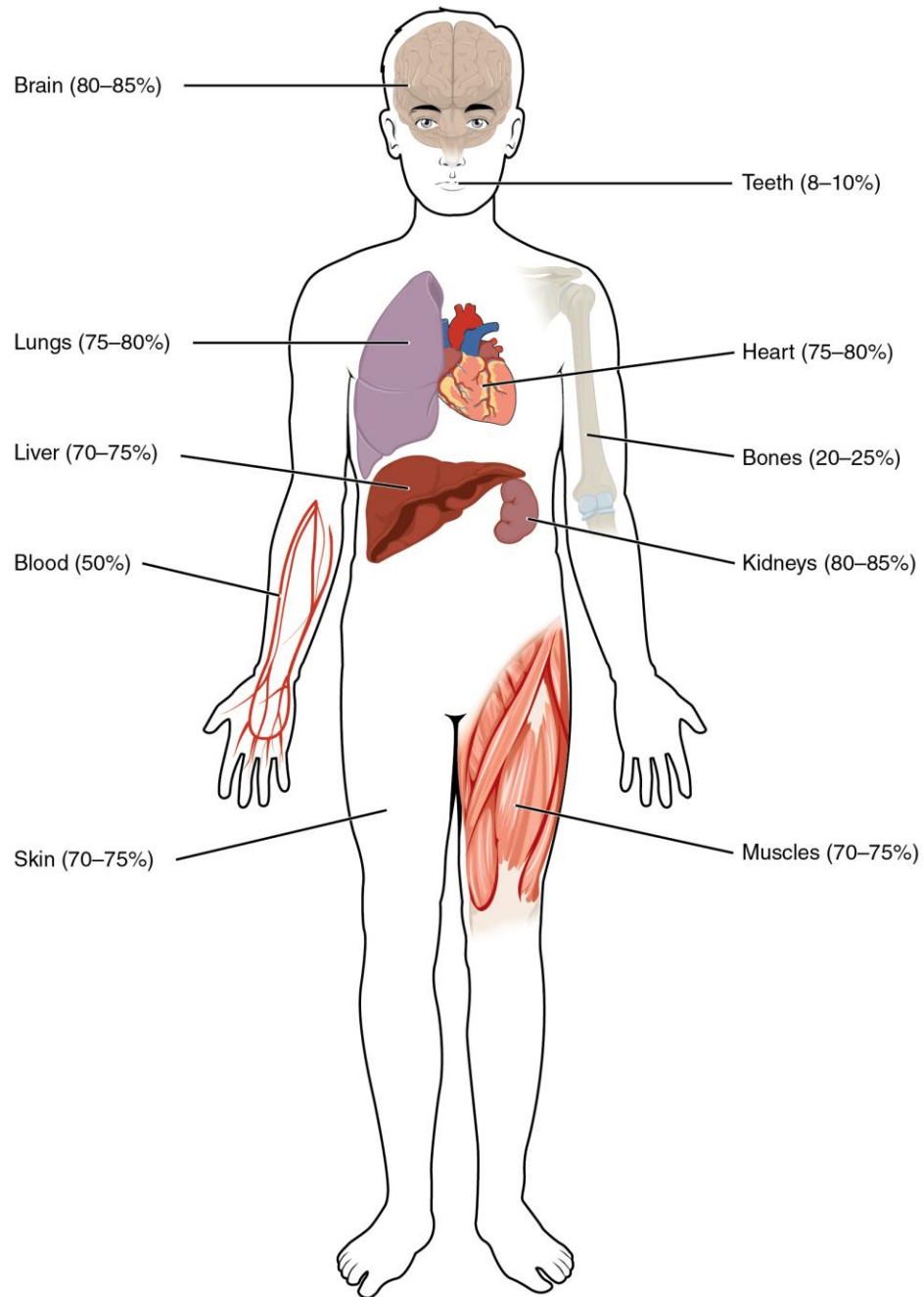
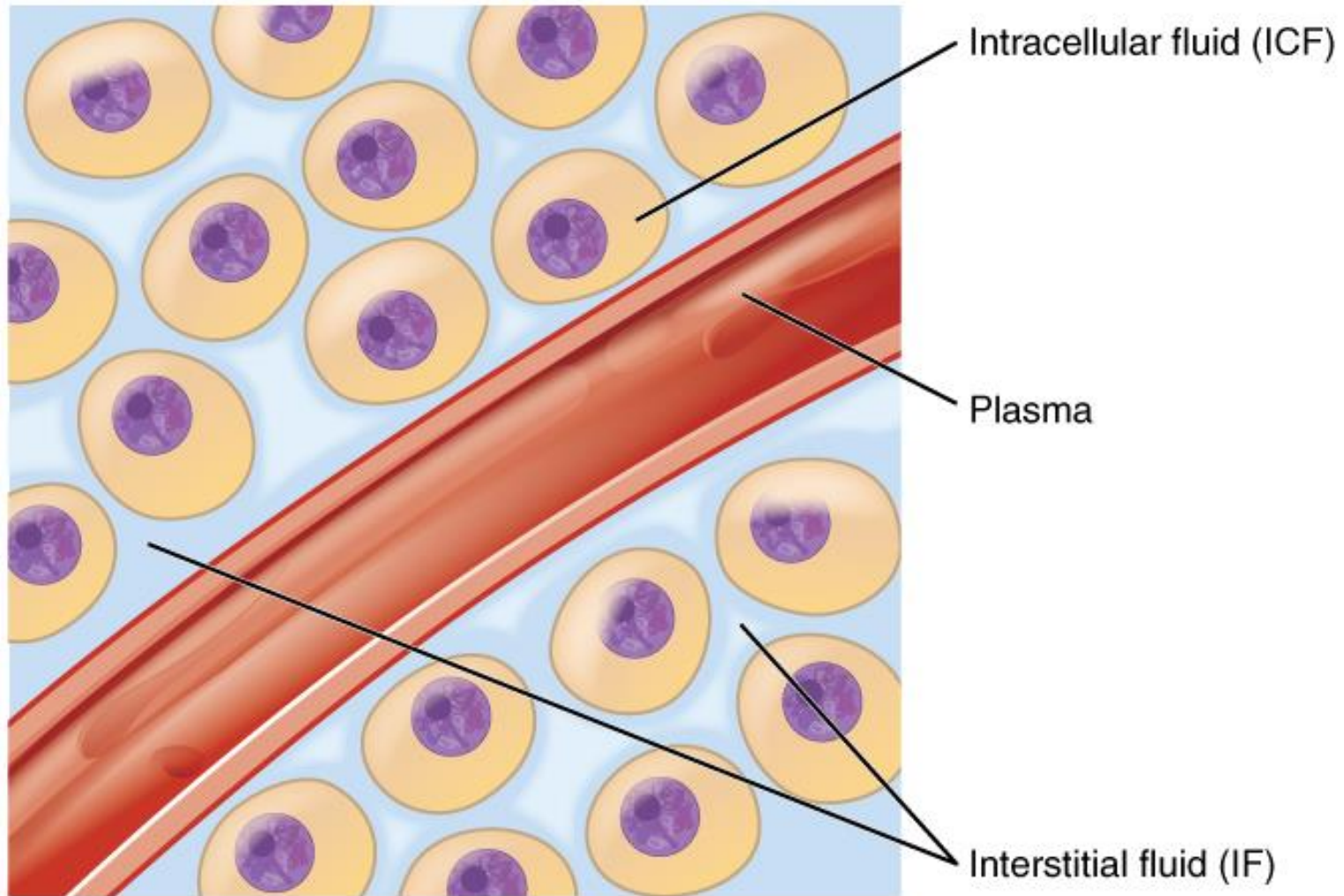


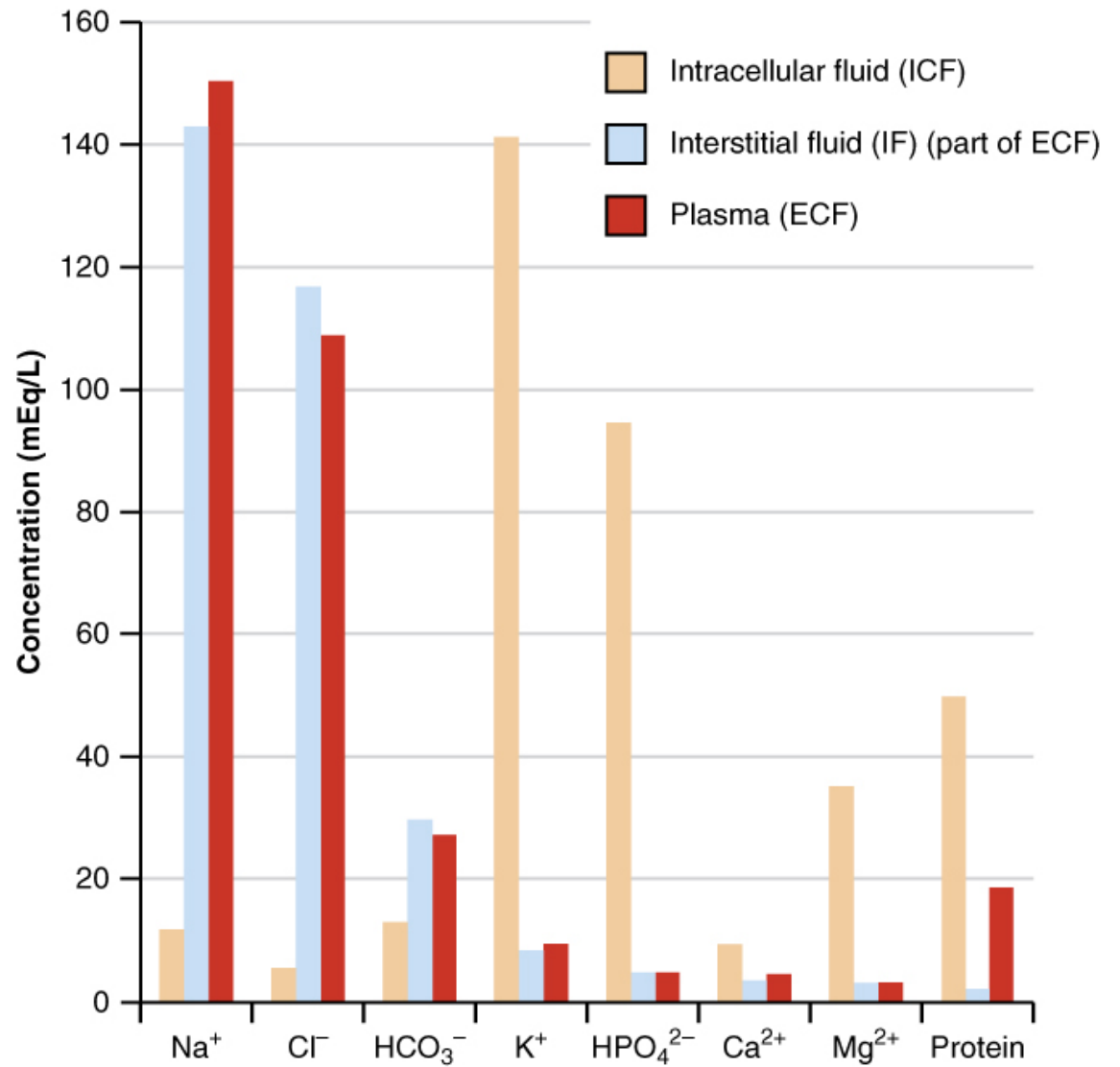
# Body Fluids and Fluid Compartment



# Fluid Compartment in the Human Body



## Composition of Body Fluid



**Filtration**

**No net movement**

**Reabsorption**

Arterial end  
net filtration pressure  
= +10 mm Hg

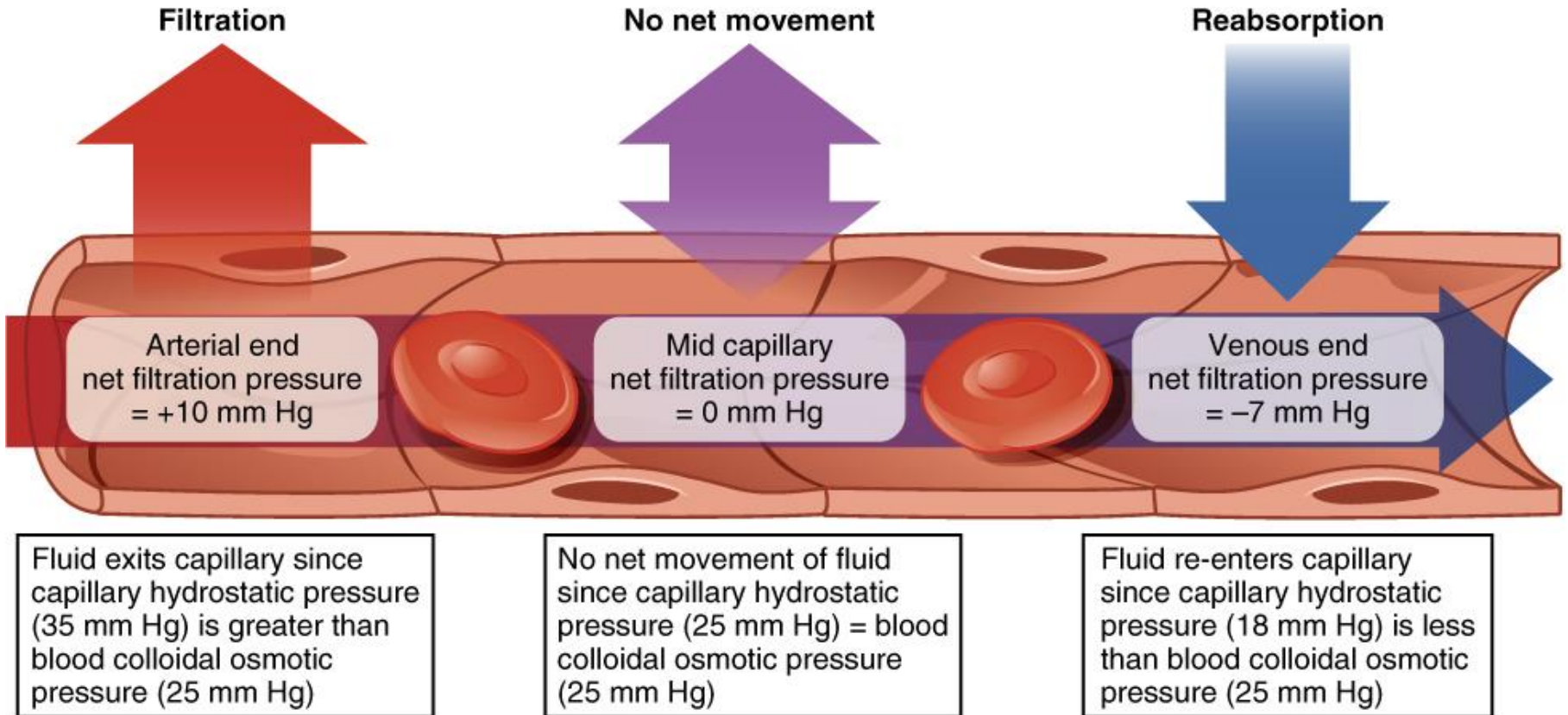
Mid capillary  
net filtration pressure  
= 0 mm Hg

Venous end  
net filtration pressure  
= -7 mm Hg

Fluid exits capillary since  
capillary hydrostatic pressure  
(35 mm Hg) is greater than  
blood colloidal osmotic  
pressure (25 mm Hg)

No net movement of fluid  
since capillary hydrostatic  
pressure (25 mm Hg) = blood  
colloidal osmotic pressure  
(25 mm Hg)

Fluid re-enters capillary  
since capillary hydrostatic  
pressure (18 mm Hg) is less  
than blood colloidal osmotic  
pressure (25 mm Hg)

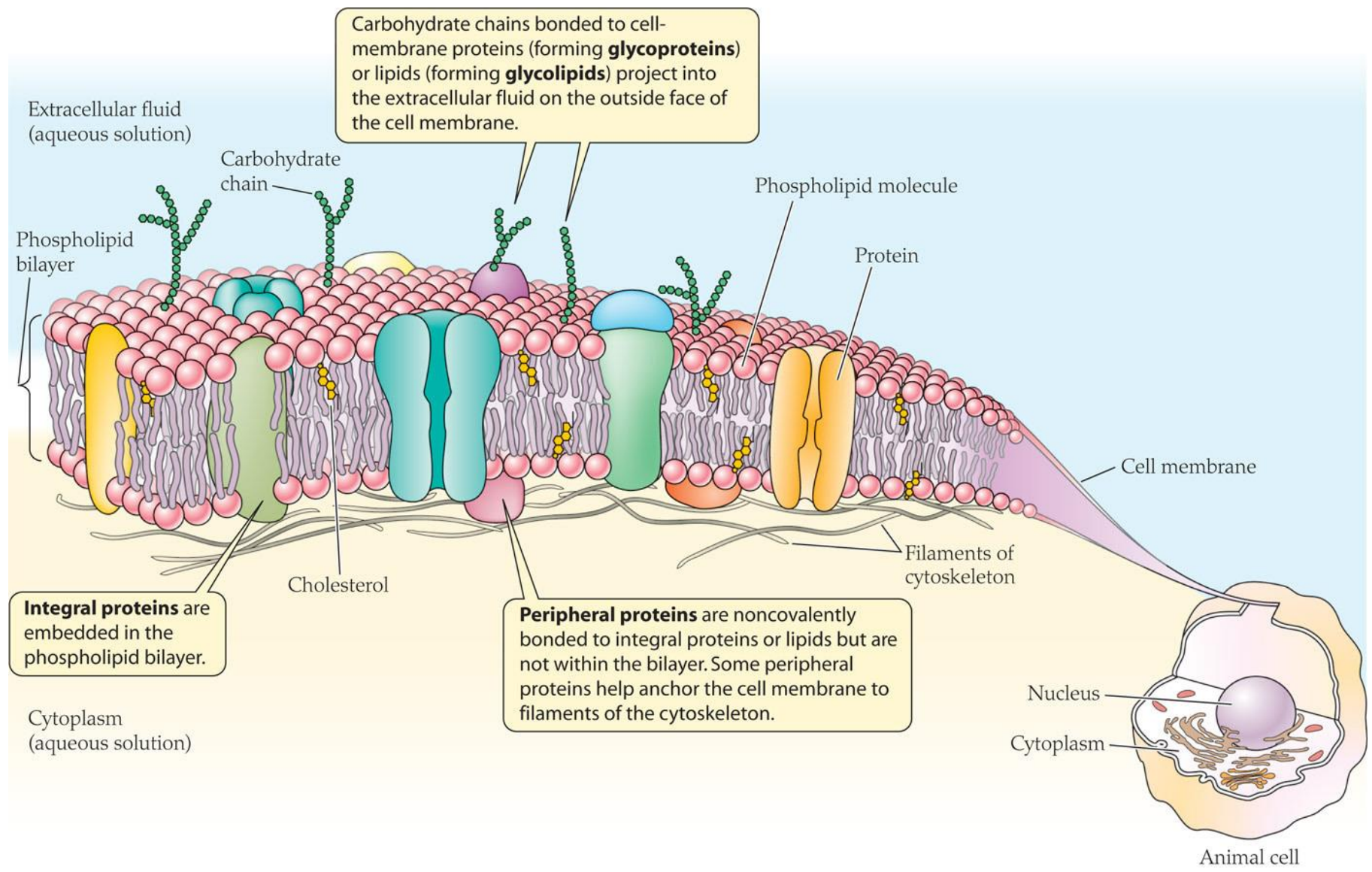


# Cell membranes and intracellular membranes

1. Cell membranes and intracellular membranes
2. Epithelia-the sheets of tissue that line body cavities and form the outer surfaces of organs



Figure 2.1 The structure of a cell membrane





# Terminology

- **Polar** – electrons are unevenly distributed
- **Nonpolar** – electrons are evenly distributed, no charge imbalance
- **Amphipathic**-each molecule consists of a polar part and a nonpolar part (head group of phospholipid)
- **Leaflets** – two layers of the membrane

# Figure 2.2 The structure of membrane phospholipid molecules

(a) A phospholipid molecule (a phosphatidylcholine)

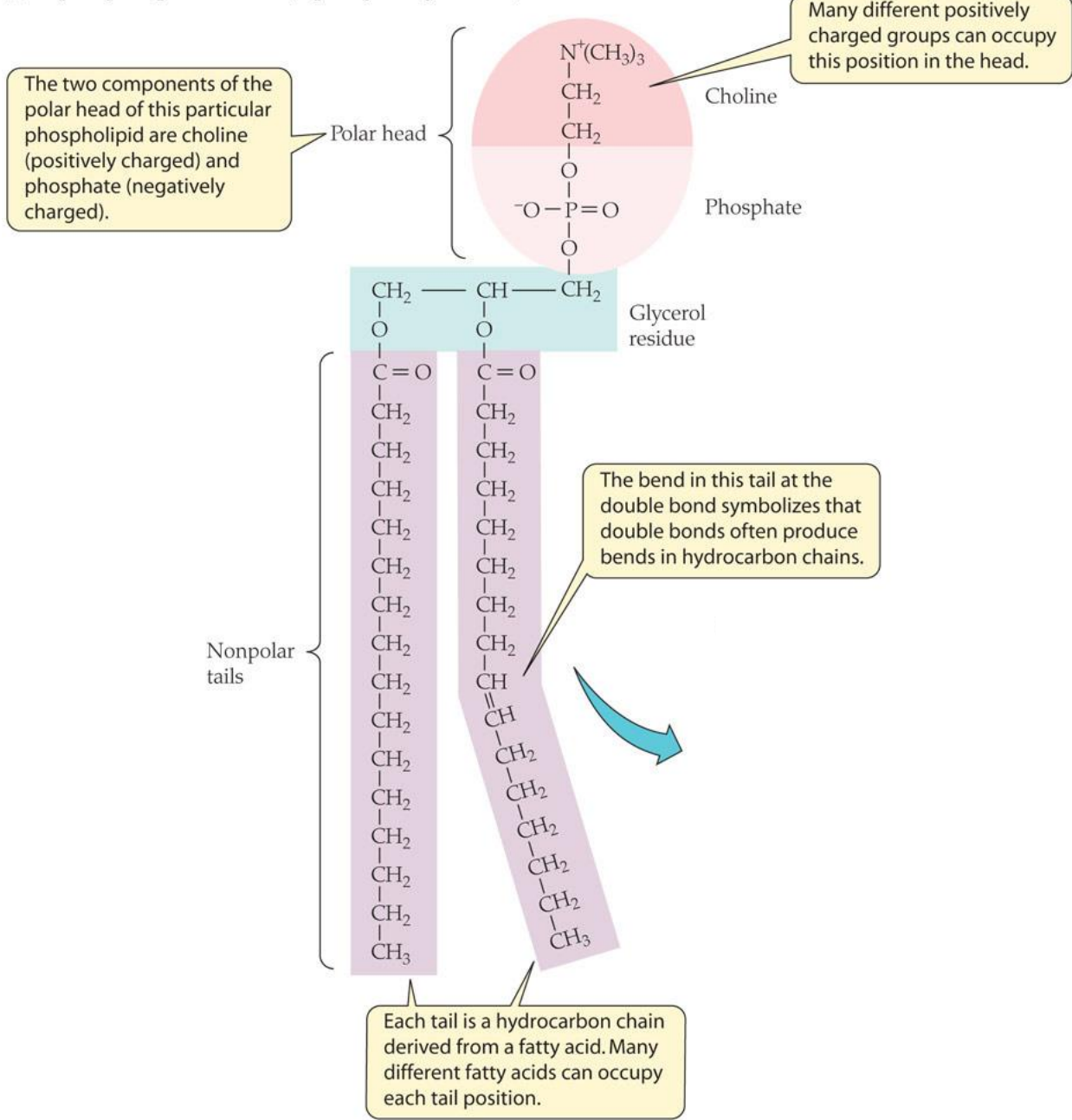
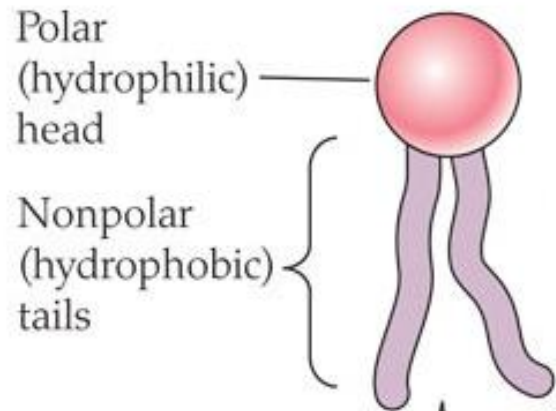


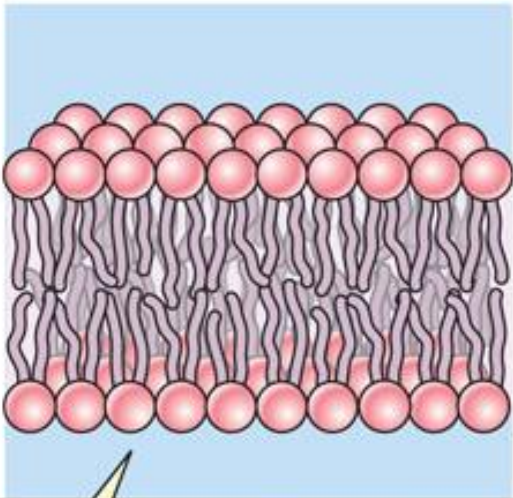
Figure 2.2 The structure of membrane phospholipid molecules

(b) Model of a phospholipid molecule



Phospholipid molecules  
are represented  
symbolically like this.

(c) Phospholipid molecules  
assembled into a bilayer  
with water on either side

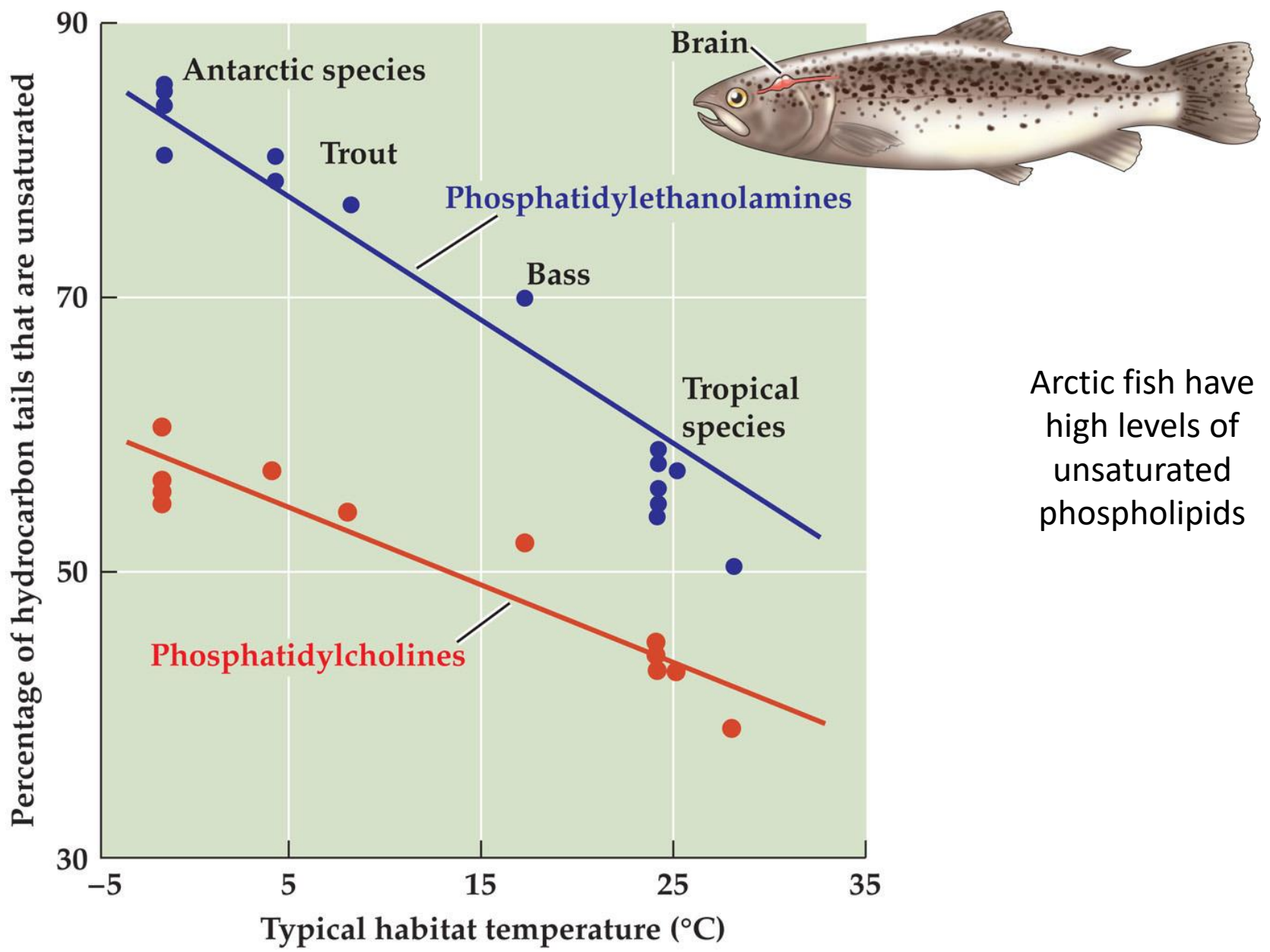


When placed in water, phospholipid molecules spontaneously assemble into a bilayer in which the nonpolar, hydrophobic tails occupy the core and the polar, hydrophilic heads occupy the two surfaces.

# Membrane Fluidity

- RBC's have >150 kinds of phospholipids
- Phospholipids can travel (by diffusion) around the whole cell in a matter of minutes
- Temperature: lower → lower fluidity
- Fluidity depends on chemical saturation of the hydrocarbons in the tails
  - Saturated-non double bonds
  - Unsaturated – one or more double bonds, creates bends → higher fluidity

Figure 2.3 The degree of chemical unsaturation of the hydrocarbon tails of brain phospholipids in fish varies with habitat temperature



# Proteins Endow Membranes with Numerous Functional Capacities

# Two *structural* designations for membrane proteins

1. **Integral** – parts of the membrane, can't be removed, usually transmembrane (span the whole membrane)
2. **Peripheral** – can be removed, bonded noncovalently to membrane components, on one side of the membrane or the other



# Five *functional* designations for membrane proteins

**TABLE 2.1** The five functional types of membrane proteins and the functions they perform

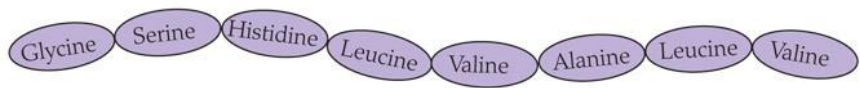
Functional type	Function performed (defining property)
Channel	Permits simple or quasi-simple <i>diffusion</i> of solutes in aqueous solution (see page 104)—or <i>osmosis</i> of water (see page 121)—through a membrane. A simplified view of a channel is that it creates a direct water path from one side to the other of a membrane (i.e., an aqueous pore) through which solutes in aqueous solution may diffuse or water may undergo osmosis.
Transporter (carrier)	Binds noncovalently and reversibly with specific molecules or ions to move them across a membrane intact. The transport through the membrane is <i>active transport</i> (see page 108) if it employs metabolic energy; it is <i>facilitated diffusion</i> (see page 108) if metabolic energy is not employed.
Enzyme	Catalyzes a chemical reaction in which covalent bonds are made or broken (see page 41).
Receptor	Binds noncovalently with specific molecules and, as a consequence of this binding, initiates a change in membrane permeability or cell metabolism. Receptor proteins mediate the responses of a cell to chemical messages (signals) arriving at the outside face of the cell membrane (see page 58).
Structural protein	Attaches to other molecules (e.g., other proteins) to anchor intracellular elements (e.g., cytoskeleton filaments) to the cell membrane, creates junctions between adjacent cells (see Figure 2.7), or establishes other structural relations.

# Fluid Mosaic Model of Membranes

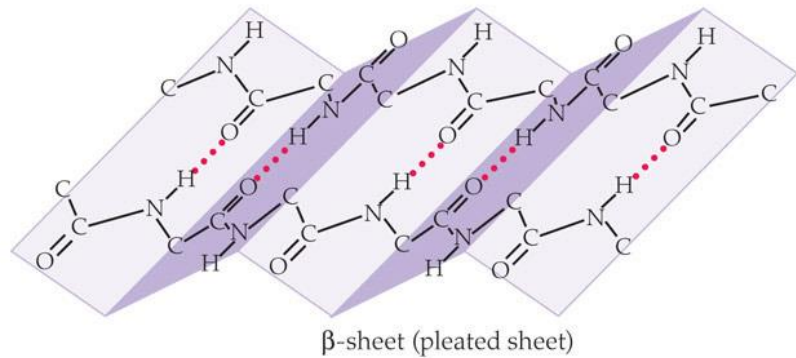
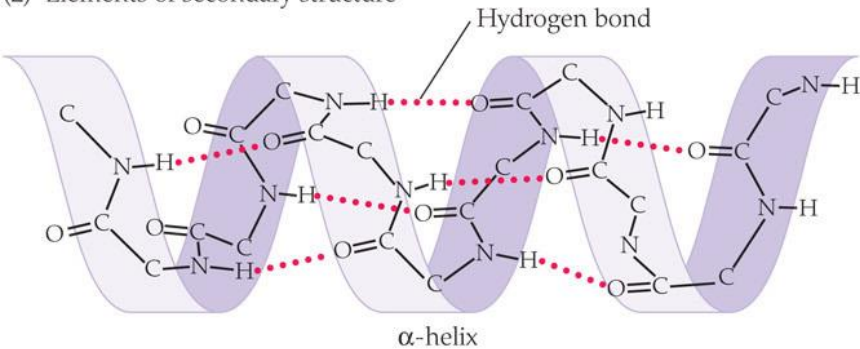
A membrane consists of a mosaic of protein and lipid molecules, all of which move about in directions parallel to the membrane faces because of the fluid state of the lipid matrix

Box Extension 2.1 Figure A The structural hierarchy of proteins

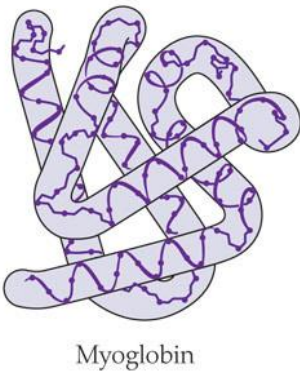
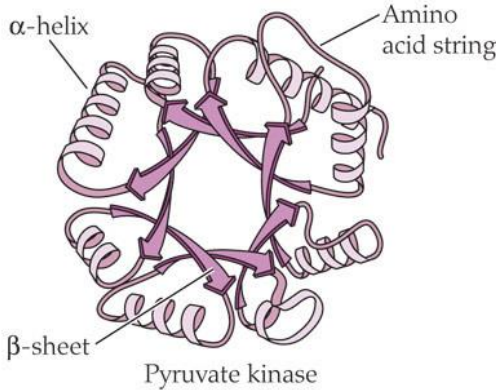
(1) Primary structure



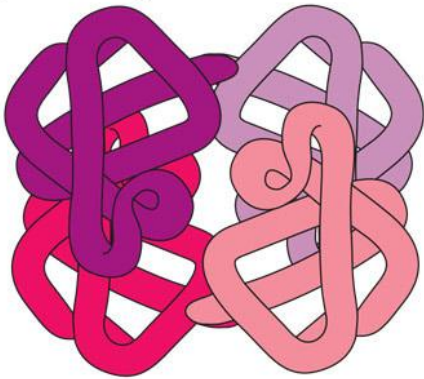
(2) Elements of secondary structure



(3) Tertiary structure drawn in two ways

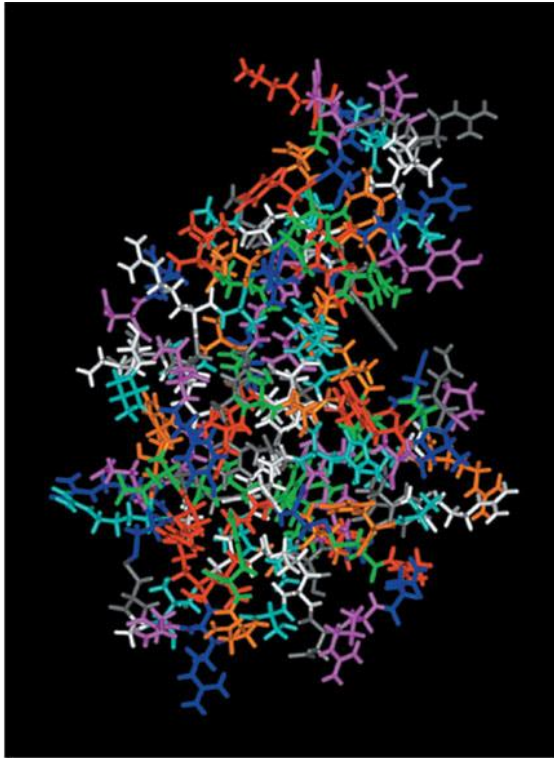


(4) Quaternary structure



This molecule consists of four separate proteins (symbolized by the four colors) bonded into a four-unit complex.

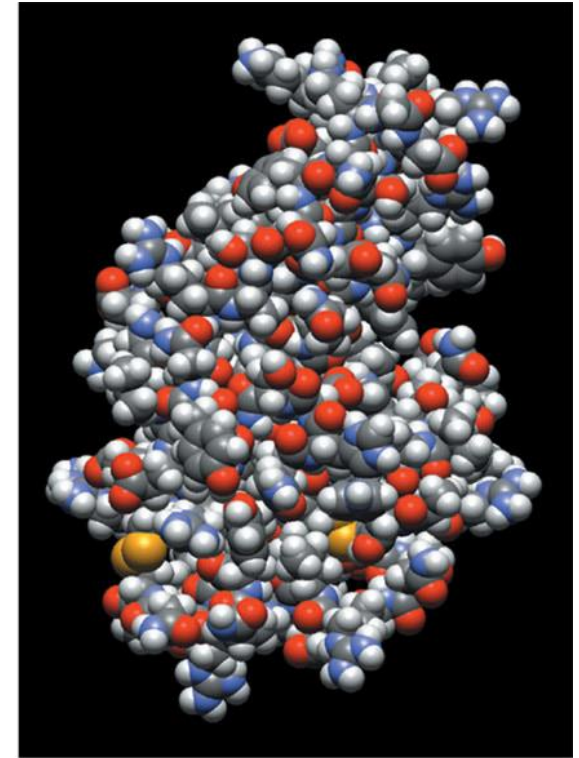
Box Extension 2.1 Figure B Three ways to diagram the tertiary structure of one protein. All three diagrams represent lysozyme



Stick model



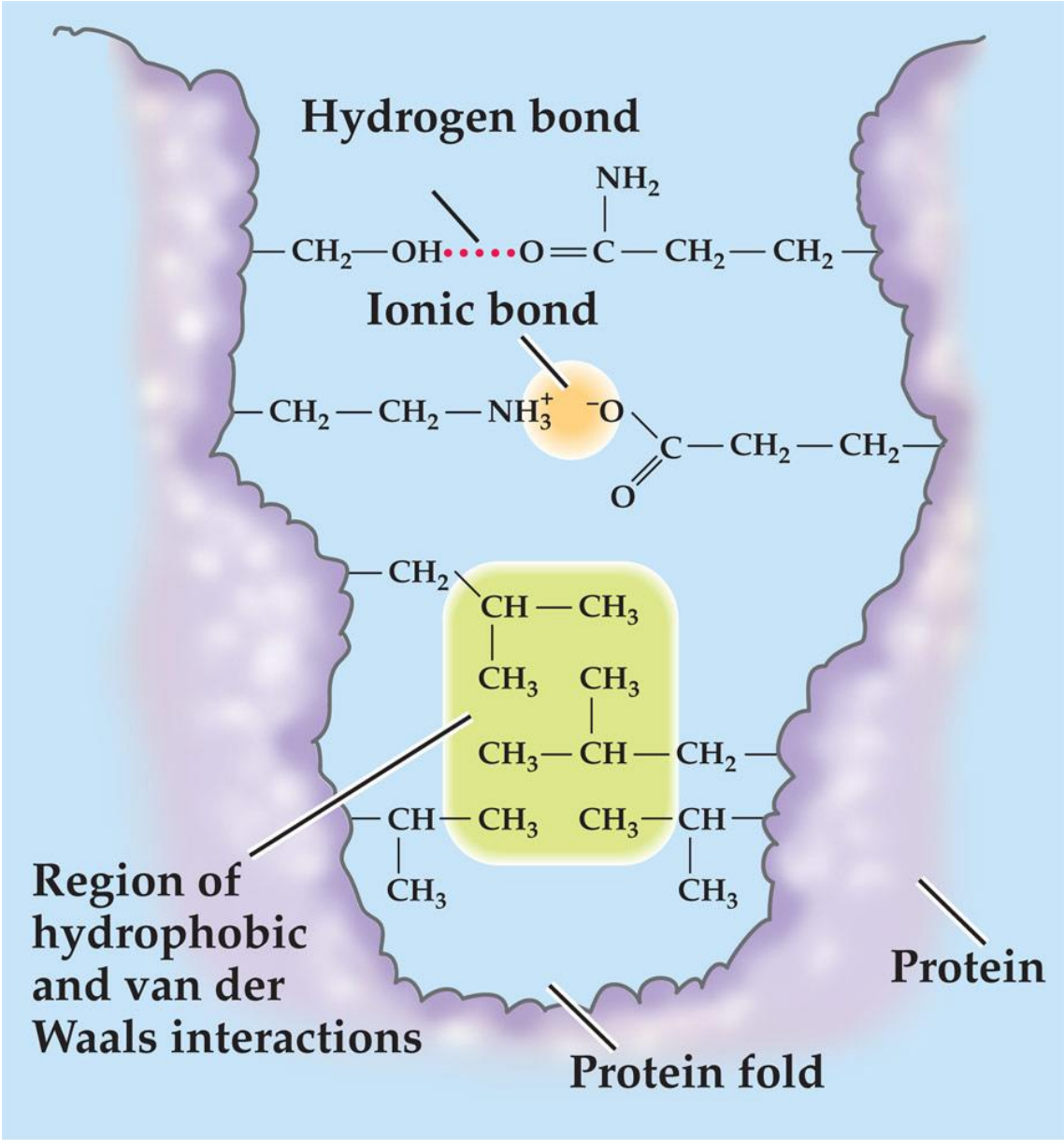
Ribbon model



Space-filling model

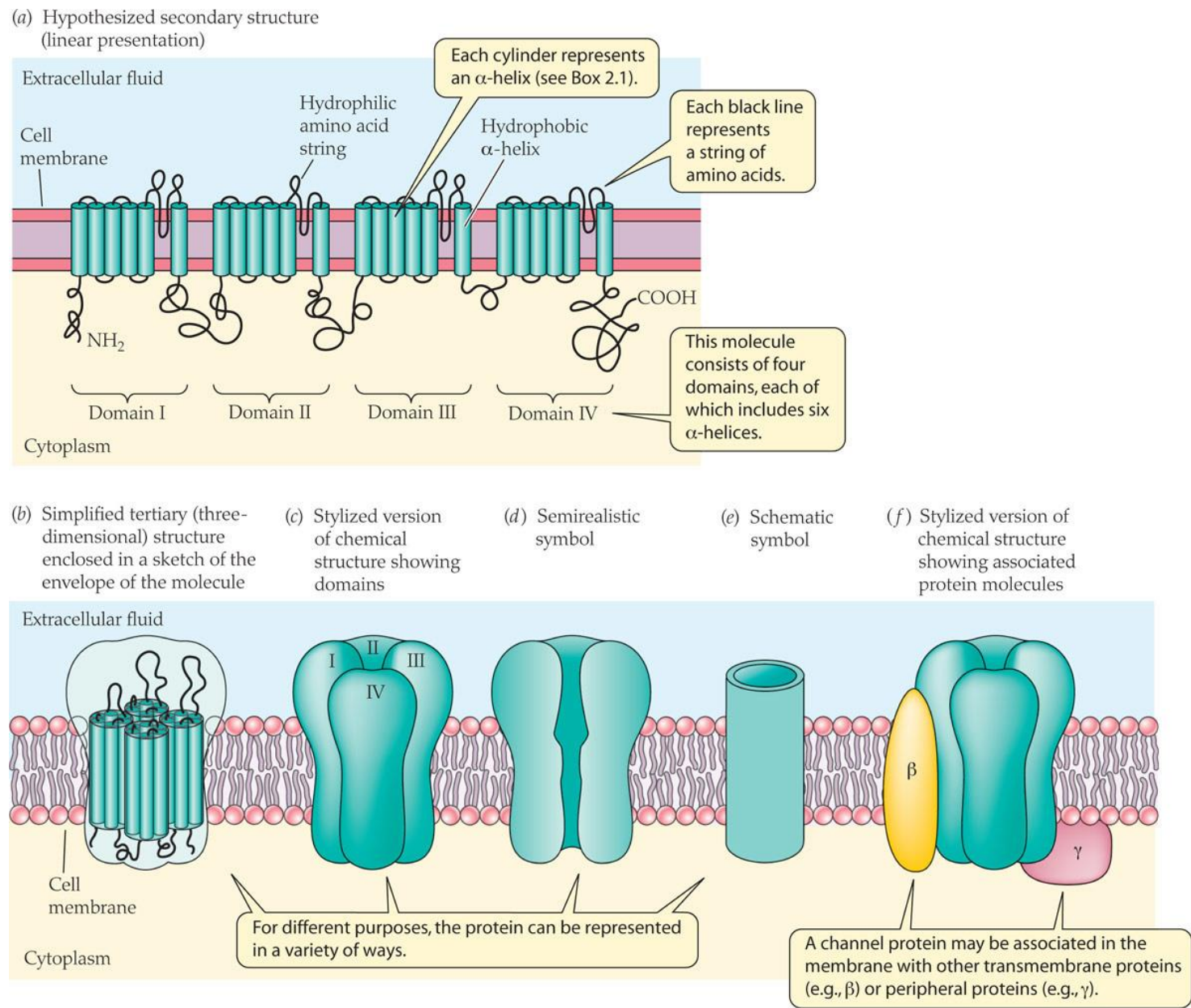


Box Extension 2.1 Figure C Types of weak, noncovalent bonds that are important in protein structure



**ANIMAL PHYSIOLOGY 3E, Box Extension 2.1 Figure C**

Figure 2.4 The structure of a transmembrane protein—a voltage-gated Na<sup>+</sup> channel—illustrating several modes of presentation



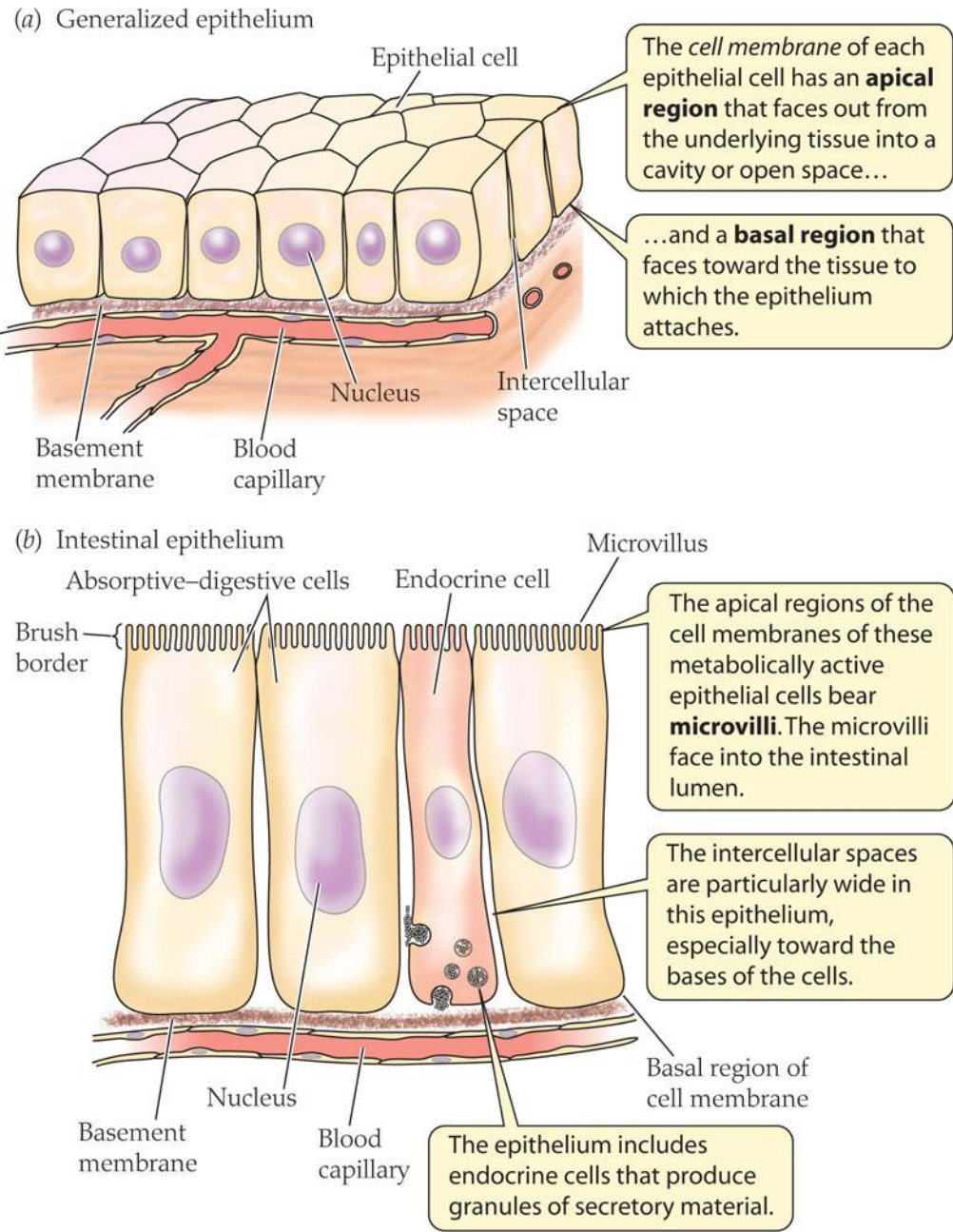
# Four Topics in Chapter 2

1. Cell membranes and intracellular membranes
- 2. Epithelia-the sheets of tissue that line body cavities and form the outer surfaces of organs**
3. Enzyme function, diversity and evolution
4. Mechanisms by which cells receive and act on signals

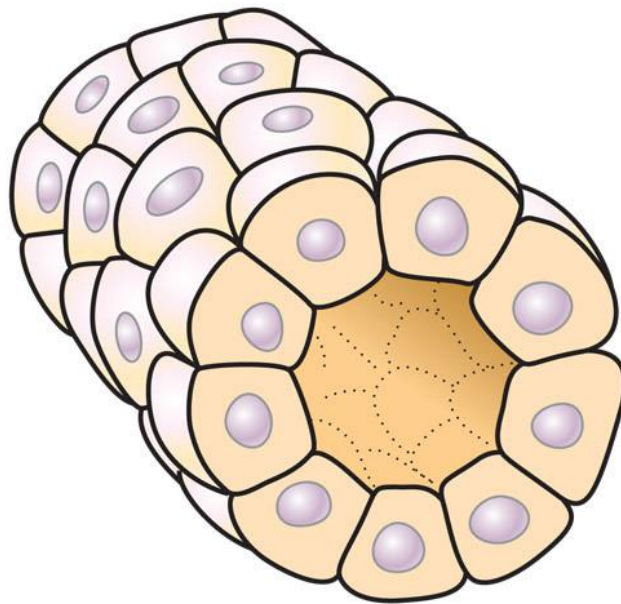


- Epithelia compartmentalize the body by forming boundaries between body regions
- Also form a boundary between an animal and its external environment

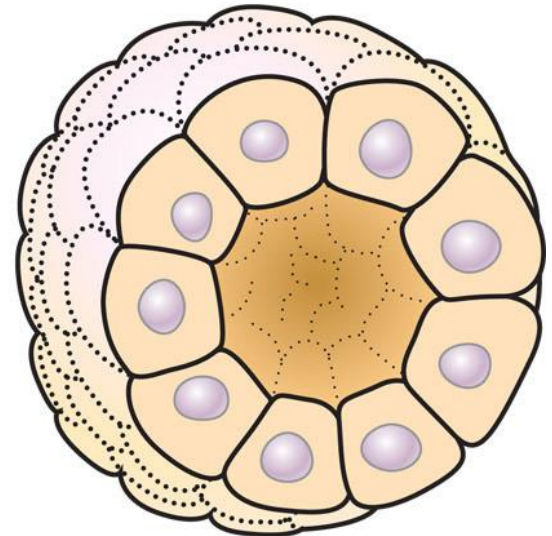
Figure 2.5 Simple epithelia-single layer of cells (squamous, cuboidal, columnar)



## (a) Epithelial cells can form tubules and follicles

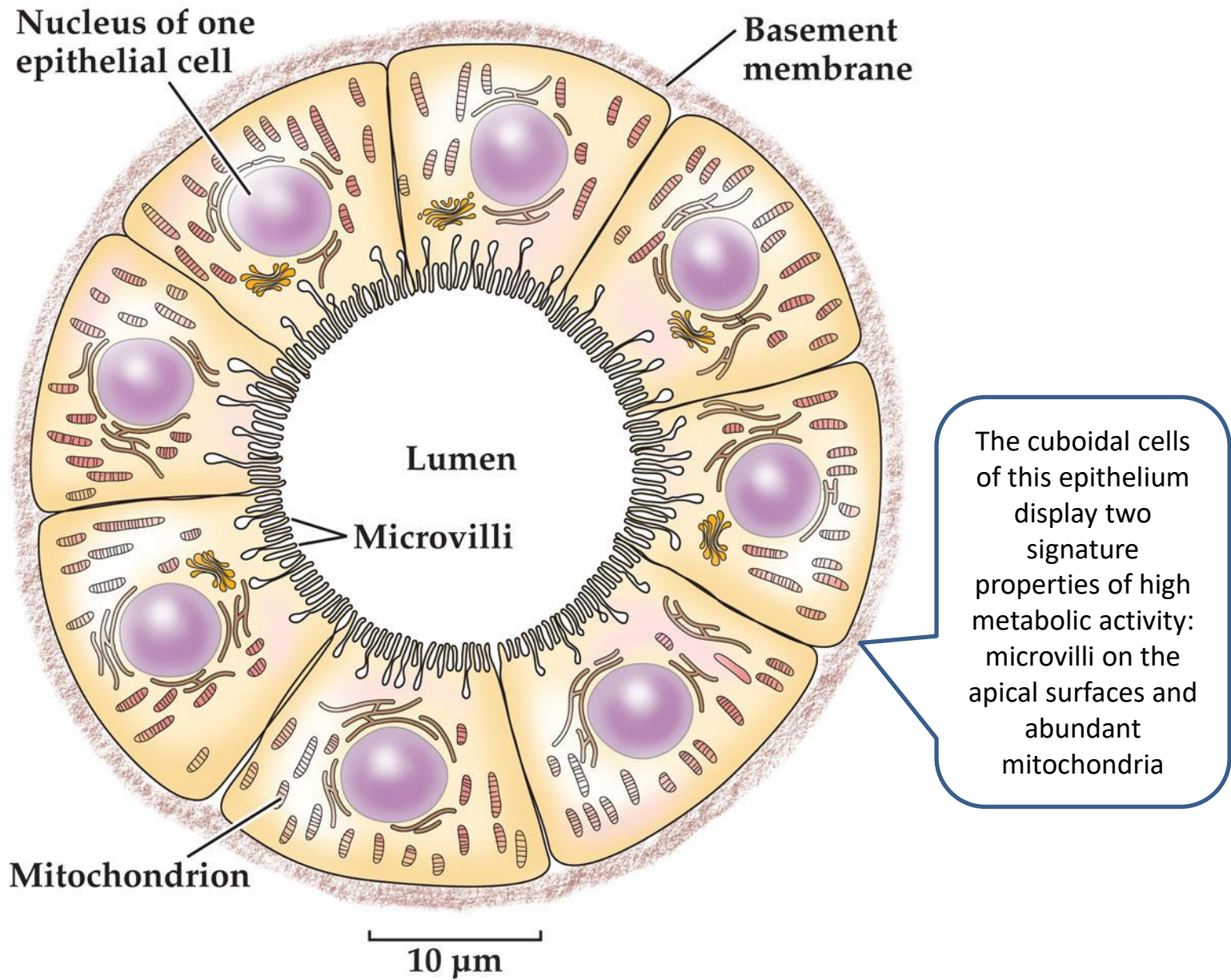


**Tubule**



**Follicle**

(b) Proximal part of a mammalian nephron (kidney tubule) in cross section





(c) Mammalian blood capillary in cross section

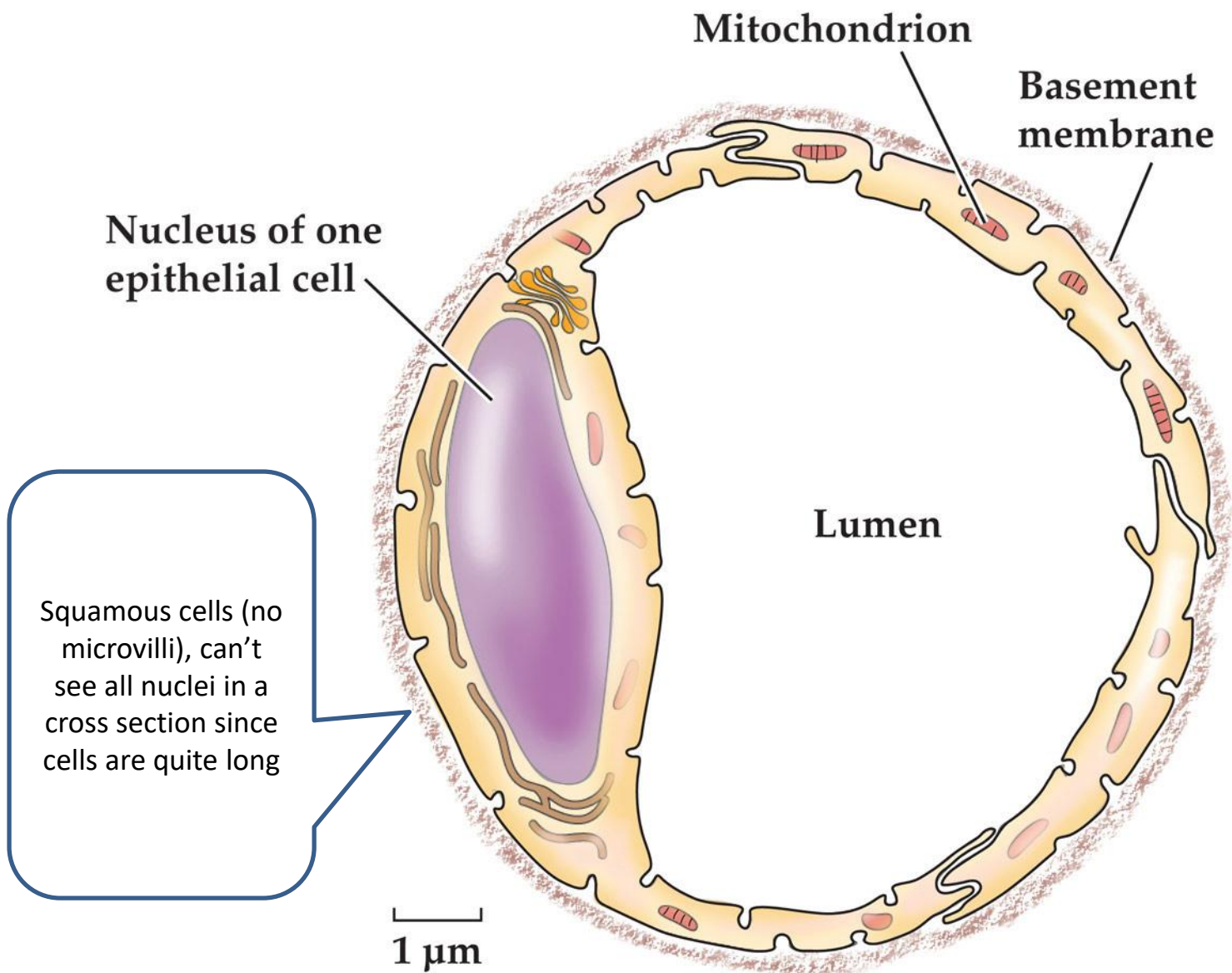


Figure 2.7 Types of junctions between cells

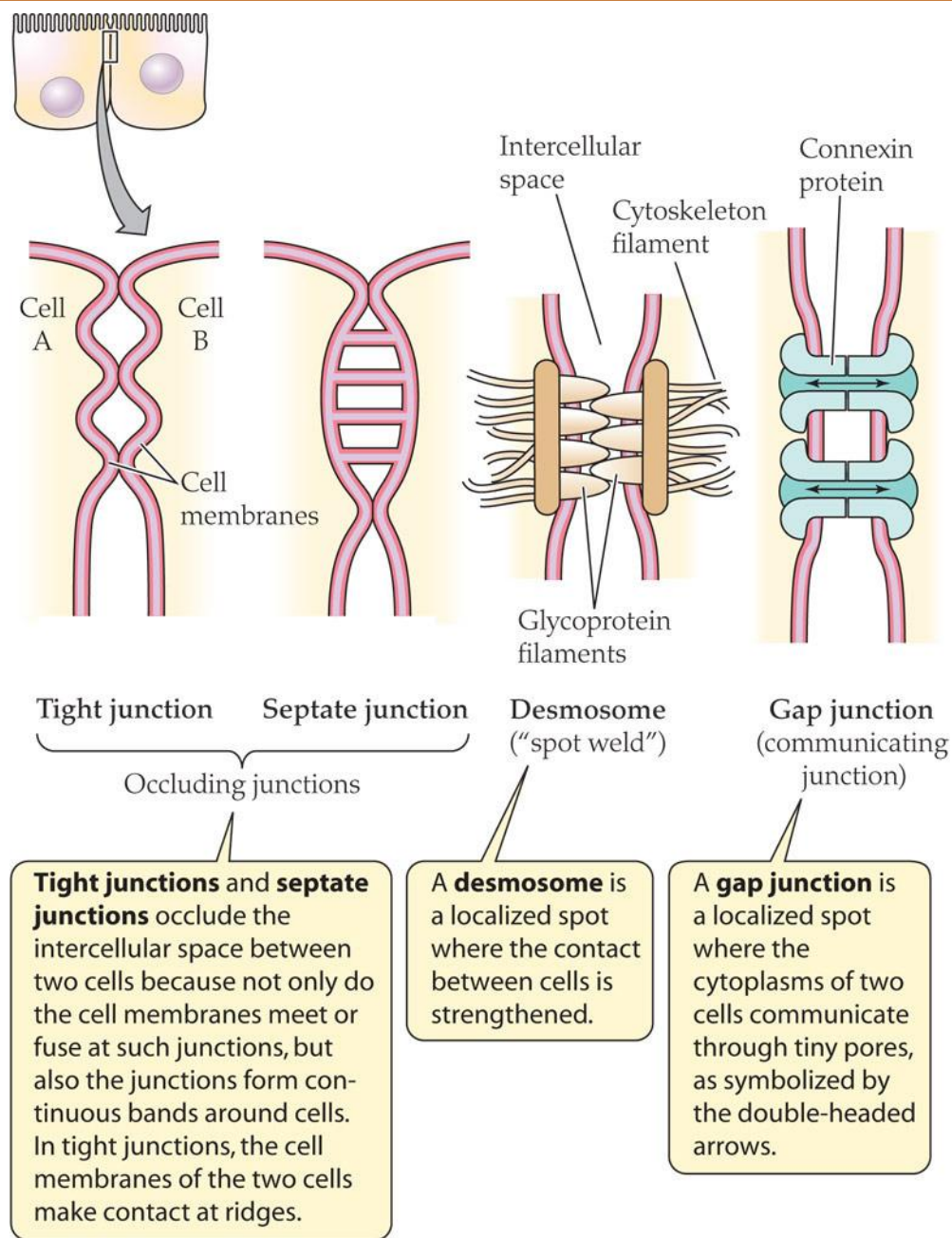
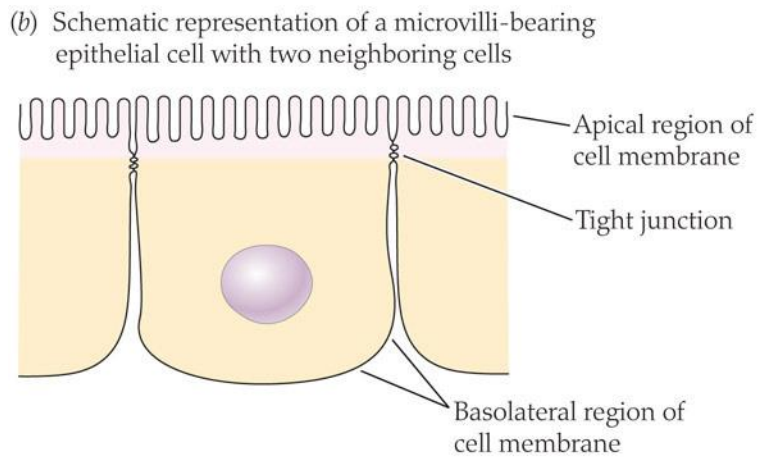
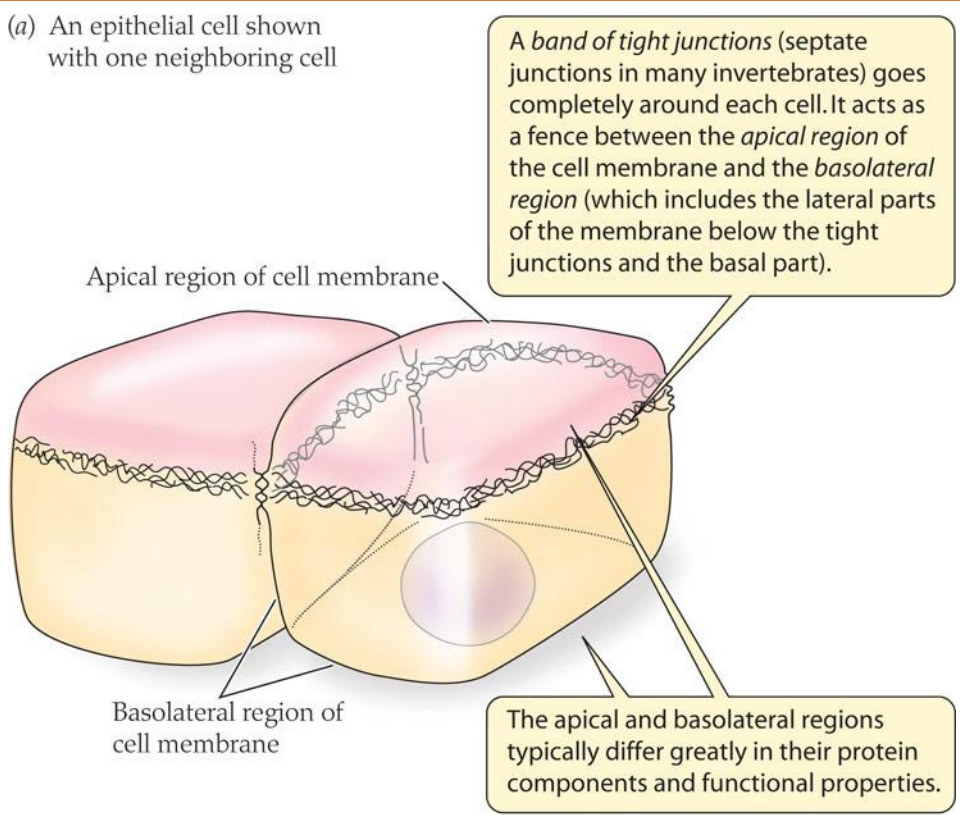


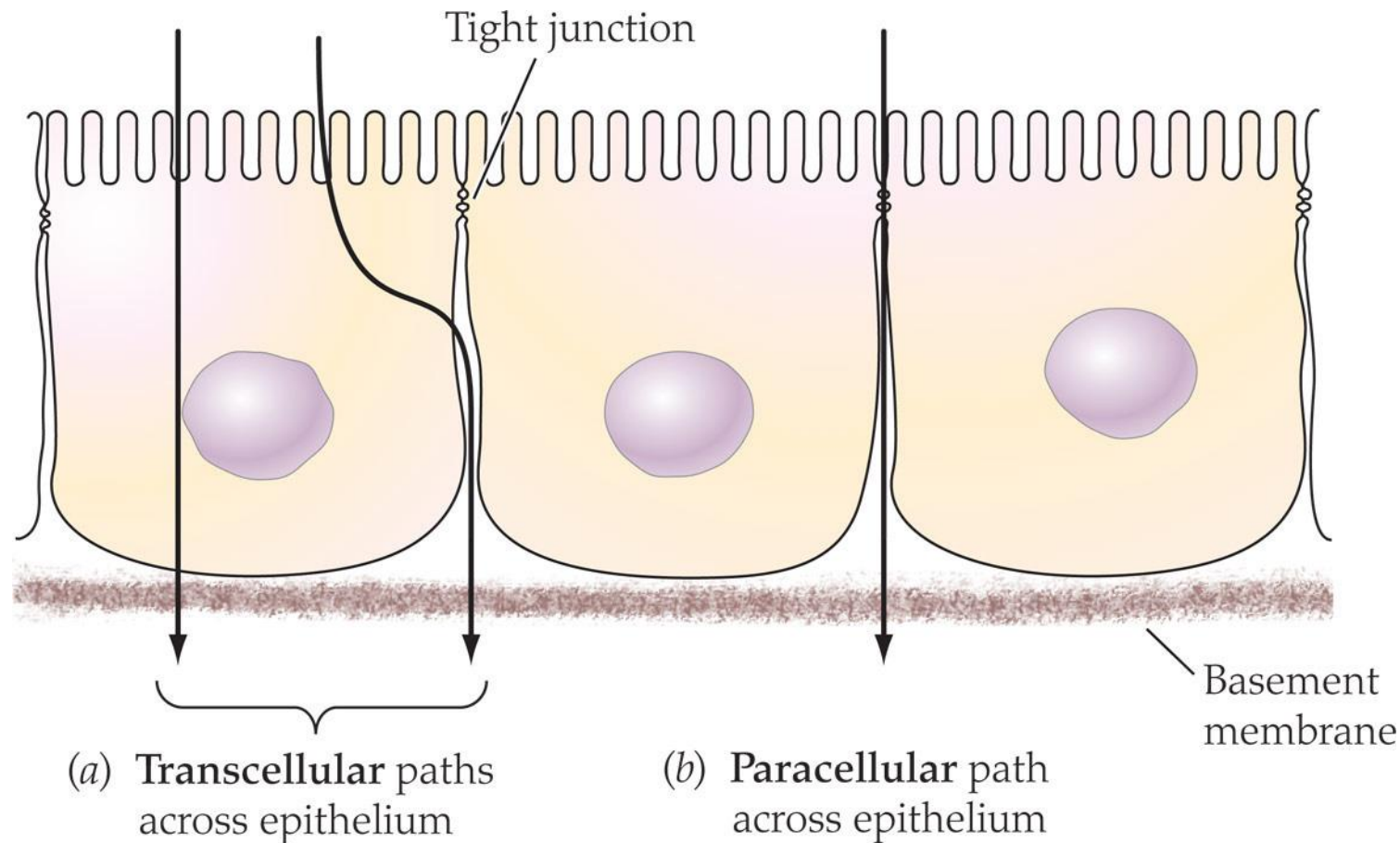
Figure 2.8 The organization of epithelial cells into apical and basolateral regions



**ANIMAL PHYSIOLOGY 3E, Figure 2.8**



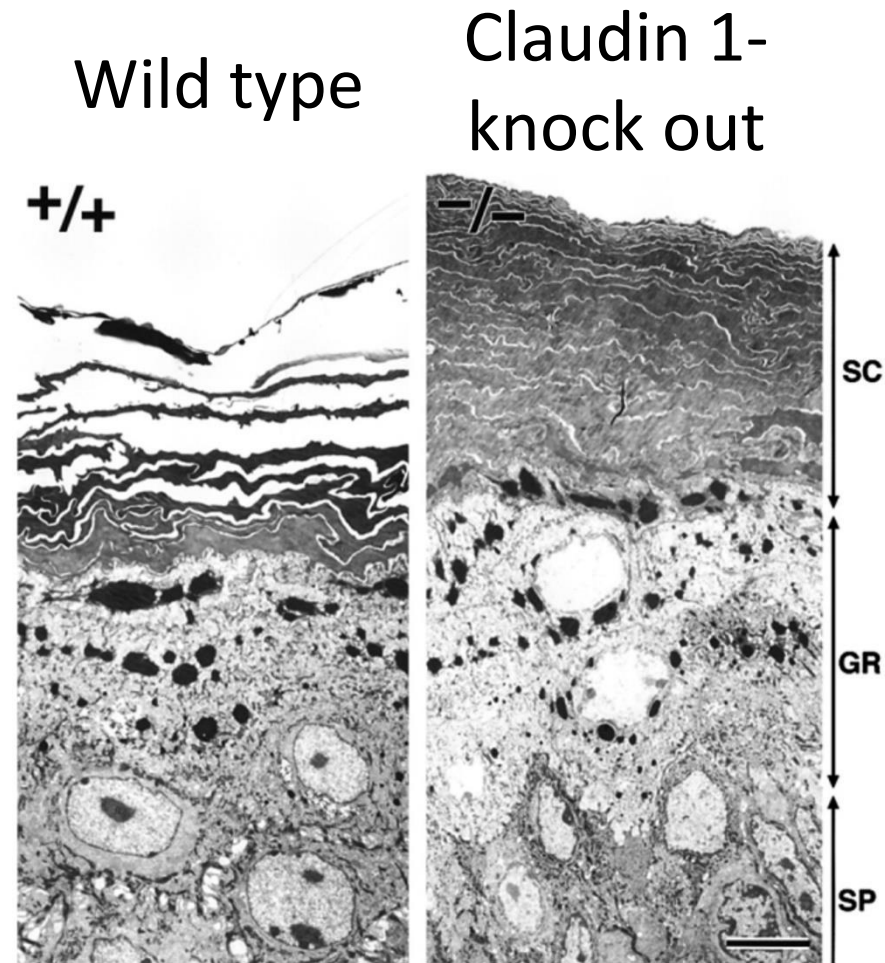
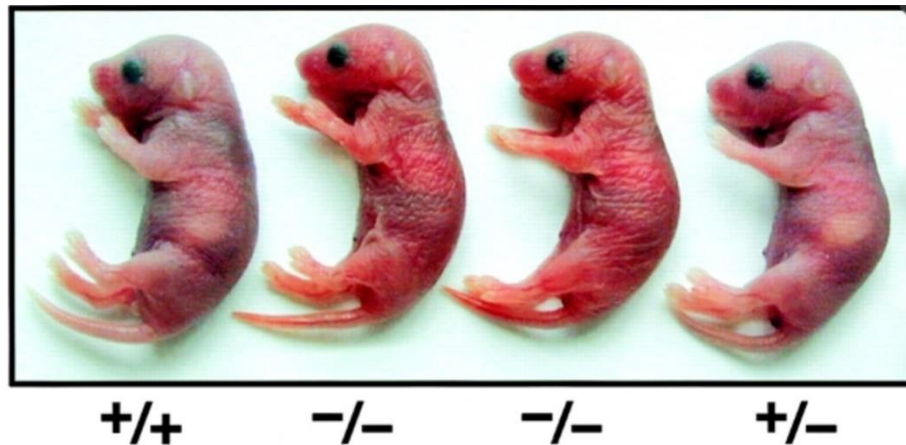
Figure 2.9 Transcellular and paracellular paths across an epithelium



Materials following a **transcellular** path must cross both apical and basolateral cell membranes.

Materials following a **paracellular** path must be able to move through the band of tight (or septate) junctions; in many epithelia, only very small molecules are able to do this, restricting the paracellular path to such molecules.

# Impairment of epidermal barrier in Claudin 1 deficient mice



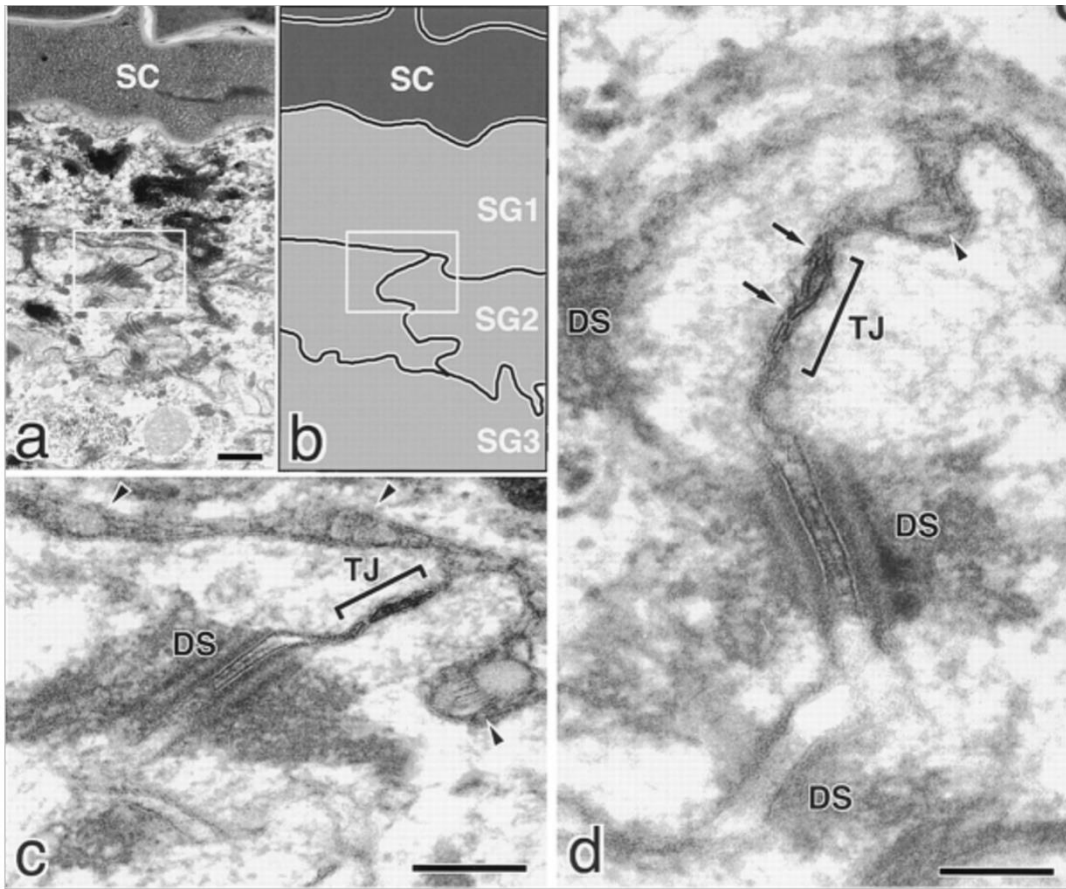
SC: stratum corneum; SG: stratum granulosum; SP: stratum spinosum

**Claudin-based tight junctions are crucial for the mammalian epidermal barrier.**

J Cell Biol. 2002 Mar 18; 156(6): 1099–1111.

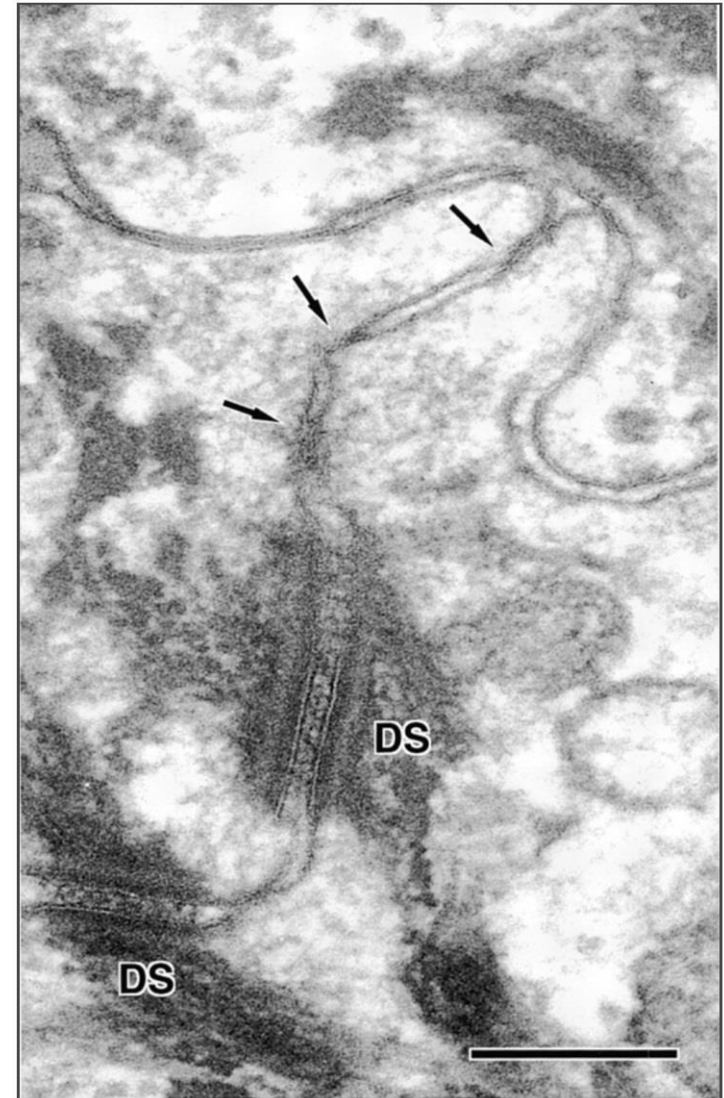


## Wild type



TJ: tight junction; DS: desmosome

## Claudin 1-knock out



## Chapter 2 Opener Two slow-moving predators that use molecular weapons to capture fast-moving prey

