the specific auditory information via the medial geniculate body. It seems likely that this is related to the well-known effects of motivation on **learning**: we remember better the material that has emotional coloring. The amygdala

Finally, there are connections from the amygdala (especially from the central nucleus) to various **brain stem nuclei**, such as the **PAG** (Fig. 31.6), parts of the **reticular formation**, and **parasympathetic** cranial nerve nuclei. These connections—together with those to the hypothalamus—are important for the autonomic and somatic expressions of emotions. The connections to the PAG are involved in eliciting **conditioned fear behavior**. The PAG sends fibers to various brain stem premotor networks (Fig. 31.6). In rats, conditioned fear includes so-called **freezing**; that is, the animal becomes completely still. The freezing reaction, which includes suppressed pain transmission, occurs typically when a rat meets a predator, such as a cat. When exposed to a sudden threat, humans also experience a similar halt of all movements, until an appropriate behavioral response is selected (flight, fight, or continued immobility).

Many behavioral changes have been produced by **electrical stimulation** of the amygdala in animals with implanted electrodes. (In such experiments, the electrodes have been inserted and fixed to the skull under general anesthesia. Afterward the electrodes cause the animal no pain or obvious discomfort.) Stimulation produces a varied pattern of somatic and autonomic responses, which appear to be parts of more complex behavioral reactions. As would be expected from the anatomic data discussed

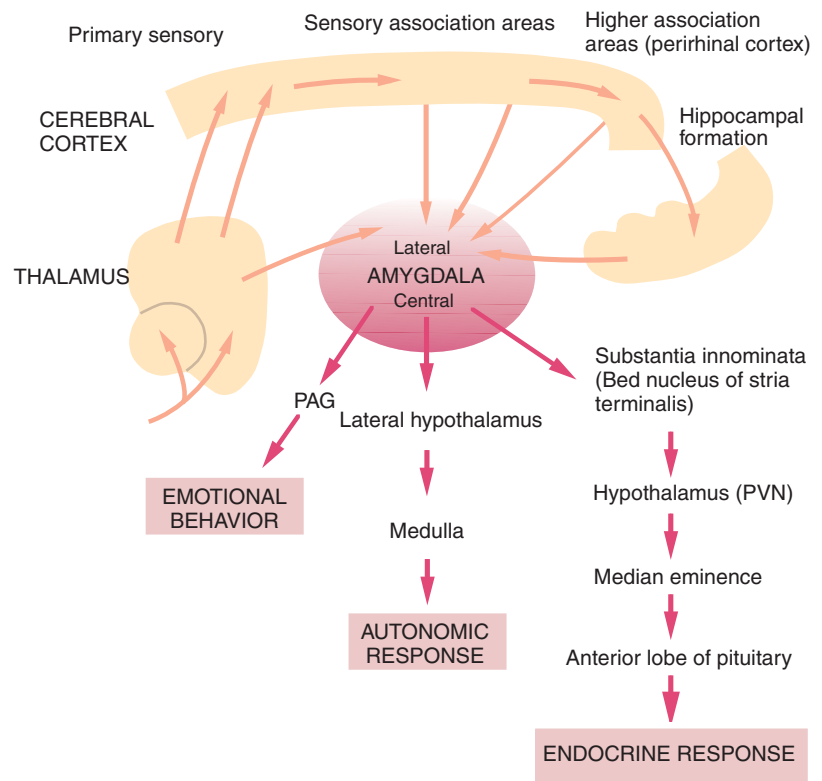


FIGURE 31.6 *The amygdala and conditioned fear.* The neural substrate of the autonomic, endocrine, and somatic responses elicited by a conditioned stimulus (sound) associated with an electric foot shock. The connections of the amygdala with the sensory association areas are necessary for discriminative aspects of stimulus analysis, while connections with the hippocampus mediate contextual conditioning. (Based on LeDoux 1995.)

in the preceding text, stimulation of medial and lateral parts of the amygdala gives different responses. Stimulation of the **corticomedial nuclear group** produces smacking, salivation, and licking and chewing movements. Emptying of the rectum and the bladder may occur, together with inhibition of voluntary movements. Stimulation of the **basolateral nuclear group** often produces **arousal** and signs of increased attention: the animal lifts its head, its pupils are dilated, and it looks around (especially toward the side opposite of the stimulating electrode). The attention of the animal appears to be directed toward something in the surroundings. As might be expected, activation of the EEG occurs along with these behavioral changes. Strong stimulation can produce dramatic effects, such as signs of fear or rage.

Amygdala and Conditioned Fear

As mentioned, the amygdala has a role in a variety of emotions and emotionally related behaviors. This, with its anatomic heterogeneity, strongly suggests that the amygdala is not a functional unit. In the search for the biologic substrate of specific behaviors, **conditioned fear** has been intensively studied (Fig. 31.6), and crucial links in the pathways that underlie this phenomenon have been “dissected” out in experimental animals. Such detailed data may help us understand how the amygdala participates in other kinds of tasks.

Situations with purely mental stress can produce a conditioned-fear reaction in rats as in humans. In rats, the conditioning may be established by letting a tone be followed repeatedly by a painful stimulus (e.g., an electric shock to the foot). After a while, the tone alone elicits fear-related behavior—that is, behavior that normally occurs in threatening or dangerous situations (such as the sight of a cat). As discussed earlier, the fear reaction of a rat includes autonomic, somatic, and endocrine responses, such as freezing and increased secretion of cortisol. When a tone is the conditioning stimulus, the necessary information about the sound is transmitted directly from the thalamus to the **lateral amygdaloid nucleus**. The auditory **cortex** is necessary for the occurrence of the fear reaction only if the animal has to discriminate two tones with different frequencies. The **hippocampus** is not necessary for the tone conditioning. It is necessary, however, for a weaker **contextual conditioning**; that is, the fear reaction can also be elicited by clues in the environment of the experimental situation (such as objects, sounds, or odors). Signals are relayed (directly and indirectly) from the lateral nucleus to the **central nucleus**. Efferents from the central nucleus to autonomic and somatic cell groups in the brain stem (among them the PAG) probably mediate main components of the fear reaction. Thus, destruction of the central nucleus abolishes both the freezing and autonomic responses in conditioned fear.

Main Tasks of the Amygdala

The tasks performed by the amygdala have been clarified by lesion and stimulation experiments in animals, and recently by numerous functional magnetic resonance imaging (fMRI) studies in humans. Further, important information comes from examination of a few persons who lack the amygdala (usually after surgical treatment of epilepsy). While the connection between the amygdala and emotions is firmly established, much investigation remains before we understand its specific contributions to emotional processing and human behavior.

Animal experiments show that a central task for the amygdala is to establish **associations between sensory stimuli and their emotional coloring**. It is crucial that we can decide quickly—before slower conscious deliberations—whether a stimulus (or a situation) is threatening or safe (punishment or reward). Accordingly, fMRI studies in humans show activation of the amygdala when viewing pictures with an emotional content. Further, bilateral lesions of the amygdala in monkeys reduce behavior elicited by emotions. For example, the animals show no fear of snakes (monkeys have an inborn fear of snakes). This can probably be explained by the removal of amygdaloid effects on the hypothalamus and on the brain stem autonomic and somatic motor centers (among them the PAG; Fig. 31.6). Sensory stimuli (such as the sight of a snake), although reaching consciousness, would not be able to elicit the normal behavioral reactions.

The sight of **faces** expressing **anger** or **fear** causes a robust activation of the human amygdala, as shown via fMRI. Correspondingly, patients with amygdaloid lesions have difficulties with **recognizing facial expressions**. Interestingly, such patients do not show the normal tendency to remember events or stimuli that have an emotional coloring better than they remember neutral ones. This has been demonstrated, for example, by showing films containing emotionally neutral material and scenes that evoke strong emotions.

Selective lesions in adult monkeys produce a pattern of behavior characterized by **social disinhibition**.³ For example, they initiate more physical contact, suggesting “. . .that heightened affiliative social interactions following amygdala lesions stems from a more general inability to properly perceive danger or threat in the environment and use such information to modulate social behavior adaptively” (Machado et al. 2008, p. 263). This would fit with the amygdala working as a sort of **alarm**—it

evaluates very rapidly a stimulus for its threatening value, and initiates appropriate behavior. However, selective lesions of the amygdala in monkeys produce fewer behavioral changes than reported in early experiments with lesions that included adjoining parts of the temporal lobe (see Chapter 34, under “Symptoms after Lesions of the Temporal Cortex”). For example, no signs of abnormal **social development** were observed the first 6 months after bilateral lesions of the amygdala in infant monkeys, in relationships with either the mother or other infants in the group. However, the infants did not exhibit the normal signs of distress to separation from the mother, presumably owing to a reduced ability to perceive danger and threatening situations.

Is the Amygdala Concerned Only with Negative Emotions?

Whereas most studies have focused on a correlation between amygdala activity and negative emotions such as fear and anger, recent studies suggest that the amygdala plays a role for the recognition of positive emotions as well. For example, a meta-analysis of human positron emission tomography (PET) and fMRI studies found that both negative and positive stimuli were associated with higher amygdala activity compared with neutral stimuli. Indeed, single-unit studies in monkeys identified distinct populations of amygdaloid neurons responding to positive and negative stimulus valence, respectively (the two kinds of neuron were not spatially segregated in the basolateral amygdala, however).

Electric Stimulation of the Human Amygdala

In humans, the amygdala has been stimulated in conjunction with brain surgery of the temporal lobe under local anesthesia. A wide spectrum of autonomic and emotional reactions has been produced in such cases, but most pronounced is a feeling of **anxiety**. Memory-like hallucinations and **déjà vu** experiences (the feeling of having experienced the same situation before) have also been reported. This is called a **dreamy state** and can occur in epileptic seizures that start in the temporal lobe. Similar effects—that is, fear and various kinds of hallucinations—have been produced by stimulation of the anterior portion of the hippocampus and the lateral cortex of the temporal lobe (in the superior temporal gyrus). This relationship can presumably be explained by the close connections between these regions and the amygdala, all being parts of a more widespread network for handling of emotions and memories.

Is Amygdala Necessary for the Experience of Emotions?

The finding that the amygdala is necessary for expression of emotions (at least some aspects) raises the question of

³ **Hypersexuality** was one of the behavioral changes reported in the early studies with large bilateral lesions of the amygdala in monkeys. However, this may be related to damage to allocortical areas near the amygdala rather than to the amygdala itself. Nevertheless, it is conspicuous that the amygdala is among the brain regions with the highest density of **receptors for sex hormones**. Conceivably, the level of sex hormones in the blood influences the activity of neurons in parts of the amygdala (the sex hormones are lipid-soluble and pass the blood–brain barrier easily).

whether the amygdala is necessary also for the subjective **experience** of emotions (such as fear or anger). A patient with bilateral destruction of the amygdala, described by the British psychiatrist R. Jacobson (1986), illustrates this point. She appeared calm and relaxed outwardly and had normal heart rate in situations in which she experienced great anxiety and wanted to run away. Presumably, the coupling between the emotions and the emotional reactions was disrupted in this patient (see Chapter 30, under “Emotions and Emotional Reactions”). Further, 20 patients with amygdaloid lesions after epilepsy surgery described their daily emotions—positive and negative—in the same manner as normal controls (Anderson and Phelps 2002).⁴

The Amygdala, Learning, and Unlearning

The conditioned-fear reaction discussed in the next subsection requires a learning process: the rat learns to associate an innocuous stimulus with something painful. Destruction of the amygdala prevents establishing the conditioned response. Indeed, induction of **LTP** occurs in certain parts of the amygdala in conjunction with development of a conditioned fear response. Thus, experiments with monkeys after a lesion restricted as far as possible to the amygdala show that they have difficulties in learning the association between objects and their meanings. They can recognize objects but cannot relate them to other kinds of information, such as whether the object was associated with a reward or something unpleasant. Many other observations support that the amygdala is necessary for the learning of associations between stimuli and their significance in terms of reward or punishment. We may say that the amygdala is crucial for the emotional coloring of experiences and sensations, and that associations are remembered. The amygdala is not alone in this respect, however. Both the amygdala and parts of the prefrontal cortex are necessary in monkeys for the learning and later retrieval of associations between visual stimuli and food rewards. The connections involved are partly direct fibers from the amygdala to the **ventromedial prefrontal cortex** and partly a pathway interrupted in the mediodorsal thalamic nucleus (MD). Experiments in rats suggest that connections between the basolateral amygdala and the **ventral striatum** are also necessary for establishing stimulus–reward associations.

4 Not all patients with bilateral damage of the amygdala exhibit a dissociation of the experienced emotion and the emotional expressions, however. Indeed, there are surprisingly large individual variations in symptoms among patients with amygdaloid lesions. Conceivably, the age at which the lesion occurred plays a decisive part: with early lesions, other parts of the brain would be expected to at least partly take over the tasks of the amygdala. Further, prior experiences and subtle difference in context may strongly influence how different subjects with lesions of the amygdala experience and respond to identical stimuli.

Connections from medial parts of the prefrontal cortex appear to be necessary for unlearning—**extinction**—of the conditioned fear response. Extinction occurs when the conditioned stimulus regularly occurs without a subsequent unconditioned stimulus, but not in rats after removal of the medial prefrontal cortex. Other data also indicate that extinction depends on an active **inhibition** of the amygdala from the prefrontal cortex—not on the disappearance of the synaptic changes underlying the associations.

Amygdala, Anxiety, and Neurotransmitters

The amygdala contains many neurotransmitters, and to sort out the functional role of each is a formidable task. For practical reasons, therefore, scientists concentrate on studying one or a few at a time, with the danger of overlooking the contributions of other transmitters. We restrict ourselves here to the transmitters involved in **conditioned fear** and **psychic stress**. As mentioned, signals pass from the lateral to the central nucleus through both direct and indirect routes.

Electrical stimulation of the lateral nucleus evokes primarily γ -aminobutyric acid (**GABA**)-mediated inhibition in the central nucleus (acting at both GABA_A and GABA_B receptors). Some inhibition occurs presynaptically by reducing the release of **glutamate**. Drugs that reduce **anxiety** (anxiolytics) may function by interfering at this level. The **benzodiazepines** (Valium and others) bind to specific sites on the GABA receptor (benzodiazepine receptors) and potentiate the effect of GABA. The density of benzodiazepine receptors is high in the amygdala and particularly high in the lateral nucleus (and one other subnucleus of the basolateral complex). Local infusion of benzodiazepines in these nuclei reduces expressions of conditioned fear in experimental animals.

Corticotrophin-releasing hormone (CRH) may also be an important transmitter in the amygdala in relation to anxiety and stress. Besides containing CRH-positive cell bodies, the central nucleus receives many CRH-containing fibers (neurons in the parabrachial area and the locus coeruleus contain CRH, for example). Injection of CRH into the cerebral ventricles increases **stress reactions** and fear-related behavior, presumably by acting in the amygdala but also in other areas. For example, noradrenergic neurons in the locus coeruleus are activated, which may contribute to arousal as part of a stress reaction. Because acute and chronic stress increases CRH in the amygdala, microinjections of CRH antagonists in the central nucleus abolish some stress reactions.

The **expectation of pain** evokes an endocrine response, as one part of a stress reaction (see Chapter 30, under “Psychosomatic Disorders”). This may be mediated by neurons in the central nucleus, which project to the paraventricular hypothalamic nucleus. CRH-containing neurons in the paraventricular nucleus project to the

median eminence. There CRH is released and reaches the anterior pituitary via the portal system (see Fig. 30.7). CRH increases the secretion of ACTH, thus increasing cortisol in the bloodstream.

Several transmitters other than CRH show changes in relation to anxiety and stress. **Neuropeptide Y (NPY)** has attracted much interest. Thus, microinjection of NPY in the amygdala evokes largely the opposite effects of CRH on stress and fear-related behavior. NPY is present in many neurons in the amygdala (colocalized with norepinephrine, GABA, or somatostatin) and in terminals of afferent axons. Animal experiments suggest that, whereas CRH is crucial in eliciting a stress reaction, NPY that is released after the reaction has started protects against overshooting.

The Amygdala and Depression

As mentioned, CRH is one likely transmitter (among several) for evoking fear-related behavior and stress reactions, and the amygdala is an important site of action. CRH also appears to be related to mood. Thus, the concentration of CRH in the cerebrospinal fluid is increased in many deeply depressed patients and victims of suicide. In the latter group, lowered density of the CRH receptor occurred in the frontal lobe (down-regulation due to constantly increased CRH available?). A transgenic mouse strain overproducing CRH has increased levels of ACTH and cortisol in the blood as expected. In addition, mice from this strain show behavior indicative of anxiety (e.g., the way they behave in novel situations). This behavioral pattern is normalized by supply of CRH antagonists.

Measurement of **regional cerebral blood flow** supports the fact that the function of the amygdala is altered in seriously depressed patients. Thus, compared with a control group, depressed patients had increased blood flow in the left prefrontal cortex and amygdala. This observation does not tell us how the amygdala is involved in depression, however—for example, whether the blood flow changes are secondary to a change of mood, or whether changes in the amygdala come first.

SOME ASPECTS OF CORTICAL CONTROL OF AUTONOMIC FUNCTIONS AND EMOTIONS

Assigning specific functions to cortical regions builds on methods that can only provide indirect answers (functional deficits after lesions, blood flow changes associated with certain behaviors, EEG, single neurons recordings, and so forth). Because distributed networks—not single areas—are responsible for the execution of complex tasks, assigning functions to specific regions must be imprecise and simplistic. Although we need “pigeon holes” and labels to aid our thinking, we should

bear in mind that our simplifying concepts have limited explanatory power. All cortical areas we mention here with focus on autonomic functions and emotions are also involved in other tasks (participate in other networks). For example, most of the cortical areas regulating autonomic functions also participate in processing of emotions. This is not unexpected, as the autonomic adjustments are an integral part of complex behavioral responses.

Autonomic Functions

Experimental and clinical data show that wide areas of the cerebral cortex influence the activity of structures innervated by the autonomic nervous system. This influence is mainly exerted via the amygdala, the hippocampal formation, and the septal nuclei, which in turn, influence the hypothalamus and brain stem nuclei. In addition, there are some direct connections from the **insula** and the **orbitofrontal cortex** to the hypothalamus. Further, neocortical areas in the frontal and temporal lobes project to the amygdala and can therefore influence the hypothalamus indirectly.

The **cingulate gyrus**—one of the limbic structures—appears to be involved in organization and initiation of various kinds of goal-directed behavior.⁵ It projects to the hippocampal formation, to the septal nuclei, and to the amygdala—all of which have connections to various parts of the hypothalamus (Fig. 31.7). Electrical stimulation of the cingulate gyrus elicits a combination of autonomic (visceral) and somatic effects. Autonomic effects include, for example, alterations of **respiration** and **circulation** (reduced rate of breathing, heart rate, and blood pressure), of the **digestive tract** (altered peristaltic movements and secretory activity), and **pupillary dilatation**. Somatic effects are expressed mainly as changes of muscle tone and often inhibition of ongoing movements.

Alterations of functions controlled by the autonomic system can be produced by stimulation of parts of the cortex other than the cingulate gyrus. Stimulation of the **orbitofrontal cortex**, the **insula**, and the **pole of the temporal lobe** produces effects similar to those obtained from the cingulate gyrus—that is, combined behavioral, emotional, and autonomic responses. Stimulation of the aforementioned neocortical regions not only produces effects on autonomic functions; somatic functions are altered as well. In contrast, alterations of autonomic functions can occur after stimulation of cortical regions that one might believe to be purely somatic, such as the motor and the premotor cortical areas. Thus, stimulation

⁵ While minor parts of the cingulate gyrus belong to the allocortex, most of it probably belongs to the oldest parts of the neocortex. The terms **limbic** or **paralimbic cortex** are often used of cortical regions that have intimate connections with limbic structures, such as the amygdala and the hippocampus.

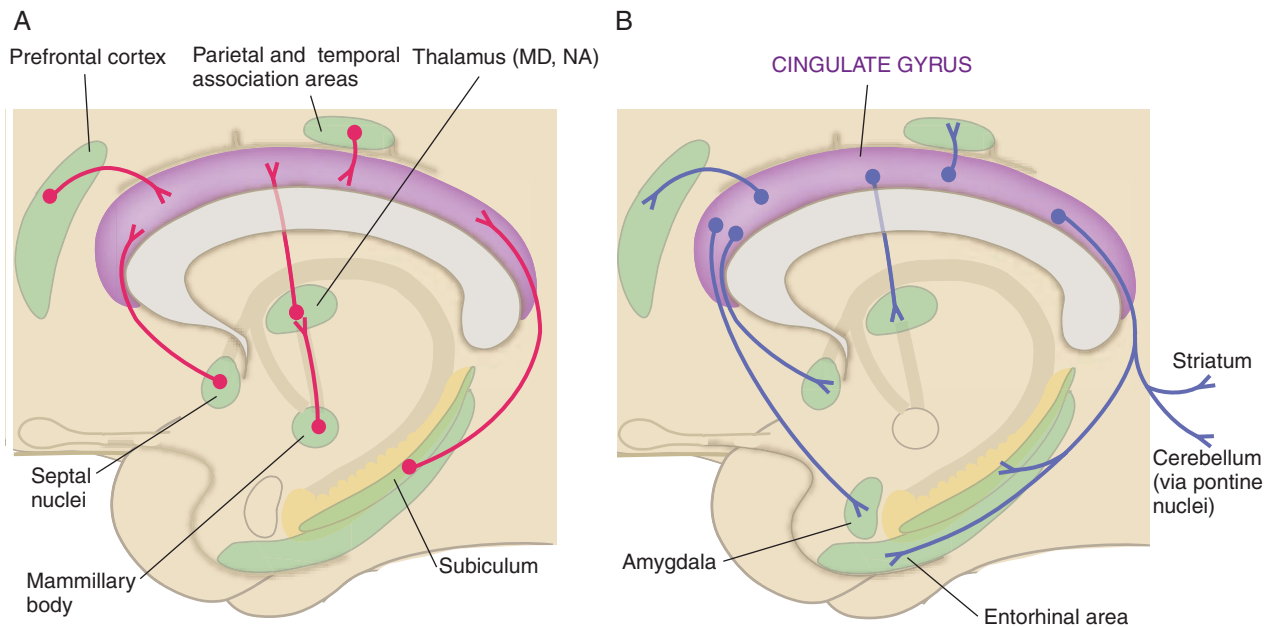


FIGURE 31.7 Main connections of the cingulate gyrus. **A:** Afferent connections **B:** Efferent connections. The cingulate gyrus has reciprocal

connections with neocortical association areas and with limbic structures and may act as a mediator between them.

of the **motor cortex** produces **vasomotor changes** (i.e., changes of the blood vessel diameter and, therefore, of blood flow) of the opposite body half. On damage to these cortical areas (as seen in patients with a cerebral **stroke**), vasomotor changes often occur in the paralyzed parts of the body. Even alterations of the heart rate and blood pressure and of the digestive tract can occur. As a final example of combined somatic and autonomic effects, stimulation of the **frontal eye field** (see Fig. 25.7) produces pupillary dilatation in addition to the more obvious conjugated eye movements.

Emotions and the Neocortex

As mentioned, cortical areas that show altered activity (as measured with PET and fMRI) in relation to emotions are more extensive than those initially included in the “limbic system” (Fig. 31.8). On the other hand, no area is solely concerned with emotional processing. The **cingulate gyrus** is a pertinent example. Even if it consists of several smaller subdivisions that differ with regard to connections, they all seem to be involved in both cognitive and emotional processing, albeit to a varying degree. The anterior part of the cingulate gyrus (**anterior cingulate cortex [ACC]**) consists of rostral part that is more concerned with affect regulation and a caudal part more concerned with cognitive task. In general, the ACC appears to be important for the choice of behavior in response to **conflicting stimuli**. The ACC also seems to monitor mental and bodily processes with

special focus on the detection of errors and conflicts.⁶ For this monitoring, emotions provide important information about values of different signals. Parts of the cingulate gyrus (both anterior and posterior parts) and the anterior **insula** (Fig. 31.8) also alter their activity in relation to emotions such as admiration and **compassion**.

Several parts of the **prefrontal cortex** show altered activity in relation to emotions in humans, and accordingly, lesions often produce emotional disturbances (see Chapter 34, under “Symptoms after Prefrontal Lesions”). Especially the **orbitofrontal** and **ventromedial** parts seem important for emotional regulation (Fig. 31.8). As mentioned, these parts have reciprocal connections with the amygdala. The orbitofrontal cortex may integrate competing, emotionally colored signals to provide an appropriate response. For example, a study compared the behavior of normal and orbitofrontal-lesioned monkeys in a situation where a snake occurred between the monkey and a piece of food. Presumably, amygdala informs about the values of the signals (snake and food), whereas the orbitofrontal cortex is necessary for evaluation and appropriate action. Such studies strongly suggest that the orbitofrontal cortex is important for behavioral **flexibility**—that is, the ability to alter behavior when needed and to choose among conflicting choices.

⁶ Stimulation of the ACC in monkeys produces, for example, **aggressive reactions**, whereas bilateral removal makes the animals tamer. They may also become **socially indifferent**—that is, they appear to have lost interest in other members of their group and do not try to make contact.

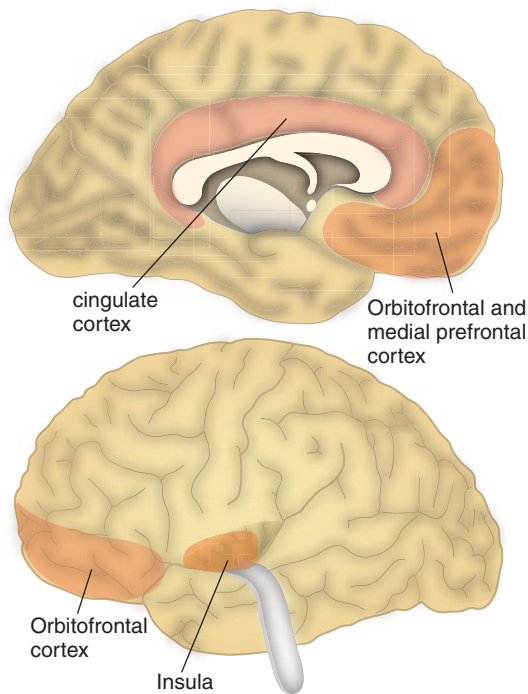


FIGURE 31.8 Parts of the cerebral cortex involved in processing of emotions.

Finally, an interesting hypothesis proposes that a network comprising the amygdala, the nucleus accumbens, the orbitofrontal cortex, the insula, and the anterior cingulate cortex monitors the **energetic demands** involved in performing a task (Boksem and Tops 2008). In so doing, the network would need to weigh the potential reward against the energetic cost of a task to reach a decision on whether to go on with it or not. Presumably, this process would require integration of emotional (motivational), and cognitive signals. The authors furthermore suggest that **mental fatigue** "...can best be considered as an adaptive signal that the present behavioural strategy may no longer be the most appropriate . . . fatigue may present the cognitive system with a signal that encourages the organism to lower present goals . . ." (p. 133).

Neocortex and emotions are discussed further in Chapter 34.

NEURONAL GROUPS IN THE BASAL PARTS OF THE HEMISPHERES: THE BASAL FOREBRAIN

Below and medial to the well-defined basal ganglia many neurons are spread out rather diffusely. Unfortunately, the nomenclature for this region is not consistent among authors. Thus, different names are often applied to the region or parts of it without attempting to describe more precisely what is meant. The old anatomists named it the **substantia innominata** (the region without a name),

presumably reflecting that nothing was known about its connections and functions. The **basal forebrain** (basal prosencephalon) is usually used synonymously with the substantia innominata. Only recently are its connections, cytochemistry, and functions being clarified, showing that the basal forebrain is not an anatomic or functional unity. It is now commonly accepted to divide the basal forebrain into three overlapping regions, differing regarding neurotransmitters and connections: the **ventral striatopallidum**, the **extended amygdala**, and the **basal nucleus** (Figs. 31.3 and 31.9; see Fig. 10.1). The ventral striatopallidum was discussed in Chapter 23. We discuss here only certain aspects of the basal forebrain, starting with a large group of cholinergic neurons that are found in the basal nucleus and some adjacent cell groups.

Cholinergic Neurons Projecting to the Cerebral Cortex

Cholinergic neurons of the basal forebrain form a thin disc close to the basal surface of the hemisphere—the **basal nucleus**—and are furthermore found in the **septal nuclei** and the diagonal band of Broca that extend dorsally close to the midline (Figs. 10.1, 31.9, and 31.10). The septal nuclei form the most well-defined anatomic entity among these cell groups, and lie just in front of the anterior commissure (Fig. 31.1; see Fig. 23.15).⁷ The **diagonal band of Broca** forms a transition between the basal nucleus and the septal nuclei.

The cholinergic neurons of the basal forebrain project to the cerebral cortex (both to the allocortex, including the hippocampus, and the neocortex).⁸ The septal nuclei and the diagonal band send fibers mainly to the hippocampal formation, whereas the basal nucleus projects to the neocortex with a rough topographic order.

Although the **afferent** connections of the cholinergic cell groups of the basal forebrain are incompletely known, important inputs arise in the brain stem, notably in the **locus coeruleus** and in **cholinergic cell groups in the dorsolateral pons** (such connections are involved in the ascending activating system discussed in Chapter 26). Further, afferents come from the **hypothalamus**, the **cingulate gyrus**, and the **nucleus accumbens** of the ventral striatum.

⁷ The region containing the septal nuclei is called the precommissural part of the septum (located anterior to the anterior commissure, see Fig. 6.26). The postcommissural part is the **septum pellucidum**, which contains no neurons (see Fig. 6.29).

⁸ Not all neurons in the basal forebrain projecting to the cortex are cholinergic. Some neurons, mingled with the cholinergic ones, are **GABAergic**. Some contain the neuropeptide **galanin**, partly colocalized with acetylcholine. In humans the diagonal band of Broca has many **somatostatin**-containing neurons, although it is not known whether they project to the cortex. Thus, although apparently the majority of the neurons in the basal forebrain projecting to the cortex are cholinergic, it is not a transmitter-specific system. This is of relevance when trying to explain the symptoms of diseases with loss of neurons in the basal forebrain (notably Alzheimer's disease, discussed in Chapter 10).

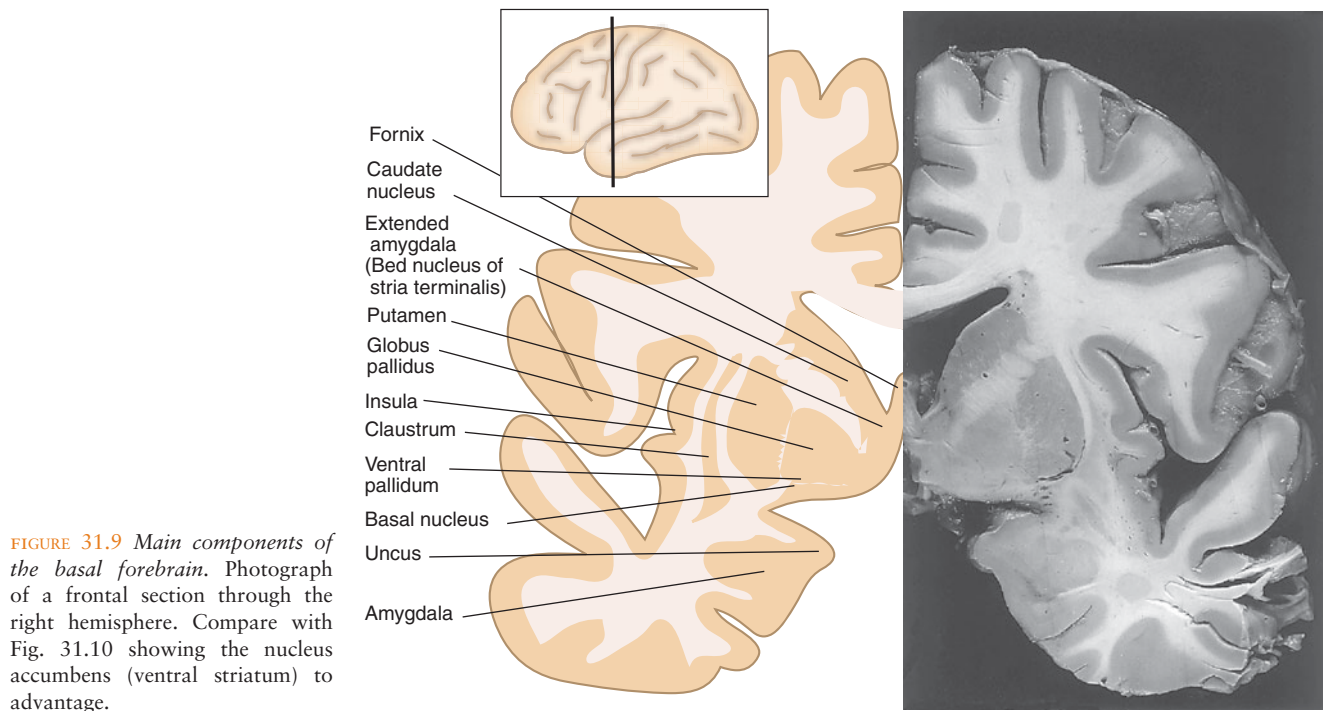


FIGURE 31.9 *Main components of the basal forebrain.* Photograph of a frontal section through the right hemisphere. Compare with Fig. 31.10 showing the nucleus accumbens (ventral striatum) to advantage.

Among the cholinergic cell groups, the **septal nuclei** first attracted interest because early lesion and stimulation experiments showed that they influence autonomic functions, emotions, and behavioral reactions. For example, lesions of the septal nuclei in animals alter sexual and foraging behavior: aggressive behavior appears to be reduced (as stimulation of the septal nuclei can produce aggression). The effects are similar to those produced by lesions of the amygdala and the anterior parts of the cingulate gyrus and presumably can be explained by the connections of the septal nuclei with these parts and the hypothalamus. Symptoms specific to the septal nuclei, constituting the so-called septal syndrome, have not been proved convincingly.

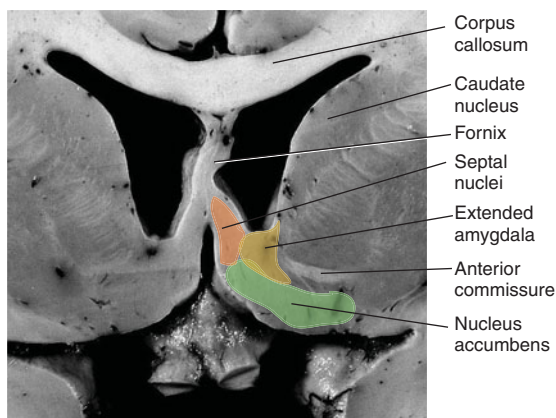


FIGURE 31.10 *The basal forebrain.* The positions of three main nuclear components are indicated with different colors in the photograph of a frontal section through the hemisphere (cf. Fig. 23.15).

Functional Roles of the Basal Forebrain Cholinergic Neurons

Later studies turned to the role of the cholinergic neurons in **attention** and **memory** mechanisms. Particularly seminal in this respect was the discovery of cell loss in the **basal nucleus** in patients with **Alzheimer's disease**. Some studies in monkeys suggested that the septal nuclei, the diagonal band of Broca, and the basal nucleus all must be destroyed to produce memory impairments. More detailed experimental studies with injection of substances that destroy the cell bodies (but not passing fibers) suggest that the septal nuclei and the diagonal band of Broca are especially important for memory (presumably because of their connections with the hippocampus), whereas the basal nucleus is more concerned with maintaining and perhaps focusing attention. Further, there are several reports of patients with small lesions in the anterior parts of the cholinergic cell groups who exhibited clear-cut memory loss (both for recent and past events). One such patient, for example, had a lesion affecting primarily the diagonal band of Broca, as judged via MRI. The difficulties with the exact localization of the damage in such patients warn against firm conclusions, however.

Other Components of the Basal Forebrain: The Ventral Striatopallidum and the Extended Amygdala

The ventral striatopallidum—discussed in Chapter 26—consists of the ventral striatum (including the nucleus accumbens) and the ventral pallidum. These parts of the

basal forebrain show similarities with the basal ganglia (dorsal striatum and dorsal pallidum) regarding cytoarchitectonics, cytochemistry, and connections (e.g., a dense dopaminergic innervation from the mesencephalon). In contrast to the dorsal striatum (the caudate nucleus and the putamen), which receives the main afferent input from the neocortex, the ventral striatum receives main inputs from the allocortex and the amygdala. A further characteristic of the ventral striatum is that it projects to the hypothalamus and brain stem nuclei, such as the periaqueductal gray (PAG) and the motor vagus nucleus. The ventral pallidum projects to the mediodorsal thalamic nucleus (MD) that sends fibers to the prefrontal cortex (cf. the projection of the dorsal pallidum to the VL/VA thalamic nuclei that project mainly to the premotor areas).

Another rather diffuse cell group in the basal forebrain, continuous with the ventral pallidum, is called the **extended amygdala** because it forms a rostral extension of the medial amygdala (Figs. 31.9 and 31.10). Most of it is made up of the **bed nucleus of the stria terminalis** (Figs. 31.6 and 36.9). The stria terminalis is a bundle of efferent fibers from the amygdala to the septal nuclei and the hypothalamus (Fig. 31.5). The bed nucleus lies medial to the pallidum at the same anteroposterior level and further anterior to the anterior commissure (i.e., close to the septal nuclei). When this part

of the basal forebrain is lumped with the amygdala, it is because they share many transmitters and connections. The anatomic distinction between the extended amygdala and the nucleus accumbens is not sharp, however, and both receive, for example, dopaminergic fibers from the mesencephalon.

The Medial Forebrain Bundle

Many fibers interconnecting the various limbic structures are located in a diffusely delimited, parasagittal fiber mass in the basal part of the hemisphere. This ill-defined structure is called the **medial forebrain bundle** and passes through the lateral parts of the hypothalamus. It extends from the region of the anterior commissure anteriorly and into the mesencephalon posteriorly. Most of the fibers are short, interconnecting nuclei found close to each other, such as the septal nuclei, other nuclei in the basal forebrain, various hypothalamic nuclei, and the PAG in the mesencephalon (see Fig. 31.4). Fibers from the **monoaminergic** cell groups of the brain stem pass through the medial forebrain bundle on their way to forebrain structures, such as the cortex (including the hippocampal formation). Functionally, the medial forebrain bundle is heterogeneous, and lesions of it cannot be expected to reveal the function of any particular cell group or fiber tract.

32 | The Hippocampal Formation: Learning and Memory

OVERVIEW

The hippocampus and nearby areas in the parahippocampal gyrus (the dentate gyrus and the entorhinal area) comprise the **hippocampal formation**, which plays a crucial role for certain kinds of learning and memory. The quantitatively dominating **afferent inputs** to the dentate gyrus and the hippocampus arise in the **entorhinal area**. The entorhinal area receives afferents from both nearby areas in the temporal lobe and cortical association areas. Thereby, it receives highly processed sensory information—that is, about all important events. The hippocampal projection neurons send signals via intermediaries to several areas, notably back to the entorhinal area and to the mammillary nucleus. Thus, the hippocampus acts largely back onto the areas from which it receives information.

Bilateral damage of the hippocampal formation leads to **amnesia**—impaired memory without intellectual reduction. Lesions restricted to the hippocampus proper produce amnesia, although it is much less severe than when the whole hippocampal formation has been damaged.

We can roughly distinguish two kinds of memory: one kind concerns the memory of events and facts and is called **declarative** or **explicit** memory; the other concerns skills and habits and is called **nondeclarative** or **implicit** memory. Only declarative memory depends critically on the integrity of medial parts of the temporal lobe. For nondeclarative memory, the basal ganglia, cerebellum, and parts of the neocortex appear to be most important.

Certainly, the hippocampal formation is not the only part of the brain that is important for memory. Parts of the parahippocampal gyrus outside the hippocampal formation appear to play an independent role in memory formation and retrieval. Further, amnesia has also been reported after lesions of the **medial thalamus**, and cholinergic cell groups of the basal forebrain also have a role (presumably by way of their connections with the hippocampus). Finally, **amygdala** is crucial for learning of associations between stimuli and their emotional value.

The hippocampal formation appears to be important for memory only for a certain time after an event. Thus, isolated damage of the hippocampal formation does

not usually abolish recall of older memories, although it prevents the learning of new material. This is probably because, after a certain time, the memory traces are stored in many parts of the cerebral cortex.

THE HIPPOCAMPAL FORMATION

Macroscopic Appearance and Constituent Parts

The **hippocampal formation** consists of the **hippocampus** and nearby regions in the temporal lobe: the **dentate gyrus**, the **subiculum**, and the **entorhinal area** (located in the parahippocampal gyrus; Figs. 32.1 and 32.2). Whereas the interest formerly was directed mainly to the hippocampus itself, it is now realized that the function of the hippocampus can be understood only in conjunction with the nearby structures in the parahippocampal gyrus, which are closely interconnected with each other and with the hippocampus. Although hippocampus is included among the limbic structures (see Fig. 31.1), its functional role is distinct from that of the amygdala, the septal nuclei, and the cingulate gyrus. The amygdala and the hippocampal formation are, for example, both crucial for learning and memory, but for different kinds.

The **hippocampus** (Figs. 32.1 and 32.3; see Figs. 6.31 and 31.2) forms an elongated bulge medially in the temporal horn of the lateral ventricle, produced during early development by invagination of the ventricular wall by the hippocampal sulcus (see Fig. 9.12). The hippocampus is easily recognized in thionine-stained microscopic sections by its conspicuous layer of pyramidal cells (Fig. 32.2). Along the medial aspect of the hippocampus, the **dentate gyrus** forms a narrow, notched band (Fig. 32.1; see Fig. 31.2). Microscopically, the narrow, dark layer of small granule cells is characteristic (Fig. 32.2). The hippocampus and the dentate gyrus belong to the allocortex and have a simplified laminar pattern compared with the neocortex. Nevertheless, they are far from simply built, with several different cell types and precisely organized, complex patterns of connections. Figure 32.2 shows that the hippocampus (the CA1 field) continues into the **subiculum** but with a marked change in the thickness and organization of layers (both have three neuronal layers). The transition

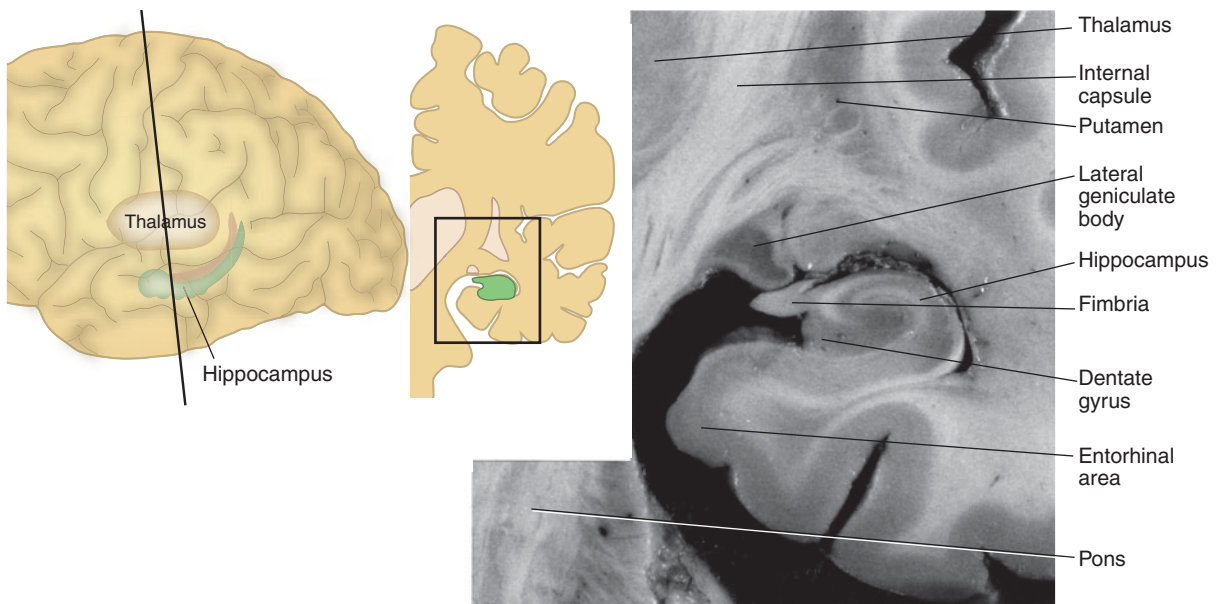


FIGURE 32.1 *The hippocampus.* Photograph of a frontal section through the left hemisphere. The hippocampus forms a continuation of the temporal cortex, as an invagination of the temporal horn of the

lateral ventricle. Figure 22.2 shows the whole section. Compare Figs. 31.1 and 31.2.

between the subiculum and the **entorhinal cortex** is marked with the appearance of a six-layered cortex.

Two Main Sets of Connections

Two aspects of the connections of the hippocampal formation are, presumably, crucial for the understanding of its functional roles: first, the extensive, two-way connections with various cortical association areas and, second, the direct and indirect connections with the amygdala,

cingulate gyrus, and septal nuclei (Fig. 32.4). As for neocortical connections, the hippocampus obviously processes large amounts of information. The parallel increase in the size of the hippocampus and the neocortex during evolution furthermore indicates that its main functions are related to the neocortex. A large number of commissural fibers connect the hippocampus of the two sides, indicating a close cooperation between them.

Afferent Connections of the Hippocampal Formation

The main afferents to the **dentate gyrus** arise in the entorhinal area (Figs. 32.4 and 32.5). Because the dentate gyrus sends its efferents to the hippocampus, the **entorhinal area** is the quantitatively dominating deliverer of information to the hippocampus. Smaller but functionally important contingents to the hippocampal formation come from the **septal nuclei** and **monoaminergic cell groups** in the brain stem (the locus coeruleus and the raphe nuclei). In addition, some fibers come from the **hypothalamus** and several **thalamic nuclei**. Finally, the **amygdala** projects to the subiculum and the entorhinal area. The latter connections most likely contribute to the well-known effect of emotions on learning, as discussed in Chapter 31 (see “Amygdala, Learning, and Unlearning”).

To understand the nature of the information processed by the hippocampus, we must know the afferent connections of the **entorhinal area**. Recent studies with retrograde transport in monkeys have shown that most **association areas** of the neocortex are likely to influence the entorhinal area. Signals reach the entorhinal

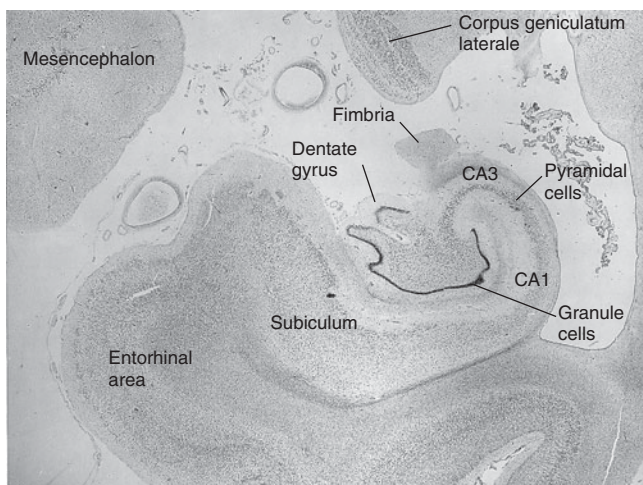


FIGURE 32.2 *The hippocampal formation.* Photomicrograph of thionine-stained frontal section through the human temporal lobe. The temporal horn of the lateral ventricle with some of the choroid plexus is seen above and to the right of the hippocampus. The mesencephalon with the crus and the substantia nigra is seen to the left. Compare with Fig. 32.1.

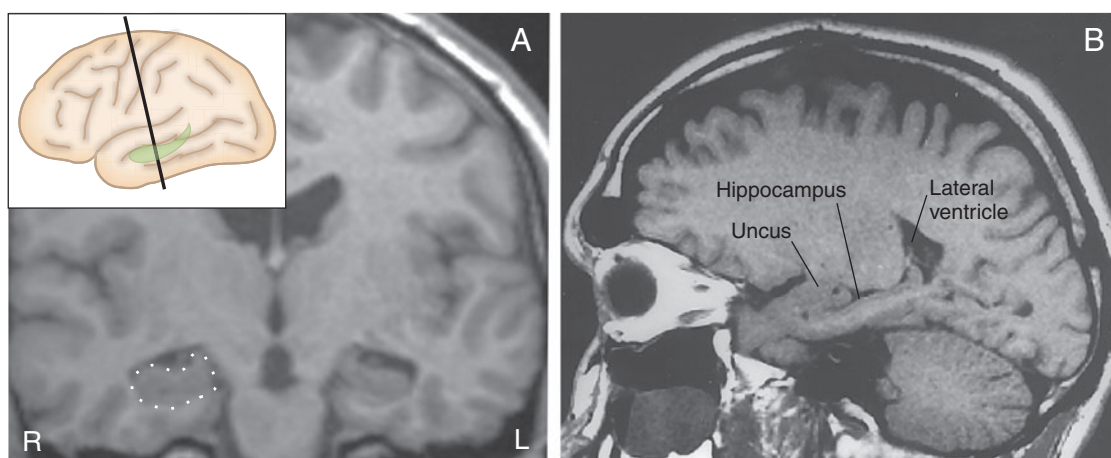


FIGURE 32.3 Magnetic resonance images (MRIs) showing the hippocampus. **A:** Frontal plane. The hippocampus is positioned medially in the temporal lobe (stippled outline on one side). Inset shows approximate position of the hippocampus in the temporal lobe and

the plane of sectioning. **B:** Parasagittal plane through the medial part of the temporal lobe. (Courtesy of Dr. S. J. Bakke, Rikshospitalet University Hospital, Oslo, Norway.)

area directly, or indirectly by means of fibers to other areas in the parahippocampal gyrus, which, in turn, project to the entorhinal area. Thus, the majority of direct entorhinal afferents arise in adjacent parts of the **parahippocampal gyrus** (see Figs. 6.26 and 31.1) and in the **perirhinal cortex** (located around the rhinal sulcus; see Fig. 19.3). These areas form a continuous cortical region, although different parts are not identical regarding connections. Together, the region receives **visual** information from extrastriate areas in the inferior part of the temporal lobe, **auditory** information from the superior part of the temporal lobe, **somatosensory** information from the posterior parietal cortex, and

information from **polysensory** association areas—that is, areas that integrate several sensory modalities. Finally, afferents arrive from the **cingulate gyrus**, the **insula**, and the **prefrontal cortex**. Together, the entorhinal area receives highly processed sensory information. We assume, for example, that information about words comes to the hippocampus regardless of whether we see (read), hear, or read by touch (Braille writing).

The cortical areas that provide the entorhinal area (and thus the hippocampus) with its main inputs project to other areas as well. For example, both the **perirhinal cortex** and the **subiculum** project to the **mediodorsal thalamic nucleus** (MD), which projects to the prefrontal cortex and parts of the cingulate gyrus. This might explain why **amnesia** (memory loss) is more severe after damage of the perirhinal cortex and the areas neighboring the entorhinal area than after a lesion restricted to the entorhinal area and the hippocampus (we return to this point later).

Modulatory Afferents Increase Hippocampal Plasticity

Although hippocampal afferents from the **septal nuclei** (Fig. 32.4) are not numerous, they have an important functional role. Many of the septohippocampal neurons release **acetylcholine** as a modulatory transmitter, increasing the excitability of the hippocampal pyramidal cells. The hippocampal afferents from the **raphe nuclei** (serotonin) and the **locus coeruleus** (norepinephrine) have modulatory effects on the hippocampal cells. Thus, there is evidence that hippocampal **long-term potentiation** (LTP; see later) is reduced after pharmacologic removal of monoamines from the hippocampus. These modulatory pathways may mediate the effects of attention and motivation, which we know have profound influences on both learning and memory.

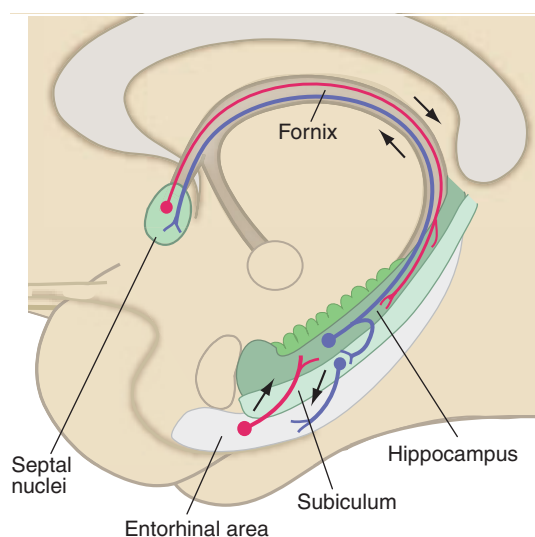


FIGURE 32.4 Some main connections of the hippocampus. The connections with the entorhinal area mediate specific, highly processed sensory information, whereas the connections with the septal nuclei are modulatory.

Efferent Connections of the Hippocampal Formation

Comparison of the efferent connections with the afferent ones shows that the hippocampal formation has largely **reciprocal** connections with subcortical and cortical areas. Thus, we must assume that whatever the hippocampus is doing, it requires a constant exchange of information with many other areas. The parallel evolutionary increase of the hippocampus and the cerebral hemispheres also suggests that hippocampal functions are most closely related to the neocortex.

There are **three parallel pathways** out of the hippocampal formation to cortical areas but they all eventually reach primarily association areas in the temporal and frontal lobes. The major pathway goes from the entorhinal area to adjacent areas in the parahippocampal gyrus and the perirhinal cortex, and from there to more distant areas, notably the tip of the temporal lobe, medial and orbital parts of the prefrontal cortex (including the cingulate gyrus), and polysensory areas in the superior temporal gyrus. A parallel pathway goes directly from the entorhinal area to the same areas that receive the indirect connections, while a third pathway goes directly from the CA1 and the subiculum.

In addition to these neocortical projections, the subiculum sends many fibers to the **mammillary body** (passing in the fornix), which then influences the **cingulate gyrus** via the anterior thalamic nucleus (see Fig. 31.6).¹ Some fibers in the fornix pass to the **nucleus accumbens** and the **ventromedial hypothalamus**. The subiculum also sends fibers to the **amygdala**. Thus, signals from the hippocampus can influence neuronal groups that are related to emotions and motivation.

The connections from the hippocampus to the subiculum, and from the subiculum to other areas, are **topographically organized**. For example, a longitudinal subicular zone close to the dentate gyrus projects to allocortex and nucleus accumbens, whereas a longitudinal zone farther from the dentate gyrus projects to the cingulate gyrus. Thus, as different parts have different connections, symptoms after lesions may be expected to vary with their exact localization within the hippocampal formation. Some controversies among authors regarding the effects of hippocampal lesions on memory are probably due to disregard of such anatomic facts.

1 Until the mid-1970s it was believed that most efferents from the hippocampus were directed to the mammillary body, passing in the fornix (fibers destined for the mammillary nucleus comprise the majority of the fornix fibers). This was based on the erroneous assumption that all axons in the fornix came from hippocampal pyramidal cells. Papez and the postulated circuit interconnecting the limbic structures presumably influenced this view on hippocampal efferents (see Chapter 31, under "The Circuit of Papez"). With the introduction of methods using axonal transport of radioactively labeled amino acids, however, it was shown that the fornix fibers originate in the subiculum.

Hippocampal Architecture and Intrinsic Connections

Most of the neurons of the dentate gyrus are the small **granule cells**, whereas the only well-defined hippocampal cell layer consists of large **pyramidal cells** (Fig. 32.5). Above and below the pyramidal cell layer are layers that contain the pyramidal cell dendrites and incoming axons. There are also various kinds of **interneurons**, notably the GABAergic **basket cells**, which inhibit the pyramidal cells. The hippocampus can be divided into three **longitudinal zones**, named CA1 to CA3 (Figs. 32.2 and 32.4). The granule cell axons contact the pyramidal cell dendrites, and the pyramidal cells send axons out of the hippocampus.

Even though the internal architecture of the hippocampus is rather complicated, with several neuronal types with complex interconnections, a relatively simple main transmission route from input to output appears to exist (Fig. 32.5). This pathway starts in the entorhinal area and has three synaptic interruptions. Neurons in the entorhinal area send their axons, forming the so-called **perforant path**, to the hippocampus, where many end in the dentate gyrus. The axons of the granule cells of the dentate gyrus, called **mossy fibers**, end primarily on the apical dendrites of the pyramidal cells of CA3. The CA3 pyramidal cells send so-called **Schaffer collaterals** to the apical dendrite of the CA1 pyramidal cells. From CA1 a significant part of the signal traffic goes to the subiculum and from there to the entorhinal area, thus closing the circuit that passes from the entorhinal area through the hippocampus and back to the entorhinal area. All links in this pathway are excitatory, using **glutamate** as neurotransmitter. Figure 32.5 gives the impression that signal transmission is confined to a plane perpendicular to the long axis of the hippocampus. Thus, the hippocampus would seem to be organized in numerous **lamellas**, each lamella presumably representing a functional unit.

While the signal pathway shown in Figure 32.5 appears to be central to hippocampal information processing, the conditions are more complex. For example, the fibers of the perforant path not only contact granule cells of the dentate gyrus but also end directly on the hippocampal pyramidal cells. Thus, several parallel pathways enter the hippocampus. Further, anatomic investigations performed in monkeys show that the efferent fibers from a narrow transverse zone of the entorhinal area extend for a considerable distance longitudinally in the hippocampus. Thus, each entorhinal efferent neuron can presumably contact neurons in many hippocampal lamellas. Further, the collaterals of the hippocampal pyramidal neurons extend not only in the plane of the lamellae (as shown in Fig. 32.5) but also longitudinally. In fact, it is still not clear what should be regarded as a functional unit within the hippocampus and the degree of functional localization present.

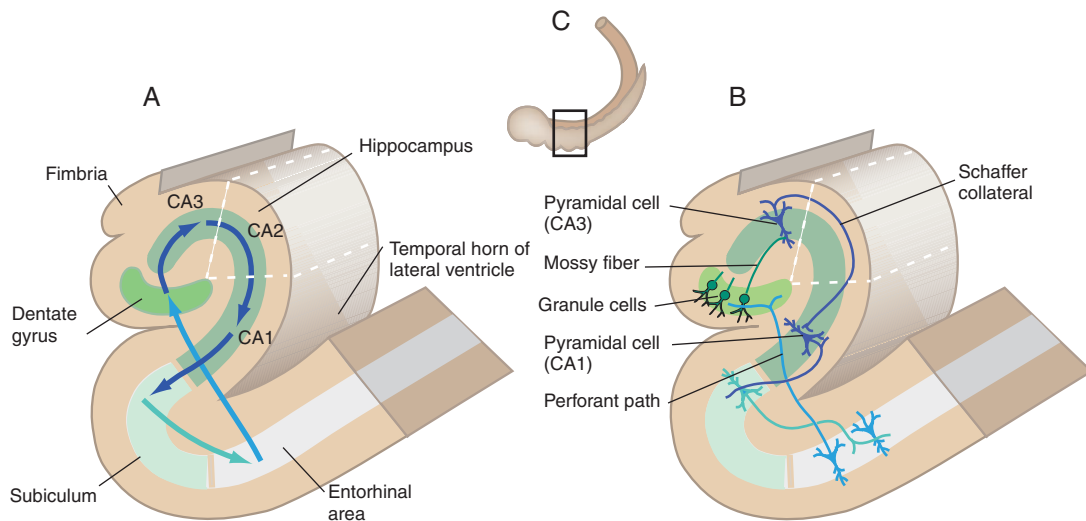


FIGURE 32.5 *Signal pathways through the hippocampus.* Schematics of a frontal slice through the hippocampus, as shown in C (cf. Fig. 31.2). **A:** The flow of signals through the hippocampal formation. The pathway emanates from and returns to the entorhinal area, which has

widespread connections with cortical association areas. **B:** Major kinds of neuron in the hippocampus and the course of their axons (not all collaterals are shown).

FUNCTIONAL ASPECTS

Different Kinds of Learning and Memory

Before discussing further the functions of the hippocampal formation, some words about learning and memory may be useful. The following is a very superficial treatment of a large and important field in psychology. Other classifications than those used here exist, and the different forms of memory may not be as independent as this presentation may suggest.

Much of our learning is **associative**: we learn from experience that two phenomena occur together. This holds for **classic conditioning** (learning of a conditioned response) when one stimulus always follows another, and for **operant conditioning** in which we learn that a certain behavior produces a certain response. An example of operant conditioning is a hungry rat that learns by trial and error to press a lever to obtain food. In the same situation later, the rat knows what the appropriate behavior is. Not all learning is associative, however. With constant exposure to a stimulus, the response is altered—for example, by ceasing to notice a loud sound we have heard many times. This is called **habituation**. The response may also increase with repeated exposure, as with painful stimuli (**sensitization**). Other, more complex, kinds of learning are **nonassociative**, such as learning by **imitation** elements of language and movement patterns. Often, however, learning includes several of these forms at the same time.

Another way to characterize learning and memory relates to **what** is learned, rather than to how it takes place. We can then distinguish two main kinds of memory: **explicit** or **declarative** memory and **implicit**, **procedural**,

or **nondeclarative** memory. Declarative memory depends on the integrity of the medial temporal lobe (including the hippocampus)² whereas nondeclarative memory depends primarily on the basal ganglia, the cerebellum, and parts of the cerebral cortex (see Chapter 22, under “The Motor Cortex and Learning,” Chapter 23, under “What Activates the Nigrostriatal Neurons?” and “Functions of the Basal Ganglia,” and Chapter 24, under “Mossy and Climbing Fibers Mediate Different Kinds of Information” and “The Cerebellum and Motor Learning”).

Episodic, Semantic, and Autobiographic Memory

Declarative memory is of two main kinds. **Episodic memory** concerns episodes and events from one’s own life (“Where did I put the scissors?”). Afterward we can recall the episode, often in the form of pictures, which may be described verbally (memory of what). **Semantic memory** concerns general knowledge about the world we live in (the capital of Italy, the first president of the United States, the location of the post office, the letters of the alphabet, and so forth). **Autobiographic** memory—that is, the ability to remember events from one’s own life—usually combines episodic and semantic memory but also involves memory processes dependent on the

2 One striking piece of evidence comes from patients with **Alzheimer’s disease** (affecting medial parts of the temporal lobe early in the disease process) who have markedly impaired declarative memory whereas the ability to learn new skills is relatively preserved. Also, observations of patients with memory deficits after lesions of the medial temporal lobe (without dementia) indicate that they learn and remember new movements better than new faces, words, places, and so forth.

amygdala (emotional coloring) and several parts of the cortex (Fig. 32.6). Common to declarative memories is that we as a rule must consciously “search our minds” to recall them. In contrast, **nondeclarative** memory is needed to learn and perform **skills** (riding a bicycle, dress, use a knife and fork, etc.). Habits and attitudes also largely fall in this category. The learning leads to altered behavior, but not so that the stored material can be subject to a conscious analysis. With skills like playing an instrument or arithmetic, the stored information becomes accessible only by performing the skill (memory of how). In ordinary teaching situations, much of the learning is implicit—for example, the acquisition of attitudes of which both the teacher and the student are unaware.

Relationship between Memory and Long-Term Potentiation

The cellular basis of brain plasticity is discussed in Chapter 4, under “Synaptic Plasticity”). Because the hippocampus is involved in learning and memory, it is of special interest that the synapses are plastic at several steps in the circuit shown in Figure 31.4. Thus, their efficacy can be increased for a long time after intensive stimulation. Indeed, **long-term potentiation** (LTP) was first discovered in the hippocampus, although it was later found in many other areas of the brain, among them the cerebral cortex and the amygdala. LTP is produced whenever the hippocampal pyramidal cells are subjected to excitatory inputs—for example, from the Schaffer collaterals—while they are in a depolarized state (i.e., caused by another excitatory input). Thus, simultaneous synaptic activation of the cell from two sources can make it “remember,” in the sense that the next time the cell is activated by the same fibers the postsynaptic effects are stronger than earlier. To demonstrate a direct

relationship between LTP and learning is difficult, however. For example, only specific subsets of synapses in widely distributed networks are likely to show LTP in a natural learning situation. To look for such altered synapses would seem like looking for “a needle in a haystack.” Nevertheless, much indirect evidence supports that hippocampal LTP is related to learning and memory. Increased synaptic efficacy has been found in the hippocampus in rats housed in an environment rich in stimuli and challenges (assuming that more learning takes place in this situation than in a standard cage). LTP-like phenomena have also been observed in the hippocampus after specific training situations. Another piece of evidence is that **N-methyl-D-aspartate (NMDA)-receptor** antagonists both prevent induction of LTP and reduce learning and memory in experimental animals. Gene-technological manipulations have produced strains of mice in which hippocampal LTP cannot be induced, and these animals also show reduced learning ability. Finally, structural synaptic changes have been observed in the hippocampus in conjunction with the induction of LTP. Even more compelling, structural synaptic changes (formation of new synapses and splitting of spines into two) have also been reported in the hippocampus in rats at the time they improve their performance in a learning situation.

Medial Parts of the Temporal Lobe Is Necessary for Declarative Memory

The belief that the hippocampus is of importance for memory goes back to the end of the nineteenth century and was based on careful examination of patients with amnesia (loss of memory) as a result of brain damage (Fig. 32.7). In particular, the importance of the medial temporal lobe for declarative memory was strikingly demonstrated in the 1950s by observations of the patient H.M. with bilateral removal of the hippocampus and surrounding regions. During the past few years, refined studies in monkeys with selective lesions, and observations with magnetic resonance imaging (MRI) and positron emission tomography (PET) in humans, have helped clarify the mutual roles of the hippocampus and other subregions of the medial temporal lobe.³

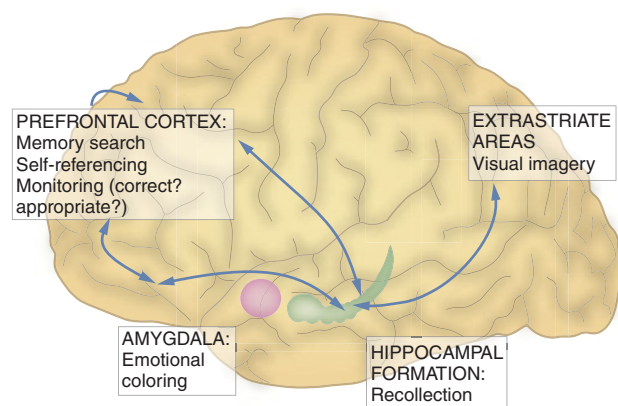


FIGURE 32.6 *Network serving autobiographical memory.* Some of the regions showing increased activity in relation to retrieval of autobiographical memories, as revealed by fMRI. (Based on Cabeza and Jacques 2007.)

³ Various memory tests are used in experiments with monkeys to study the relationship between brain structures and memory. One common test is the so-called **delayed nonmatching-to-sample** test. The monkey is briefly shown an object. After a certain time, the same object is shown with a new one. To receive a reward (e.g., orange juice) the monkey must move the new object, thus showing that it remembers which one was seen before. The experiment goes on with continually new pairs of items. With a brief interval between the first and second presentation, the performance is independent of the integrity of medial temporal structures. With intervals above 10 sec, however, the frequency of errors increases in monkeys with such lesions as a sign of failing memory. With more than 2 min intervals, the performance is no better than chance, whereas normal monkeys reduce their performance to about 80% with intervals of 3 min.

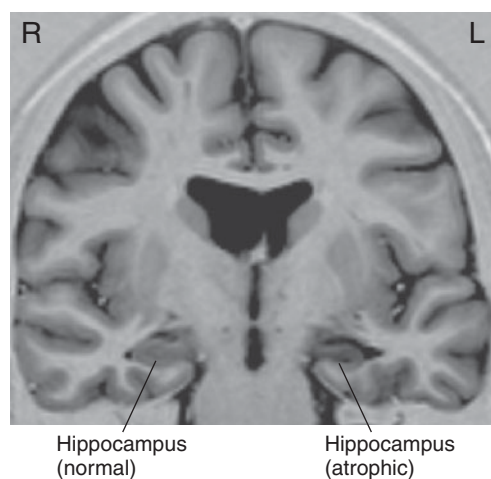


FIGURE 32.7 MRI in the frontal plane showing hippocampal atrophy on the left side. A nearby focus of epileptic activity is most likely the cause of hippocampal cell loss, causing so-called mesial temporal sclerosis. (Courtesy of Dr. S.J. Bakke, Rikshospitalet University Hospital, Oslo, Norway.)

Patients with severe **amnesia** that is not accompanied by intellectual reduction (dementia) typically have lesions that affect the medial parts of the temporal lobe (and often also the medial parts of the thalamus). Such lesions include the hippocampal formation as the most constant finding. Further, testing of patients with restricted lesions of the medial temporal lobe suggested that memory impairments concerned **long-term memory**, whereas **short-term memory** was virtually unimpaired (i.e., the ability to remember, e.g., numbers or words for up to a minute). Nevertheless, these conclusions may not be generally valid. Thus, conclusions were generally based on tests requiring the patients to retain simple motor sequences or short strings of digits or letters. Indeed, when using more natural test situations, requiring short-term memory of faces or locations, patients with restricted medial temporal lobe lesions show reduced performance.

The amnesia is **anterograde**—that is, the memory is lost for events that take place after the time of the brain damage. There are also varying degrees of **retrograde amnesia**—that is, the patient is unable to recall events that took place before the damage. Retrograde amnesia usually occurs together with anterograde amnesia (but some cases have been reported with relatively pure retrograde amnesia alone). Retrograde amnesia is usually graded, that is, it extends from the time of injury and a certain time backwards. Observations of patients with lesions restricted to medial temporal lobe indicate that the retrograde amnesia extends from about one year up to about 20 years. If the amnesia extends further back and includes childhood memories, most studies suggest that there is damage also outside the medial temporal lobe. Not all researchers agree with this view, however, and maintain that lesions of the hippocampal formation

and surrounding areas can produce an amnesia that comprises all prior material without any gradation. Further, other evidence suggests the participation of the hippocampal formation in the retrieval of remote memories. For example, most functional MRI (fMRI) studies agree that the hippocampus increases its activity in relation to recollection of autobiographic memories, independent of how remote they are. As usually in science, the disagreements among authors may depend more on how questions are posed to nature (e.g., different tests and contexts) than on erroneous results.

The Contribution of the Perirhinal Cortex

The most pronounced memory loss occurs when the hippocampal formation, the perirhinal cortex, and areas outside the entorhinal area in the parahippocampal gyrus are destroyed on both sides. Especially, the inclusion of the **perirhinal cortex** appears to increase the severity of amnesia as compared with what occurs after lesions restricted to the hippocampal formation. Thus, the perirhinal cortex and the adjoining cortex of the parahippocampal gyrus appear to contribute to memory not only by their connections with the hippocampal formation but also by their connections to other parts of the brain. For example, the perirhinal cortex projects to the thalamus (MD), amygdala, mammillary body, and parts of the prefrontal cortex. The perirhinal cortex may be uniquely involved in **stimulus recognition memory**.

Amnesia: The Famous Case of H.M.

Patients with bilateral damage to medial aspects of the temporal lobe have pronounced anterograde (and often less severe retrograde) amnesia, as described previously. Scoville and Milner (1957) described the most famous case of this kind in the 1950s. This patient, called H.M. in the voluminous literature that deals with the results of tests to which he had been subjected, underwent surgery to remove bilaterally the medial aspects of his temporal lobes (the purpose was to cure his severe epilepsy). His lesion most likely comprised the hippocampal formation, additional parts of the parahippocampal gyrus, the uncus, and the amygdala. The epileptic seizures became less frequent, but, unfortunately, he acquired severe, permanent anterograde amnesia. Initially, he also had considerable retrograde amnesia that gradually improved to just 1 year before the operation. He remembered well the address of the place he lived before the operation, but he never learned the new address when moving afterward. He could easily recall songs he had learned before the operation but not those he heard for the first time afterward. He never learned where the lawn mower was kept in his new house. Shortly after eating dinner, he could start on a new meal without

remembering that he had just eaten. Nevertheless, his intelligence, as measured with various tests, was unaltered, compared with what it was before the operation, and his capacity for abstract reasoning was normal. Interestingly, his ability to learn new movements was much better than that for learning new faces, words, and so forth. The memory deficits described all concern events that took place some time ago—that is, long-term memory. His short-term memory was not correspondingly impaired, however. For example, he could recognize a word among nine presented to him 40 sec earlier. Once H.M. described his life as follows: “Every day is alone, regardless of the pleasures I have had or the sorrows I have had.” Without long-term memory, we lose the continuity in our lives.

Can Destruction of the Hippocampus Alone Produce Anterograde Amnesia?

Lesions restricted to the hippocampus (in contrast to the hippocampal formation) are difficult to obtain. Nevertheless, recent experiments in monkeys with stereotaxic lesions, or with controlled global ischemia, come close to the ideal situation. Thus, global ischemia of limited duration may kill hippocampal pyramidal cells selectively. Such experiments show that selective hippocampal lesions produce impaired memory in monkeys, although the impairment is much less severe than when other parts of the hippocampal formation are included. The following observations, published by Zola-Morgan and coworkers in 1986, strongly suggest that isolated hippocampal damage produces amnesia in humans, too. The patient, called R.B., had an episode of brain ischemia during cardiac surgery. Afterward, he had moderate anterograde amnesia that lasted until his death some years later. He was not appreciably reduced intellectually. Even though his amnesia was much less severe than that of H.M., he had grave problems remembering the events of the day before. During his visits to his doctor, he would repeat the same story at short intervals. Histological examination of the brain after his death showed that the most pronounced alterations were in the hippocampus and, most interestingly, restricted to CA1 on both sides. Within the CA1, there was an almost total loss of pyramidal cells. That the CA1 field of the hippocampus is particularly vulnerable (to ischemia) was suggested in the nineteenth century based on observations of patients with epilepsy. The mechanism in such cases may be excessive release of glutamate that activates the NMDA receptors (see Chapter 11, under “Ischemic Cell Damage and the Glutamate Hypothesis”). As a possible explanation of the marked symptoms of R.B. caused by a seemingly minor damage to the hippocampus, Zola-Morgan et al. suggest that destruction of the CA1 field (in its entire length) effectively interrupts signal transmission through the hippocampus and thus isolates it from the rest of the brain.

Making Memory Traces Permanent: The Hippocampal–Cortical Dialogue

As mentioned, the amnesia after lesions of the hippocampus and surrounding structures is predominantly anterograde (although retrograde amnesia for several years may occur). This has been taken as evidence that representations of events are only temporarily stored in the hippocampus, before permanent storage in other parts of the brain. Other clinical observations of patients with damage in various parts of the brain indicate that there is no specific “memory center,” but well-consolidated information is stored in a distributed fashion. When a memory is consolidated—presumably as long-term synaptic changes—the hippocampal formation no longer seems necessary for storage and recall. The time required to reach this stage is not known, but based on observations of patients with damage to medial parts of the temporal lobe it probably takes a year or more. During this period, we imagine a gradual **consolidation** of the memory traces in the relevant parts of the cortex (and probably in subcortical structures). It seems unlikely that the information is first held in the hippocampus for a certain period and then transmitted out to the permanent stores in a finally processed form. More likely, the consolidation takes place by a continuous **dialogue** between the hippocampus and other parts of the cortex. **Sleep** may be of special importance for the dialogue, as witnessed by specific patterns of synchronized activity in the hippocampus and in cortical areas that were particularly active during the learning phase (see Chapter 26, under “Dreaming”). Further, it appears that every time a memory is recalled it becomes **labile** and can be modified before renewed consolidation occurs.

As discussed previously, however, it is not quite clear whether the hippocampus ever ceases entirely to participate in storage and retrieval of episodic memories (most would agree that retrieval of semantic knowledge—much of it acquired during childhood—becomes independent of the hippocampus and nearby regions).

Amnesia Caused by Lesions Outside the Medial Temporal Lobe, and Korsakoff’s Syndrome

Clinical observations suggest that amnesia can arise after lesions of the **medial thalamus**, the **mammillary body**, and the connections from the latter to the thalamus (the **mammillothalamic tract**). Loss of memory also has been reported after lesions of **cholinergic cell groups in the basal forebrain**. Whether isolated damage to any one of these structures causes amnesia is not finally settled, however. As to the relation between the site of a lesion and ensuing functional disturbances, PET studies of glucose uptake in a group of patients with severe amnesia are illuminating. These patients all showed altered glucose uptake in the hippocampus,

thalamus, cingulate gyrus, and ventral parts of the prefrontal cortex, although their lesions as identified with MRI were differently placed. It seems reasonable to conclude that declarative memory, as usually tested, depends on the integrity of several cell groups that are mutually interconnected. When one part of the network is damaged, it has consequences for the functioning of the other parts. This illustrates the problems inherent in assessing the functional role of a cell group or a tract only from the symptoms caused by their destruction.

As to **thalamic lesions**, inclusion of the MD nucleus (see Fig. 33.8) seems necessary to cause amnesia, although destruction of the anterior nucleus and the internal medullary lamina close to the MD may contribute (perhaps due to interruption of fibers to and from MD). Severe amnesia was described in a patient who, as judged from a computed tomography (CT) scan of the brain, had a small lesion confined to the anterior and medial parts of the thalamus on the left side. (In this particular case, the verbal memory was more severely affected than the visual. The patient could with some difficulty remember objects he had seen some time ago, whereas words heard were completely forgotten.) The MD sends efferents to the prefrontal cortex, and this is involved in various aspects of memory (cf. Chapter 34, under “Frontal Association Areas”).

The role of the **mammillary body** in amnesia has attracted much interest because of its involvement in **Korsakoff's syndrome**, in which the patient suffers from severe anterograde and retrograde amnesia (among other cognitive impairments). Cell loss is also found regularly in the medial thalamus, however. Accurate testing of patients with Korsakoff's syndrome shows that they also have symptoms suggestive of prefrontal dysfunction. The disease is usually due to chronic alcohol abuse. The retrograde amnesia may be very pronounced—for example, more than 25 years. This in itself suggests that the pathology in these patients does more than interrupt connections between the medial temporal lobe and other parts of the cerebral cortex. Whether a lesion of the mammillary bodies alone causes amnesia has been a matter of controversy. In monkeys, however, bilateral destruction of the mammillary bodies produces a moderate memory loss. Further, amnesia has been reported after bilateral interruption of the **fornix** in epileptic patients (cutting the pathway from the hippocampal formation to the mammillary body). In any case, it seems unlikely that the memory traces are stored in the thalamus or the mammillary body.

Amnesia with Confabulation

A peculiar form of amnesia occurs together with **confabulation**: that is, the patient invents stories (without knowing that they are not real). Most of these patients have a lesion involving the substantia innominata, the medial hypothalamus, and the orbitofrontal cortex

(usually caused by a ruptured aneurysm of the anterior cerebral artery). The often-bizarre stories can usually be traced back to real events, although they consist of various, unrelated fragments from memory. It seems that the patient is unable to suppress irrelevant associations and cannot check them against reality. These are faculties usually associated with the prefrontal cortex.

What Is the Unique Contribution of the Hippocampus to Learning and Memory?

There is little doubt that the hippocampus is of crucial importance for certain kinds of learning and memory, as discussed earlier. Nevertheless, its specific contribution is not entirely clear, and neither is the division of labor between the hippocampus and the other components of the hippocampal formation.⁴ One central task of the hippocampus—in animals and humans—seems to be **spatial orientation** and **navigation**. Indeed, imaging studies of London taxi drivers suggested correlation between the size of the hippocampus and duration of navigational training (animal experiments have found the same effect). Properties of single hippocampal cells, as first described by O'Keefe and Nadel (1978), are of particular interest in this connection. Thus, the firing of single hippocampal neurons—called **place cells**—changes with the position of the animal in relation to its surroundings: for example, the firing pattern changes with the location of the animal in different corners of the cage. The hippocampus obviously receives information about starting position, the direction of movement, and the distance moved. Based on such experiments, O'Keefe and coworkers proposed that the hippocampal neurons together form a **cognitive map** of our surroundings. Necessary information may be integrated in the entorhinal cortex, where neurons are topographically arranged according to their spatial receptive fields. Together, such **grid neurons** produce a systematic map of the surroundings. Indeed, the activity of a few neurons can code for the position of a rat with a few centimeters accuracy. Accordingly, lesions of the hippocampus in rats severely reduce their ability to find their way back to previous locations. Further, monkeys with lesions of the hippocampal formation have difficulties with remembering where an object was located—the association between **objects** and **space**.

4 Studies of the development in children with early damage to the hippocampus shed some light on these questions. Three children suffered hippocampal lesions without signs of damage to the surrounding cortical regions at birth, and at the age of four and nine, respectively. They were examined at the age of 14, 19, and 22 years. As expected, they suffered from anterograde amnesia. However, the amnesia was much more severe for episodic than for semantic memory. Their language, reading abilities, and general knowledge were a little below average for their age groups but they followed ordinary school with normal progression (Vargha-Khadem et al. 1997). This suggests that the hippocampus is necessary for episodic memory, whereas areas around the hippocampus can take care of semantic memory in the absence of the hippocampus.

One of the great challenges in understanding the hippocampus is to combine its role in spatial navigation and spatial memory with its undeniable importance for declarative memory. Recent animal experiments suggest that both tasks may be carried out simultaneously: neuronal populations may at the same time signal *where* something happens and *what* is going on. Such a double role might—loosely considered—fit with the everyday experience that recall depends strongly on **context**. For example, experiments with divers show that a series of numbers learned under water is better recalled under water than on dry land. Similarly, when unable to recall why we went into a room, it may help to go back to the place where we first got the idea to go into the room.

“Knowledge Systems of the Brain”

In a wide sense, memories represent our knowledge of the world and the actions that are necessary to relate

successfully to it. Not all this knowledge is accessible for conscious “inspection” and analysis (nondeclarative or implicit memory, as described). Nevertheless, all memories presumably have as their substrate synaptic changes in specific parts of the central nervous system. Damasio and Tranel (1992) use the term “knowledge systems of the brain” about the widespread networks dedicated to specific tasks. Examples are knowledge systems dealing with social interactions, faces, objects, language, or ourselves. Although a part of the temporal lobe cortex is particularly important for recognition of faces, this does not necessarily mean that all information about faces is stored there. More likely, this part is unique because it has access to face-related information stored in many other areas. Presumably, the memory of faces fails after a stroke in the temporal lobe because there is “no one” to retrieve and process all the relevant information, not because the storehouse is empty.

VIII

THE CEREBRAL CORTEX

THE human cerebral cortex consists of about 20 billion neurons and constitutes more than half of all gray matter of the central nervous system. This gives a rough impression of its functional importance. In lower vertebrates, there are only very modest primordia of the cerebral cortex (allocortex), and it is first in mammals, particularly anthropoid apes and humans, that the cerebral cortex comes to dominate the rest of the nervous system quantitatively. This enormous increase in the volume of the cortex has necessitated a marked folding of the surface of the hemisphere. The outer layer of the human cerebral cortex is around 0.2 m^2 , but only one-third of this is exposed on the surface. The **neocortex** constitutes most of the cerebral

cortex in higher mammals, and this part deals only with the neocortex. In Chapter 33, we deal mainly with the structure and connections of the cerebral cortex from a functional perspective. In Chapter 34 we discuss the complex tasks attended to by the cortical association areas, which make up the bulk of the human cerebral cortex. In previous chapters we treated the role of the cerebral cortex in sensory processes (Chapters 14–19), control of body and eye movements (Chapters 22 and 25), autonomic functions and emotions (Chapter 31), and memory (Chapter 32). The role of the cerebral cortex in relation to sleep and consciousness is briefly discussed in Chapter 26.

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The Cerebral Cortex: Intrinsic Organization and Connections

OVERVIEW

In this chapter, we address two levels of organization. The first concerns information processing in a small volume of cortex; the second level concerns the interconnection of functionally different cortical units with long association and commissural fibers. These connections are essential parts of distributed, task-specific cortical networks.

The neurons of the neocortex are arranged in **six layers** parallel to the cortical surface. The layers differ with regard to afferent and efferent connections. In general, layers 2 and 4 are receiving, whereas layers 3 and 5 are mainly efferent. The bulk of afferents from the thalamus end in lamina 4, whereas layer 5 gives origin to subcortical tracts to the cord, brain stem, and basal ganglia. Layers 2 and 3 receive and send out most of the corticocortical fibers, interconnecting various parts of the cortex.

The cortex is divided into numerous **cytoarchitectonic** areas that differ with regard to connections and functional specializations. At a microstructural level, the cortical neurons are arranged in smaller **modules**, often in the form of **columns** perpendicular to the cortical surface. Each column, containing some thousand neurons, represents a computational unit. The neurons of the columns communicate with each other and with neurons in neighboring and distant columns.

There are two main kinds of cortical neuron: the **pyramidal cells** with long axons destined for other parts of the cortex or subcortical targets, and **interneurons** (with numerous subtypes) with short axons remaining in the cortical gray matter. The pyramidal cells are **glutamatergic**; whereas all interneurons are **GABAergic** (γ -aminobutyric acid [GABA] is colocalized with different combinations of neuropeptides). In addition, **modulatory transmitters**, such as dopamine and acetylcholine, regulate the cortical excitability level and the signal-to-noise ratio of cortical neurons.

Cortical connections fall into four groups. One group of afferents consists of precise, topographically organized connections from the **specific thalamic nuclei**; each thalamic nucleus supplies one particular part of the cortex. Another group consists of diffusely organized connections from the **intralaminar thalamic nuclei**

and several other **subcortical nuclei** (releasing modulatory transmitters). The two final groups—making up the majority of all cortical connections—consist of corticocortical fibers; that is, **association fibers** and **commissural fibers**. Association fibers are precisely organized connections linking cortical areas within the same hemisphere, while commissural fibers pass in the corpus callosum and connect areas in the two hemispheres. The **efferent** connections of the cerebral cortex can also be divided into **subcortical** and **corticocortical** ones (association and commissural connections). The subcortical fibers are destined for the thalamus, the striatum, various brain stem nuclei (among them the pontine nuclei projecting to the cerebellum), and the spinal cord. The corticocortical connections are for the most part **reciprocal**—that is, an area receives fibers from the same areas to which it sends fibers.

STRUCTURE OF THE CEREBRAL CORTEX

Levels of Organization

To understand the relationship between the performance of the cerebral cortex and the underlying neural processes, we need to address two levels of organization. The first concerns information processing in a small volume of cortex; we may term this **intracortical** synaptic organization. The second level concerns the interconnection of functionally different cortical units with long association and commissural fibers, which we term **interareal** synaptic organization. How are signals arriving in a small bit of the cortex treated before “answers” are sent to other parts of the cortex or subcortical nuclei? How are the many different cortical areas interconnected to form networks designed for solving specific tasks? The enormous number of neurons and synaptic couplings in the human cerebral cortex explain why we do not have complete answers to such questions. For example, about 100,000 neurons reside below 1 mm² of cortical surface (rodent SI), and each neuron receives synaptic contacts from at least some hundred other neurons. Further, the tasks of the cerebral cortex are the most complex of all. To understand the relationship between the “machinery” of the

brain and mental functions such as personality, memory, thought, and feelings is indeed the most formidable and exciting challenge of modern neuroscience.

The Neocortex Consists of Six Layers

All parts of the neocortex have a common basic structure, with the neurons arranged in six layers, or **laminae**, oriented parallel to the surface of the cortex (Figs. 33.1 and 33.2). Another general feature is the arrangement of the neurons in rows or **columns** oriented perpendicular to the cortical surface (Fig. 33.2). Both kinds of cellular aggregation relate to functional specializations among the neurons, as we discuss below. Figure 33.1 shows the main features of the layering. It can be seen that the laminar pattern arises because cells of similar shape and size are collected in more or less distinct layers. The density of cell bodies also differs among the layers.

About two-thirds of the neurons are cortical **pyramidal cells** (the name refers to the triangular shape of their cell bodies). A typical pyramidal cell has a long **axon** arising from the base of the pyramid and a long **apical dendrite** that extends toward the cortical surface; it thus extends through several layers superficial to the

layer in which the cell body is located (see Fig. 1.1). The large pyramidal cells lie in layers 3 and 5, but many of the smaller cells in the other layers are also pyramidal. The large number of **dendritic spines** further characterizes the pyramidal cells (see Fig. 1.1). The rest of the cortical neurons constitute a heterogeneous group whose neurons have in common that their cell bodies are not pyramidal; such neurons are therefore lumped together as **nonpyramidal cells**. Their shape and size vary considerably, but all of them are most likely interneurons.¹

The most superficial cortical layer, **layer 1**, the **molecular layer**, is rich in fibers but has few neurons (Figs. 33.1 and 33.2). Apart from axons, it contains the apical dendrites of pyramidal cells in the deeper layers. **Layer 2**, the **external granular layer**, contains densely packed, small cell bodies.² **Layer 4**, the **internal granular layer**,

1 Some of the nonpyramidal cells are multipolar and are called **stellate cells**. Others are called **basket cells** because their axonal branches form a wickerwork around the cell bodies of pyramidal cells. In addition, several other varieties of interneurons are given names that, as a rule, reflect the shape of the neuron.

2 Formerly, the term **granule cell** was used of the small cortical neurons, explaining the names of layers 2 and 4, which contain mainly small cell bodies. Many of the so-called granule cells are, in fact, small pyramidal cells, and the term “granule cell” should therefore not be used of cortical neurons.

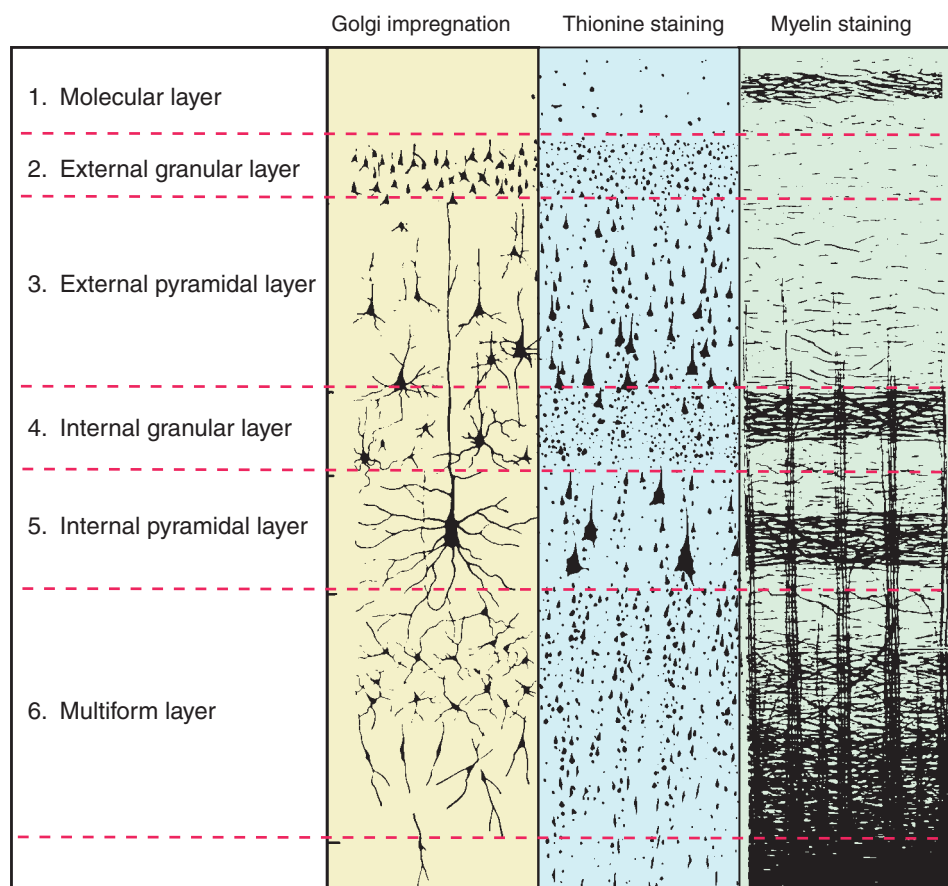


FIGURE 33.1 The basic six-layered structure of the neocortex. The three columns show sections perpendicular to the cortical surface subjected to different staining methods. **Left:** Appearance in Golgi-impregnated sections, in which the cell bodies and some of the dendrites can be seen. **Middle:** A thionine-stained section, in which only the cell bodies are visible. **Right:** Appearance after myelin staining—that is, the main pattern of the myelinated axons is evident, with perpendicular bundles of fibers entering the cortex, and horizontal bundles of fibers interconnecting nearby parts of the cortex. Compare Figs. 33.6 and 33.7.

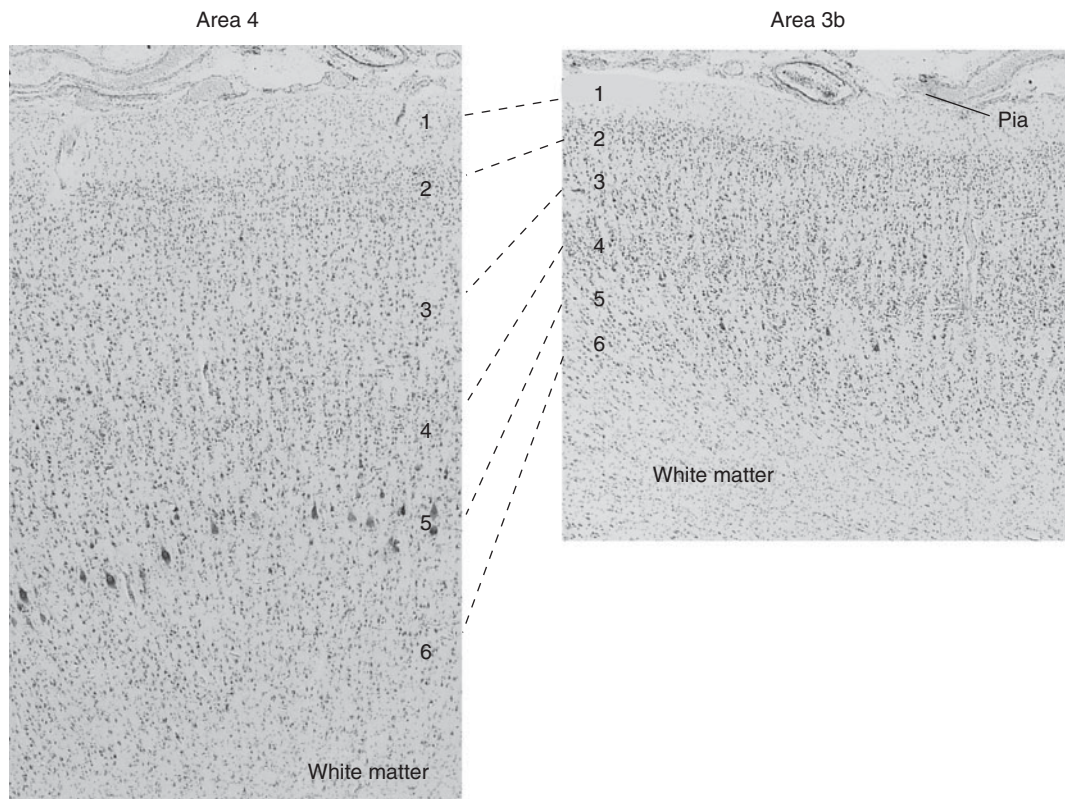


FIGURE 33.2 *Cytoarchitectonics of the cerebral cortex.* Photomicrograph of a thionine-stained section through the central region of the human brain. The section is perpendicular to the direction of the central sulcus. The six-layered structure is evident in area 4 (MI) and in area 3b (SI), but development and appearance of the various layers are different in the two areas, especially with regard to layers 4 and 5. The motor cortex in the precentral gyrus is much thicker than the

somatosensory cortex. This is not because there are more neurons but because of more extensive dendritic trees, more axonal branches and boutons, and perhaps more glial cells. There is a tendency for the cells to be arranged in vertically oriented rows or columns. See also Fig. 12.3 with a corresponding section from the monkey. Magnification, $\times 170$.

is similarly built. In primary sensory cortical areas, layer 4 is especially well developed. In the striate area, lamina four exhibits a further laminar subdivision (layer 4A, B, and C; see Fig. 33.5). **Layer 3, the external pyramidal layer**, can be recognized by its content of medium-sized pyramidal cells, whereas **layer 5, the internal pyramidal layer**, also contains many large pyramidal cells. **Layer 6, the multiform layer**, contains many cells with spindle-shaped cell bodies.

Considering only certain salient features, we can say that layers 2 and 4 are mainly **receiving**—being most developed in the primary sensory areas—whereas layers 3 and 5 are mainly **efferent** and send their axons out of the part of the cortex in which they are located. The pyramidal cells of layer 5 send their axons primarily to **subcortical nuclei**, and this layer is particularly well developed in the motor cortex (MI) (Fig. 33.2; see Fig. 22.4). The layer 3 pyramidal cells send their axons primarily to other areas of the **cortex** (association and commissural fibers). Layer 6 is also largely efferent and sends many axons to the **thalamus**.

Cortical Microstructure: Columns

We mentioned that the cortical neurons appear to be organized into small units formed by cylinders or **columns** of tissue extending perpendicular to the surface of the cortex. A tendency for cortical neurons to be arranged in perpendicular rows is evident from thionine-stained sections (Fig. 33.2). Figure 33.3A gives a simplified three-dimensional view of a column. Vernon B. Mountcastle first observed the fact that neurons within such a column share functional properties that differ from those in adjacent columns (in the monkey SI). He found that when a microelectrode was inserted perpendicular to the cortical surface, the cells encountered as the electrode traversed the thickness of the cortex had similar receptive fields. They were furthermore similar with regard to their stimulus specificity. In contrast, when the microelectrode was inserted obliquely, the receptive fields of the neurons moved with the advancement of the electrode. Based on such experiments, it was proposed that neurons in the somatosensory

cortex (SI) with similar receptive fields and modalities are grouped together in columns with a diameter of some hundred micrometers. A similar columnar arrangement of neurons has been observed in the motor cortex (MI), but with respect to muscles rather than to receptors (neurons within one column act on one or a few synergistic muscles). Interestingly, a pyramidal cell with all its dendrites and recurrent collaterals is contained within a cortical tissue cylinder with a diameter of about 350 μm (Fig. 33.3A). Thousands of other neurons are present within the same cylinder.

Modular Organization of the Cerebral Cortex

Module is a more general term for columns and other assemblies of neurons that share salient properties (in the cortex and elsewhere; see Chapter 16, under “Modular Organization of the Visual Cortex”). It is striking how connections and cytochemical markers show a **patchy distribution** all over the cortex, strongly suggesting some kind of modular pattern as a basic principle in the organization of the cerebral cortex. For example, afferent projections—from the thalamus or from other parts of the cortex—end in many, regularly spaced patches in the cortex rather than continuously (Fig. 33.3B). This means that inputs from different sources may converge systematically on cortical neurons. Similarly, projection neurons with a common

target tend to be lumped together. There is, furthermore, a striking correspondence between the diameters of cortical dendritic trees and of the terminal patches formed by afferent fibers.

It may not be feasible to find a definition that fits modules in all parts of the cortex, however. Indeed, different criteria are used to define modules in the cortex, and, depending on the criteria used, their shape and size vary greatly. Further, the distribution neurons with different properties is less schematic than the modular concept might imply. In the visual cortex, in which the segregation of functionally different neurons has been most thoroughly investigated, cells sharing functional properties are arranged in **bands** rather than in cylinders (see Fig. 16.23). Another problem with the columnar concept is that neurons in different layers—for example, of the striate area—are not functionally identical (e.g., cells that are color-specific and cells that are movement-specific are located in different layers). Thus, the modules may not always extend through the depth of the cortex but may be limited to one or a few layers (cf. color-specific “blobs” in the striate area; see Fig. 16.24). To apply the columnar concept to such cases confuses rather than clarifies.

The **biologic significance** of modular organization in the cortex and elsewhere (e.g., in the striatum and the cerebellum) is not fully understood. The modular pattern gives each cortical neuron a varied afferent

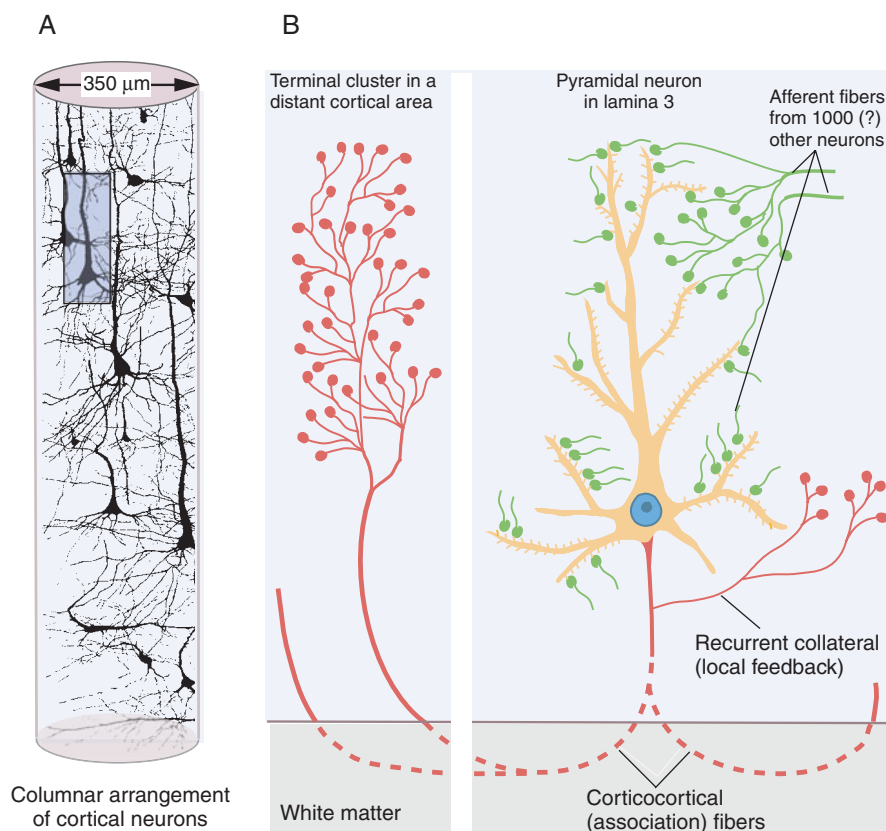


FIGURE 33.3 Columnar arrangement of cortical neurons and clustering of cortico-cortical terminal areas. **A:** A cylindrical volume of cortical tissue forms a functional unit (column), for example, in the somatosensory cortex. Golgi-impregnated neurons (from Conel 1939) show position of cell bodies and the main orientation of the dendrites. Note, however, that only a fraction of the total neuronal population within a column is shown in the figure. **B:** Schematic. A pyramidal neuron in cortical layer 3 (inset in A) and its main connections: recurrent collaterals and corticocortical fibers. A patch of termination in a distant cortical area is also shown. Each neuron forms many such patches in several areas. The number of neurons forming synapses on each layer 3 pyramidal neuron is indicated.

input, with each neuron receiving samples of information from several sources (several patches contact each neuron). In the motor cortex, for example, a neuron might need to integrate information from the thalamus (perhaps from different subgroups in the VL nucleus), from SI, and from the premotor cortex. Another advantage of modular organization may be that it permits shorter intracortical connections and thus saves nervous tissue. Further, modules may be advantageous during development by making it easier for growing axons to find their target. Some claim, however, that modular organization is merely a byproduct of how the nervous system develops and that it lacks an intrinsic functional meaning.

Afferents from Different Sources End in Different Cortical Layers

The cortical layers differ with regard to the origin of their extrinsic afferents. Thus, there is some degree of specialization among the laminae with regard to what kind of information they process. As we discuss later, however, connections between the laminae ensure integration of their outputs. Schematically, pathways that convey **precise sensory information** end primarily in **layer 4**. This includes thalamocortical fibers from the somatosensory relay nucleus, VPL, and from the relay nuclei of the visual and auditory pathways, the lateral and medial geniculate bodies. **Association fibers** (i.e., from other cortical areas) end preferentially in **layers 2 to 4**. Subcortical afferents with **modulatory effects** end in several layers but especially in **layer 1**. Such fibers arise in the intralaminar thalamic nuclei, in several brain stem nuclei (the raphe nuclei, nucleus locus coeruleus, and dopaminergic cell groups in the mesencephalon), and in the basal nucleus (acetylcholine).

The Cerebral Cortex Can Be Divided into Cytoarchitectonic Areas

Even though all parts of the neocortex consist of six cell layers, the thickness and structure of the various layers vary from one area to another. This is observed most easily in sections stained to visualize the cell bodies only (Figs. 33.1, 33.2; see Fig. 22.4). Such **cytoarchitectonic** differences form the basis of the subdivision of the entire cortex into cytoarchitectonic areas, as done around the turn of the century by Brodmann (1909) and others (Fig. 33.4). This is briefly described in Chapter 6, and several of the cytoarchitectonic areas are mentioned in previous chapters. The development of cytoarchitectonic areas is briefly discussed in Chapter 9 (“Specification of Cortical Cytoarchitectonic Areas”).

The main importance of the division of the cortex into cytoarchitectonic areas is that these areas have

proved in many instances to differ functionally, even though they were initially defined solely on the basis of the size, shape, and arrangement of the neuronal cell bodies. In many cases, a cytoarchitectonically defined area is unique with regard to its afferent and efferent connections and the physiological properties of its cells.

Although the borders between different cytoarchitectonic areas are sometimes easy to identify, as exemplified in the section of the visual cortex shown in Fig. 33.5, more often the differences are rather subtle. Thus, it should not come as a surprise that authors often disagree with regard to the parcellation of the cortex into areas. Today we try to define a cortical area not only based on cytoarchitectonics but also by additional criteria, such as fiber connections, cellular markers, recordings of single-cell activity, and the behavioral effects of stimulation or ablation of the area in question.

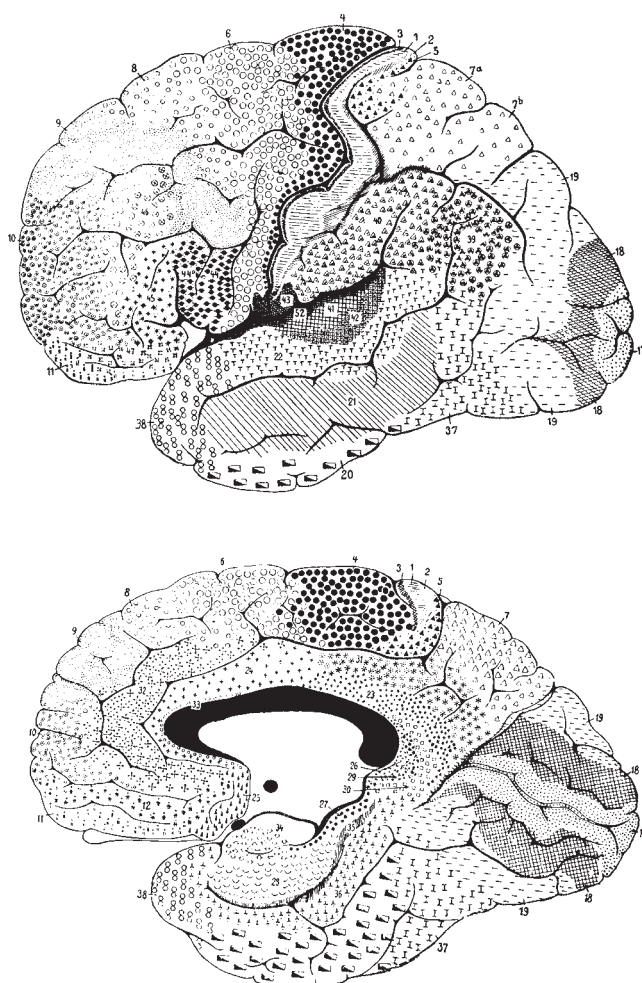


FIGURE 33.4 Brodmann's cytoarchitectonic map of the human brain. The various areas are labeled with different symbols and numbers. (From Brodmann 1909).

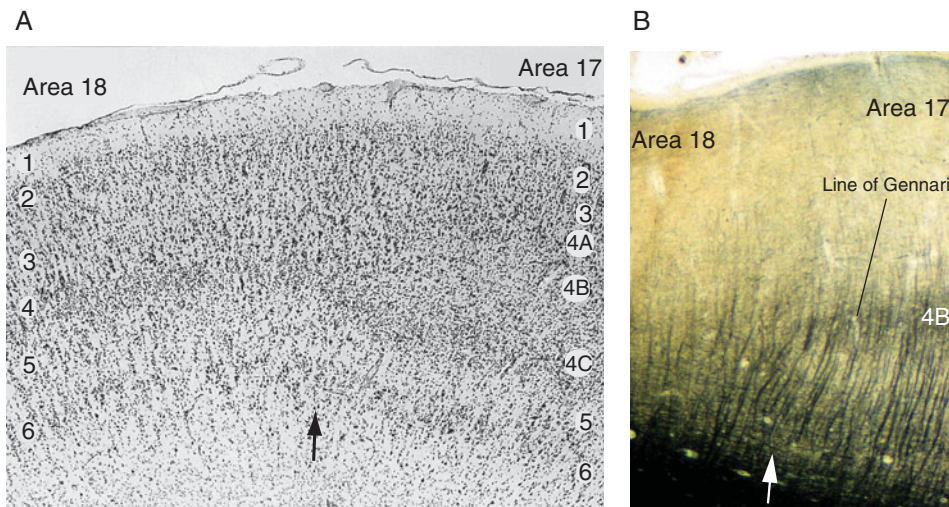


FIGURE 33.5 *The transition between area 17 (the striate area) and area 18.* **A:** Thionine-stained section from the human visual cortex. The various layers change in cell size, cell density, and thickness at the transition between the two cytoarchitectonic areas. Lamina 4 is particularly well developed in area 17, and is subdivided into sublayers (4A–4C). **B:** Section from the same region stained to show myelinated

nerve fibers. Note the myelinated fibers running parallel with the cortical surface in lamina 4B (the line of Gennari). See Fig. 16.17, which shows the macroscopic appearance of the line of Gennari. Most areal borders are less clear cut than the border between areas 17 and 18.

We return to connections and functions of different cortical areas later in this chapter.

With regard to the **size of cortical areas**, there are surprisingly large **individual variations**. For example, the volume of the striate area (the most easily identified one) varies by a factor of three among adult humans, and similar differences have been documented for the somatosensory cortex, the auditory cortex, and prefrontal cortical areas. Because the volume of the hemispheres does not show similar variations, this implies that in a brain with a large striate area, other areas are relatively smaller. Whether such anatomic differences also have functional significance is unknown, but conceivably, they may contribute to the large differences between humans in mental and other capacities.

Intracortical Connectivity: Interneurons and Pyramidal Cell Collaterals

The **pyramidal cells** send their axon toward the white matter to reach other parts of the cortex or subcortical cell groups. Thus, they are the **projection neurons** of the cortex (Fig. 33.3B and 33.6), which most likely constitute more than two-thirds of all cortical neurons. The projection neurons send **recurrent collaterals** before the axon leaves the cortex and can thus influence the level of activity among the cortical neurons in their vicinity. The recurrent collaterals may, for example, excite inhibitory interneurons and thereby limit the activity of the parent cell and other pyramidal cells.

All cells belonging to the other main type of cortical neurons, the **nonpyramidal cells**, have locally branching axons that do not reach the white matter—that is, they are the cortical **interneurons** (Fig. 33.7). Such interneurons may be classified into **three main types** on the basis of the course and branching pattern of their axons. The first type has an axon that forms numerous terminal branches close to the cell body; it mediates influence mainly to neighboring neurons within the lamina in which the cell body is located. The second type has an axon coursing perpendicularly or vertically toward either the cortical surface or the white matter, giving off collaterals on its way; this enables the interneuron to influence neurons in several layers. The third type sends its axon in a horizontal direction (parallel to the cortical surface).

Even though this division into three kinds of cortical interneurons is an oversimplification, it shows the main features of the intracortical connections, which permits interactions among neighboring neurons, among neurons within different layers, and among neurons located at some distance within the same layer. **Vertically oriented axons** (from either interneurons or pyramidal cell recurrent collaterals) ensure the communication among neurons within a narrow cortical cylinder or **column** (Fig. 33.3A). The **horizontal** axonal branches mediate communication among neurons in different columns. Such horizontal influences can be inhibitory or excitatory. In sensory cortical areas, the horizontal intracortical connections mediate **lateral inhibition**, which

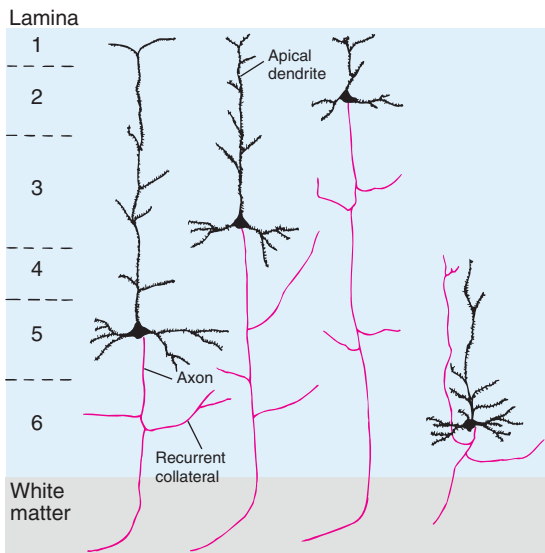


FIGURE 33.6 *Cortical projection neurons.* Schematic. The axons (red) give off several recurrent collaterals on their way to the white matter. The dendrites have numerous spines. The largest neurons, with the largest cell bodies and the thickest axons, are located in layer 5. Drawing based on studies by Jones (1988) of the central region of the monkey with the Golgi method.

increases the spatial resolution of the sensory information (lateral inhibition is present at several levels of the sensory pathways; see Fig. 13.4). They may also contribute to cortical **plasticity** by modulating the size of the receptive fields of cortical neurons.

The majority of horizontal intracortical axons appear to be collaterals of pyramidal cells rather than of interneurons. They are generally longer in association

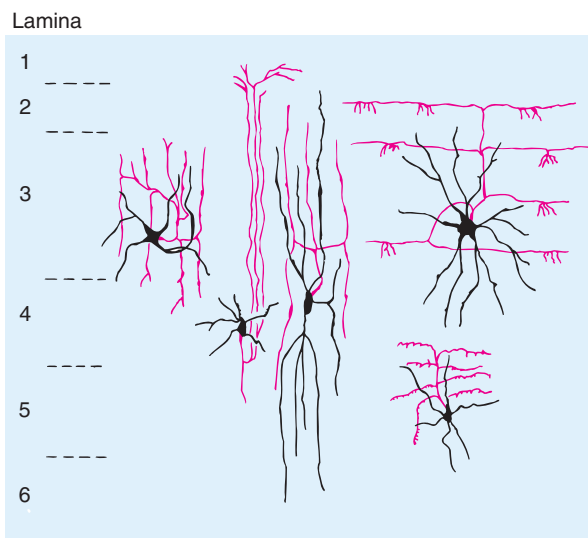


FIGURE 33.7 *Main kinds of cortical interneurons.* There are three patterns of axonal distribution (in blue): in the immediate vicinity of the cell body, horizontally in the layer of the cell body, and vertically spanning several layers. (Based on Jones 1987.)

areas than in primary sensory areas—measuring up to 9 mm in monkey posterior parietal cortex, although the majority of them are probably less than 1 to 2 mm. Thus, cortical columns farther apart than this must communicate by means of projection neurons with an axon coursing in the white matter—that is, association fibers (Fig. 33.3B). Such connections are discussed later in this chapter.

Fast Excitation and Inhibition in the Cortex: Glutamate and GABA

Most cortical neurons receive inputs from the thalamus and other parts of the cortex with fast, **excitatory** synaptic actions (**glutamate** acting on amino-methylisoxazole propionic acid [AMPA] receptors) and fast **inhibition** from interneurons (**GABA** acting on GABA_A receptors). Such synapses are responsible for precise and specific information transfer in topographically organized connections. In addition, cortical neurons are influenced by slow, **modulatory** synaptic actions (we return to that later). Most or perhaps all **projection neurons**—those sending their axons to subcortical nuclei and those forming association or commissural connections—use **glutamate** as a neurotransmitter. Accordingly, physiological studies show that they have fast, excitatory synaptic actions (however, there are also metabotropic glutamate receptors in the cortex). Glutamate released from pyramidal-cell collaterals can also bind to *N*-methyl-D-aspartate (NMDA) receptors, which are thought to be involved in plastic changes related to learning and restitution after brain damage.

A subgroup of cortical interneurons—**spiny stellate cells**—is most likely also **excitatory**, as judged from, among other things, the kind of synapse they form (asymmetrical synaptic thickening, in contrast to the symmetrical thickening found at most inhibitory cortical synapses). As the name implies, this kind of interneuron has numerous dendritic spines, and in this respect it resembles the (excitatory) pyramidal cells. The rest of the interneurons are termed **aspiny** because they have few spines. These are **inhibitory**, with **GABA** as the predominant neurotransmitter. GABA-containing neurons constitute 20% to 25% of all cortical neurons. Most of the GABAergic interneurons also contain one or several **neuropeptides** (such as substance P, somatostatin, vasoactive intestinal peptide [VIP], cholecystokinin, and neuropeptide Y).

There is some experimental evidence that, in some cases, **epilepsy** may be related to selective loss of GABAergic cortical interneurons.

Modulatory Synaptic Effects in the Cerebral Cortex

Modulatory influences on cortical neurons are partly due to activation of **metabotropic glutamate receptors**

and GABA acting on GABA_B receptors. In addition, **neuropeptides** colocalized with GABA exert modulatory effects, although we do not know their functional roles. Most studied, however, are modulatory actions mediated by diffusely organized fiber systems from several brain stem and basal forebrain cell groups. We mentioned such connections in Chapter 5 (under “Modulatory Transmitter ‘Systems’”) and in Chapter 26 (under “Pathways and Transmitters Responsible for Cortical Activation”). The following neurotransmitters are involved: **acetylcholine**, **norepinephrine**, **serotonin**, **dopamine**, and **histamine**. In general, they act to improve the precision of cortical signal transfer—for example, by improving the **signal-to-noise ratio**. Such effects are probably important in relation to **arousal**, **focused attention**, and **motivation**. Acetylcholine, for example, brings layer 5 pyramids in the motor cortex from a state with low-frequency burst firing to single-spike firing. In the latter state, the frequency of single spikes depends on the degree of depolarization: that is, the intensity of synaptic excitatory inputs to the neuron from, for example, the premotor cortex and the thalamus.

Intracortical Signal Traffic and Information Processing

Afferent fibers that end in a small volume of the cortex make excitatory synapses with a large number of projection neurons and interneurons. Thus, one afferent fiber from the thalamus has been estimated to contact about 5000 cortical neurons. The synaptic contacts are established in certain layers only, but the excitation is propagated to other layers by the pyramidal cell recurrent collaterals and **excitatory interneurons** (spiny stellate cells). At the same time, activation of numerous **inhibitory interneurons** serves to focus the excitatory signals and to limit the activity of the projection neurons. In addition, the inhibitory interneurons inhibit other inhibitory interneurons, with resulting **disinhibition**. **Horizontal** axonal collaterals propagate both inhibition and excitation laterally from the focus of cortical excitation. This does not occur at random but so that functionally related neurons are interconnected. Finally, the integrated signals are issued—especially from layer 3 and 5 pyramids—to other parts of the cortex and subcortical cell groups (among them, the motoneurons).

As we see, the activity of each cortical neuron (i.e., its firing frequency and firing pattern) depends on the activity of the numerous other neurons with which it is synaptically connected. Such connections reach a cortical neuron from subcortical nuclei and other parts of the cortex and from cells in its immediate vicinity (within a radius of a few millimeters in the horizontal direction). One cortical neuron, such as a pyramidal cell of the MI, integrates information from perhaps 600

nearby cortical cells and has been estimated to receive about 60,000 synapses (monkey).

The enormous number of neurons and their complex interconnections within even a small volume of cortical tissue explain why we still do not understand the basic rules underlying intracortical information processing. Promising advances have been made, however, especially in the visual cortex (see under “Intracortical Signal Traffic in the Visual Cortex”).

Intracortical Signal Traffic in the Visual Cortex

Regarding intracortical signal traffic, detailed studies have been performed in the visual cortex with the use of methods enabling the recording of single-cell activity in relation to specific stimuli and subsequent intracellular injection of horseradish peroxidase. Thus, the dendritic and axonal patterns of individual, functionally characterized neurons can be determined. Successful attempts have also been made to abolish the activity of neurons in specific layers and then study how the properties of neurons in other layers are changed. As expected from the known terminal pattern of axons from the lateral geniculate body, neurons are first activated in **layer 4** after visual stimuli (in addition, neurons in other layers with dendrites extending into layer 4 can be influenced). From layer 4, the excitation is propagated to **layers 2** and **3**, and from there to **layers 5** and **6**. Some cells in layer 6 send axons upward to layer 4. Presumably, at every step in such a pathway through the cortex some processing of the sensory information takes place, such as integration by one neuron of the signals from other functionally different neurons. In accordance with this assumption, the functional properties of neurons in different layers vary, as shown with microelectrode recordings after natural stimulation of receptors. A projection neuron in layer 5 has quite different properties than a cell in layer 4; for example, the receptive fields of the layer 5 cells are larger (as a sign of convergence of signals from several neurons in layer 4). Other properties of layer 5 neuron also suggest that signals from functionally different layer 4 neurons converge on layer 5 cells (via processing in layers 2 and 3).

Cortical Neurons Are Coincidence Detectors

Many cortical neurons react primarily when information about two events reaches them simultaneously, like cells in the visual cortex that respond poorly to signals from one eye only but vigorously to simultaneous signals from both eyes (binocular cells). Like a good detective who has a special eye for **coincidences** (events occurring simultaneously) and disregards numerous trivial bits of information, the cortical neurons respond preferentially to certain coincidences of stimuli that have a survival value. This is the basis for association

learning; that is, learning the relationship between cause and effect. We know that a novel or unexpected stimulus, or one occurring in an unusual context, causes arousal and improved retention of new material. Often, synaptic changes occur when a neuron receives simultaneous a specific input about a stimulus or the context and a modulatory input (e.g., acetylcholine from the basal nucleus) signaling the salience of the specific input (see Fig. 4.10). This characteristic property of cortical cells is probably built into the inborn wiring pattern (“hardware”) of the brain, but it also needs proper use to be further developed and maintained (see later, “The Parietal Lobe and the Development of the Ability to Integrate Somatosensory and Visual Information”).

Glutamatergic **thalamocortical fibers** appear to act through **AMPA** receptors on cortical neurons but not through **NMDA** receptors (the latter being related to induction of long-term potentiation [LTP]). The recurrent pyramidal cell collaterals, however, act also on **NMDA** receptors. Some experimental evidence shows that simultaneous activation of a cortical neuron from the thalamus (AMPA) and from other cortical neurons (NMDA) can induce LTP. In the monkey motor cortex, LTP has been established during the learning of new motor skills. Thus, not only are the cortical neurons especially sensitive to coincident inputs, they may also be the cellular basis of associative learning in the cortex.

Horizontal Integration and Cortical Plasticity

The extensive horizontal intracortical connections appear to be crucial for the working of the cortex. They help explain why the response of so many cortical neurons depends on the **context** of a stimulus (cf. Chapter 16, under “Color Constancy”). As mentioned, horizontal connections ensure that the **receptive fields** of cortical neurons are not static but subject to modification by inputs from their neighbors. For example, single cells in the visual cortex have smaller receptive fields and react more strongly when the **attention** of the animal is directed to the visual stimulus. Horizontal connections most likely also contribute to well-known **psychophysical phenomena**, such as the filling in of missing lines in otherwise meaningful visual images. When blocking GABA receptors (and thus inhibitory horizontal connections) the receptive fields of cortical neurons enlarge immediately. After blocking GABA transmission, stimulation of a small peripheral spot activates an area in the SI that is much larger than before. After **amputation** of a finger in monkeys, the area in SI activated from the neighboring fingers enlarges immediately. These examples show that there must exist excitatory connections in the cortex that are **suppressed** under ordinary conditions. Further, due to such modifiable connections there is a **dynamic balance**

between the cortical representations of various body parts.

Training of a sensory task can alter the cortical representation of the trained part (e.g., training roughness discrimination with the index finger). It seems likely that such examples of **cortical plasticity** are due, at least partly, to changes in the synaptic efficacy of horizontal connections. The same mechanism probably operates during **recovery** after brain damage that affected the cerebral cortex or its connections.

CONNECTIONS OF THE CEREBRAL CORTEX

The connections of the cerebral cortex with subcortical structures are described in several of the previous chapters, where we deal with the terminal regions of the major sensory pathways and the areas that give origin to the descending pathways involved in motor control. Here we describe general aspects of connections between the **thalamus** and the cerebral cortex and of the **cortico-cortical connections** (association and commissural connections). Such knowledge is a necessary basis for the following treatment of the cortical association areas and their functional roles.

A Brief Survey of Cortical Connections

We can classify the **afferent** connections of the cerebral cortex as follows:

1. Precise, topographically organized connections from the “**specific**” **thalamic nuclei**; each thalamic nucleus supplies one particular part of the cortex
2. Diffusely organized connections from the **intralaminar thalamic nuclei** and several other **subcortical nuclei** (the raphe nuclei, the nucleus coeruleus, dopaminergic cell groups in the mesencephalon, and cholinergic cell groups in the basal forebrain); such connections do not respect the cytoarchitectonic borders in contrast to the connections from the specific thalamic nuclei
3. **Association fibers**—that is, precisely organized connections linking cortical areas within the same hemisphere
4. **Commissural fibers**—that is, precisely organized connections between areas in the two hemispheres

The **efferent** connections of the cerebral cortex can also be divided into **subcortical** and **corticocortical** ones (association and commissural connections). The subcortical fibers are destined for the thalamus, the striatum, various brain stem nuclei (among them the pontine nuclei projecting to the cerebellum), and the spinal cord. The corticocortical connections are for the most part **reciprocal**—that is, an area receives fibers from the same areas to which it sends fibers.

Specific Thalamocortical Connections

The thalamus has been mentioned in several contexts (see Chapters 6 and 14 for descriptions of its gross anatomy and main subdivisions). The thalamus supplies all parts of the neocortex with afferents. Each part of the cortex receives fibers primarily from one of the **specific thalamic nuclei**. The main features of this topographic arrangement of the thalamocortical projection are shown in Fig. 33.8 (as we believe it is organized in humans). Figures 14.6 and 24.16 show the projections from the ventral thalamic nucleus in more detail, based on experimental studies in monkeys with various axonal tracing methods. Figure 16.20 shows the projection from the lateral geniculate nucleus to the visual cortex.

Conditions are more complex than shown in the diagrams, however. In the first place, the topographic arrangement is much more fine-grained. Thus, the projection from one thalamic nucleus is precisely arranged, with subdivisions of the nucleus supplying minor parts only of the large fields shown in Fig. 33.8. As described in Chapters 14 and 16, the thalamocortical

connections from the (VPL) and the lateral geniculate body are somatotopically and retinotopically organized, respectively, with a precision that enables the cortex to extract information about minute spatial details. Further, each thalamic nucleus sends fibers to more than one cortical area. We can take the **mediodorsal nucleus (MD)** as an example. In the scheme, it is depicted as sending fibers to the prefrontal cortex (without any topographic arrangement). In reality, the MD sends fibers to other parts of the cortex, too, such as parts of the cingulate gyrus. The MD furthermore consists of several subdivisions, each supplying a different part of the prefrontal cortex. The main point emerging from such knowledge is that the MD is not a functional unit. A small lesion, for example, must be expected to produce quite different effects, depending on its exact location within the MD.

Some of the specific thalamic nuclei are relay stations in the pathways for signals from **sensory receptors** to specific cortical areas (vision, hearing, cutaneous sensation, etc.). The **ventral posterolateral nucleus (VPL)** receives afferents from the somatosensory pathways and projects to SI (areas 3, 1, and 2; see Figs. 14.2,

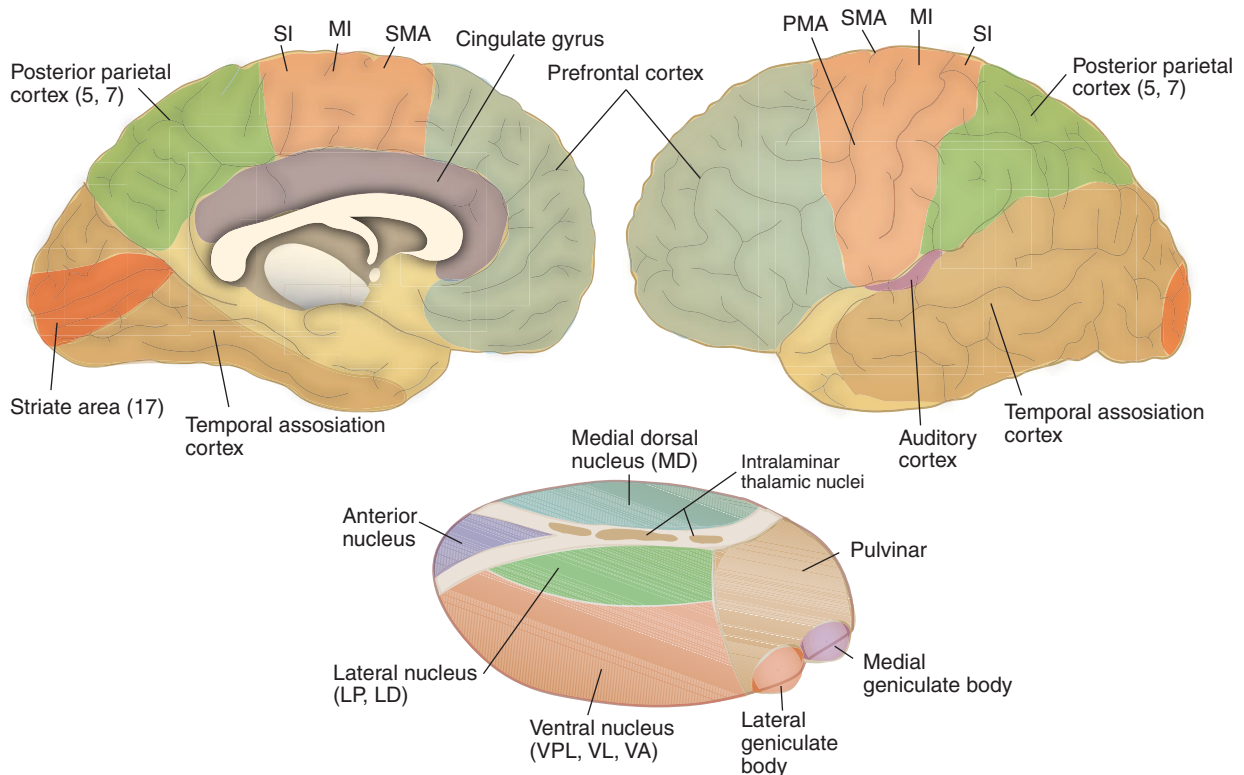


FIGURE 33.8 *The thalamocortical projection.* Highly simplified scheme showing the main features of its topographic organization.

Overlap exists between the cortical terminal regions of different thalamic nuclei but is not shown in the figure.

14.4, and 14.6); the **lateral geniculate body** or nucleus receives afferents from the retina and projects to the striate area (see Figs. 16.14 and 16.20); the **medial geniculate body** is the last subcortical station in the auditory pathways and it sends efferents to AI (see Figs. 17.9 and 17.11). Other specific thalamic nuclei, the **ventrolateral nucleus** (VL) and the **ventral anterior nucleus** (VA), are relay stations in the pathways from the cerebellum and the basal ganglia to the motor and premotor cortical areas (see Fig. 24.16). Other thalamic nuclei relay signals from limbic structures: the **anterior thalamic nucleus** (A) receives afferents from the mammillary nucleus (which receives its main input from the hippocampal formation) and projects to the cingulate gyrus; and the **mediodorsal nucleus** (MD; Fig. 33.8) can relay signals from the amygdala to the frontal lobes. The posterior parietal cortex (areas 5 and 7) receives fibers from the posterior part of the thalamus, the **lateral posterior nucleus** (LP), and parts of the **pulvinar** (Fig. 33.8; see Figs. 6.21 and 6.22). Other parts of the pulvinar projects to the temporal lobe. The LP and the pulvinar receive afferents from nuclei related to vision and eye movements, such as the superior colliculus and the pretectal nuclei, and may relay such information to the posterior parietal cortex (see also Chapter 25, under “Cortical Control of Eye Movements”). However, the pulvinar is not primarily a relay nucleus but a link in a cortico–thalamic–cortical circuit, as we will return when discussing corticothalamic connections.

The Intralaminar Thalamic Nuclei

Modern tracer studies have shown that the cortical projections from each of the intralaminar nuclei end in certain parts of the cortex only; for example, the **contralateral nucleus** (CL) sends fibers predominantly to the parietal cortex. Nevertheless, the projections are considerably more widespread and diffuse than those from the specific thalamic nuclei and do not respect areal borders (the intralaminar nuclei were formerly termed the “unspecific thalamic nuclei”). Physiologic studies indicate that the intralaminar nuclei exert general effects on the **excitability** of cortical neurons. Thus, electrical stimulation produces a so-called **recruiting response** in extensive parts of the cortex, which resembles the EEG changes associated with **arousal** (desynchronization; see Chapter 26). The coactivation of cortical neurons by signals from the specific thalamic nuclei and the intralaminar nuclei may be important for the **binding** of specific stimuli with their salience—that is, a form of **coincidence detection** that perhaps may be necessary for awareness of the stimulus.

The tasks of the intralaminar nuclei are related not only to the cerebral cortex, because they have even stronger connections with the **striatum** (see Chapter 23,

under “Thalamostriatal Connections”). Such connections are precisely organized (another fact speaking against the use of the term “unspecific thalamic nuclei”).

Extrathalamic, Modulatory Connections to the Cerebral Cortex

Apart from the major thalamocortical connections, several subcortical nuclei provide sparser cortical inputs without synaptic interruption in the thalamus (the raphe nuclei, the locus coeruleus, the mesencephalic ventral tegmental area [VTA], the basal nucleus, and the tuberomammillary nucleus in the hypothalamus). These nuclei supply most of the central nervous system with modulatory inputs and are involved in a number of functions, as discussed in previous chapters. Briefly stated, the fibers from the aforementioned nuclei exert a modulatory control over the **excitability** level of cortical neurons, with relation to **wakefulness** and **phases of sleep**. In addition, they probably control more specifically selected cortical neuronal groups when our attention is focused on relevant, novel stimuli.

All of these transmitter-specific nuclei project to large parts of the cortex with no distinct topographic pattern. Nevertheless, recent studies in monkeys show that each nucleus (and thus fibers with a particular transmitter) projects with a higher density to some than to other parts of the cortex. For example, **dopaminergic fibers** from the VTA end with highest density in the prefrontal and temporal neocortex, whereas **noradrenergic fibers** from the locus coeruleus innervate especially the central region (MI, SI). Further, fibers from the various nuclei end in somewhat different cortical layers. A striking feature of the fibers is that, after having entered the cortex, they run **horizontally** for a considerable distance (in contrast to the vertical organization of the afferents from the specific thalamic nuclei). Further, their actions are partly mediated by **volume transmission**. (See Chapter 2, under “Modulatory Transmitter ‘Systems’” and Chapter 16, under “Signal Pathways and Transmitters for the Activation of the Cerebral Cortex”).

The Corticothalamic Connections

All of the thalamic nuclei receive massive **back-projections** from the cerebral cortex. In fact, the number of corticothalamic fibers is much higher than the number of thalamocortical ones; for example, the relationship for VPL has been estimated to 7:1. Yet, the corticothalamic connections have until recently received relatively little attention. The largest thalamic nuclei, with weak or no inputs from peripheral sense organs such, have reciprocal connections with association areas in the parietal, temporal, and frontal lobes

(e.g., the pulvinar and the MD.³ These nuclei should presumably be viewed as dialogue partners for the cerebral cortex rather than relay stations in pathways from subcortical nuclei.

In general, the nuclei receive afferents from the cortical areas to which they send their efferents—that is, the thalamocortical and the corticothalamic connections are **reciprocal**. The corticothalamic projections are also precisely, topographically organized. As mentioned, corticothalamic neurons have their cell bodies mainly in **layer 6**, whereas projections to other subcortical nuclei arise mainly in layer 5. This indicates that the information received by the thalamus is not simply a copy of information sent from the cortex to other regions.

In view of the massive corticothalamic connections, the function of the thalamus cannot be limited to mediating information from subcortical cell groups to the cortex. Even for “relay” nuclei such as VPL, merely quantitative considerations show that crosstalk with the cerebral cortex must be of major importance. All thalamic nuclei receive and presumably process vast amounts of information from the cortex and is therefore intimately involved in processes taking place in the cortex itself. That the corticothalamic fibers really influence the information processing in the thalamus is witnessed by several observations. For example, the receptive fields of neurons in the VPL increase dramatically if their

synaptic input from the SI is removed. The same holds true for neurons in the lateral geniculate body and their input from the striate area. Thus, the cerebral cortex exerts strong top-down control of the signals it is going to receive. Equally important, the corticothalamic connections also influence the firing pattern of the thalamocortical neurons (burst or single-spike firing; see Chapter 26, under “Thalamic Neurons Have Two States of Activity”). For example, the feedback from the cortex governs the **synchronization of spindle oscillations** (burst firing) in various parts of the thalamus during early stages of **sleep**.

Inhibition in the Thalamus: Interneurons and the Reticular Thalamic Nuclei

Inhibitory interneurons—the majority **GABAergic**—probably constitute one-fourth or more of all neurons in some of the thalamic nuclei. In the thalamic sensory relay nuclei, the inhibitory interneurons may contribute to the enhancement of stimulus contrasts (by lateral inhibition) and to selection of certain kinds of stimuli by suppressing other kinds (e.g., in relation to transmission of signals from nociceptors). **Opioid peptides** and other neuropeptides (such as **substance P**) are present in several of the thalamic nuclei, including those relaying nociceptive signals.

The **reticular thalamic nucleus** is unique among the thalamic nuclei because virtually all neurons are **GABAergic**. The nucleus, which forms a thin shell at the lateral aspect of the thalamus (Fig. 33.9), sends its efferent fibers in the medial direction to end in the other thalamic nuclei (and not to the cortex, differing also in

³ Some authors use the term **first-order thalamic nuclei** of those primarily driven from peripheral sense organs (e.g., VPL and the lateral geniculate body), which serve as relay stations in the large sensory pathways. Nuclei not receiving such direct sensory inputs (such as the pulvinar and the MD) are termed **higher order thalamic nuclei** (Sherman 2005).

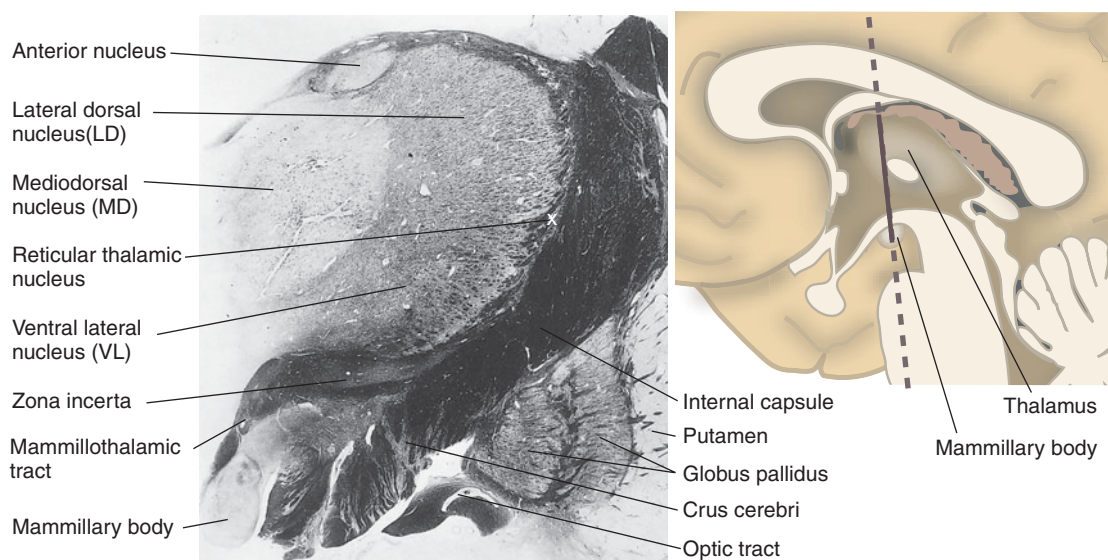


FIGURE 33.9 *The thalamus.* Frontal section through the middle part of the human thalamus. Schematic; myelin stained. (See also Figs. 3.24 and 3.27.)

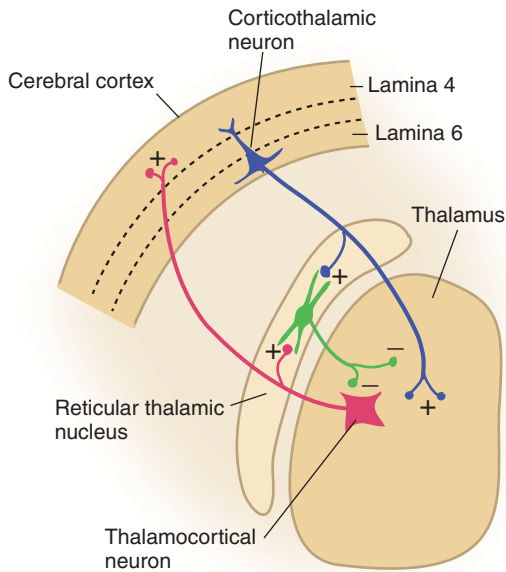


FIGURE 33.10 *Reciprocal thalamic connections with the cerebral cortex and the reticular thalamic nucleus.* Thalamocortical and corticothalamic fibers give off collaterals to the reticular thalamic nucleus. The neurons in this nucleus are GABAergic, and influence significantly the functional state of the thalamocortical neurons.

this respect from the other thalamic nuclei). Both the corticothalamic and thalamocortical fibers pass through the reticular nucleus on their way to and from the thalamus and give off numerous collaterals that end in the nucleus (Fig. 33.10). Further, the reticular nucleus receives afferents from the **mesencephalic reticular formation** and has reciprocal connections with the **PAG** (which, among other things, is related to the control of transmission of signals from nociceptors).

The function of the reticular thalamic nucleus is not fully understood, but its connections indicate that it can influence the activity of thalamocortical neurons and thus, indirectly, the activity of the cerebral cortex. Indeed, electrophysiological studies show that the reticular nucleus can **synchronize** the activity of neurons throughout the thalamus. Further, its activity correlates closely with changes of the **EEG** during sleep and wakefulness. Its relation to **attention** and **control of sensory information** transfer to the cortex may explain why lesions of the reticular nucleus reduce learning and memory (of spatial tasks).

Association Connections of the Cerebral Cortex

Vast numbers of association fibers ensure the cooperation among the cortical areas. The **shortest** association fibers connect minor parts within one area (so-called U fibers), whereas somewhat longer fibers link together

neighboring areas (Fig. 33.3). For example, there are ample connections between the SI and area 5 posteriorly and the MI anteriorly. MI furthermore receives association fibers from the premotor area (PMA) and supplementary motor area (SMA). The **longest** association fibers interconnect functionally related areas in different lobes: for example, connections from the extrastriate visual areas and the posterior parietal cortex lead to the premotor and prefrontal areas.

When we follow the association connections “outward” from the **primary sensory areas** (SI, VI, and AI), signals first reach **unimodal** association areas—that is, areas that process only on sensory modality (Figs. 33.11 and 33.12). From there, association fibers pass to **polymodal** areas—that is, areas that integrate sensory information of different modalities. One area integrates somatosensory and visual information, another visual and auditory, and so forth. The areas outside the primary sensory areas (such as the areas of the posterior parietal cortex and the extrastriate areas) send their efferent projections not only to their immediate neighbors but also to distant areas in the frontal lobe (premotor and prefrontal areas) and to “limbic” cortical areas (the cingulate gyrus and the hippocampal gyrus) (Figs. 33.11 and 33.12).

As a rule, each cortical area establishes association connections with many other areas; that is, there is a

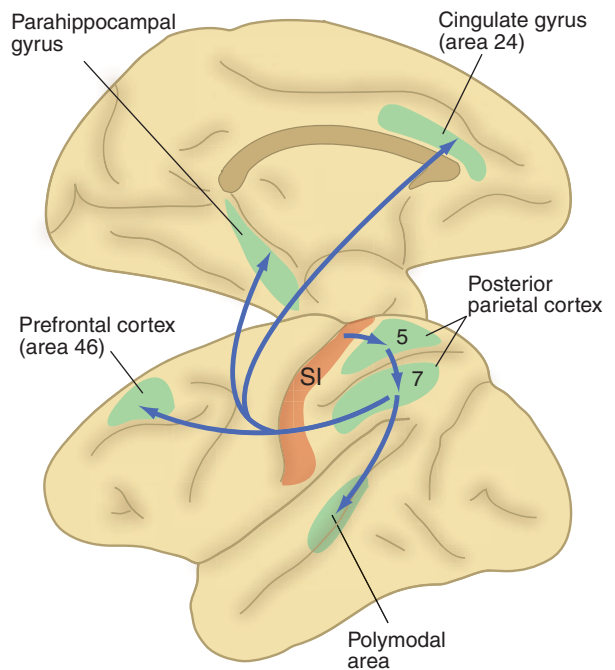


FIGURE 33.11 *Association connections of the somatosensory cortex (monkey).* Only some fiber connections are shown, to illustrate the flow of information progressing from SI to the posterior parietal cortical areas, and from there to polysensory areas in the temporal lobe, to the prefrontal cortex, and to limbic cortical regions. (Based on Jones and Powell 1970.)

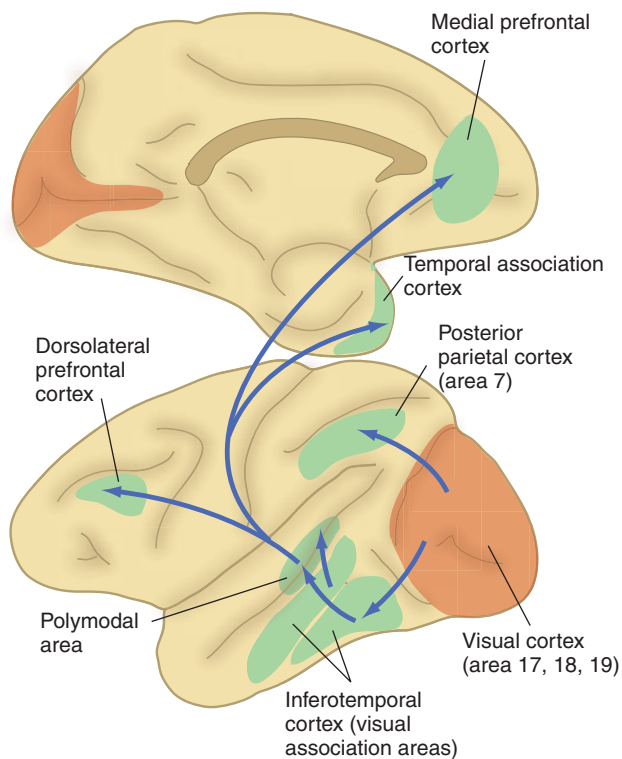


FIGURE 33.12 Association connections of the visual cortex. Compare with Figure 33.9 and note similarities with regard to the progression of information outward from the striate area. Only some connections are shown. (Based on Jones and Powell 1970.)

considerable **divergence** of information. At the same time, there is also considerable **convergence**, as each area receives afferents from several other areas (Figures 33.11 and 33.12 give very simplified accounts). Together, the many areas of the cortex are extensively interconnected, forming complex networks specialized for specific tasks. Complex functions—such as control of attention and speech—are carried out by many areas in conjunction with each other by way of their association connections. At the cellular level, the extensive association connections enable neurons to integrate many, different pieces of information and to act as coincidence detectors, as discussed in the preceding text.

Outward and Backward Corticocortical Connections

The discussion so far may give the impression that the flow of information between various cortical areas goes only in one direction, but this is not the case. Generally, the association connections are **reciprocal**, enabling areas to exert mutual influences. For example, there is not a one-way traffic outward from the striate area through successive stages of extrastriate areas, as at all levels areas send fibers back to each other and to the striate area. Differences exist, however, between the

connections going in opposite directions between two areas, notably with regard to laminar origin and destination. This has been most studied in visual cortical areas. Fibers going **outward**—also called **feedforward** connections—from the striate area to the extrastriate areas arise mainly in layer 3 and end primarily in layer 4. Fibers going **backward**—**feedback** connections—arise in deep layers (layer 6) and end mainly in the most superficial layers. The outward connections are precisely, topographically organized, whereas in general, the backward connections show a less fine-grained topography. Based on their anatomic characteristics, the outward connections from the sensory areas appear to be concerned with **segregation** and specific **convergence** of sensory information. For example, the striate area distributes different features of a visual image to extrastriate areas that are specialized for analysis of color, motion, form, and so forth. The backward connections are probably responsible for **context-dependent modifications** of sensory processing at the earlier stages—for example, the well-known effect of context on receptive fields of neurons in primary sensory areas. The backward connections may also provide a **prediction** of the sensory input expected to arise from a motor command, enabling the brain to compare the expected and actual results.

Commissural Connections of the Cerebral Cortex

Most commissural fibers pass in the **corpus callosum** (see Figs. 6.26 and 6.27). A small fraction courses in the **anterior commissure** (in humans, 1%–2%). The latter fibers are believed to belong primarily to the olfactory pathways. (In the monkey, the anterior commissure contains about 5% of all commissural fibers, mainly linking corresponding parts of the temporal lobes.)

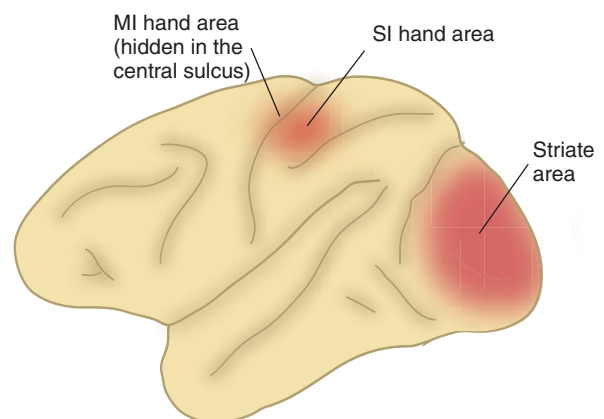


FIGURE 33.13 Total distribution of commissural connections (monkey). Almost all parts of the cerebral cortex send and receive commissural connections, except the representation of the central visual field in the striate area and the SI and MI hand regions (shown in red).

Generally, commissural fibers interconnect **corresponding areas** in the two hemispheres. The density of such fibers varies considerably among regions, however. Some regions are almost or totally devoid of commissural fibers (Fig. 33.13), especially the **striate area** and parts of the **MI** and **SI** that represent **distal parts of the extremities** (the regions representing the hands and feet). Thus, regions of the cortex dealing with parts of the body that usually work in a symmetrical fashion (such as the two halves of the back) are amply interconnected, whereas parts that usually work independently (such as the hands) have few commissural fibers. A corresponding organization is present in the somatosensory and the

motor pathways; connections related to distal body parts are entirely crossed, whereas connections related to proximal parts are bilateral. It seems possible that commissural connections between, for example, the hand areas, would disturb the independent control of the hands (nevertheless, information about motor commands and sensory signals pertaining to the distal body parts of one side can reach the other hemisphere by other cortical areas and their commissural connections).

We return to the corpus callosum and the commissural connections in Chapter 34 when dealing with lateralization of functions—that is, tasks that are the division of tasks between the two hemispheres.

34 | Functions of the Neocortex

OVERVIEW

This chapter deals primarily with functional aspects of the cortical **association areas**, that is, parts of the cortex that neither receive direct sensory information through the major sensory pathways nor send direct fibers to subcortical motor nuclei. All of the cerebral lobes contain association areas, according to this definition. In Chapter 33, we discuss the connections of the association areas in general. Here we look at the main groups of association areas and their possible functions. It should be realized, however, that the functional roles of a group of association areas, for example, those of the frontal lobe, can be understood only if considered as part of distributed, task-specific networks involving other association areas and subcortical nuclei.

Typically, the association areas receive and **integrate** various kinds of information. Some are specialized for integration of two or more sensory modalities; others integrate highly processed sensory information with information about intentions and goals. The integrative capacity depends on proper use of sensory information, and must be learned in early childhood.

The **posterior parietal cortex** integrates visual and somatosensory information, and is responsible for, among other functions, the control of **visually guided behavior** and **spatial orientation**. From the parietal areas, signals pass to premotor and motor areas. The posterior parietal cortical areas also have reciprocal connections with the cingulate gyrus and the prefrontal cortex, mediating the influence of emotions, attention, and motivation on behavior produced by somatosensory and visual stimuli.

The **prefrontal cortex**, situated in front of the frontal lobe motor areas, receives information about all sensory modalities and also about the motivational and emotional state of the individual. It is of special importance for planning and initiation of **goal-directed behavior**. More specifically, the prefrontal cortex is important for attention, for selection of a specific behavior among several possible ones, and for suppression of unwanted behavior. These functions are of particular importance for social behavior.

The **temporal association areas** are particularly concerned with high-level processing of auditory and visual information. The cortex of the superior temporal gyrus is characterized by its connections with the auditory cortex, whereas the inferior part of the temporal

lobe—the **inferotemporal cortex**—is dominated by processed visual information important for object recognition. The **medial parts** of the temporal lobe are of special importance for learning and declarative memory, as discussed in Chapter 32.

The **insula**, hidden in the lateral sulcus, receive all kinds of sensory information and is involved in pain perception, homeostasis, and bodily awareness.

Language and speech depends on a distributed network with two special hubs, one in the frontal lobe (**Broca's area**) and one at the temporoparietal junction (**Wernicke's area**). The anterior (frontal) area is particularly important for the expressive aspects of speech, while the posterior area is more concerned with the sensory aspects.

The left hemisphere is **dominant** for speech in 95% of the population. We also discuss further examples of hemispheric specialization (lateralization). The **corpus callosum** ensures that the two halves of the brain keep each other updated and cooperate to obtain common goals.

Finally, we discuss possible biologic bases of **cognitive sex differences**.

ASSOCIATION AREAS

For some of the cortical areas, functional aspects are discussed in previous chapters. Here we address primarily functions that can be ascribed to the so-called **association areas** of the cerebral cortex. This term is not precisely defined but is traditionally used for parts of the cortex that neither receive direct sensory information through the major sensory pathways nor send direct fibers to subcortical motor nuclei. All of the cerebral lobes contain association areas, according to this definition.

The connections of the association areas indicate that they are able to integrate information from sensory and “limbic” parts of the cortex (i.e., the cingulate gyrus, parts of the prefrontal cortex, and the hippocampal region) and thereafter issue commands to motor cortical areas and (indirectly) to the hypothalamus.

Comparison of the cerebral hemispheres of humans and monkeys shows that the association areas occupy a much larger fraction of the total in humans than in monkeys. Comparison of monkeys with other mammals, such as cats and dogs, shows again that the main

difference between them with regard to the cerebral cortex is the relative size of the association areas. These parts of the cortex are of importance for what we may loosely call **higher mental functions**, as we discuss here. The association areas are not “centers” for specific mental faculties, however. First, several areas—often widely separated—participate in one task or function, and, further, one area participates in more than one function. This is witnessed by the high degree of divergence and convergence of the connections of the association areas, as discussed in the preceding text. Second, the operations of the association areas cannot be understood if considered in isolation; the intimate connections between the association areas and subcortical cell groups, such as the thalamus, the basal ganglia, the amygdala, and the hippocampus, are essential for their normal functioning.

Measurements of **regional cerebral blood flow** and metabolism during the performance of various **cognitive tasks** indicate that large parts of the cortex participate in all higher mental functions. When a person is asked to **imagine** that she is walking from one place to another in a city she knows, the activity increases in the extrastriate visual areas, in the posterior parietal cortex, in parts of the temporal lobe, and in several prefrontal areas. Solving a **mathematical** problem activates many of the same cortical areas but with certain differences, and a **verbal** task activates multiple areas that partly coincide with and partly differ from those activated in the spatial and the mathematical tasks.

In Chapter 16, we discussed the cortical substrate of visual imagery (under “Consciousness and Visual Experience”).

Cognition and Cognitive Functions

The word **cognition** stems from the Latin word *cognitio*, meaning “acknowledge, come to know.” According to the *Encyclopedia Britannica*, cognition “includes every mental process that can be described as an experience of knowing as distinguished from an experience of feeling or of willing. It includes, in short, all processes of consciousness by which knowledge is built up, including perceiving, recognizing, conceiving, and reasoning.” Among neuroscientists today, the word is often used more broadly to include the **affective** aspects of higher mental functions. For example, the scholarly book *The New Cognitive Neurosciences* (2000) edited by Michael Gazzaniga deals not only with consciousness, language, memory, attention, and similar phenomena but also with emotions. This presumably reflects the realization that there is no sharp distinction between brain structures that govern rational thought and actions on the one hand and those that underlie emotions and subconscious drives on the other.

Integration of Different Sensory Modalities Depends on Learning

The ability to integrate somatosensory and visual information and to use visual information to guide voluntary movements is not inborn but learned in infancy and early childhood. Persons who were born blind but gained their vision back as young adults do not manage to coordinate the visual information with the other senses and therefore cannot use the “new” sense. They usually continue to use tactile sensation to “see” objects, and, in fact, the visual information can be more confusing than helpful (this is described by Oliver Sacks in *An Anthropologist on Mars* 1995).

Studies by Hyvärinen (1982) and coworkers exemplify how meaningful use of the systems is necessary for sensory integration to take place during early development. He studied how the properties of single cells in the parietal cortex change during the phase in which an infant monkey learns to combine **visual** and **somatosensory information**. Monkeys were prevented from seeing from birth (by suturing the eyelids) until they were between 6 months and 1 year old. At that time, few cells in area 7 responded to visual stimuli, in contrast to the normal situation at that age. An abnormally large fraction of the cells were activated by passive somatosensory stimuli and by active movements. Most striking was the almost total absence of cells that could be activated by both somatosensory and visual stimuli. Even 2 years after reestablishment of normal vision, the single-cell properties of area 7 remained virtually unaltered, with very few cells responding to visual stimuli. Properties of cells of the extrastriate cortex (area 19) were also altered in the visually deprived monkeys; for example, some cells were activated by somatosensory stimuli (which never occurred in normal monkeys), and there were fewer than normal visually driven cells. Behaviorally, the monkeys were blind after opening of the eyes, and no improvement occurred during the next month. They bumped into obstacles, fell off tables, and were unable to retrieve food by sight alone. Threatening faces did not frighten them, in contrast to normal monkeys at the same age. One monkey that was observed for 3 years improved to some extent, but it never regained the full use of vision. Brain-imaging studies of **humans born blind or deaf** also indicate that considerable reorganization takes place, as compared with normal persons. For example, in persons who have been blind since birth, the visual cortex is activated by somatosensory stimuli during Braille reading.

In conclusion, the data indicate that the functional properties of neurons in the association areas, and thus the capacity of the areas to contribute to certain tasks, are determined to a large extent in early childhood. Further, there is only a limited possibility of regaining the proper function of these regions at a later stage

(see Chapter 9, under “Sensitive (Critical) Periods”). The experiments also show that when one kind of sensory information is lacking during an early stage of development, other sensory modalities take over parts of the cortex not normally used for processing that kind of sensory information. This has been shown also after early lesions of the auditory system.

Lesions of the Association Areas: Agnosia and Apraxia

Lesions of association areas typically disturb higher-level aspects of sensory and motor functions. **Agnosia** is usually defined as the lack of ability to recognize objects (when not due to reduced sensation or dementia). **Apraxia** is used similarly about the loss of the ability to do certain, formerly well-known skilled actions (such as dressing, using household tools, copy a drawing, etc.). There is no clear-cut distinction between these two categories of symptoms, however. Loss of the ability to copy a drawing—called **constructional apraxia**—for example, may be due to the patient not being able to perceive more than a small piece of the drawing at a time. Thus, she is not able mentally to put several pieces together.

Agnosia or apraxia seldom occurs as the only symptom, however, and even less frequent are cases with isolated subcategories. Such cases are nevertheless of great theoretical interest because they shed light on how the brain works to solve specific tasks. Depending on their site and size, cortical lesions can produce a wide spectrum of difficulties with perception. This concerns not only objects (strictly defined), but also all aspects of higher processing of sensory information. The word “agnosia” is therefore used more widely than the definition may suggest. For example, we use the term **visual agnosia** for loss of the ability to recognize objects and persons by sight, whereas **tactile agnosia** means loss of the ability to recognize objects by touch.

Visual agnosia is the most common kind and has been studied the most. It may appear in several varieties, each with a specific name. For example, **prosopagnosia** means inability to recognize familiar faces; **autotopagnosia** is inability to recognize one’s own body parts; **simultanagnosia** is the inability to perceive more than one object at a time, and so forth. Inability to recognize letters, **alexia**, is regarded by some as a special kind of agnosia. To some extent, the different kinds of agnosia can be attributed to lesions of specific parts of the cortex. For example, several kinds of visual agnosia are associated with lesions of specific parts of extrastriate areas (see Chapter 16, under “Further Processing of Sensory Information outside the Striate Area”). **Tactile agnosia** has been described after lesions at the parieto-temporal junction (see Chapter 14, under “Lesions of the Somatosensory Areas”). Disturbances of the **body**

image and **self-awareness** are typical of lesions in the posterior parietal cortex (see later).

There are numerous forms of apraxia (about 30 are listed by Petreska and coworkers in a comprehensive review). For example, **ideational apraxia** is used when it is not the execution of movements that is impaired (e.g., the use of a toothbrush) but the objects are used inappropriately (e.g., to eat with the toothbrush). **Ideomotor apraxia** is characterized by impaired execution of a movement rather than the conceptualization of its purpose. Basically, apraxia seems to “. . . result from a specific alteration in the ability to mentally evoke actions, or to use stored motor representations for forming mental images of actions” (Petreska et al. 2007, p. 64). Although apraxia is most frequently observed after parietal or frontal lesions, lesions in other parts of the cortex and even subcortical ones may produce apraxia too.

If in the term “agnosia” we include the inability to recognize **complex sounds** (like music, spoken words, and laughter), the distinction with speech disturbances after brain damage (aphasia) becomes blurred. Further, there are elements of apraxia in aphasia when it includes (as often happens) the inability to write (**agraphia**). Patients with lesions of the association areas often have symptoms belonging to several categories mentioned above—that is, elements of aphasia, agnosia, and apraxia. This can only partly be explained by the fact that many lesions are large and affect several specialized regions. It also reflects that, although they are anatomically separated, association areas are extensively interconnected and their normal functioning requires that they cooperate.

Parietal Association Areas

Usually **areas 5 and 7**—located in the upper and lower parietal lobules, respectively—are considered to constitute the parietal association cortex (Fig. 34.1; see Fig. 33.4). The term **posterior parietal cortex** is also used of this region. Both areas 5 and 7 can be further subdivided into parts differing in connections and functional properties.¹ These areas are intercalated between the visual cortical areas in the occipital lobe and the somatosensory cortex in the anterior parietal cortex. Functionally, as one might expect from this location, areas 5 and 7 process and integrate somatosensory and visual information. From these areas, signals are conveyed to premotor and motor areas (Fig. 34.1), explaining why a parietofrontal network is activated during

1 There is some disagreement in the literature with regard to the parcellation of the posterior parietal cortex in humans. Brodmann (Fig. 33.3) placed areas 5 and 7 in the superior parietal lobule, whereas the inferior lobule contained areas 39 and 40. Others, however, describe area 5 as located in the superior parietal lobule and area 7 in the inferior—that is, corresponding to the situation in monkeys.

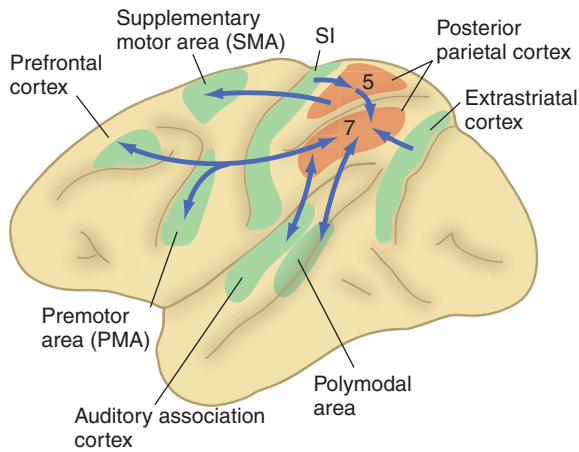


FIGURE 34.1 Association connections of the posterior parietal cortex (monkey). Connections with the limbic cortical areas are not shown (see Fig. 33.11). Visual and somatosensory information converge in area 7. Connections are reciprocal.

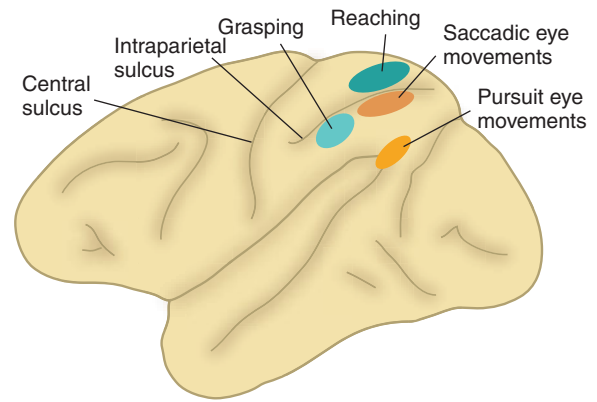


FIGURE 34.2 Subregions within the posterior parietal cortex with relation to planning of specific kinds of movement. (Based on Andersen and Buneo 2002.)

most voluntary movements. The posterior parietal cortical areas also have ample connections (both ways) with the **cingulate gyrus** and the **prefrontal cortex**. These connections are assumed to mediate the influence of emotions, attention, and motivation on behavior produced by somatosensory and visual stimuli.

Experiments with recording of **single-cell** activity in **area 5** of the monkey indicate that this area is essential for the proper use of somatosensory information, for goal-directed voluntary movements, and for the manipulation of objects. This fits well with the symptoms that arise in humans after damage to the posterior parietal cortex, as discussed next. Single-cell recordings indicate that **area 7** has an important role in the integration of visual and somatosensory stimuli, which is essential for the coordination of the eye and the hand—that is, for visual guidance of movements. Area 7 is also involved in the control of **eye movements**. Studies in monkeys suggest that certain subregions of the posterior parietal cortex are specialized for reaching movements, grasp, saccades, and smooth-pursuit eye movements (Fig. 34.2). It further appears that activity in these subregions is closely linked with the **intention to move**²—that is, the posterior parietal cortex contains a “map of intentions.”

Lesions of the posterior parietal cortex in humans can cause different symptoms, depending on which

parts are most affected. Some of the symptoms can be summarized as difficulties with the transformation of sensory stimuli into adequate motor actions. This can probably be explained by lack of parietal influence on the premotor areas. The understanding of the meaning of sensory stimuli is seriously impaired (but usually not the mere recognition of a stimulus). This **agnosia** concerns especially the recognition of the form and spatial position of objects. A typical symptom after right-sided lesions is a tendency to **neglect** the opposite side of the body and the visual stimuli from the opposite side. Patients may suffer from **apraxia**—that is, they are unable to use well-known tools and objects. They may further have problems with **visually guided movements** (such as stretching out the arm to obtain an object).

Even though the symptoms mentioned here are most often seen after damage of the posterior parietal cortex, most of them have been described after lesions in other parts of the brain, too, especially of the prefrontal cortex, the thalamus, and the basal ganglia, all of which have connections with the parietal cortex. These structures take part in a **distributed network** responsible for, among other functions, the control of visually guided behavior and spatial orientation.

Properties of Single Neurons in the Posterior Parietal Cortex

Studies of monkeys with permanently implanted electrodes have demonstrated a wide repertoire of properties among neurons in areas 5 and 7. In general, a task-related increase in firing frequency occurs only when a stimulus is relevant and the **attention** of the animal is directed toward the stimulus. Thus, many neurons are virtually impossible to activate when the

2 Stimulation of parietal and premotor cortical areas in awake patients undergoing surgery corroborates the importance of the parietal cortex for movement intention and awareness of own movements. Stimulation of the Brodmann's areas 7, 39, and 40 (Fig. 34.4) provoked an intention to move, and with increasing stimulus strength, the patients reported that they had actually performed the movements (although no movement occurred). Stimulation of the premotor cortex, on the other hand, produced overt movements but the patients were not aware of them. Thus, “Conscious intention and motor awareness... arise from increased parietal activity before movement execution.” (Desmurget et al. 2009).

animal is drowsy and inattentive. Some cells respond to stimulation of proprioceptors, but their response is much more vigorous when a movement (stimulating the proprioceptors) is **self-initiated** by the monkey than when the joint is passively manipulated by the examiner. In area 5, many neurons change their firing frequency in relation to manipulatory hand movements. Other neurons increase their firing in relation to reaching movements, but only when the hand is moved toward an object the monkey wants to obtain (such as an orange). The increase of firing in such neurons starts at the time the animal discovers the object—that is, before the arm movement starts—and is therefore not a result of proprioceptive stimulation. The American neurophysiologist Vernon B. Mountcastle, who first described such neurons, suggested that they might function as **command neurons** for the target-directed **exploration** of our immediate surrounding extrapersonal space. Such neurons appear to respond to the **coincidence** of two events: a sensory stimulus (e.g., the sight of an orange) and a signal that depends on motivation (whether the monkey is hungry and wants the orange; see Fig. 4.10).

More about Symptoms after Lesions of the Posterior Parietal Cortex

The most marked symptom produced by bilateral parietal lesions is the inability to **grasp** and to **manipulate** objects. Thus, the patient may be unable to move the hand toward an object that is clearly seen, even though there are no pareses and no visual defects. Movements that do not require visual guidance—such as buttoning, bringing an object to the mouth, and so forth—are performed normally. When the patient is asked to pour water from a bottle into a glass, he pours the water outside the glass over and over again, even though he can see clearly both the bottle and the glass. Such patients also have severe difficulties with the **appraisal** of **distances** and the **size** of objects. Further, to **fix the gaze** becomes difficult, especially to direct the gaze toward a point in the periphery of the visual field. The **identification of objects** is difficult, because of the inability to attend to more than one detail at a time (such as seeing a cigarette but not the person who smokes it). This may be a fundamental defect after parietal lobe lesions, perhaps also explaining the difficulties mentioned earlier with pouring water into a glass (the patient is unable to locate in space the bottle and the glass at the same time).

Patients with parietal lobe lesions typically have difficulties with **drawing** an object or a scene; again the inability to perceive more than one feature at a time is the probable basic defect. The parts of an object are drawn separately, without the proper spatial relations, or the drawing gives an extremely simplified representation.

This symptom occurs most often after damage to the right parietal cortex, in cases of unilateral lesions. The **use of tools** is also difficult or impossible: for example, the patient no longer knows how to use a hammer (**apraxia**).

Unilateral lesions of the right parietal lobe typically produce negligence—**neglect**—of the opposite body half and visual space. Such a patient behaves as if the left part of his body does not exist. He dresses only the right side, shaves only the right half of the face, and so forth. He may deny that the left leg belongs to him and claim that it belongs to the person in the adjacent bed, for example. A similar symptom is denial of the disease and the functional loss, called **anosognosia**. The patient may deny that the limb is paralytic or that he is blind. Thus, certain aspects of body knowledge no longer exist in the mind of the patient, but the loss is not consciously perceived. When drawing a face, for example, the right side is drawn normally, whereas the left side is vague or not included in the drawing.

A peculiar constellation of symptoms, the **Gerstmann syndrome**, can occur after lesions of the parietal lobe at the transition to the temporal lobe (usually of the left hemisphere). The symptoms are as follows: **finger agnosia** (the patient cannot recognize and distinguish the various fingers on her own or other people's hands), **agraphia** (inability to write), sometimes **alexia** (inability to read), **right-left confusion**, and, finally, **dyscalculia** (reduced ability to perform simple calculations, especially to distinguish categories of numbers such as tens, hundreds, and so forth). The most distinctive feature of the syndrome is the finger agnosia, which can occur in isolation. That finger agnosia can be the only symptom of a parietal lobe lesion indicates that a disproportionately large part of the human parietal cortex is devoted to the hand. Thus, the hand has a unique role as an exploratory sense organ and as a tool, and, further, it has a special place in our inner, mental, body image. The British neurologist M. Critchley (1953, p. 210) expressed it as follows: “The hand is largely the organ of the parietal lobe.”

Frontal Association Areas

In this context we use the term “association cortex” only about the **prefrontal cortex**—that is, the parts of the frontal lobe in front of areas 6 (premotor area [PMA], supplementary motor area [SMA]) and 8 (the frontal eye field) (Fig. 34.3; see Fig. 33.4). The prefrontal cortex consists of several cytoarchitectonic areas, each with a specific set of connections. Together, the prefrontal areas receive strong **afferent** connections from areas in the occipital, parietal, and temporal lobes and, in addition, from the cingulate gyrus (Fig. 34.3). Thalamic afferents come from the **mediodorsal nucleus**, MD (see Fig. 33.8), which, in turn, receives afferents

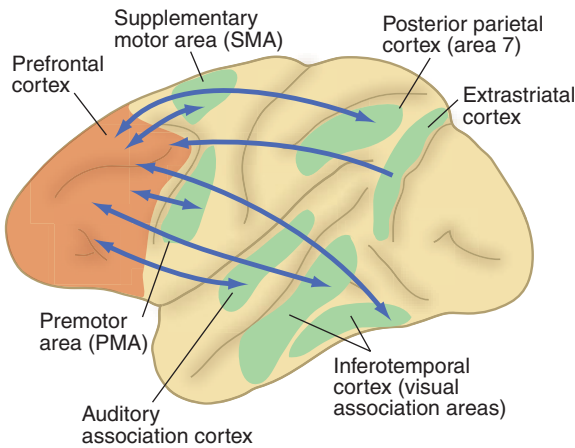


FIGURE 34.3 Association connections of the prefrontal cortex (monkey). Note the convergence of all kinds of processed sensory information and the connections with PMA and SMA. Connections with limbic cortical areas are not shown.

from the **amygdala** and the **ventral pallidum** (among other places). In sum, the prefrontal cortex appears to receive information about all sensory modalities and also about the motivational and emotional state of the individual.

The prefrontal cortex sends **efferents** back to most of the areas from which it receives afferents, among them the SMA and the PMA. In addition, many prefrontal efferents reach the caudate nucleus of the **striatum** (see Fig. 13.6). Finally, some efferents reach the **amygdala** and the **hypothalamus**.

Animal experiments, observations in brain-damaged humans, and brain imaging in normal persons all give a fairly consistent picture of the major tasks of the prefrontal cortex. Figure 34.4 shows some tasks associated with particular prefrontal subdivisions. The prefrontal cortex is obviously of crucial importance for **planning** and **initiation** of **goal-directed behavior**. More specifically, the prefrontal cortex is important for **attention**, for **selection** of a specific behavior among several possible, and for **suppression** of unwanted behavior. With regard to selection, the prefrontal cortex cooperates with the basal ganglia, as discussed in Chapter 23, under “Functions of the Basal Ganglia.” Further, the prefrontal cortex is important for certain aspects of **memory**—both for working memory and for the long-term establishment of memory traces. Working memory enables us to retain a stimulus long enough for its evaluation and linking with ongoing processes and memory. We discussed in Chapter 32 that medial parts of the temporal lobe, including the hippocampus, are necessary for declarative memory. Functional magnetic resonance imaging (fMRI) studies indicate, however, that we remember words or pictures only if their presentation also activates the

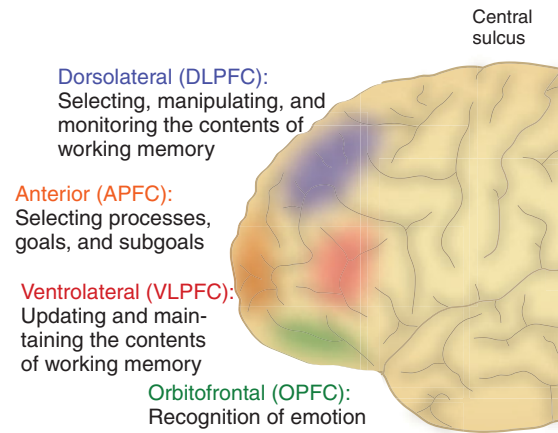


FIGURE 34.4 Some functional specializations within the prefrontal cortex. Regions that appear to be related to different aspects of memory, as judged from functional neuroimaging studies (PET and fMRI). The orbitofrontal cortex is activated in association with explicit identification of emotional facial expressions. (Based on Fletcher and Henson 2001, and Adolphs 2002.)

prefrontal cortex. Probably, the prefrontal cortex tells the hippocampal region about the emotional coloring and the context of information that is transmitted to the hippocampus from sensory areas.

Learning of rules by association appears to be a central task of the prefrontal cortex. This may be performed by neurons that associate behaviorally relevant but otherwise dissimilar bits of information—such as that a red traffic light means stop. Appropriate behavior requires that we are able to learn such rules, but equally important is that we can replace them quickly with new ones. Both faculties suffer after damage to the prefrontal cortex.

In conclusion, the prefrontal cortex is of crucial importance for our ability to organize our own lives and to function socially. Indeed, the tasks mentioned previously, such as planning and choice of behavior, choosing between signals with different emotional coloring, suppression of unwanted behavior, and so forth, are indispensable for **social adaptation**. Another important factor in social functioning is **empathy**,³ which also seems to depend on the integrity of the prefrontal cortex. However, the prefrontal cortex is not the only part of the cortex showing empathy-related activity. It appears that empathy activates the same network that is activated by the person experiencing the painful or

³ **Empathy** is used with somewhat different meanings. Hein and Singer (2008) refer to empathy as “...an affective state, caused by sharing the emotions and sensory states of another person.” They distinguish empathy from **sympathy**, which they describe as emphatic concern or compassion. By emphasizing that empathy is an affective state, they also distinguish it from the understanding of other persons’ beliefs, intentions, and desires, which derives from reasoning (i.e., cognitively). Indeed, understanding another person’s intentions is not the same as sharing them.

distressing situation. The exact distribution of activity would therefore vary with the specifics of the emotion experienced by the suffering person.

Symptoms after Prefrontal Lesions

The prefrontal cortex consists of several subregions, which differ with regard to their connections and functional properties. The symptoms of lesions that occur in this area in humans are correspondingly varied, and, further, the individual differences are fairly large even after seemingly identically placed lesions. In general, the symptoms are compatible with the functions discussed earlier.

Prefrontal lesions typically produce changes of **mood** and **personality**, distinguishing them from lesions of other parts of the cortex. Commonly, large lesions produce apathy, indifference, and emotional leveling-off. The patient appears to be uncritical compared with before the damage. For example, he may behave in a complacent and boastful manner, which he would never have done before. The ability to **alter the behavior** on the basis of experience from previous actions appears to be reduced. Clear-cut symptoms occur usually only after bilateral damage to the prefrontal cortex. Occasionally, however, the first signs of a **frontal-lobe tumor** are changes of behavior and personality.

More about Symptoms after Lesions of the Prefrontal Cortex

A striking defect after bilateral lesions of the **dorsolateral prefrontal cortex** (DLPFC) is the lack of the so-called **delayed response**. A monkey sees that food is put into one of two bowls. Then the sight of the bowls is blocked for up to 10 min before the monkey is allowed to choose one of the two. In contrast to normal monkeys, the lesioned ones do not remember which bowl contained the food (even though they do not show reduced performance in other more complicated memory tests). The dorsolateral parts of the prefrontal cortex thus appear to be necessary for the ability to form and retain an inner conception of the existence of an object in time and space when the object is no longer seen. Interestingly, humans manage a similar test first at the age of about 1 year. Before that age, everything that is not seen or felt is presumably nonexistent for the infant.

A characteristic symptom in humans with prefrontal lesions is the inability to **alter the response** when the stimulus changes; they continue to make the same response even though it is no longer adequate. This phenomenon is called **perseveration**. The **Wisconsin card sort test** is often used to reveal such a defect. The person is asked to sort cards in accordance with certain general rules, such as color, number, shape, and so forth. The correct rule to be applied is indicated by the

response given by the examiner to the first attempts at sorting the cards. The rules can be changed without warning. Normal persons understand fairly quickly that the rule has been changed and alter their responses accordingly, whereas patients with prefrontal lesions continue sorting in accordance with the first rule, in spite of repeated warnings that they are making mistakes.

Emotional and **personality** changes after frontal lobe lesions are most common when the lesion includes the **orbitofrontal parts**. Such symptoms are difficult to evaluate, and the premorbid personality of the patients appears to play a decisive role. Nevertheless, a general tendency is to become less emotional and to show reduced emotional reactions to events. They also have difficulties in extracting the salient features from a complex situation, making their responses unpredictable and often inappropriate. A test designed for such symptoms uses drawings of complex situations, in part with a dramatic content, such as a man who has fallen through the ice on a lake and is in danger of drowning. Patients with frontal lesions usually attend only to details, saying, for example, "Since there is a sign saying 'Careful!' on the beach, there may be a high-voltage cable nearby." This kind of reduction results in inability to foresee the consequences of one's own actions and poor insight into other people's circumstances. This leads to poor **social adaptation**, with isolation as the final result.

The reduced capacity to retain inner conceptions is most likely the reason such patients have increased **distractibility**, with reduced ability to perform tasks that require continuous activity and attention. Motor hyperactivity, which can be a symptom, may perhaps result from the increased distractibility. Thus, in monkeys with prefrontal lesions, the hyperactivity disappears when they are placed in an environment with few stimuli.

In humans with frontal lobe tumors (e.g., a glioma or a metastasis from a malignant tumor elsewhere), a **depressive disorder** has been observed, but this may rather be a condition of de-emotionalization and social isolation.

The Temporal Association Cortex

A unitary functional role is even less evident for the temporal association areas than for those in the parietal and frontal lobes. Apart from the auditory cortex (areas 41 and 42) (see Fig. 17.11) and the phylogenetically old parts at the medial aspect (Fig. 31.1), the temporal lobe consists largely of Brodmann's areas 20, 21, and 22, which here are considered the association areas (Fig. 34.4).

The cortex of the **superior temporal gyrus** is characterized by its connections with the auditory cortex,

whereas the inferior part of the temporal lobe—the **inferotemporal cortex**—is dominated by processed visual information from the extrastriate visual areas. In addition, there are strong connections with the **hippocampal formation** (through the entorhinal area) and the **amygdala**. Electrical **stimulation** of the temporal association cortex in humans evokes recall of memories of past events or the experience of dreamlike sequences of imagined events (such observations were first made by the Canadian neurosurgeon Wilder Penfield, who stimulated the temporal lobe and other parts of the cortex in patients in whom the cortex was exposed under local anesthesia for therapeutic reasons). Finally, long association fibers interconnect the temporal association areas with the prefrontal cortex (Fig. 34.3).

The **inferotemporal cortex** is important especially for the interpretation of **complex visual stimuli**, as judged from experiments in monkeys. Thus bilateral removal of these regions makes the monkeys unable to recognize and distinguish complex visual patterns.

These and other observations led to the conclusion that the inferotemporal cortex is of special importance for the **categorization** of visual stimuli. In monkeys, some neurons of the inferotemporal cortex respond only when the monkey sees, for example, a **face** or a **hand**. Some neurons respond preferentially to one particular face, whereas other neurons respond to any face. A neuron that responds briskly when the monkey is shown a drawing of a face may stop firing when important features are removed, such as the mouth or the eyes. Whether monkeys and humans—highly dependent on the ability to recognize faces and interpret facial expressions—have developed a separate system for face recognition is not settled. Selective loss of face recognition—**prosopagnosia**—sometimes occurs after lesions of the temporal lobe and would seem to suggest the existence of a separate “face” system. Further, fMRI studies show that activation in the **fusiform gyrus** in the inferotemporal cortex is associated with face recognition in humans (see Fig. 16.26). Other data, however, are more compatible with a general network for object identification that is used also for face recognition. Thus, objects other than faces can activate the fusiform gyrus, and, conversely, face recognition is associated with activation of several sites outside the fusiform gyrus. Because facial recognition is so important for social interactions and is used intensively from birth, presumably a larger proportion of neurons in temporal association areas become specialized for faces than for identification of other objects.

The **medial temporal lobe** and its importance for learning and declarative memory are discussed in Chapter 32. In addition, **lateral parts** of the temporal lobe is necessary for **semantic memory**—that is, knowledge about facts, meaning of words, objects, and so forth, which was acquired some time ago.

Symptoms after Lesions of the Temporal Cortex

Bilateral damage of the temporal lobes produces a syndrome dominated by pronounced **amnesia**.⁴ The amnesia can be ascribed largely to the destruction of the hippocampal formation and neighboring areas in the parahippocampal gyrus. In addition, certain emotional changes are presumably caused by the concomitant destruction of the amygdala located in the tip of the temporal lobe. These aspects are discussed in Chapter 31. In addition, the patients become very **distractible**: they have difficulty maintaining their attention on a certain stimulus or task. Finally, psychic blindness or **visual agnosia** is a typical symptom of temporal lobe lesions that affect the **inferotemporal parts**. The patient is unable to recognize objects and persons she sees, even though her vision is normal. As for other association areas, it is the interpretation of sensory information that is deficient, not the sensory experience as such. Information about size and shape of objects may nevertheless be available to the posterior parietal cortex to be used in movement control, as discussed in Chapter 16, under “Consciousness and Visual Experience.”

The Insula

The insula, sometimes called the fifth cerebral lobe, is mentioned in several chapters. This is because the insula, as evident from positron emission tomography (PET) and fMRI studies, participates in a wide range of cortical networks. Thus, activity changes in the insula occur in relation to somatic and visceral sensory processes, emotional regulation, and aspects of bodily awareness. This part of the neocortex is hidden at the bottom of the lateral fissure (see Figs. 6.29, 6.30, 14.9, and 31.9), and consists of several cytoarchitectonic subdivisions. Anatomically, the insula is characterized by receiving all kinds of sensory information and by its extensive corticocortical connections with major parts of the cortex. Further, there are ample connections among the subdivisions of the insula. It resides at the junction of the frontal, parietal and temporal lobes and has reciprocal connections with all three. This concerns, for example, the orbitofrontal cortex, the cingulate gyrus, the parahippocampal gyrus, the temporal pole, the superior temporal sulcus, premotor areas, SII, and the posterior parietal cortex—that is, regions involved in a wide specter of behaviors and mental processes.

4 The constellation of symptoms that occur after bilateral destruction of the temporal lobes is named after the two American neurosurgeons Klüver and Bucy (1937) who first described it in monkeys. Besides amnesia, the animals lack emotional responses and aggressive behavior (increased tameness), and they withdraw from social contact. Furthermore, the syndrome includes visual agnosia, a tendency to examine all objects by mouth, a tendency to pay attention to all visual stimuli, an irresistible urge to touch everything, and hypersexuality.

The insula receives afferents from several thalamic nuclei (Ventral anterior, Ventral posteromedial, Centromedian, VPM, CM, and some other nuclei). The insula also has reciprocal connections with the amygdala.

We discuss the insula in Chapter 14 because it is among the cortical regions activated by noxious stimuli and is an essential part of the network responsible for the experience of **pain**. Further, the anterior insula is activated in conjunction with strong **emotions** (especially disgust) and is presumably a part of a network for regulation of affect.

The insula receives not only sensory signals from somatic structures, but is consistently activated both by nonpainful and painful enteroceptive stimuli. For example, nonpainful distension of hollow organs such as the stomach and the esophagus activated the insula associated with the subjective feeling of fullness. Further, the awareness of one's own heartbeats is associated with activation of the insula. Indeed, the insula may thus play a particular role in our **awareness** of the state of our internal organs. However, its role does not seem to be limited to monitoring the internal organs and evoking subjective feelings referred to them. Thus, the awareness of voluntary movements, and especially the feeling of **body ownership**, involves a network that probably includes the insula (see Chapter 18). The contribution of the insula in this respect presumably depends on its integration of **proprioceptive**, **vestibular**, and **motor signals**.

Finally, the anterior part of the insula receives **olfactory** and **gustatory** signals and presumably contributes to the integration of these modalities, and their further integration with other enteroceptive signals.

Task-Specific Networks Integrate and Analyze Information

The brain receives innumerable pieces of information, which, to a large extent are treated in separate systems. For example, separate neuronal populations encode different features of objects, such as color, form, movement, surface texture, heaviness, and so forth. With regard to representation of space, the brain appears to possess several “maps.” Yet, we experience ourselves and our surroundings as entities, not as isolated fragments. How can this paradox be explained? This **binding problem**—that is, how various bits of information represented in different parts of the cortex are integrated in the brain—is closely linked with the problem of consciousness (see also Chapter 16, under “Consciousness and Visual Experience,” and Chapter 26, under “Neurobiological Basis of Consciousness”). The brain must possess the ability to integrate, almost instantaneously, the activity in numerous specialized neuronal groups, each representing different features of, for example, a visual scene. To be useful, the integrated

“picture” must furthermore be put into a meaningful context to form the basis for appropriate actions.⁵ Imagine, for example, the continuous stream of changing information that must be evaluated and acted upon when driving a car in heavy traffic. No area appears to receive all necessary information. Rather, it seems likely that neuronal groups in many parts of the cerebral cortex are interconnected in task-specific networks. The vast number of association fibers must obviously be essential in this respect. There is now much evidence that **synchronization** of activity in large-scale networks may be the substrate for the binding together of related pieces of information—and thus for our conscious experience of our environment and ourselves. It should be emphasized that while the anatomic connectivity forms the “hardwiring” of the networks, their functioning depends on dynamic, moment-to-moment fluctuations in synchronized activity. Presumably, engagement of specific networks shifts in the time scale of milliseconds. For example, we know from everyday experience how our attention shifts instantaneously. Another example concerns viewing of ambiguous pictures; the experience (e.g., duck or rabbit) changes with no time delay in spite of unchanged sensory input.

The “Default-Mode Network”

Among the cortical networks so far identified, the so-called **default-mode network** seems to have a special position. It is characterized by consistently *decreased* activity during goal-directed tasks, while it is active when the person's attention is not directed to any specific task. The network comprises lateral parts of the parietal cortex and regions on the medial aspect of the hemisphere (posterior cingulate gyrus and adjoining posterior regions, and parts of the medial prefrontal cortex). The activity of the default-mode network is thought to be related to **introspection**. In support of this assumption, an fMRI study showed increased activity in the default-mode network when the subject contemplated a moral dilemma (requiring minimal cognitive engagement) whereas a color-word interference task produced deactivation (Harrison et al. 2008). Nevertheless, the moral dilemma situation produced a pattern of activity within the network that differed from the activity in an eyes-closed resting state.

Disturbances of the default-mode network-activity have been found in fMRI studies of several groups of patients (e.g., chronic pain, attention-deficit/hyperactivity disorder [ADHD], and mental diseases).

⁵ It takes several hundred milliseconds from the arrival of sensory signals at the cortical level to perception, as first shown by Libet (1991) with stimulation experiments in humans. This might perhaps reflect that in this situation it takes some time to bring the necessary networks in a state of synchronized activity.

For example, chronic pain patients show less than normal deactivation related to task performance.

The Cerebral Cortex and Mental Disease

It is to be expected that diseases affecting complex functions such as emotions, personality, sense of reality, and thought would involve alterations in many parts of the brain, as well as in many neurotransmitters. Indeed, attempts to explain mental illnesses by malfunction in a single brain “center” or of one transmitter have not been successful.

Evidence of changed structure, metabolism, or neurotransmitters has been found most consistently in the **prefrontal cortex** and several **limbic structures** in patients with mental disorders. However, altered neuronal activity in the prefrontal cortex (or any other structure) does not tell us that the primary pathology is there. Thus, the alterations in one area may be secondary to changed activity in other areas with which it is connected. Considering the many connections between the prefrontal cortex and the amygdala, ventral striatum, and hippocampal formation (among others), it seems likely that even if the primary pathology should arise in only one of these structures, the symptoms would be due to malfunctioning of the whole **network**. Interestingly, computational models of schizophrenia suggest that a basic problem may be that networks (especially prefrontal ones) are unstable due to a low signal-to-noise ratio. The latter may be due to faulty dopamine actions, as dopamine normally would stabilize networks by increasing signal-to-noise ratio (by acting on D_1 receptors). Presumably, the final symptomatology in a disease such as schizophrenia would reflect both the dysfunction caused by neuronal pathology and the attempts by the rest of the brain to cope with the disturbed functions. This would be so, regardless of whether schizophrenia turns out to be due to defective receptor genes, a prenatal disturbance of neuronal migration, or (most likely) a combination of many factors.

Ever since the first accurate description of schizophrenia, it has been postulated that the disease is caused by alteration of the frontal lobes. This assumption was partly based on similarities between the symptoms in schizophrenia and in cases of frontal lobe damage. Measurements of regional blood flow lend support to the theory that the prefrontal cortex may be involved in some manner. Thus, many schizophrenics have abnormally low blood flow in the prefrontal cortex at rest, and various tasks that give increased flow in normal people failed to do so in these patients. In homozygous twins discordant for schizophrenia, the affected twin was found to have slightly smaller volume of the cerebrum and of the hippocampus than the other, and significantly less activation of the dorsolateral prefrontal

cortex during cognitive tasks. One may ask, however, whether the symptoms are caused by the abnormal prefrontal activity, or whether the low activity is due to other parts of the brain disturbing the execution of cognitive tasks in the prefrontal cortex? In contrast to some other studies, no signs of abnormal brain asymmetry or cortical structure were found in this twin study. The role of the ventral striatum—especially the nucleus accumbens—as a target for antipsychotic drugs was discussed in Chapter 23, under “The Ventral Striatum, Psychosis, and Drug Addiction.” The fact that the drugs commonly used for the treatment of psychotic disorders are antidopaminergic (primarily D_2 antagonists) gave rise to the hypothesis that schizophrenia is caused by disturbances of dopamine actions pre- or postsynaptically. Recent studies have strengthened this theory—for example, by showing an increased level of dopamine receptors in the brains of schizophrenic patients. Other monoamines may be involved in schizophrenia as well, and PET studies suggest that untreated patients have reduced glutamate concentrations in the prefrontal cortex. Especially altered *N*-methyl-D-aspartate (NMDA)-receptor-mediated transmission has been reported in several studies. Needless to say, the interpretation of such findings is not straightforward.

Many questions remain about the etiology and pathophysiology of schizophrenia. Do defects of neurotransmitters and receptors produce structural abnormalities, or are abnormal brain networks the primary cause? There is evidence to suggest that the primary defect in schizophrenia is neurodevelopmental, but a hypothesis of neurodegeneration (based on evidence of slight but progressive loss of brain tissue) also has its proponents. We know that environmental factors must contribute, although we know little about their nature and how they interact with genetic predispositions.

LANGUAGE FUNCTIONS AND “SPEECH AREAS” OF THE CEREBRAL CORTEX

Clinical observations in the nineteenth century led to the identification of two so-called speech areas in the left hemisphere (Fig. 34.5). The anterior area is named after the French physician Paul Broca, who in 1861 described loss of speech—*aphasia*—caused by a lesion in the left frontal lobe just in front of the motor face area. The posterior speech area is named after the German neurologist Carl Wernicke, who discovered in 1874 that one of the clinically observed kinds of aphasia was associated with a lesion in the posterior part of the superior temporal gyrus. Aphasia is **defined** as loss (or disturbance) of speech due to a brain lesion. That speech depends almost entirely on only one hemisphere—in most people, the left—is the most marked and best-known example of **lateralization** (of function)

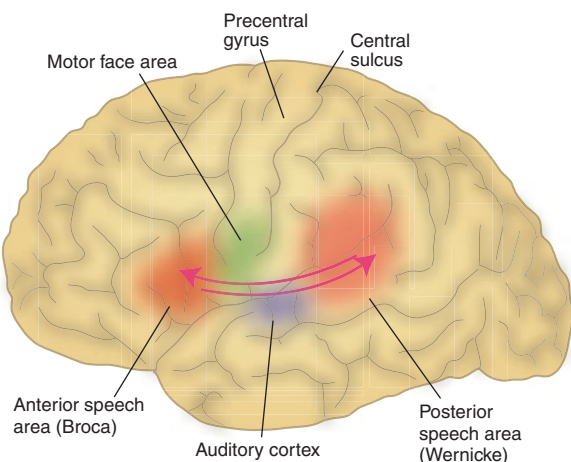


FIGURE 34.5 *Speech areas of the human brain.* Lesions of the anterior speech area (of Broca) produce predominantly motor aphasia, whereas lesions of the posterior speech area produce mainly sensory aphasia. The posterior speech area as shown here comprises parts of the inferior parietal lobule that were not included in the area described by Wernicke. Long association fibers interconnect the two areas, and these connections explain why a lesion between the two speech areas can produce aphasia (so-called conductance aphasia). Additional areas of the cerebral cortex are also involved in language processing, as shown with PET and fMRI.

or **hemispheric dominance**. We return to the topic of lateralization later.

What are termed “speech areas” should, more correctly, be termed “areas of aphasia,” since we know that their destruction produces various disturbances of language functions (aphasia), but we know less of how these areas contribute to the normal production of language and speech. Further, what rather broadly is referred to as the “anterior” and “posterior” speech areas (Fig. 34.5) consist in reality of functionally different subregions.

fMRI and PET show that various tests of language functions activate regions corresponding roughly to the areas of Broca and Wernicke, but also that other parts

of the cortex participate. This may not be surprising since language depends on several different processes, such as storage of words in short-term memory, phonologic (sound) and semantic (meaning) processing in relation to the “lexicon” in long-term memory, arranging words into sentences, and the issuing of commands to motor areas about sound production. Silent reading or repetition of words activates primarily the anterior region (Fig. 34.6). Specific tests for **semantic** analysis activate extensive areas in the temporal, prefrontal, and inferior parietal cortices, including Broca’s and Wernicke’s areas (primarily in the left hemisphere). Tests of **phonologic** analysis (like choosing matching words by sound similarity) activate partly the same and partly different areas than semantic tests. As for other complex mental functions, the main tendency is that extensive regions of the hemisphere participate in an extensive network for language and speech, and that different tasks activate overlapping regions.

Different parts of the **network** for language processing were suggested by Shalom and Poeppel (2008, p. 125) to specialize as follows: “. . . memorizing (learning new and retrieving stored primitives) in the temporal lobe, analyzing (accessing subparts of stored items) in the parietal lobe, and synthesizing (creating combinations of stored representations) in the frontal lobe.” Further, there is evidence of a specialization within each region. For example, in Broca’s area (or region), the superior part (closest to the ventral premotor area) appears to specialize in phonological processing, the midregion in syntactic processing, and the inferior part in semantic processing. A functional subdivision of Broca’s area would agree with anatomic data, since it consists of several cytoarchitectonic areas with different connections.

Are There Brain Systems that Are Specific for Language?

A central question is whether language is produced by networks specialized for only this function, or by a

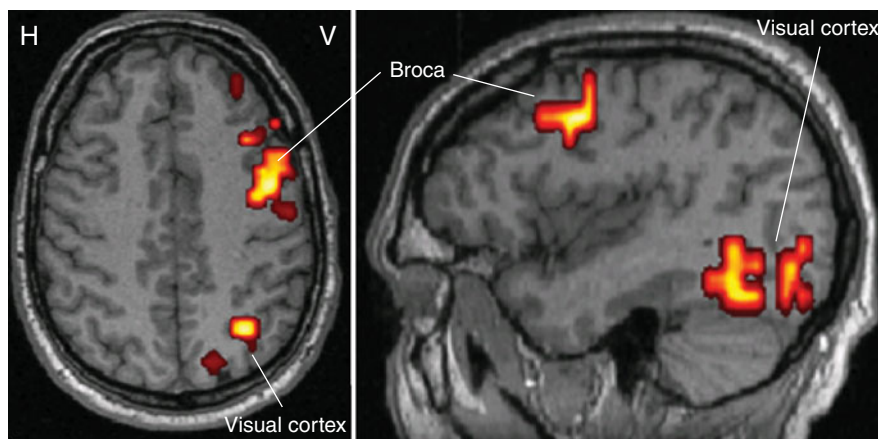


FIGURE 34.6 *Activation of Broca’s area as demonstrated with fMRI.* An anatomic (T1-weighted) MRI is produced first, thereafter color-coded activity data are superimposed on the brain image. Activity in the visual cortex occurs because the test of language functions starts with reading of word. (Courtesy of Dr. S.J. Bakke, Rikshospitalet University Hospital, Oslo, Norway.)

more general system taking part also in other cognitive functions. Although rare, over the years many patients have been described with selective loss of language functions without other cognitive defects, and vice versa. Such observations have been taken to support the **specificity hypothesis**. Still, PET and fMRI data show considerable overlap among cerebral regions activated by language tasks and by other cognitive tasks. Further, connectionist-modeling (computer-based models of neural networks) shows that the **multipurpose hypothesis** is not incompatible with the evidence of selective defects from brain-damaged patients. Models can simulate language learning, such as learning to read words. Although such networks do not have segregated pathways for different functions, partial damage may produce selective visual or semantic defects (somewhat similar to a child with dyslexia). Such modeling experiments challenge “the commonly held assumption that the fractionation of behaviour reflects an underlying fractionation of the brain systems that control such behaviour” (Nobre and Plunkett 1997). Thus, a specific impairment after a brain damage does not mean that the brain possess a specific module responsible for the lost function.

Aphasia

There is an enormous literature dealing with the disturbances of speech and language, and there are numerous classifications of aphasias. Also, there is no lack of hypotheses for the cause, whereas understanding of the basic mechanisms underlying language and speech is still limited. This should come as no surprise, however, considering the complexity of the function in question.

Very schematically, there are two main types of aphasia. The simplest type is the so-called **motor aphasia**. This occurs most often after destruction of Broca’s area (the anterior speech area). The patient more or less completely loses the ability to speak and typically produces only single words in a sort of telegraphic style. The few words used may also be applied wrongly. Other names used for this type are **nonfluent aphasia** (because the speech becomes stuttering) and **expressive aphasia**. The understanding of language is usually preserved, whereas the production of speech is deficient. Nevertheless, there are no signs of pareses of the muscles involved in speech production. Often, motor aphasia is combined with **agraphia**—the inability to express language in writing.

In patients with **sensory or receptive aphasia**, the lesion usually affects more posterior parts of the hemisphere at the junction between the parietal, temporal, and occipital lobes—that is, in or close to the Wernicke’s or the posterior speech area (Fig. 33.5). Typically, the comprehension of language is most severely affected. The various elementary sounds are not properly put

together to form meaningful words and sentences. Words that are heard cannot be repeated. In contrast to motor aphasia, spontaneous speech is fluent, but sounds are often put together into meaningless words, and proper words lack relation to each other (“word-salad”). Usually, sensory aphasia is combined with **alexia**—the inability to read.

In reality, the pure forms of aphasia are very seldom encountered; in most patients there is a mixture of motor and sensory symptoms, with one or the other dominating. Often there are other symptoms as well, since lesions of the hemispheres are rarely confined to the speech areas. Further, similarities between word blindness (alexia, dyslexia) and visual agnosia may be caused by lesions of the parietal lobe and the inferotemporal cortex. Alternatively, elementary symptoms of aphasia—such as agraphia or alexia—may sometimes occur in isolation or in combination with other kinds of symptoms (see Gerstmann’s syndrome, described previously).

The relationship between specific aphasic symptoms and the anatomic location of a brain lesion is not absolute. Virtually all forms of aphasia have been described after lesions in unexpected parts of the brain. Thus, we can only say that the probability of speech disturbances is highest when the lesion affects one or both of the areas depicted in Fig. 33.5 (or the association fibers interconnecting the anterior and posterior speech areas).

THE DIVISION OF TASKS BETWEEN THE HEMISPHERES

The fact that the two hemispheres are connected by afferent and efferent fiber tracts with the opposite body half is proof that there *is* a division of tasks between the hemispheres. We have mentioned speech as a function that is largely taken care of by one hemisphere (the left), and here we discuss other examples of lateralization of functions. This does not mean, of course, that the two hemispheres are independent units: the **commissural connections** ensure that information reaching one hemisphere also reaches the other and that commands issued to lower parts of the brain also are known to both hemispheres. Complex functions are usually carried out by cooperation of the two hemispheres, and, as discussed, lesions of the association areas often have to be bilateral to produce clear-cut functional deficits.

Are the Two Hemispheres Anatomically Different?

Many differences have been reported but most of them are small and, further, based on small samples. This, together with the large individual variations in brain anatomy (e.g. with regard to size of cortical areas and total brain size) might explain why many reported

differences have not been confirmed by later studies. In general, there are no anatomic data—with regard to neuronal numbers, synaptic densities, size of areas, and so forth—that “explain” why one hemisphere performs certain tasks better than the other (hemispheric specializations).

The most robust anatomic difference between the left and the right hemisphere in humans concerns the upper face of the temporal lobe. The **temporal plane** (planum temporale) in the vicinity of the auditory cortex is reported to be more extensive on the left than on the right side in about 70% of the population (see Chapter 17, under “Asymmetrical Organization of the Auditory Cortex in the Temporal Plane”).⁶ The difference appears to be present before birth, and recent studies indicate that the lateralization of language functions is determined prenatally. It is well known that early brain damage (in infancy, before language has been acquired) has less severe effects on language functions than lesions that occur later. Because, obviously, in such early cases of brain damage the right hemisphere takes over the language functions, it was concluded that language function is not initially lateralized. However, a more likely interpretation is that even though there is an inborn tendency for lateralization of speech, at an early stage the right hemisphere has not been fully occupied by other tasks and therefore can substitute for the left hemisphere. At later stages, both hemispheres are fully used and specialized for specific tasks and therefore are unable to take over new complex functions. Incidentally, the capacity of the right hemisphere to develop normal language in young children favors the view that language is not dependent on a language-specific, genetically determined network.

Function of the Commissural Connections: The Corpus Callosum

Here we mention a few examples of the significance of the commissural connections. When the optic nerve fibers that cross in the optic chiasm are cut (see Fig. 16.16), signals from one eye reach only the hemisphere of the same side. After such an operation, monkeys are trained in a **visual discrimination task** (to distinguish a triangle and a circle to obtain a fruit reward) with a patch that occludes vision in the left eye. The learning must depend on processes taking place in the right hemisphere, which receives visual information from the right eye. When the monkey has learned the task with the right eye, the occlusion is reversed. Nevertheless, even when using the left eye, the monkey

solves the task as well as when using the right eye. Thus, the left hemisphere also has learned the task. This must be dependent on an effective transmission of visual information from one hemisphere to the other, as can be demonstrated by cutting the corpus callosum (and the anterior commissure) before the discrimination training starts. Then the animal learns the task only with the right hemisphere when the left eye is occluded. The transmission of visual information from one hemisphere to the other in this experiment must depend on commissural connections between the extrastriate and probably the inferotemporal areas, since the striate area lacks commissural fibers (Fig. 33.13).

Corresponding experiments have shown that **tactile** and **kinesthetic** signals are also transferred through the corpus callosum. A monkey with transection of the corpus callosum (and the anterior commissure) is trained to open a box with the right hand only (without being allowed to see the box). After some training, the opening is performed swiftly. If the monkey is then prevented from using the right hand, the task has to be learned over again with the left hand—no learning had taken place in the right hemisphere (which controls the left hand). A monkey with an intact corpus callosum uses both hands with identical dexterity to solve this task, even though only one hand was used during the training period.

There is a certain **topographic arrangement** of the commissural fibers within the corpus callosum. As one might expect, the posterior parts are necessary for the transfer of visual information, whereas the anterior and middle parts are necessary for transfer of somatosensory signals. (See Chapter 26, under “Neurobiological Basis of Consciousness” with a description of a patient with damage of the middle part of the corpus callosum and abolished transfer of tactile information.)

The ample commissural connections make it possible for the two hemispheres to **specialize** and to share tasks between them. Both are not required to be equally good at all tasks. Nevertheless, they keep each other constantly informed (just as one would expect of hospital specialists sharing the responsibility for one patient). The brain, even though in a sense consisting of two anatomic parts, functions as a unit for the whole body and our extrapersonal space. The degree of hemispheric specialization or lateralization of functions can be studied only after eliminating the callosal transfer of information.

Cerebral Lateralization and Dominance

Data showing that the two hemispheres are different have received much attention since the early 1970s, and many simplistic statements about the functions of the right and the left hemispheres have been put forward.

⁶ In more than 90% of persons, the language function depends on the left hemisphere; thus, the relationship between speech lateralization and anatomic asymmetry is not absolute.

Recent studies provide a much more complex picture, however. A specialist in the field of cerebral lateralization wrote a few years ago: “The time has come to put the brain back together again.” Another scientist launched the expression **dichotomania** about the urge to equate the many examples of duality in human nature with the two halves of the brain (left–right): scientist–artist, conscious–subconscious, rationalism–mysticism, masculine–feminine, and so forth. The truth is that both hemispheres take part in most functions; the differences concern mainly how efficient they carry out individual processes, like elements of language functions, emotional processing, visual object recognition, and so forth. For example, many studies show that the **left hemisphere** usually is superior with regard to **analytical and logical thinking** as expressed verbally and in numbers, while the **right hemisphere** is superior with regard to **spatial abilities**, the comprehension of complicated patterns, and **drawing**.

Studies of Split-Brain Patients

A wealth of information on the topic of hemispheric specializations (lateralization) has been provided by the study of so-called **split-brain** patients—patients in whom the **corpus callosum** has been transected (this is done in severe cases of epilepsy, to prevent spread of the abnormal discharges from one hemisphere to the other). The American Roger Sperry was awarded the Nobel Prize in 1981 for his pioneering studies of split-brain patients. Even though lateralization is probably most marked in humans, there is much evidence that it also occurs in animals (e.g., the ability to sing depends on cell groups in the left side of the brain in birds).

Split-brain patients manage well in everyday life, mainly because visual information reaches both hemispheres (because we move the gaze constantly) and there are some bilateral sensory and motor pathways. These patients get into trouble, however, if, for example, somatosensory information is not supplemented with visual information. The preceding example of the commissurotomy monkey and tactile learning is relevant for split-brain patients, too. In some situations, conflicts may arise between commands issued from the two hemispheres; for example, the left hemisphere may command the right hand to start dressing, whereas the left hand is ordered to undress.

Studies of split-brain patients raise interesting questions, such as whether the two hemispheres have independent consciousness and what the relation is between consciousness and language and between intelligence and language (see also Chapter 16, under “Visual Awareness and Synchronized Network Activity” and Chapter 26, under “Neurobiological Basis of Consciousness”). No simple answers are available to

such questions addressing phenomena in the transition zone between neuroscience, philosophy, and religion. To argue that humans consist of two personalities, one in each of the hemispheres, is a gross oversimplification, of course. The normal cooperation and interaction between the hemispheres is so intimate that our mental life and behavior are caused by their collective activities.

Lateralization of Language

When one hemisphere is most important for a certain function, we say that it is **dominant** for that function, whereas the other hemisphere is **recessive**. The most clear-cut example of such cerebral dominance—or, in other words, lateralization of function—is speech, as mentioned above. For most people, even for most left-handers, the left hemisphere is responsible for most aspects of language functions. About 95% of right-handers have left hemisphere language dominance, while the corresponding number for left-handers is about 70% (there is obviously not a strong correlation between the lateralization of speech and hand preference).⁷ Several kinds of investigation have confirmed the lateralization of language. Studies of **split-brain** patients are especially instructive in this respect. They confirm, among other things, that the right hemisphere is mute in most people (even though it may express single words when strong emotions are aroused). When a split-brain patient is asked to identify with the right hand an object that is not seen, he can easily tell the name of the object, what it is used for, and so forth. This is because the tactile information comes to the left, speech-dominant hemisphere. When the left hand is used for the same test, however, the patient is unable to name the object, because the information reaches only the “mute” right hemisphere. The patient nevertheless shows signs of appropriate emotional reactions to the object. That the right hemisphere “understands” the nature of the object is further supported by other experiments in which the right hemisphere is presented with a picture of, for example, a key. Even though the patient cannot say anything about the object, he nevertheless picks out with the left hand a key among several objects (which are not seen). Such data show that the **right hemisphere**

⁷ The numbers concerning language dominance come mainly from studies in which one hemisphere was temporarily anesthetized by injection of a barbiturate into the internal carotid artery. In general, recent brain-imaging studies agree with these data. One fMRI study found that among 50 right-handed persons, 96% had largely left hemisphere activation when performing a language task (word finding); 4% had bilateral activation, but no one had larger right than left activation. In contrast, 76% of left-handers had left hemisphere activation with the same test, 14% had bilateral activation, and 10% had right hemisphere activation. Thus, it appears that only about 10% of left-handers have right hemisphere dominance for language.

can understand concrete language. It understands both speech and writing of this kind but cannot express understanding through language—only through action. As mentioned, the right hemisphere can utter a few words especially when they are emotionally loaded, while it cannot manage abstract or rare words or grammatical analysis.

Not all aspects of language function are localized to the dominant hemisphere. The modulation and melody of the sounds of speech, **prosody**, appears to largely depend on the right hemisphere, as witnessed by several clinical reports. Thus, in some patients who suffered a right hemisphere stroke, prosody was changed or reduced without concomitant aphasia. Brain-imaging studies show activation in the right hemisphere in tests for perception of prosody, notably in the region corresponding to the Broca's area and in the superior temporal gyrus (but also some activation of the left hemisphere). Patients with loss of prosody may also be unable to judge the emotional aspects of the speech of other persons; for example, they cannot decide whether the person is sad or happy. Such an **intonational agnosia** may have serious effects on the social life of the patient. The importance of the prosody illustrates that much of what we regard as verbal communication is, in fact, nonverbal.

Lateralization of Music

With regard to the ability to appreciate and express **music**, there is no simple division of labor between the hemispheres, although it has been assumed that the right hemisphere is most important. Indeed, that the **perception of a melody** depends mainly on the right hemisphere was supported by a study using the Doppler technique to measure changes of total blood flow to the hemispheres during various tasks. The right-sided dominance was true only for nonmusicians, however: professional musicians showed left hemisphere dominance for the same task that was presented to the nonmusicians. Listening to **rhythm** activated the left hemisphere most strongly in both groups. Furthermore, left hemisphere activation was relatively larger when the person listened attentively, trying to discriminate musical elements, rather than having the music as a background. **Imagining** a familiar tune was found with PET to activate association areas around the right auditory cortex and frontal regions on both sides. The **supplementary motor area**—which is important for rhythmic and sequential movements—is activated when imagining tunes, perhaps because there is a motor element in music imagination.

Amusia most often occurs together with aphasia, but it has also been reported to occur in isolation. This suggests that language and music use largely separate parts of the cortex, although they appear to lie close together.

This is supported by PET studies of musicians practicing sight-reading (i.e., at the same time reading and playing an unfamiliar score).

Ear and Visual Field Dominance

A certain degree of “ear dominance” exists in most people, corresponding to speech lateralization—that is, the right ear is dominant for most people. This phenomenon can be studied by use of so-called **dichotic listening**. Two words are presented at the same time, one to each ear. Afterward, most people say that they heard the word presented to the right ear. **Visual field dominance** has been described in studies in which different visual stimuli are presented to the two hemispheres simultaneously. With regard to written words and letters, there is a tendency to prefer those presented in the right visual field (that is, those transferred to the left hemisphere). For **face recognition**, the reverse situation appears to exist for most people, as can be demonstrated by the presentation of so-called **chimeric portraits** composed of two left halves and two right halves, respectively. The person is asked which of the chimeric portraits most resembles the original (authentic) portrait. Most persons claim that the chimeric portrait consisting of two right facial halves most resembles the original. This is taken to suggest that the right hemisphere dominates in the analysis of faces and other complex visual patterns. Another indication of this is that when the **shape of letters** is made sufficiently ornate, the right hemisphere appears to become necessary for their interpretation.

Lateralization of Hand Function

With regard to lateralization of hand functions, the hemispheric differences are less clear-cut than for language. It is not a question of the ability to use the hand, but a matter of preference of one hand for most or all tasks. Even though hand preference is inheritable, there are also strong social factors that contribute to the final outcome of hand preference—for example, in writing. There is most likely a gradual transition with regard to the strength of hand preference, from those with a strong tendency to use the right hand for all tasks if possible (writing, drawing, use of tools, eating, and so forth) to those with an equally strong tendency to use the left hand. The latter group probably constitutes 2% to 3% of the total population. Hand preference starts to become expressed from the second year of life and is usually finally established at the age of 5 to 6.

Lateralization of Emotions

Early observations of split-brain patients suggest that the **right hemisphere** is dominant for the expression

of emotions, but further studies show that the right hemisphere does not dominate all aspects of emotional behavior. Thus, the two hemispheres appear to be specialized for specific aspects of emotions. We mentioned **prosody** as an example of right hemisphere dominance. Overall, the right hemisphere appears to be the best at perceiving emotional expressions, whereas both hemispheres are involved in the experience and expression of emotions. The left hemisphere may be the best judge of certain kinds of emotional expressions, however. Some data have been interpreted to show that the right hemisphere is dominant as to the experience of strongly negative feelings (and the left as to positive feelings). Patients with **strokes** affecting the **left** hemisphere tend to have more depressive reactions than patients with corresponding right-sided lesions (the right hemisphere understands the agony of the left?). **Right** hemisphere lesions, especially when they affect the frontal lobe, appear to have a stronger tendency to produce a somewhat inadequate elevation of mood.

Further Examples of Hemispheric Specializations

Studies with detailed analyses of specific aspects of broader categories of cerebral functions reveal that the division of labor between right and left is more complicated than is apparent from the first split-brain observations. Sophisticated studies of visual perception show that the right hemisphere is superior to perceive and remember **specific characteristics** of objects (for example, the face of a person), whereas the left is better at **categories** (a face versus other kinds of objects). A different picture emerged in a study comparing patients with lesions in the superior temporal gyrus as to their ability to **identify letters**. Those with right-hemisphere lesions had difficulties with identification of a letter when it was composed of many small ones, but they easily perceive the small letters. Those with a left-hemisphere lesion had difficulties with identification of the small letters, but they easily identified the big one. Thus, the right hemisphere is good at identifying the **overall shape**, whereas the left is good at seeing the **details**. It may seem paradoxical that the left hemisphere is specialized both for identification of broad categories and details (and that the right hemisphere is specialized for identification of specific properties and the overall shape). Most likely, however, this reflects that different principles govern lateralization of visual and semantic memory. Another lateralization of a specific function does not fit with the usual right-left dichotomy of functions. Thus, the right hemisphere is best at **measuring distance** (e.g., the distance between a dot and a line), whereas the left hemisphere excels at judging **mutual positions** (whether the dot is above or below the line).

SEX DIFFERENCES AND THE CEREBRAL CORTEX

Gender and Cortical Structure

Many studies show that, on the average, men perform better than women on certain **spatial tasks** (shown most convincingly for imaginary rotation of a figure). Women, in contrast, excel on tasks that require **verbal fluency**, perceptual speed, and some fine-motor skills (many other cognitive differences have been proposed but few have been convincingly documented). Many speculative explanations have been offered. From an evolutionary point of view, it is now common to explain such sex differences by the living conditions during early human history, along with the different roles held by men and women. Although cognitive sex differences thus may have a genetic basis, environmental influences interact with genetic predispositions to produce the final cognitive make-up of the individual. Nevertheless, the average cognitive sex differences are small, and very much smaller than the variability among individuals of the same sex. As said by the Canadian psychologist Doreen Kimura (1996, 259): “In the larger comparative context, the similarities between human males and females far outweigh the differences.”

With regard to neuroanatomic sex differences, the most obvious is the difference in **brain volume**: the brain is on average 10% heavier in men than in women. Most, but not all, of this difference can be accounted for by different body weights. The **temporal plane** (involved in language processing) has been reported to be larger in women than in men. Whether this is causally related to sex differences in verbal fluency is so far unknown, however. Studies at a more detailed level offer many data but, unfortunately, conflicting results make it difficult to draw conclusions. Yet, it seems fairly well documented that women have a slightly thicker cortex in parts of the parietal and temporal lobes (as studied with MRI). On the other hand, one study reported higher synaptic density in men than in women. A morphometric study found no difference in cortical thickness but that men had on average somewhat higher **cortical neuronal density** than women ($117,000 \pm 31,000$ and $101,000 \pm 26,000$ per mm^2 , respectively). Even if this finding should be confirmed, the large individual variation within each sex makes any inferences about causal relationships doubtful. With PET and fMRI sex differences in brain activation patterns have been looked for. For example, one study found that men and women activated somewhat different parts of the brain when they were trying to find their way out of a (virtual) labyrinth. Apart from many regions activated in both sexes, men activated the left hippocampus, whereas women activated the right frontoparietal region. Other studies also point to gender differences in activation patterns but the interpretation

of such findings is far from straightforward. It is not advisable to make firm statements as to the biologic bases of gender differences on the basis of small statistical associations.

Sex Differences and Lateralization

Cognitive sex differences have been speculatively linked with more or less convincingly demonstrated sex differences in **lateralization** and brain structure. Some observations suggest, for example, that men have stronger lateralization of, for example, visuospatial abilities, whereas women to a higher degree use both hemispheres. (It is not obvious, however, that a strong lateralization gives a higher visuospatial ability; the reverse might just as well be the case.) It was then studied whether there might be sex differences in the cross-sectional area of the **corpus callosum**, which would correlate with differences in lateralization; that is, a corpus callosum with relatively few fibers was expected to correlate with a high degree of lateralization. Indeed, some studies reported that the corpus callosum is relatively larger in cross section in women than in men. These speculations have not been confirmed by further and more comprehensive studies, however. First, the reported differences as for visuospatial abilities constitute merely a few percentages of the individual variations among members of the same sex. Furthermore, the notion that men have more marked lateralization than women cannot be accepted as generally valid. Finally, several studies have not been able to confirm the sex differences in the size of the corpus callosum. They are at best small, whereas the individual variations again are surprisingly large. In fact, an inverse relationship between **brain weight** and cross-sectional area of the corpus callosum has been reported: smaller brains, regardless of sex, would be expected to have relatively larger corpus callosum. In monkeys the number of axons in the corpus callosum varies among individuals by a factor of 2. To complicate matters, one report claimed that the corpus callosum is about 10% larger in left-handers (and persons using both hands equally) than in right-handers. This does not support a correlation between superior visuospatial abilities, strong lateralization, and a small corpus callosum.

In **conclusion**, the available data do not support that sex differences in cognitive and other abilities can be explained by differences in lateralization.

Psychological Differences and Biology

Arguably, men and women differ more in other aspects of their psychology than cognition—for example, in aggression and emotional reactions. Some differences are expressed very early, including preferred activities. Most researchers today would ascribe such differences

to the combination of biologic and environmental factors.

In which respects are the brains of men and women different? Sex differences in the corpus callosum are very small at best, and can hardly explain the observed cognitive differences. The neuroanatomic differences in the structure of the cerebral cortex lack so far explanatory power. The most obvious microanatomic differences between the brains of men and women are found in the **hypothalamus** (see Chapter 30, under “The Hypothalamus, Sexual Functions, and Sex Differences”). Although the structural sex differences in the hypothalamus presumably relate mainly to differences in sexual functions, we cannot exclude sex differences also in hypothalamic influences on other cell groups, such as the **amygdala** and the **cerebral cortex**. Further, we know that **receptors for sex hormones** are found in many parts of the brain outside the hypothalamus (e.g., the amygdala and the prefrontal cortex, just to mention two regions related to emotions and emotional reactions). Experiments in rats and monkeys clearly show that **exposure to male sex hormones** during early critical phases of development leads to certain behavioral characteristics in later life. Indirect evidence suggests that this may be so also in humans—for example, regarding **spatial abilities**. Further, some cognitive functions have been shown to vary in the same individual with variations in the level of sex hormones (in both men and women). Thus, sex differences in distribution and densities of hormone receptors, in conjunction with different hormonal makeup, may form the basis for psychological sex differences. The blood level of hormones would then modulate the excitability of specific neuronal populations. In addition, subtle differences in the cerebral wiring patterns and synaptic organizations might contribute to the sex differences, as they would contribute to differences among persons of the same sex. These differences might be genetically determined or use-dependent, or most likely both.

Nature or Nurture?

The final performance of the brain is a product of genetic and environmental influences, as discussed in Chapter 9. Environmental challenges induce structural and neurochemical adaptations in the nervous system expressed through altered behavior. There is now an intense search for genes that influence social behavior. Such genes would be expected to participate in the establishment and refinement of the many task-specific networks discussed in this book. It is striking, however, that the search for genes with decisive influence on human behavior has not been successful. Rather, it appears that personality as well as cognition and emotions are under the influence of numerous genes, each providing a very small contribution to the final phenotype.

For example, while cognitive abilities (IQ) are among the most genetically controlled traits (40%–80% of the variability can be explained by heritage) it has proved very hard to find a connection between specific genes and IQ. Considering that, presumably, each of the many contributing genes is under epigenetic influence, the number of variables that determine our final behavior becomes astounding. Indeed, as said by Story Landis and Thomas Insel in an editorial in *Science* (2008, p. 821): “Genes code for proteins, not for behaviors.” They go on pointing out “. . . genomics is not destiny. Indeed, if genomic sequence ‘determines’ anything behaviorally, it determines diversity. It is important that we be wary about extrapolating from model organisms to humans. We must also avoid using small

statistical associations to make grand claims about human nature.”

In conclusion, all development—whether genetically determined or not—involves synaptic changes, and from the time of birth there is a dynamic interplay between genes and the environment in the establishment, maintenance and functional regulation of billions of synapses. A genetically determined disposition—for example, a certain temperament—evokes a certain kind of response from other people. The response induces synaptic and neurochemical changes in the brain as a basis for behavior adaptations. To sort out the mutual roles of nature and nurture in the formation of a person’s psychology and behavior then becomes virtually impossible.

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Six issues annually. Concise reviews of the development the last few years within a special subfield. Contains comprehensive reference lists. Key references are supplied with brief, focused abstracts.

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Neuron. Cell Press

Monthly. Each issue usually features one review article (basic neuroscience) and several mini-reviews of high standard, in addition to many research articles.

Scientific American. Scientific American Inc.

Monthly. Regularly one or more articles on various aspects of neuroscience. Requires less prior knowledge than the specialized neuroscience

review journals. Well-written, easy-to-follow articles with often superb illustrations.

The Neuroscientist. Williams & Wilkins

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Trends in Cognitive Sciences. Elsevier

Monthly. High-quality reviews covering neuropsychology.

Trends in Neurosciences. Elsevier

Monthly. Brief reviews of neurobiological topics in rapid development. Written by leading scientists. Requires fairly broad prior knowledge of neuroscience.

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CHAPTER 14: CENTRAL PARTS OF THE SOMATOSENSORY SYSTEM

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CHAPTER 23: THE BASAL GANGLIA

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CHAPTER 24: THE CEREBELLUM

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CHAPTER 26: THE RETICULAR FORMATION: PREMOTOR NETWORKS, CONSCIOUSNESS, AND SLEEP

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CHAPTER 32: THE HIPPOCAMPAL FORMATION: LEARNING AND MEMORY

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CHAPTER 34: FUNCTIONS OF THE NEOCORTEX

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