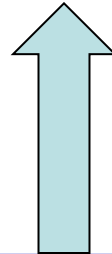


# Transport across Plasma Membranes

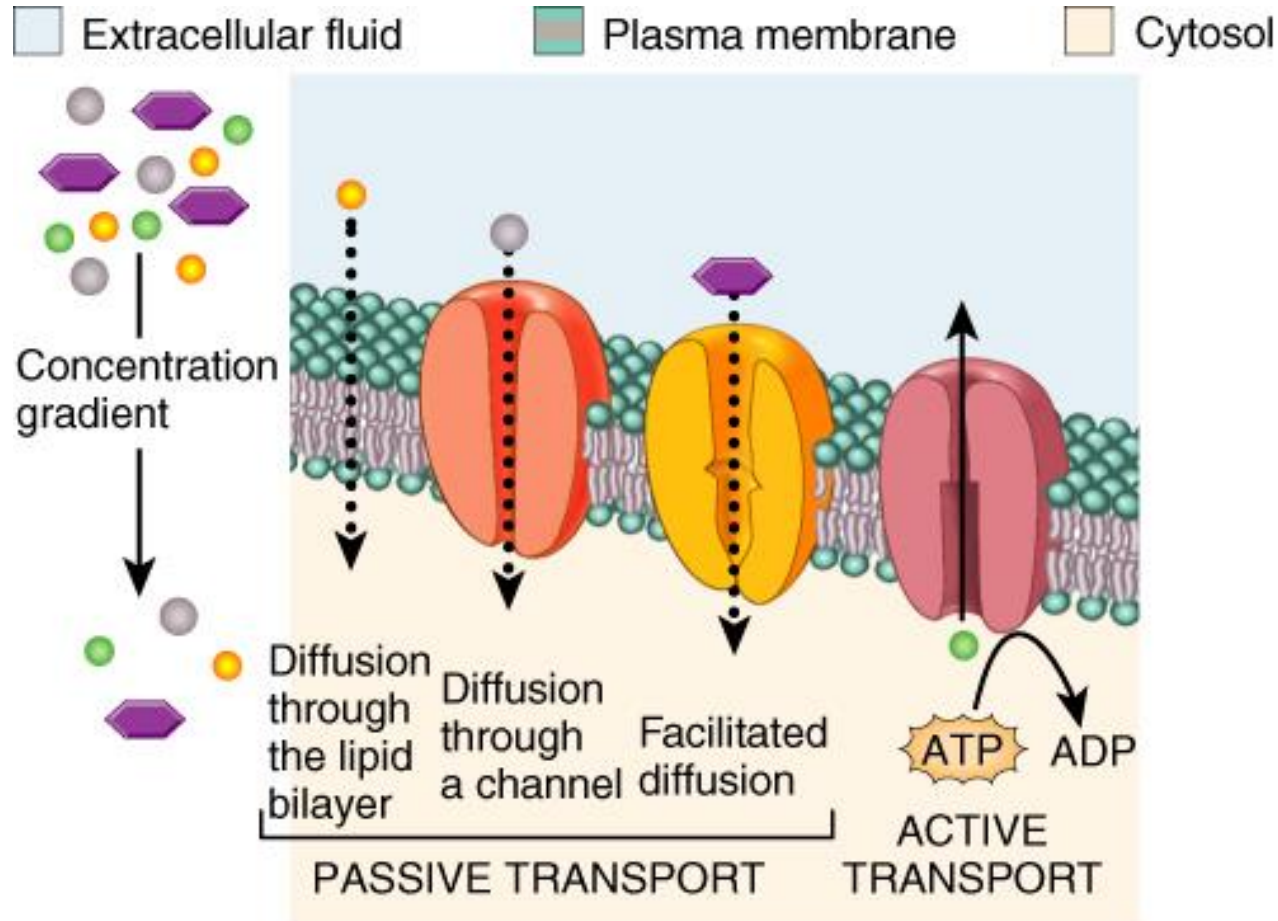
<https://www.youtube.com/watch?v=A9ihz5gYxU4>



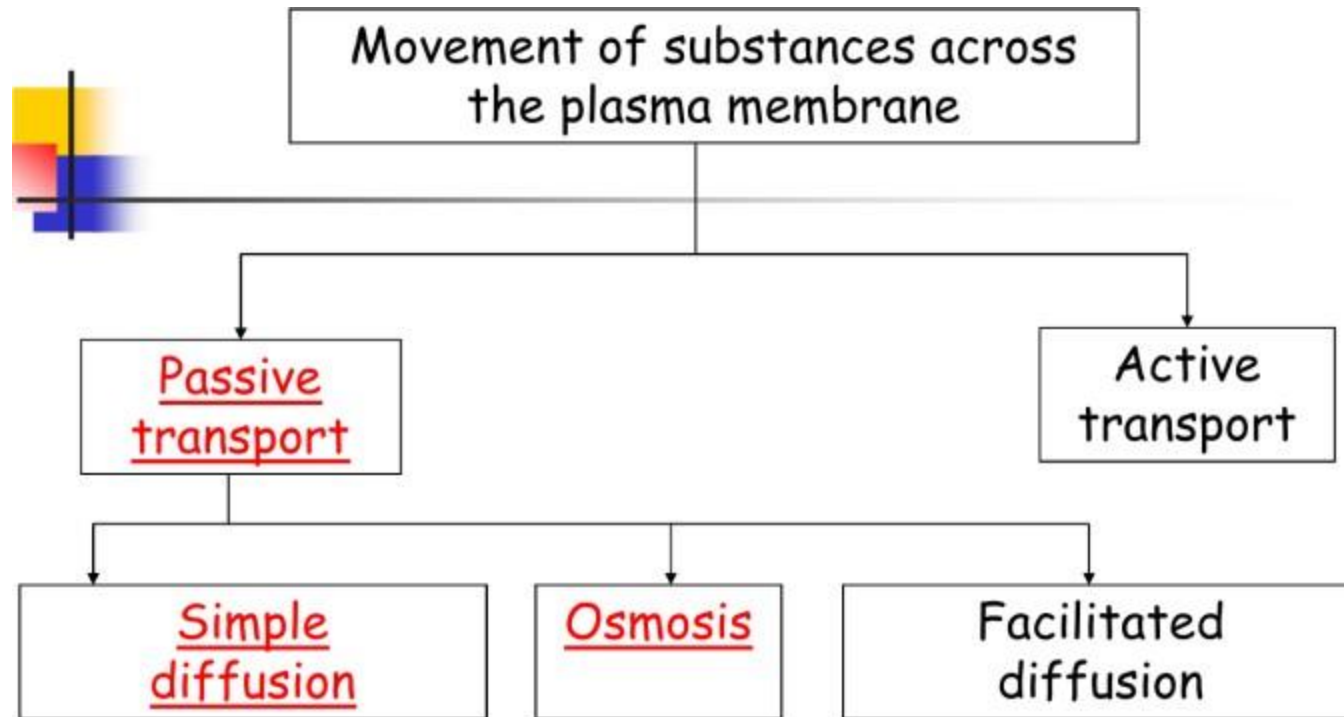
**Follow the LINK**

**Or copy and past on  
your browser**

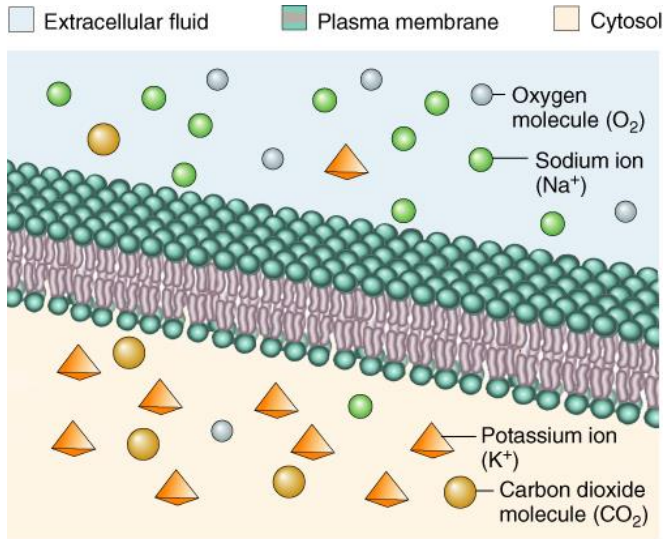
# Transport across Plasma Membranes



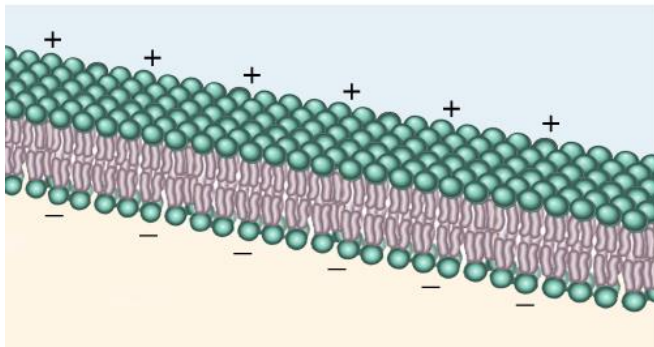
# Transport across Plasma Membranes



# Diffusion through lipid bilayer



(a) Concentration gradients

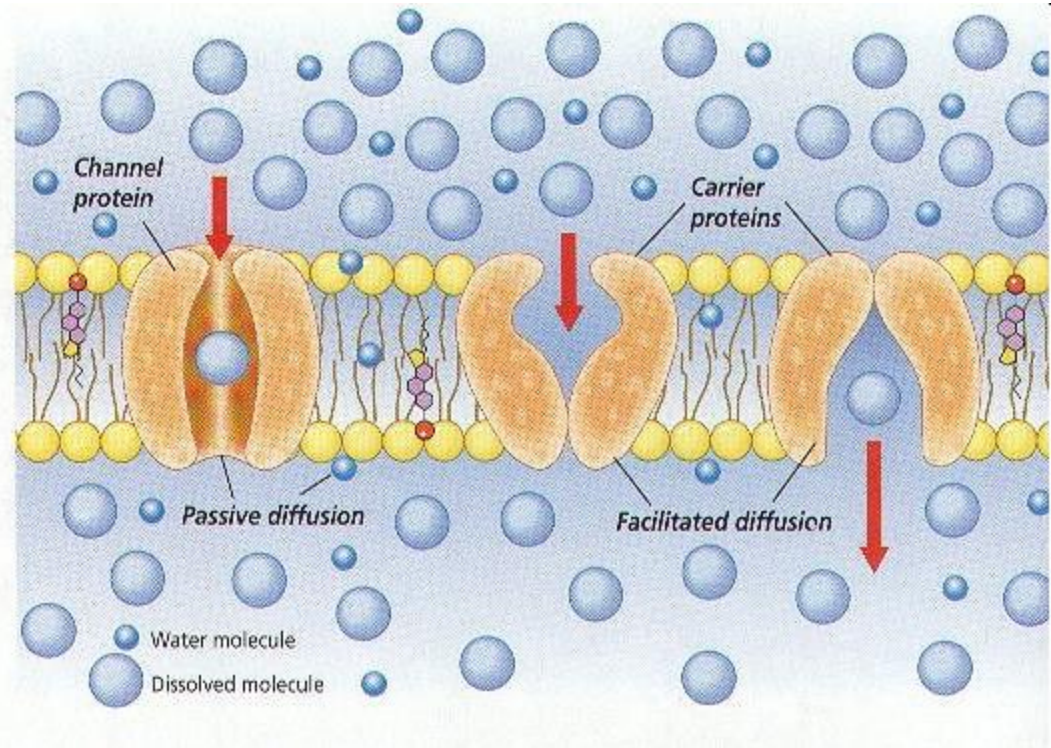


(b) Electrical gradient

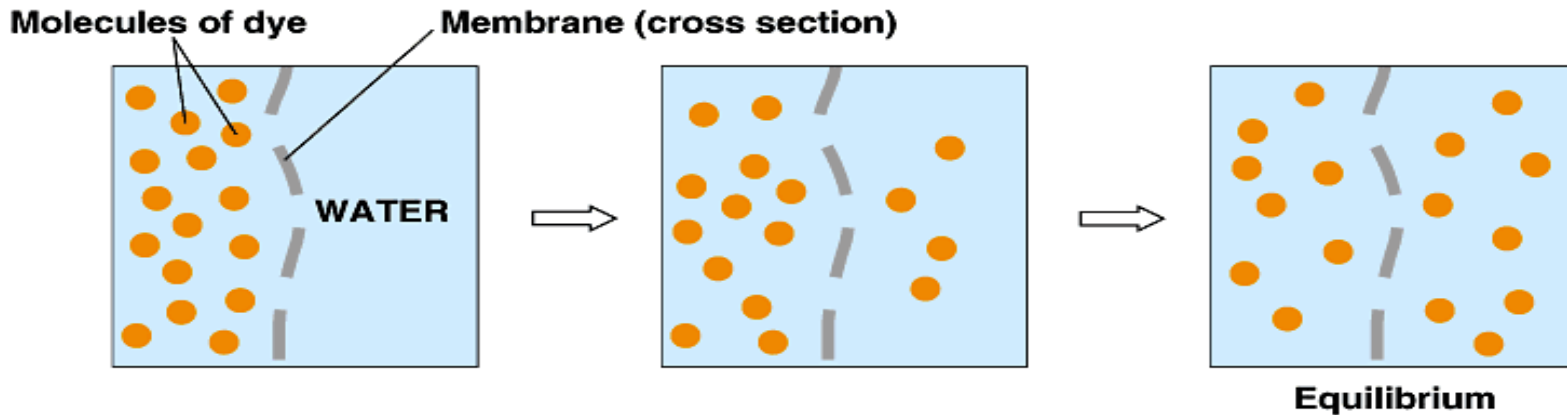
- $CO_2$
- $O_2$
- $NO$
- Steroid Hormones
- Monoglycerides

# Diffusion through channels

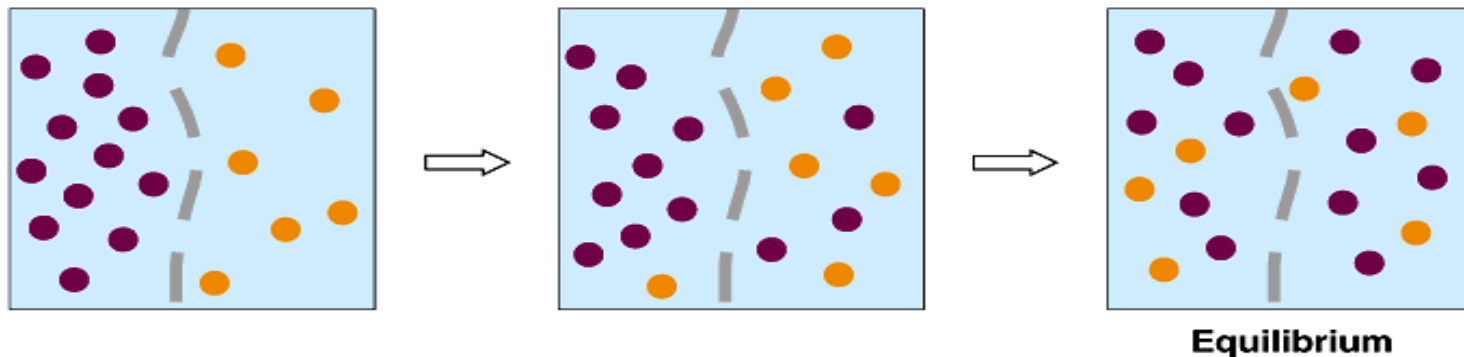
Channel proteins provide the openings through which small, dissolved particles, especially ions, diffuse by passive transport.



# The Concept of Simple Diffusion



**(a) Diffusion of one solute**

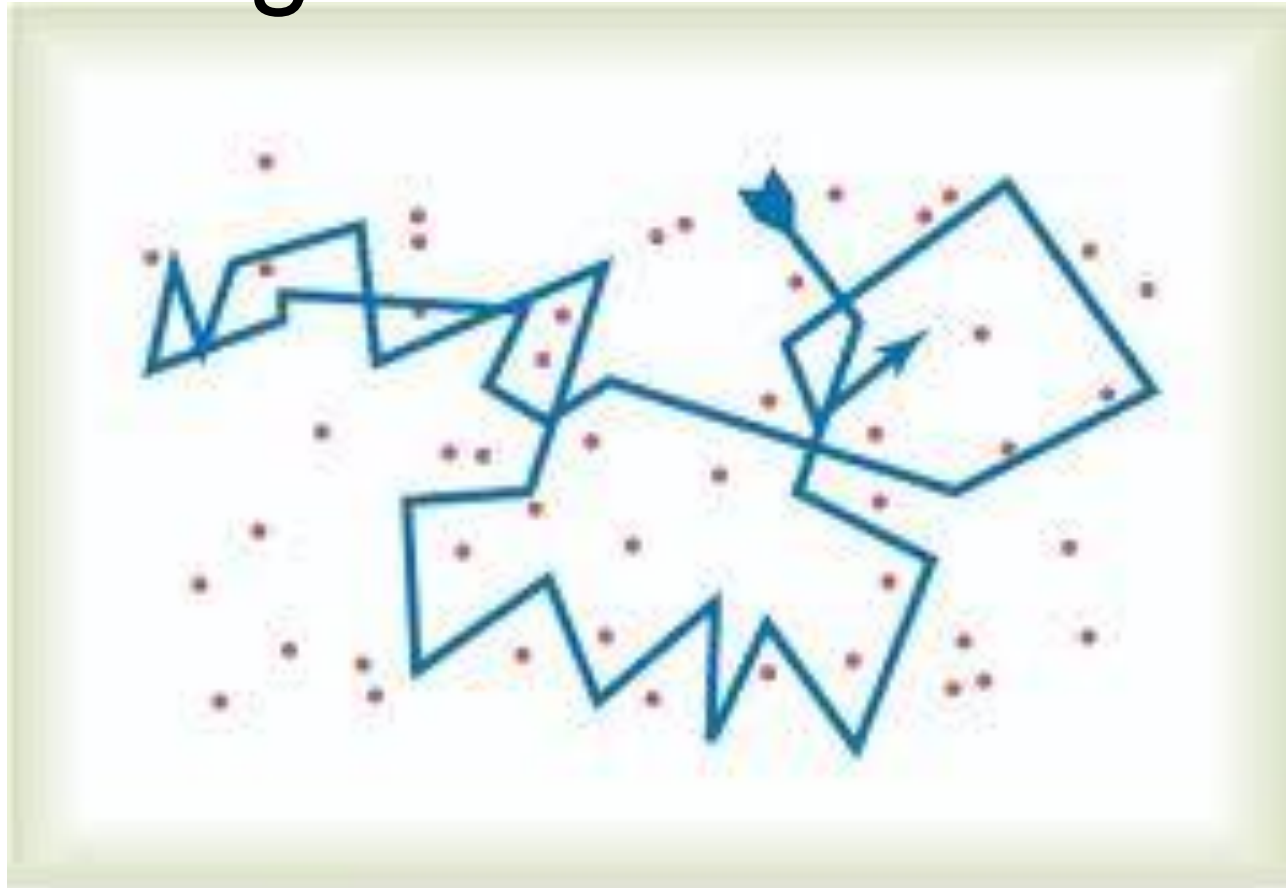


**(b) Diffusion of two solutes**

Are we needing to consume macro-energetic molecules (ATP) for diffusion across plasma membranes??



# What type of Energy is there to get the diffusion??

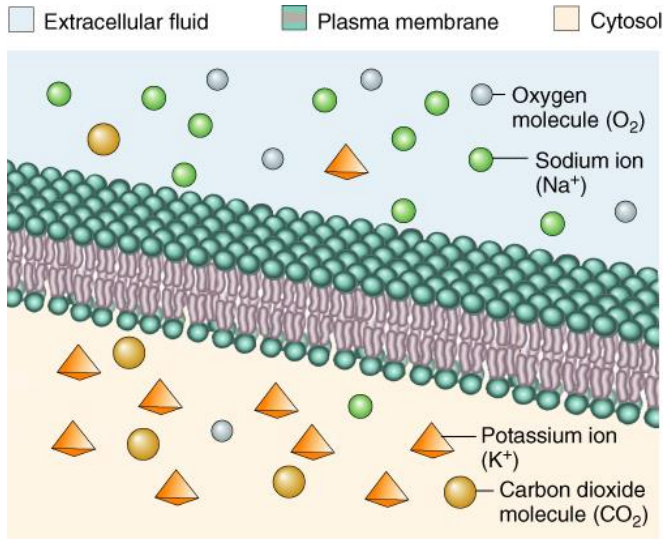


**Figure 4-3**

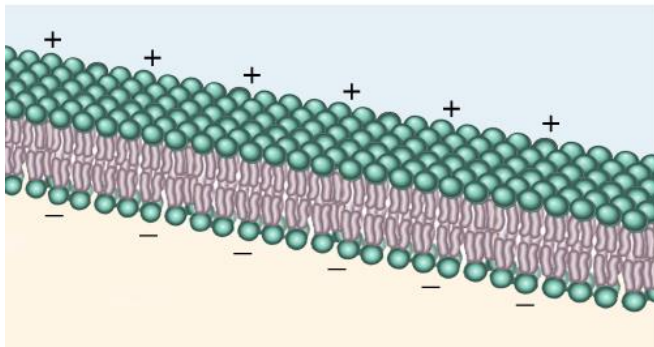
Diffusion of a fluid molecule during a thousandth of a second.



# Diffusion through lipid bilayer



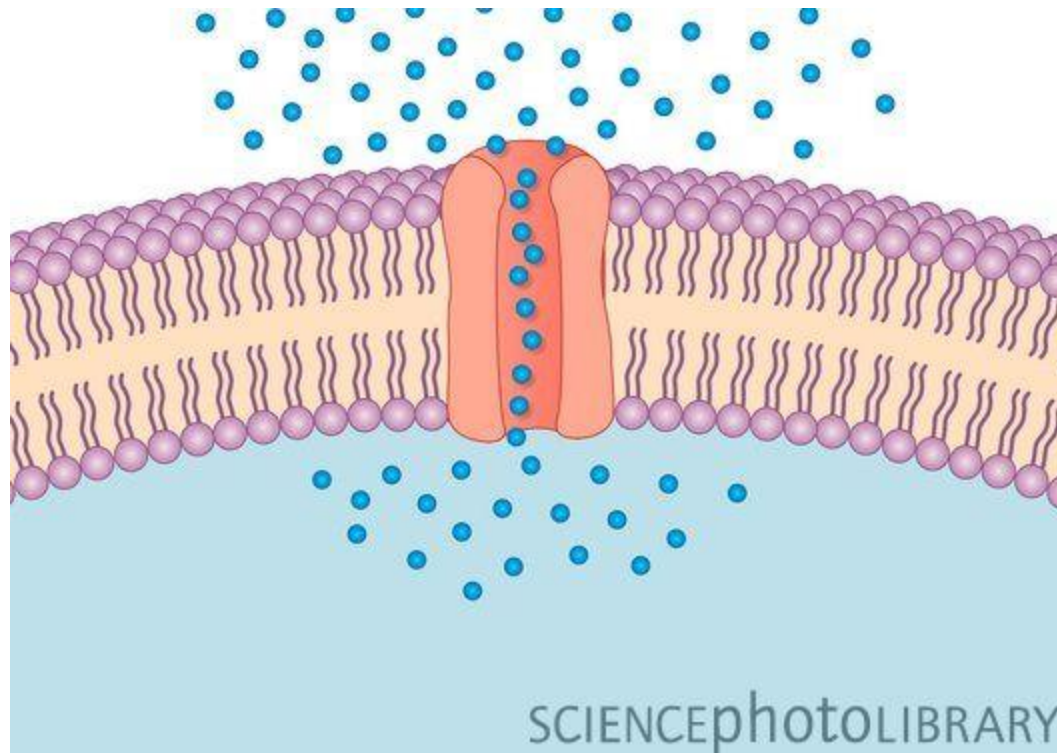
(a) Concentration gradients



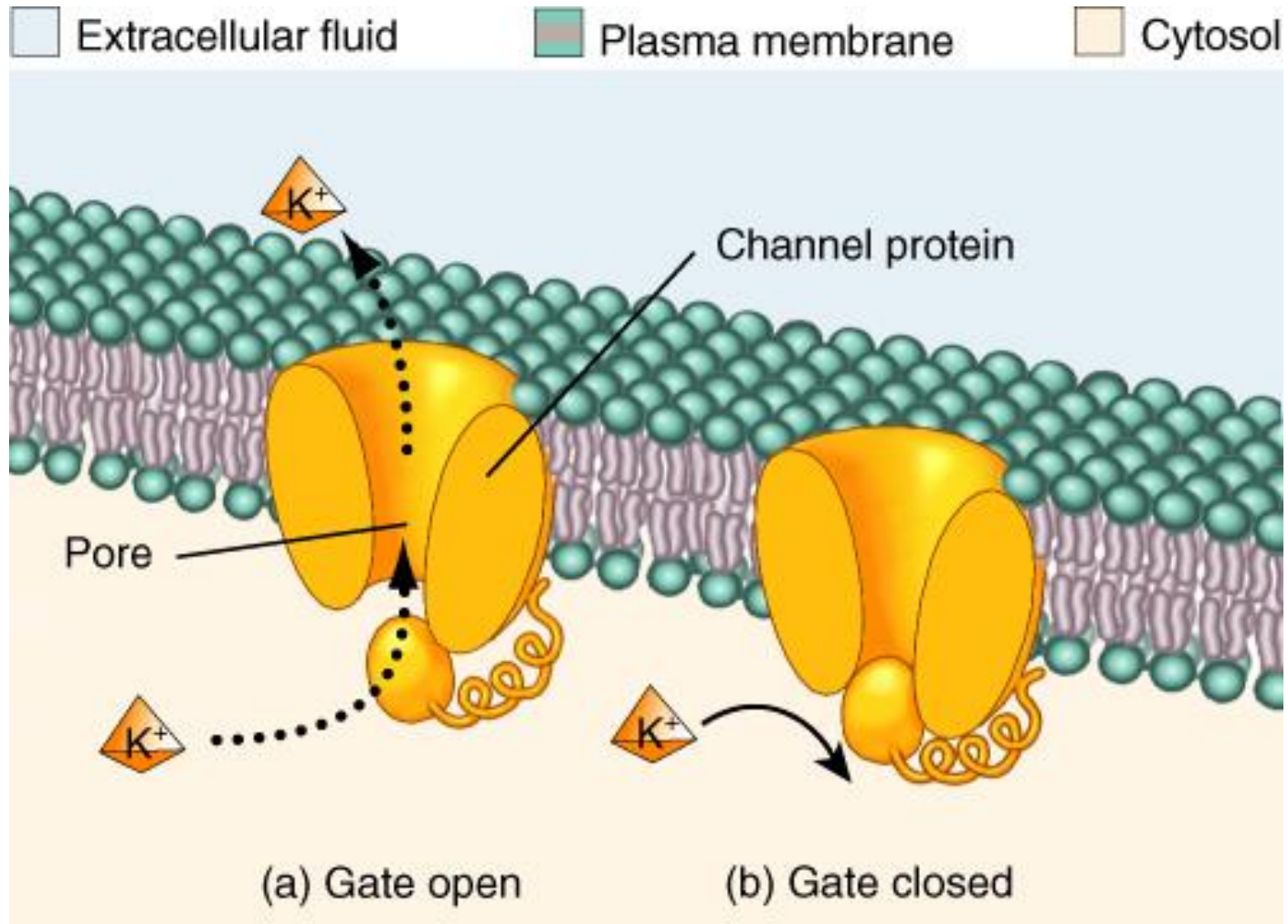
(b) Electrical gradient

- $CO_2$
- $O_2$
- $NO$
- Steroid Hormones
- Monoglycerides

# Diffusion through Channels



# Diffusion through Channels



# Fick's Law

- $J = P \cdot \Delta C$
- $P = D \cdot A / \Delta X$
- $J = D \cdot A \cdot \Delta C / \Delta X$

$J$  = Flux (Rate of diffusion)

$P$  = Permeability

$D$  = Diffusion Coefficient

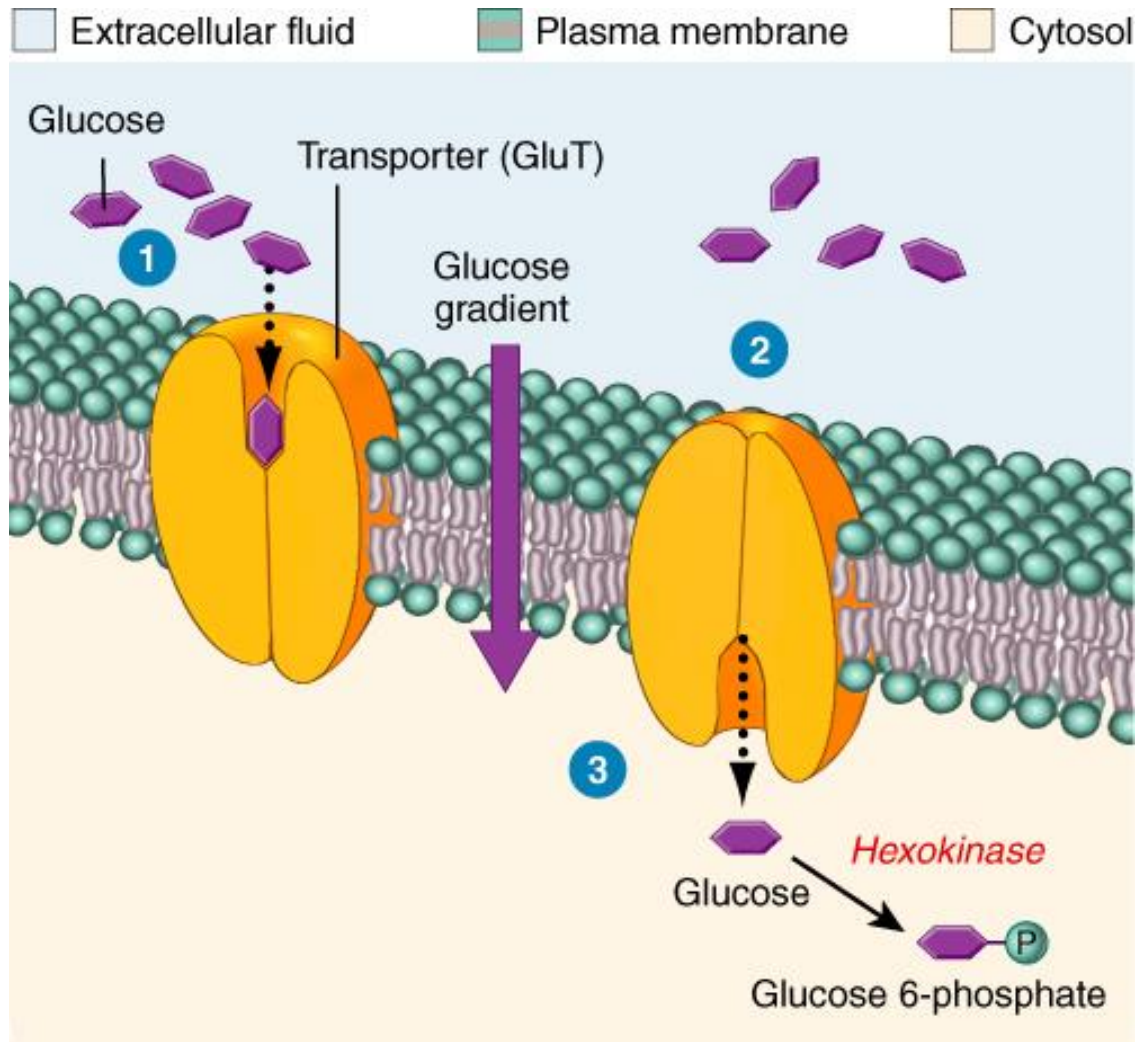
$A$  = Surface area

$C$  = Concentration

$X$  = Membrane thickness

# Facilitated Diffusion

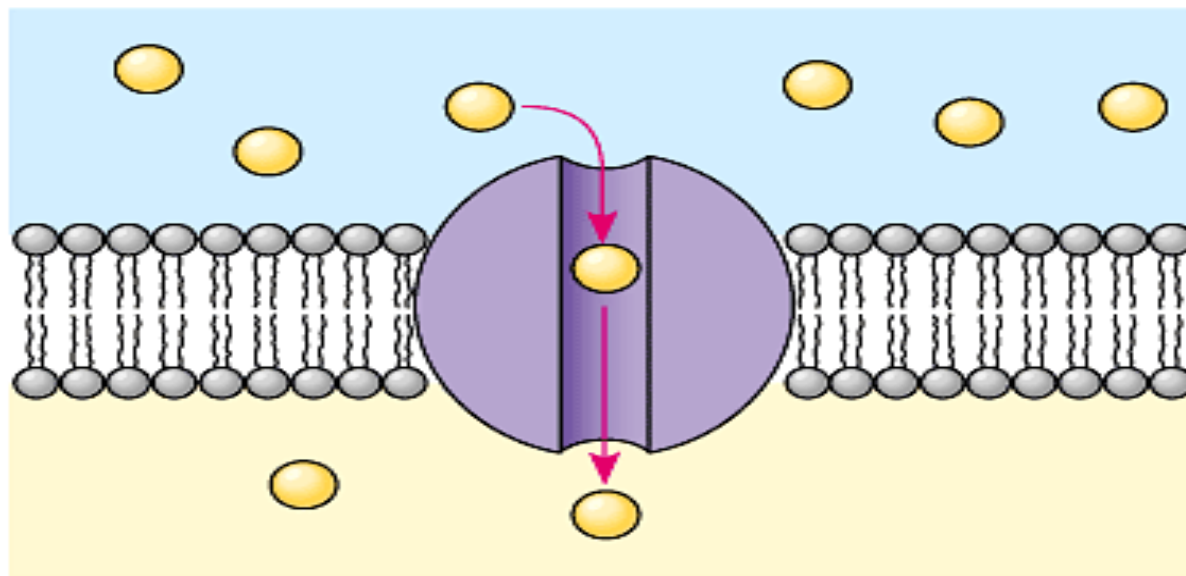
# Facilitated Diffusion



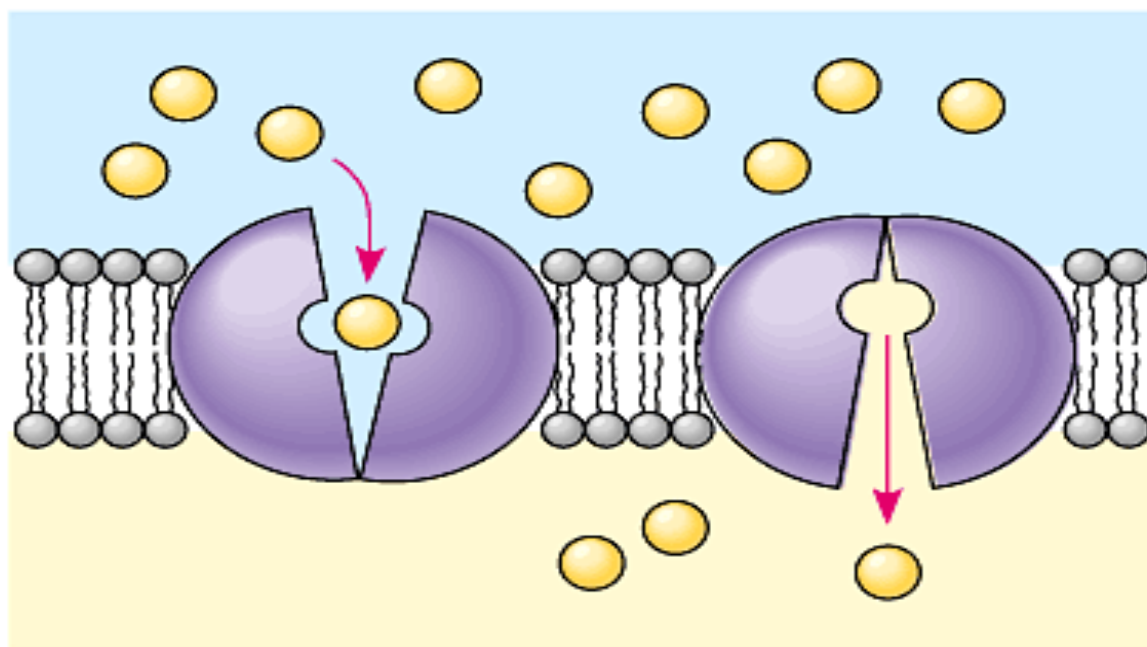
# Facilitated Diffusion

- Aminoacids
- Glucose
- Galactose
- Fructose





**(a)**

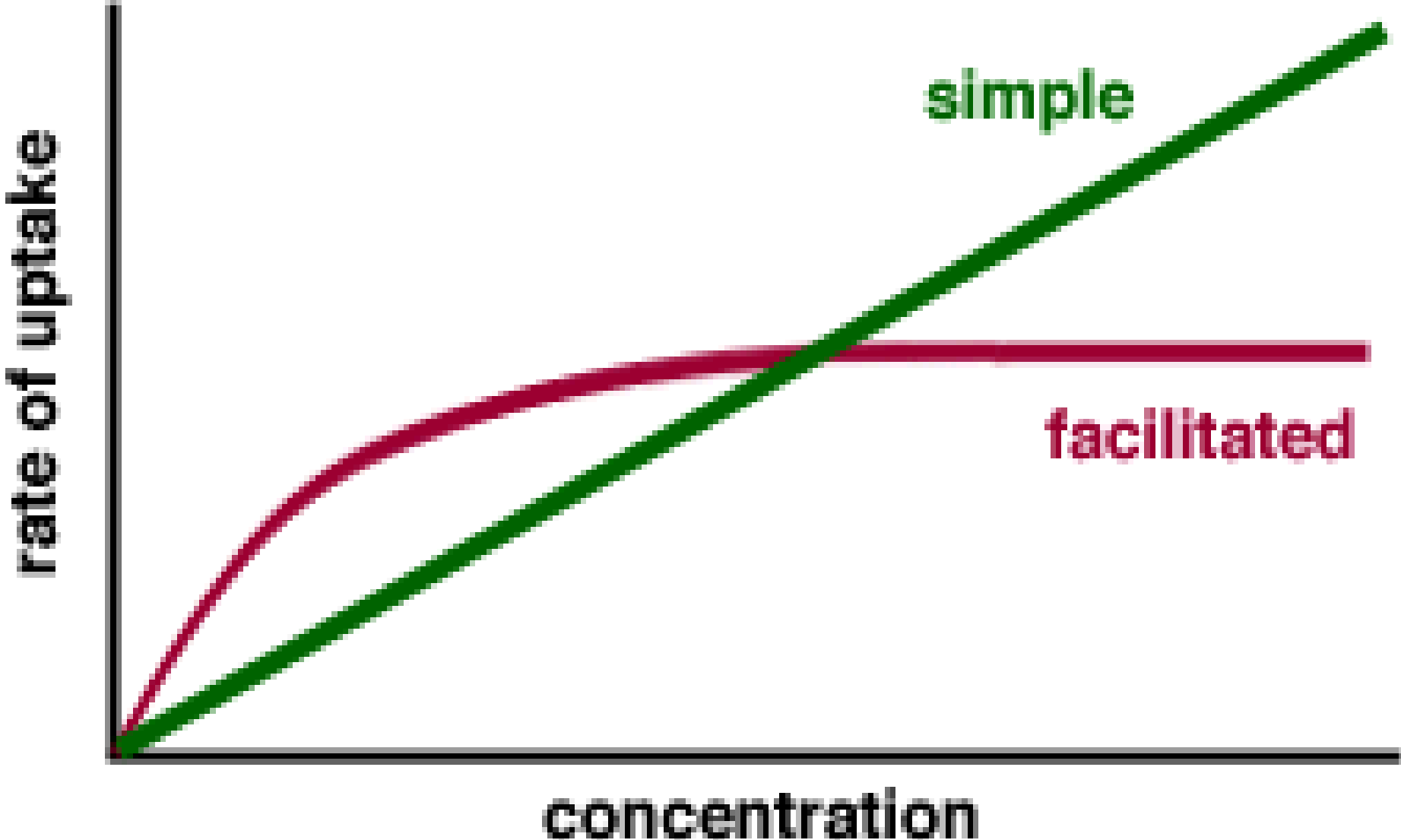


**(b)**

# Question

- Regarding the first Fick's law of diffusion,
- By activation of channels, what parameter is changed?

# Diffusion



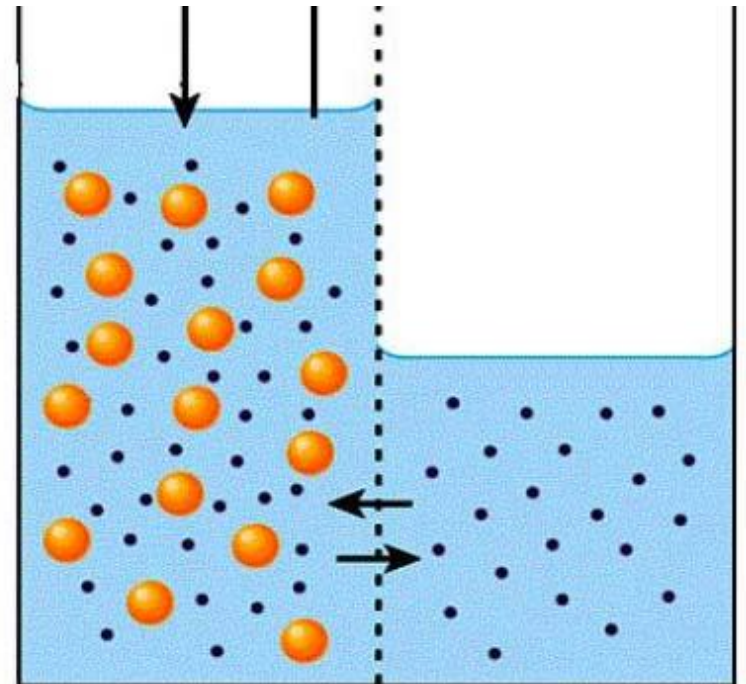
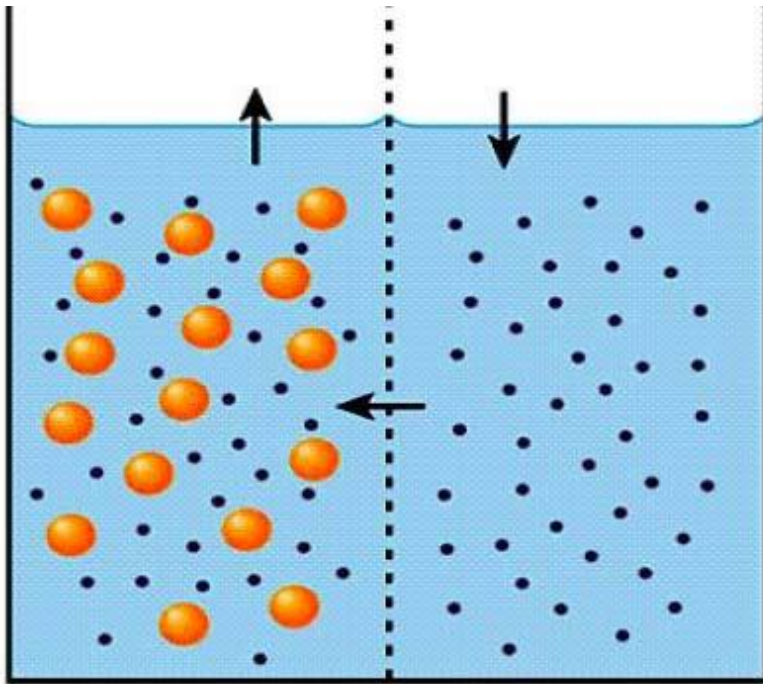
- Regarding **V max:**  
(maximum velocity of transport).

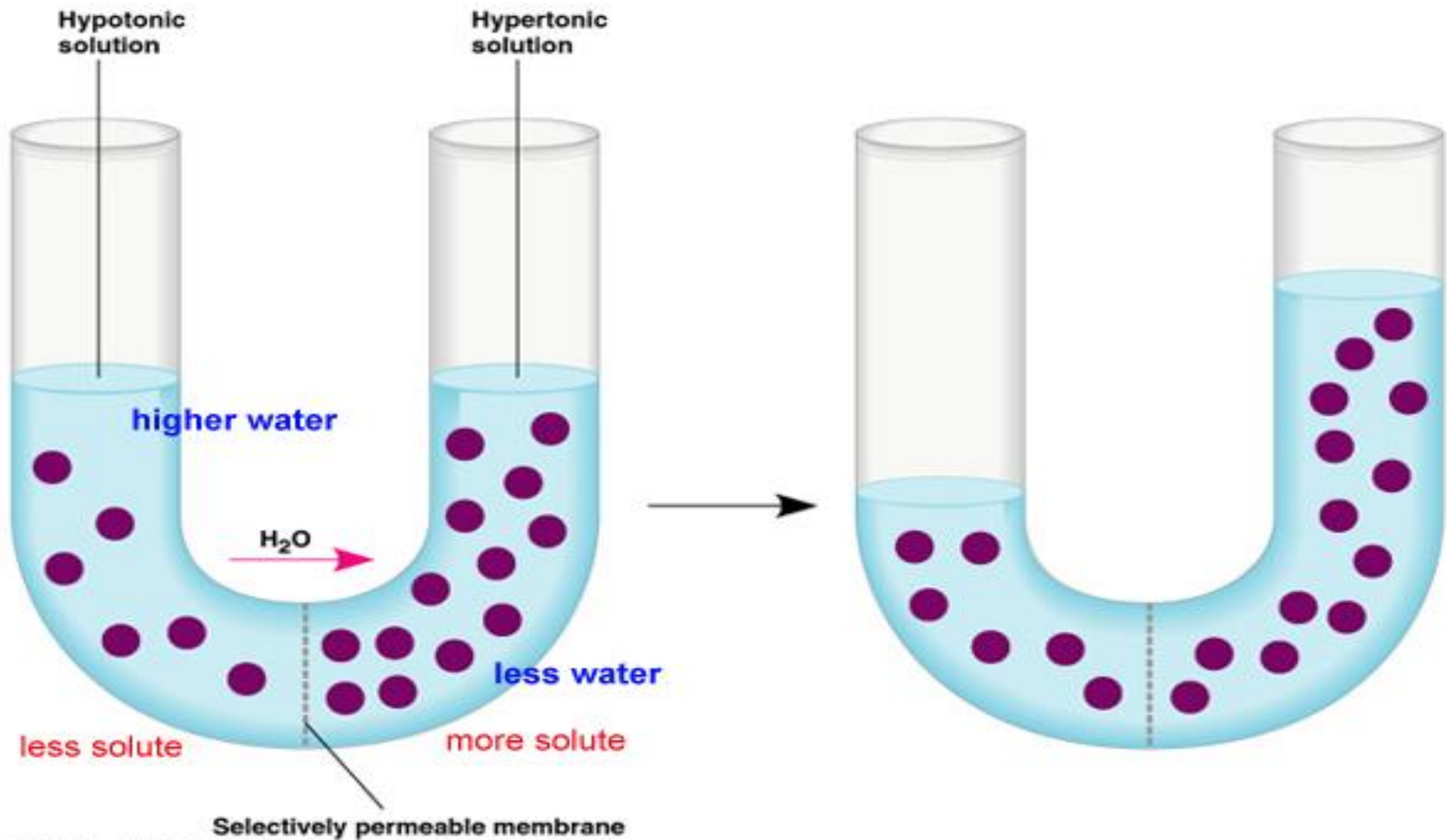
## Questions

- Is it found in Simple diffusion or Facilitated diffusion?
- Explain why having this phenomenon?

# Osmosis

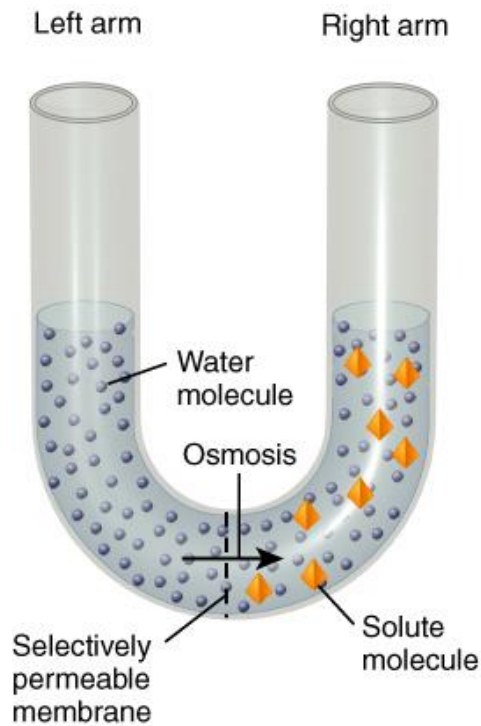
# Osmosis



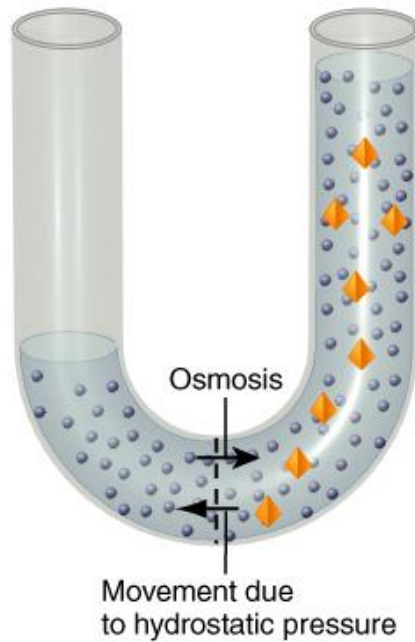




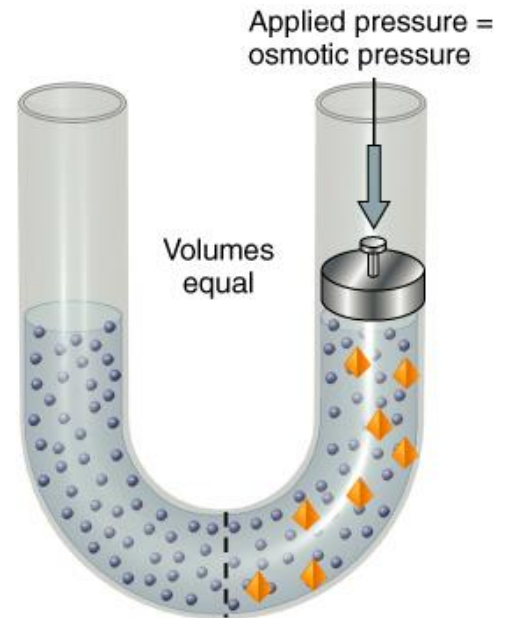
# Osmotic pressure



(a) Starting conditions



(b) Equilibrium



(c) Restoring starting conditions

# Van't Hoff's Law

$$\pi = RTC$$

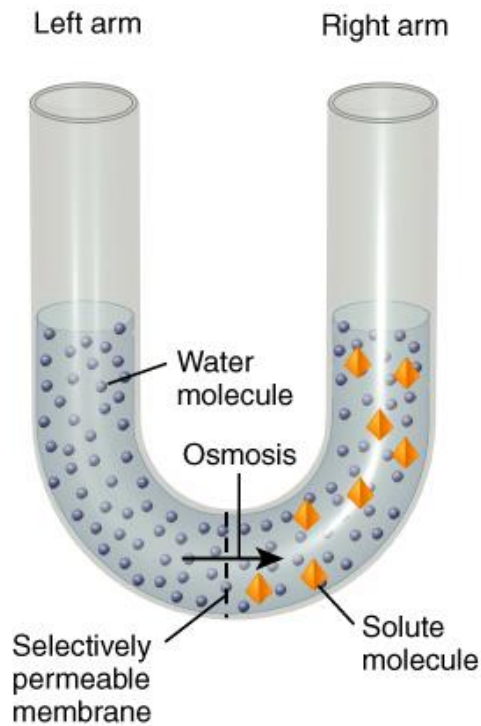
$\pi$  : osmotic Pressure

R = Gas constant

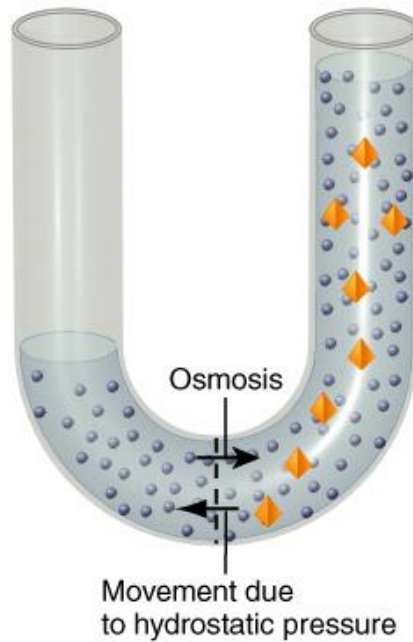
T = Absolute Temperature

C = Concentration

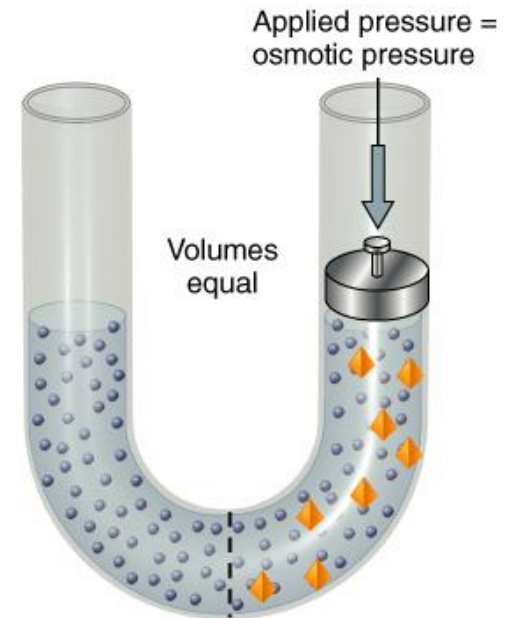
# Osmosis vs. Filtration



(a) Starting conditions



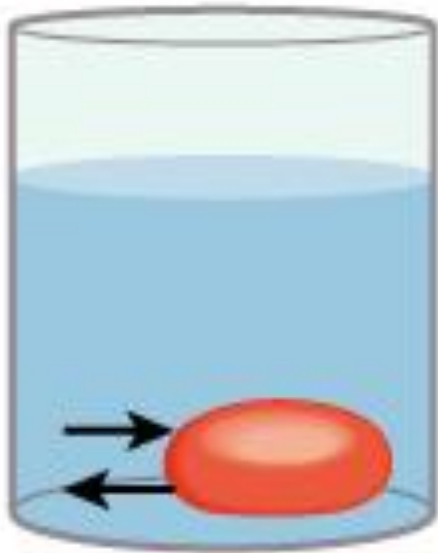
(b) Equilibrium



(c) Restoring starting conditions

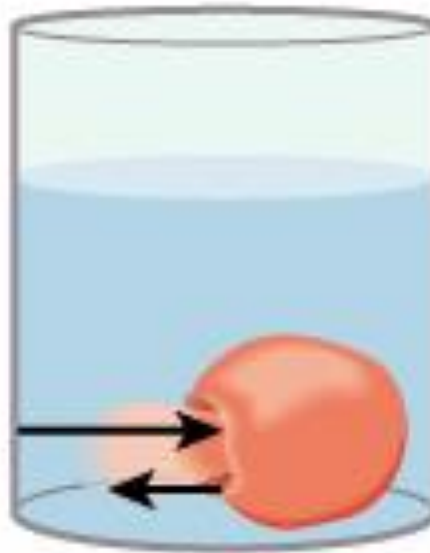
# Tonicity of solution

Isotonic solution



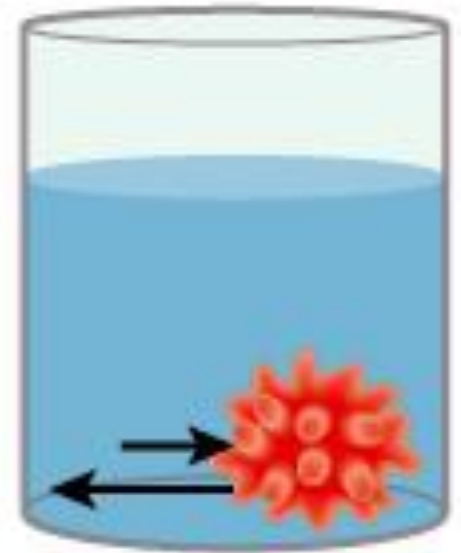
(a) Normal RBC shape

Hypotonic solution



(b) RBC undergoes hemolysis

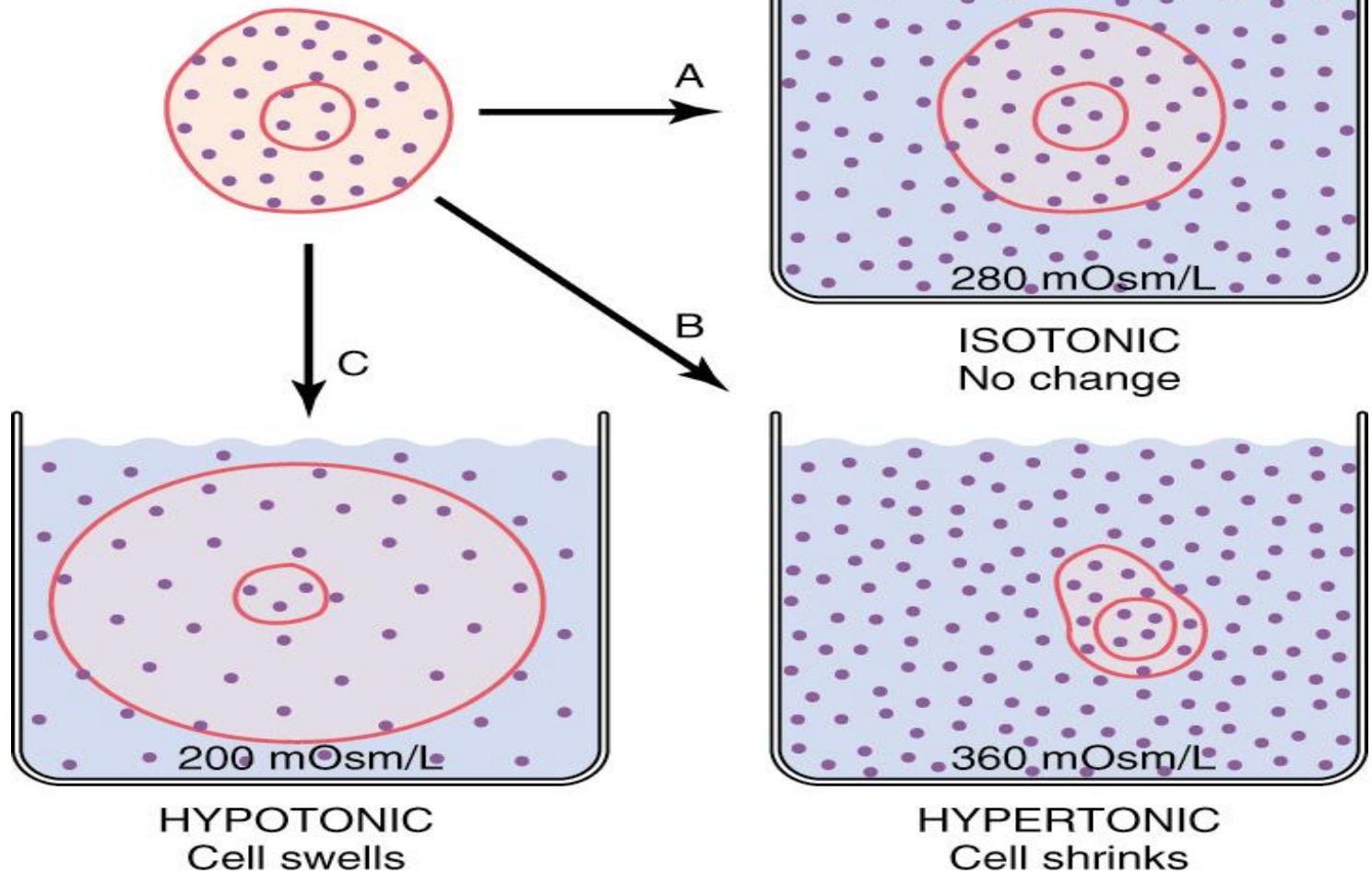
Hypertonic solution



(c) RBC undergoes crenation

Hypotonic  
(cell  
swells)

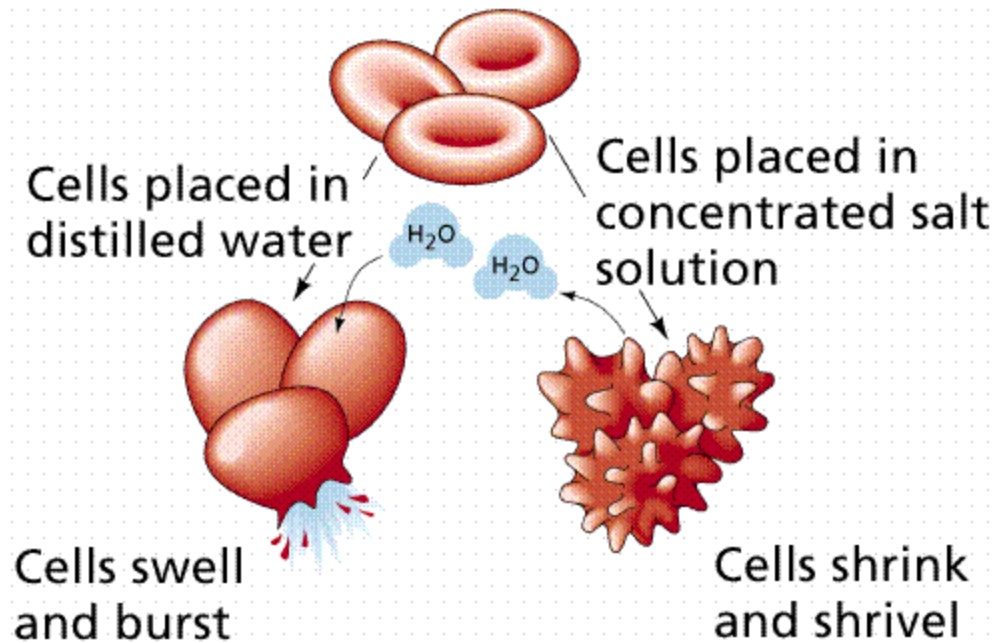
Isotonic  
(no change)



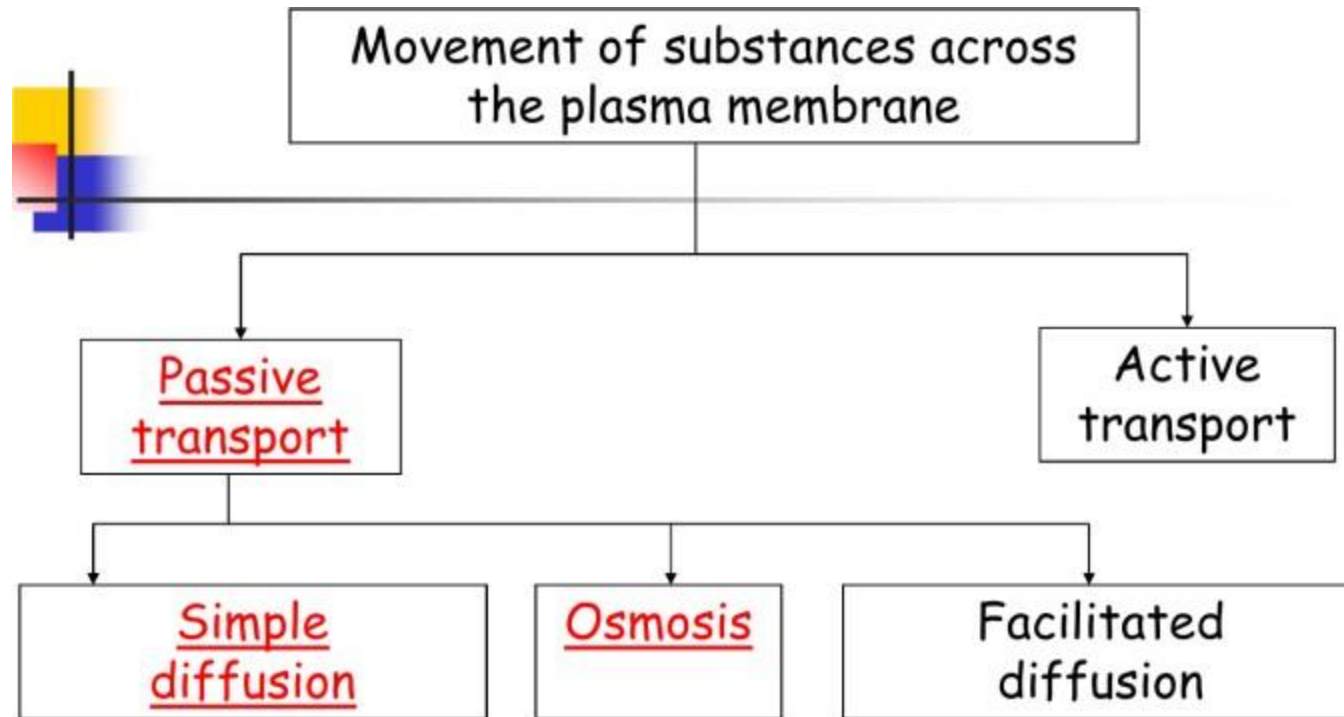
Hypertonic  
(cell  
shrinks)

Figure 25-5; Guyton and Hall

What happens by placing Red Blood Cells in Hypertonic or hypotonic solution



# Transport across Plasma Membranes





# Transport through plasma membranes

## Passive transport modalities

**-Simple diffusion:** transport through lipid bilayer, transport through channels, Ficks law of diffusion.

**-Facilitated diffusion:** by carriers

Differences in diffusion Kinetics between the previous modalities

Equivalent Concentration of particles.

# Transport through plasma membranes

## Passive transport modalities

-**Osmosis**: concept of osmotic pressure (Van't Hoff's law), Oncotic (Colloid-osmotic) pressure. Osmolarity, Osmolality  
Hydrostatic pressure and filtration

# Transport through plasma membranes

## Active transport modalities

- **Primary active transport** (ATP-ase carriers or Pumps) (functions of pumps:  $\text{Na}^+/\text{K}^+$  pump,  $\text{Ca}^{++}$  pump,  $\text{H}^+$  pump,  $\text{H}^+/\text{K}^+$  pump).
- **Secondary active transport** ( $\text{Na}^+$  dependent carriers) examples

# Transport through plasma membranes

## Active transport modalities

- **Vesicular transport:** endocytosis, phagocytosis, transcytosis, pinocytosis and exocytosis and its control in secretory cells.-

**TABLE 3.1** Passive Membrane Transport Processes

PROCESS	ENERGY SOURCE	DESCRIPTION	EXAMPLES
<b>DIFFUSION</b>			
Simple diffusion	Kinetic energy	Net movement of particles (ions, molecules, etc.) from an area of their higher concentration to an area of their lower concentration, that is, along their concentration gradient	Movement of fats, oxygen, carbon dioxide through the lipid portion of the membrane
Facilitated diffusion	Kinetic energy	Same as simple diffusion, but the diffusing substance is attached to a lipid-soluble membrane carrier protein or moves through a membrane channel	Movement of glucose and some ions into cells
Osmosis	Kinetic energy	Simple diffusion of water through a selectively permeable membrane	Movement of water into and out of cells directly through the lipid phase of the membrane or via membrane pores (aquaporins)
<b>FILTRATION</b>			
	Hydrostatic pressure	Movement of water and solutes through a semipermeable membrane (either through the plasma membrane or between cells) from a region of higher hydrostatic pressure to a region of lower hydrostatic pressure, that is, along a pressure gradient	Movement of water, nutrients, and gases through a capillary wall; formation of kidney filtrate

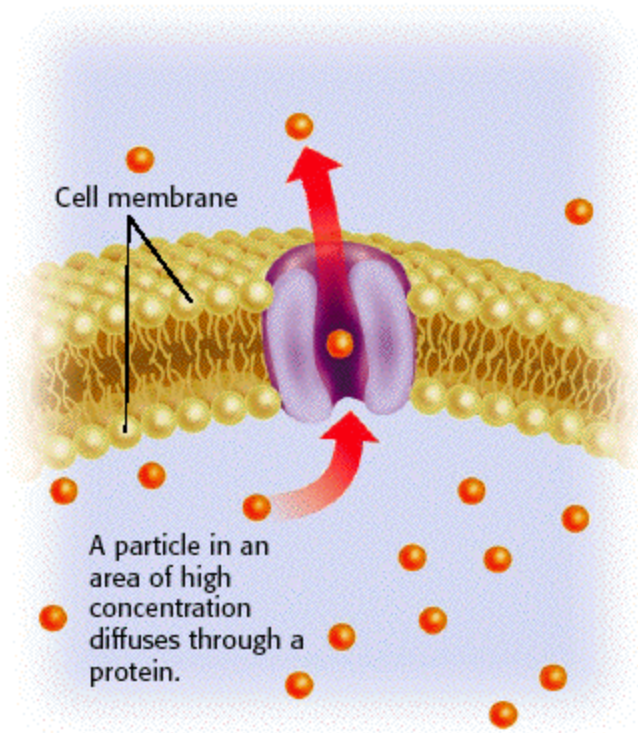
Copyright © 2006 Pearson Education, Inc., publishing as Benjamin Cummings.

# Active Transport Mechanisms

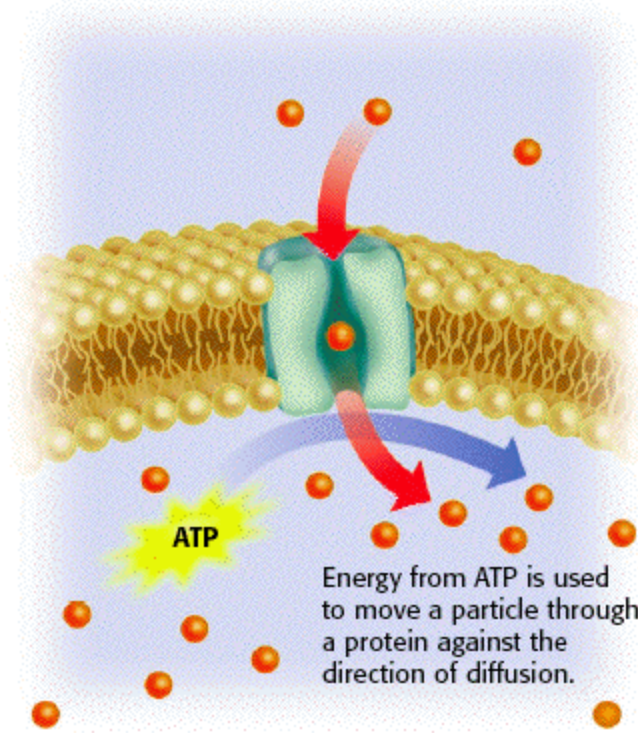
- Primary active transport
- Secondary active transport
- Vesicular transport

## Passive and Active Transport

### PASSIVE TRANSPORT



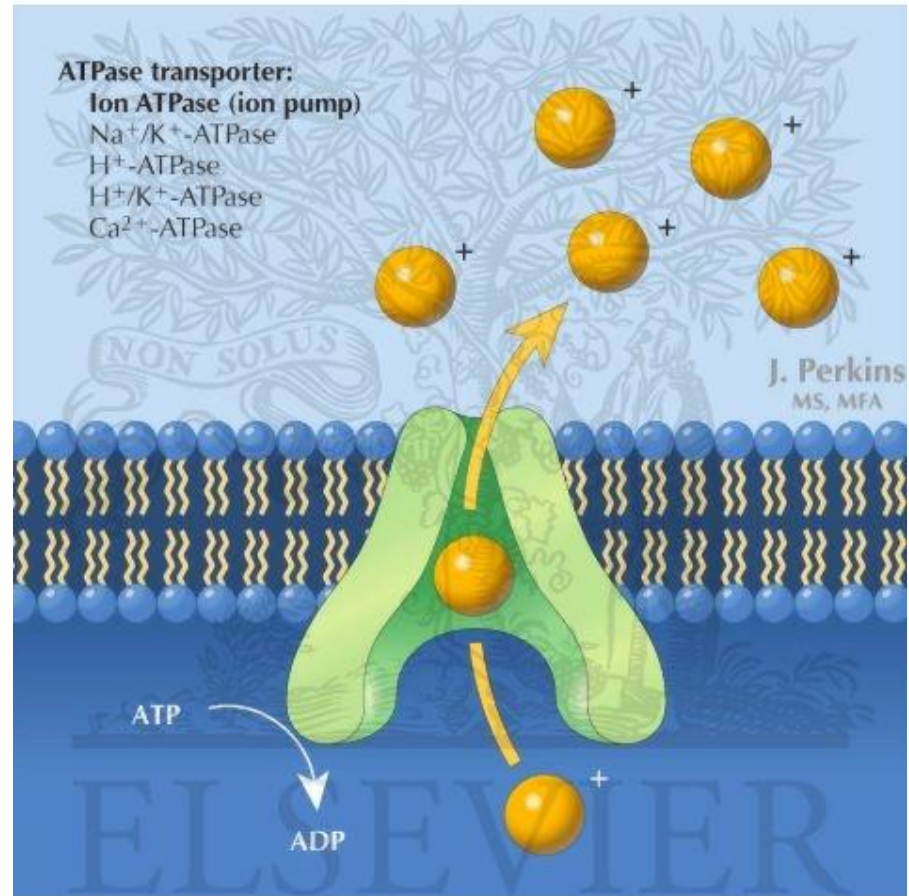
### ACTIVE TRANSPORT



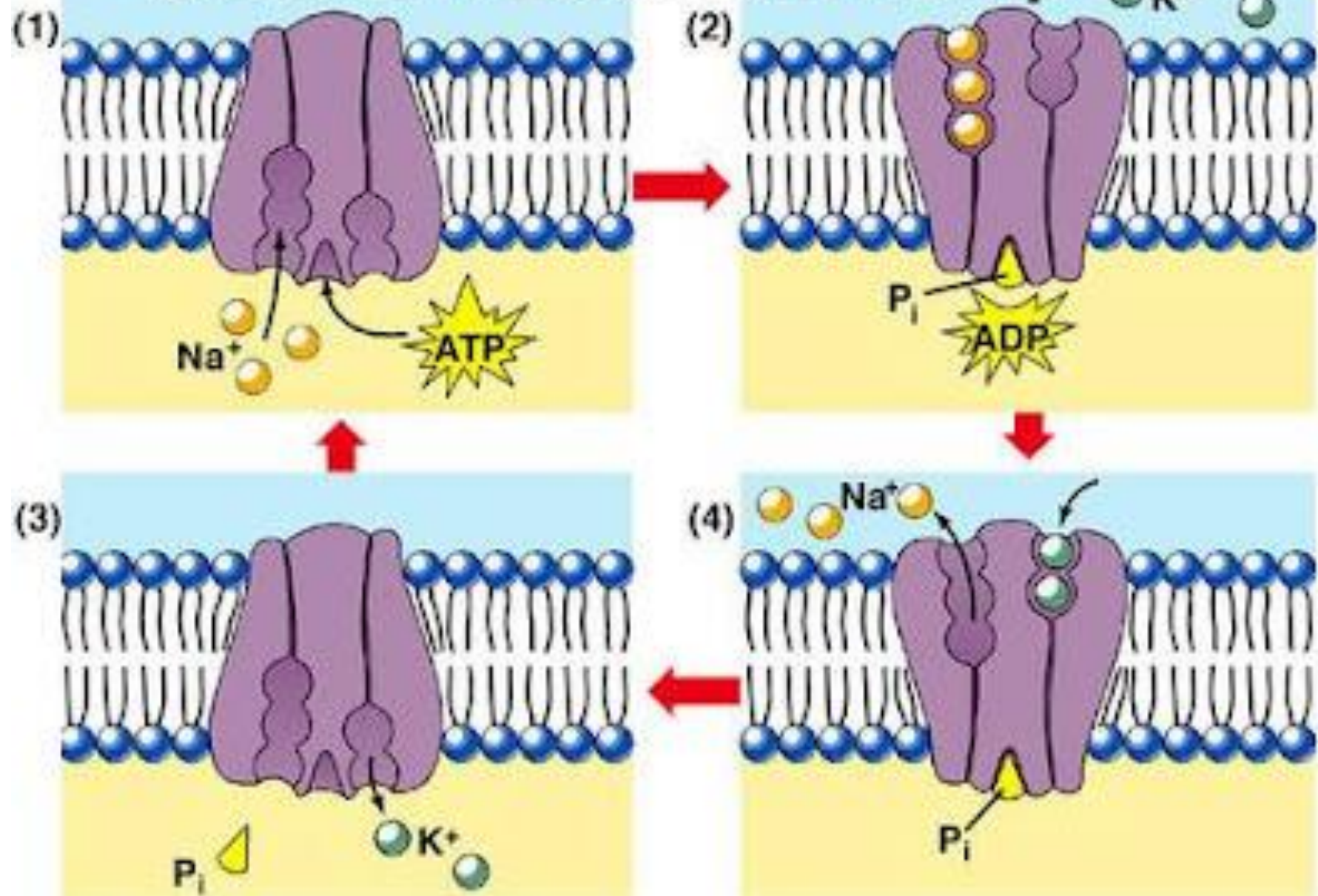


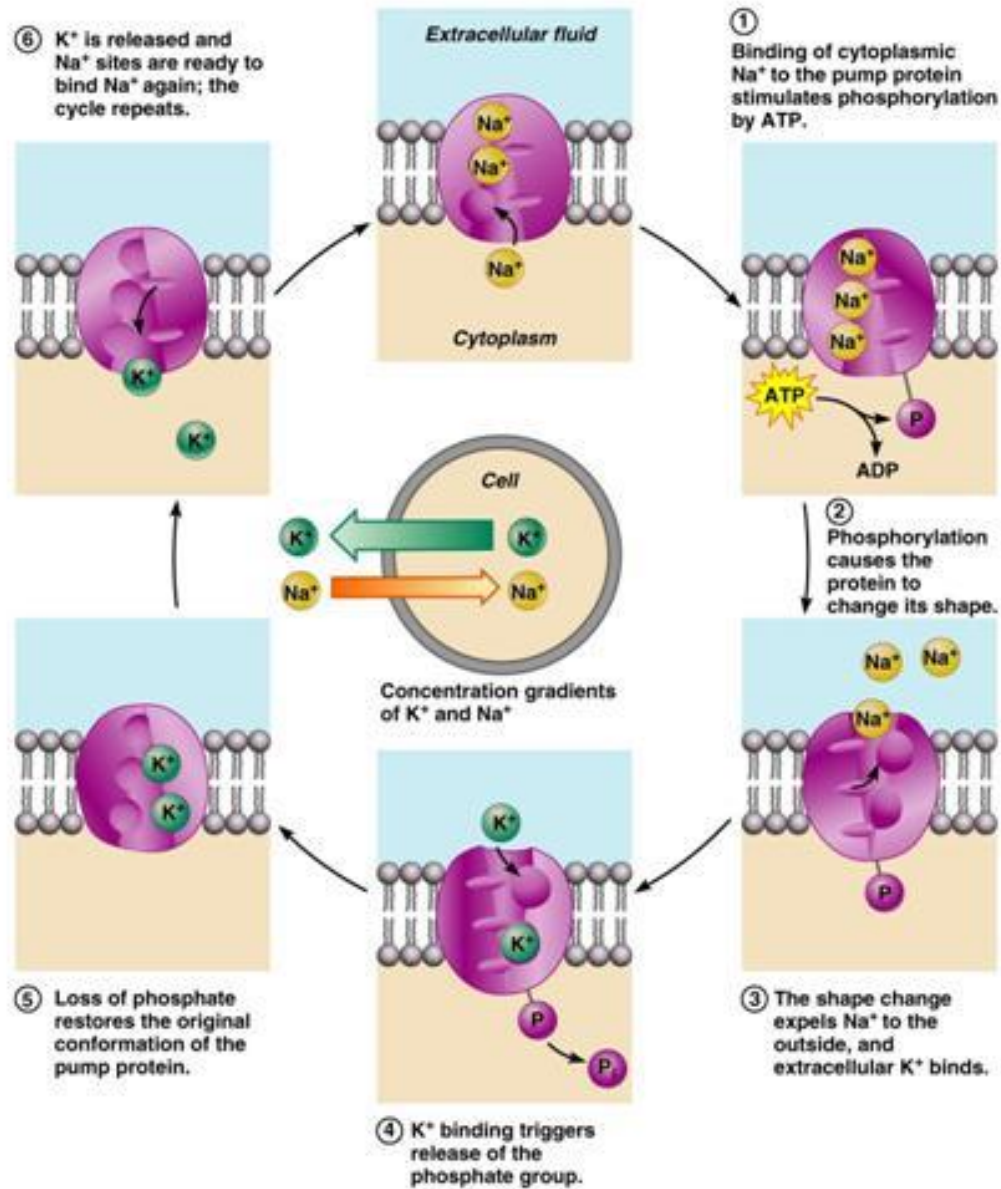
# Types of Pumps

- $\text{Na}^+/\text{K}^+$  pump
- $\text{H}^+$  pump
- $\text{H}^+/\text{K}^+$  pump
- $\text{Ca}^{++}$  pump



# Sodium-Potassium Pump

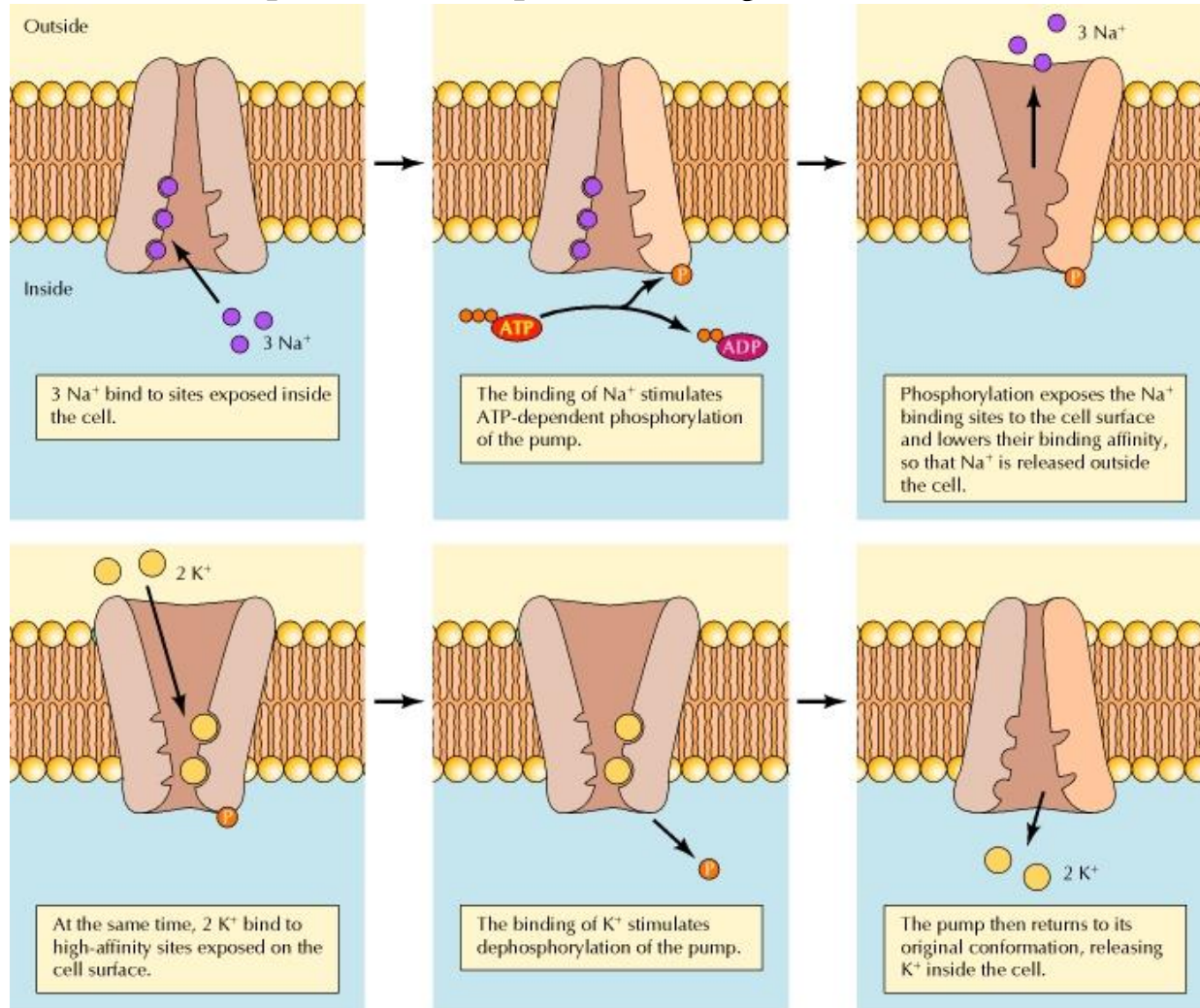




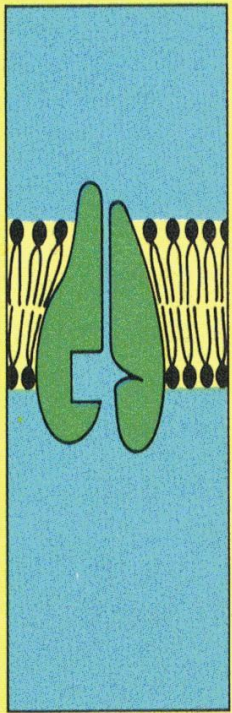
**Figure 3.10**



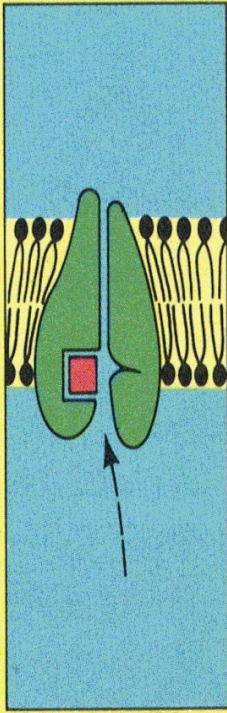
# Phosphorylation & dephosphorylation



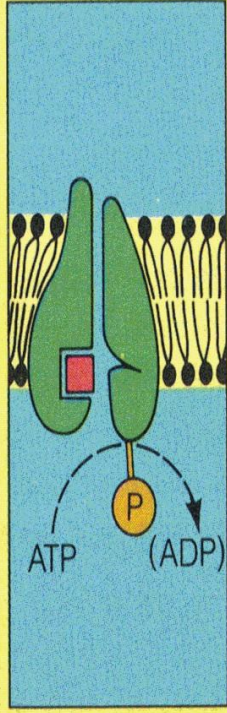




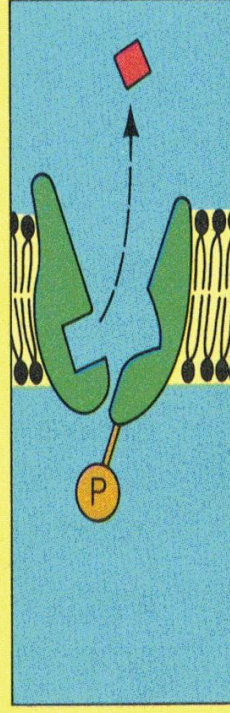
**a** Transport protein with two binding sites.



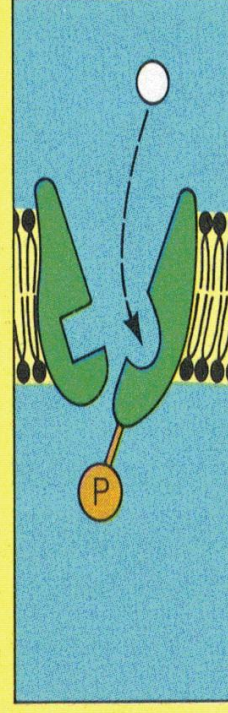
**b** Specific solute binds at one site.



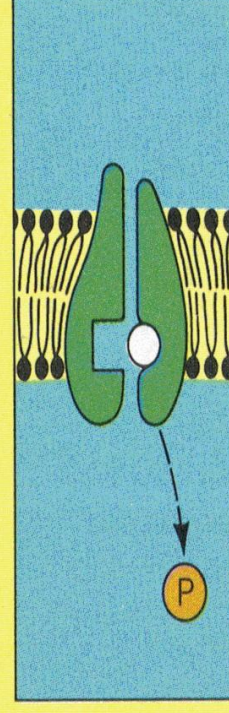
**c** Phosphate group is transferred from ATP to protein.



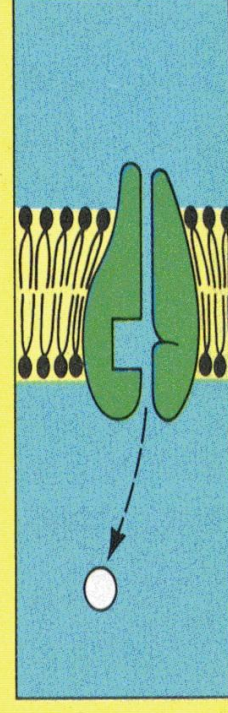
**d** Protein changes shape, pumps the solute across membrane.



**e** The other binding site is now exposed, different solute binds to it.



**f** Phosphate group is released, protein returns to original shape.

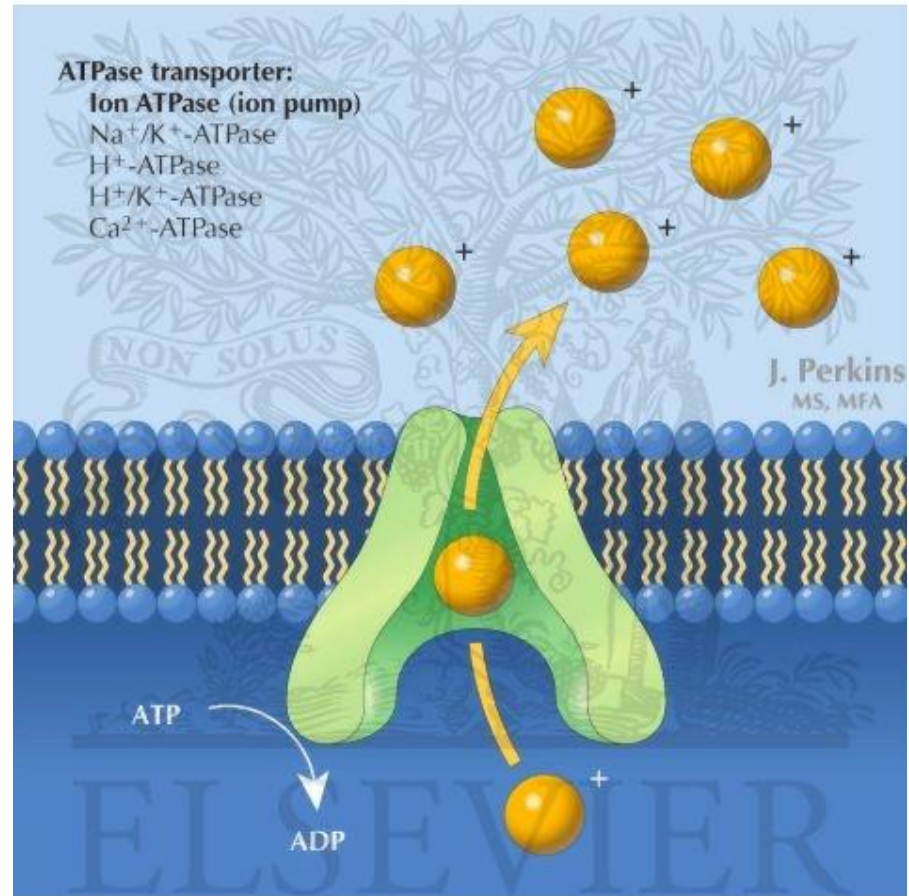


**g** The shape change causes the solute to be released.



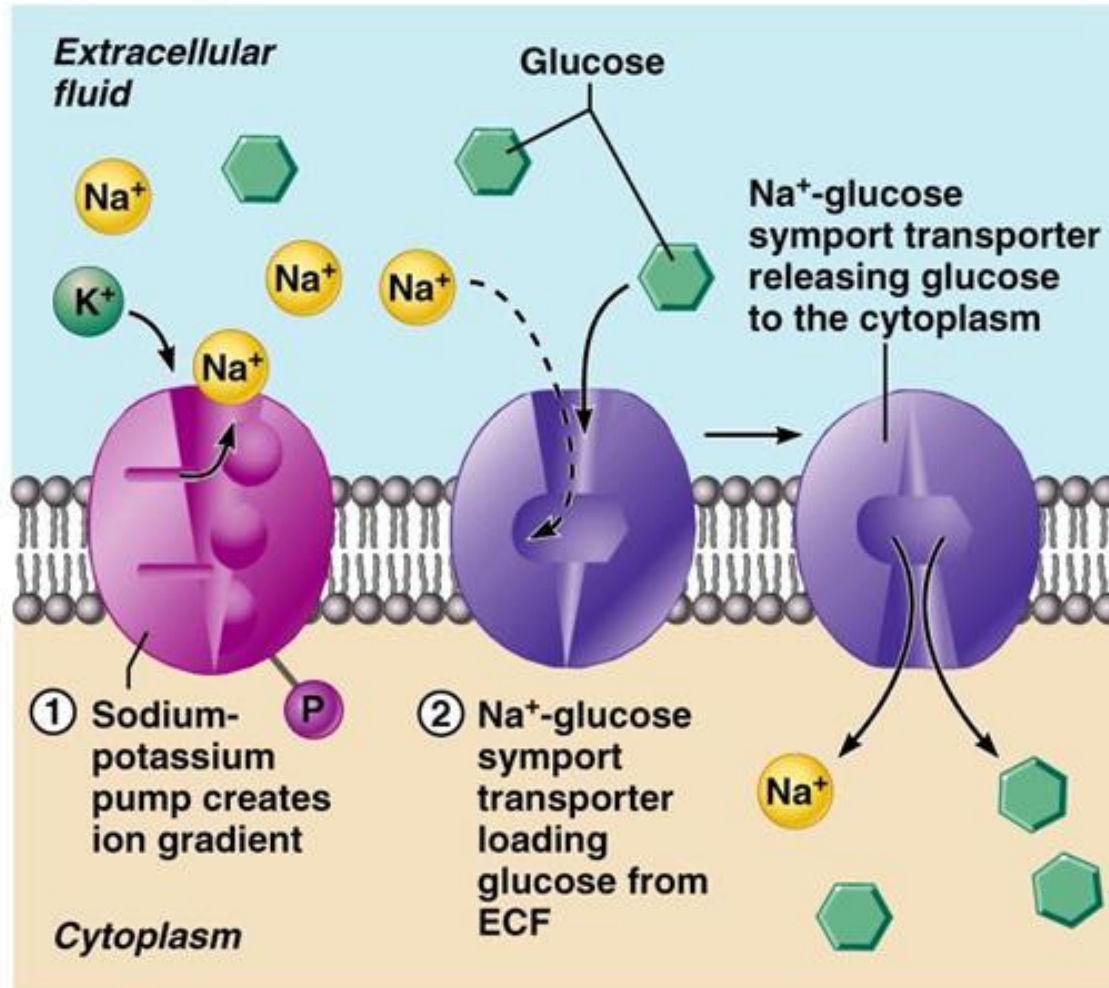
# Types of Pumps

- $\text{Na}^+/\text{K}^+$  pump
- $\text{H}^+$  pump
- $\text{H}^+/\text{K}^+$  pump
- $\text{Ca}^{++}$  pump



# The importance of pumps for cell functions

# Secondary Active Transport



Copyright © 2006 Pearson Education, Inc., publishing as Benjamin Cummings.

Figure 3.11



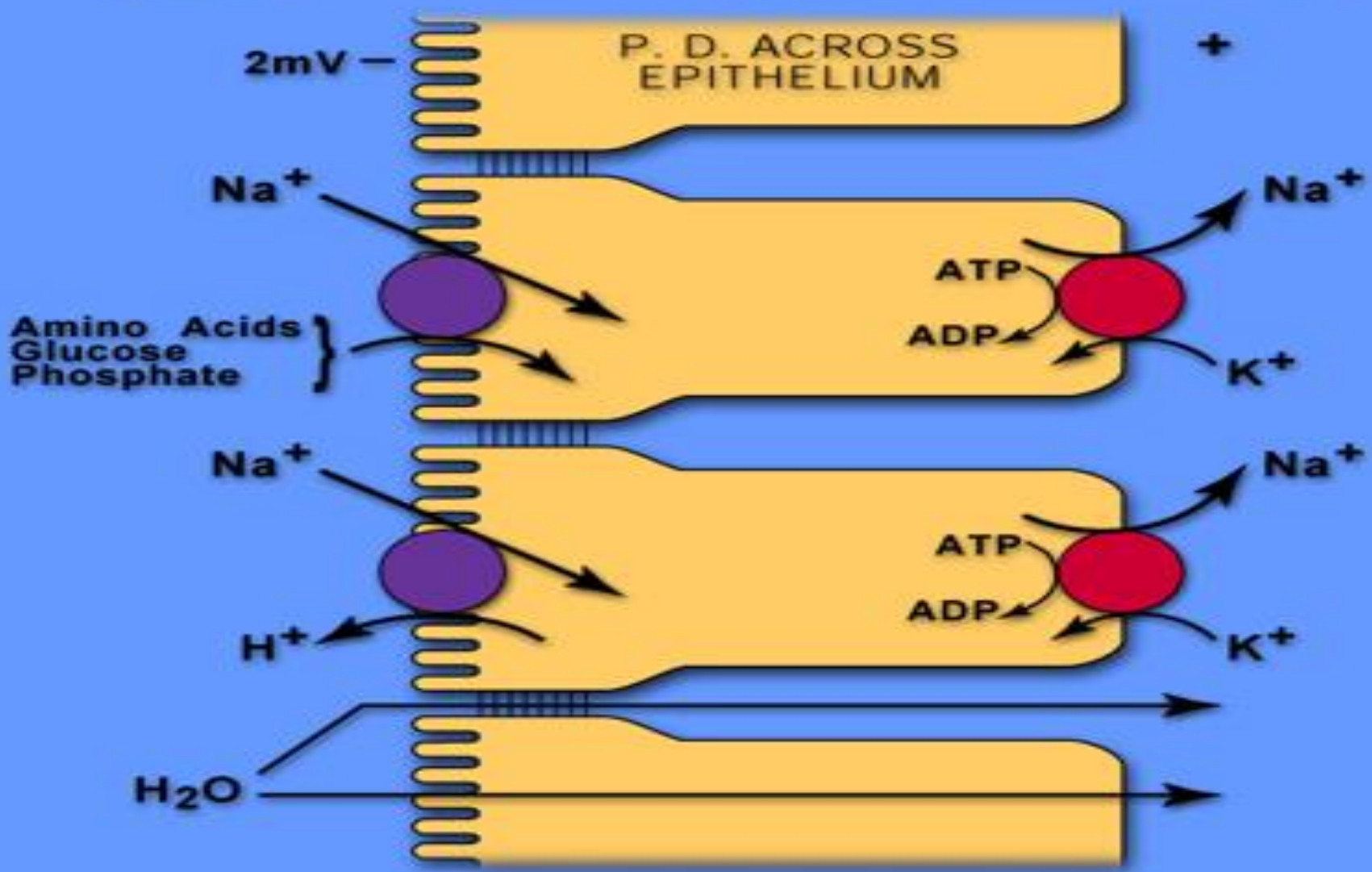
# Na<sup>+</sup> ENTRY INTO PROXIMAL TUBULAR EPITHELIAL CELLS

## EARLY PROXIMAL

TUBULAR LUMEN

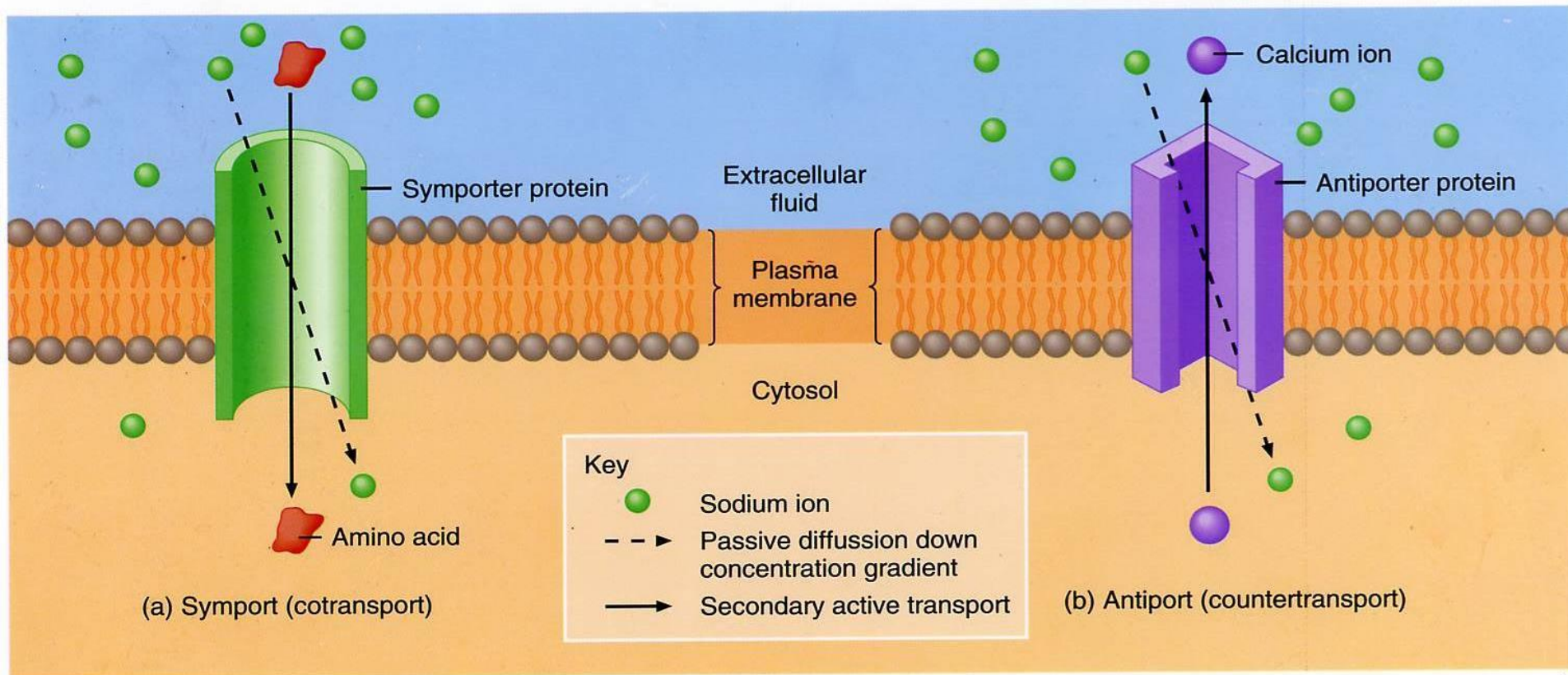
EPITHELIAL CELL

RENAL ISF



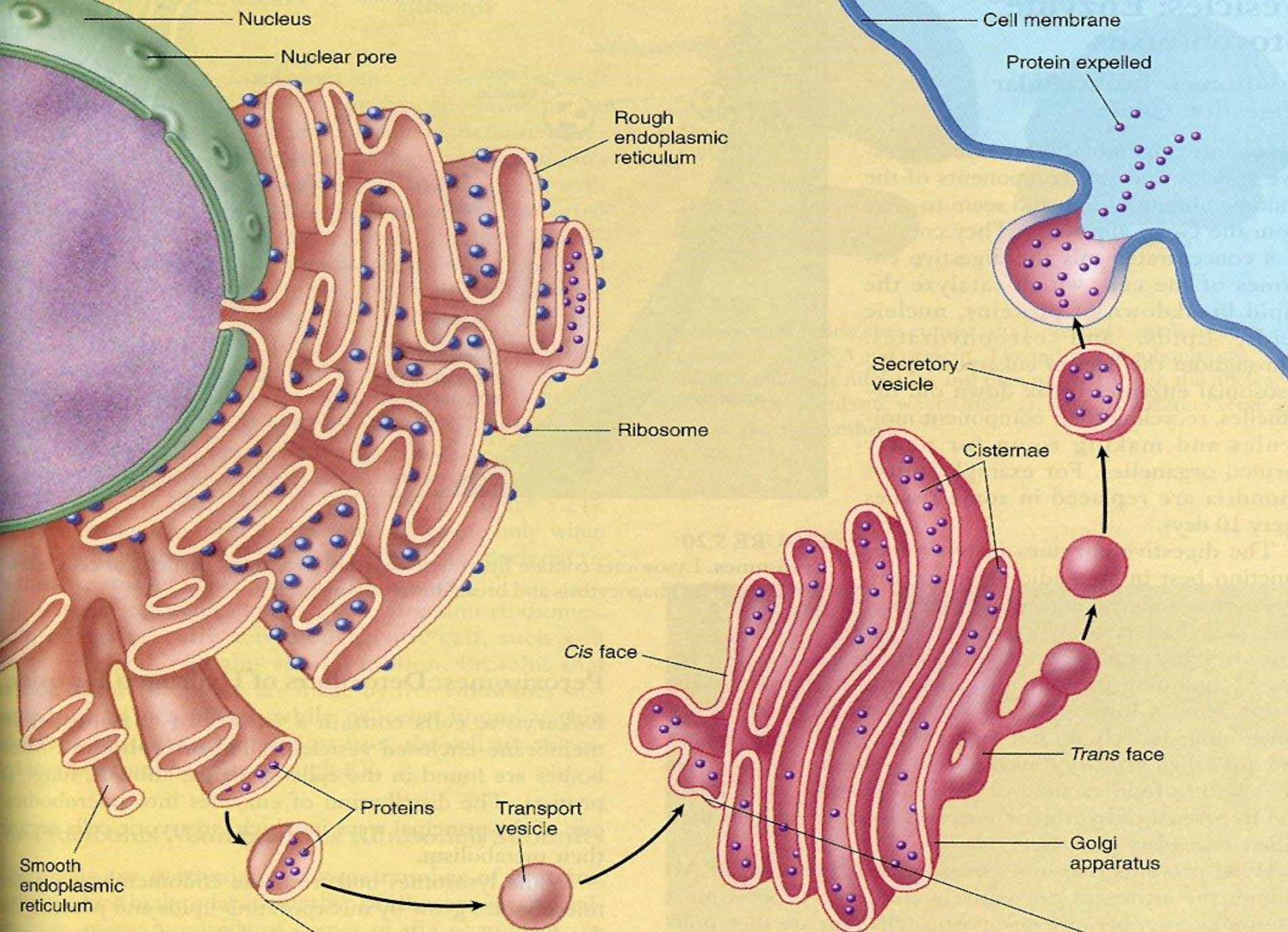
# Secondary Active Transport

- Co-transport
- Counter transport



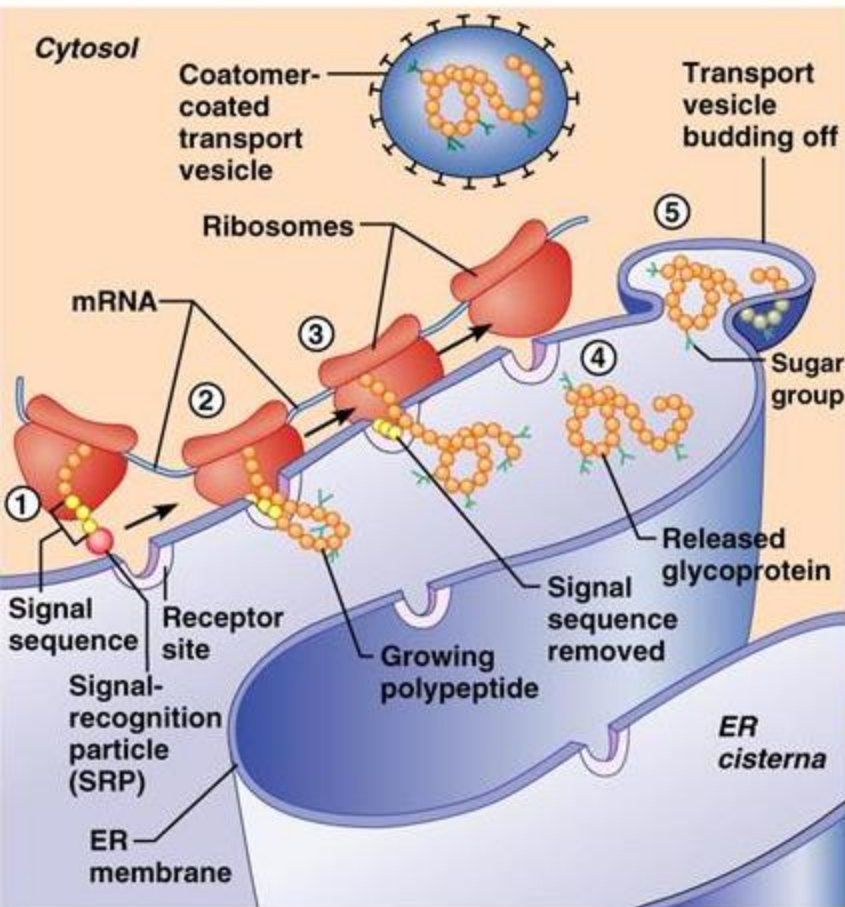
# Vesicular Transport







# Formation of vesicles at ER



Copyright © 2006 Pearson Education, Inc., publishing as Benjamin Cummings.

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.

## Rough ER and Protein Synthesis

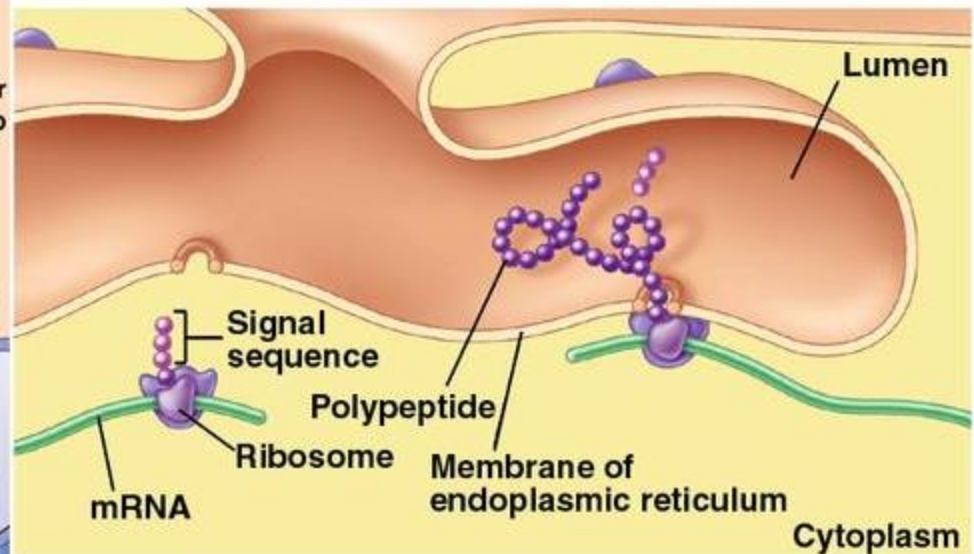
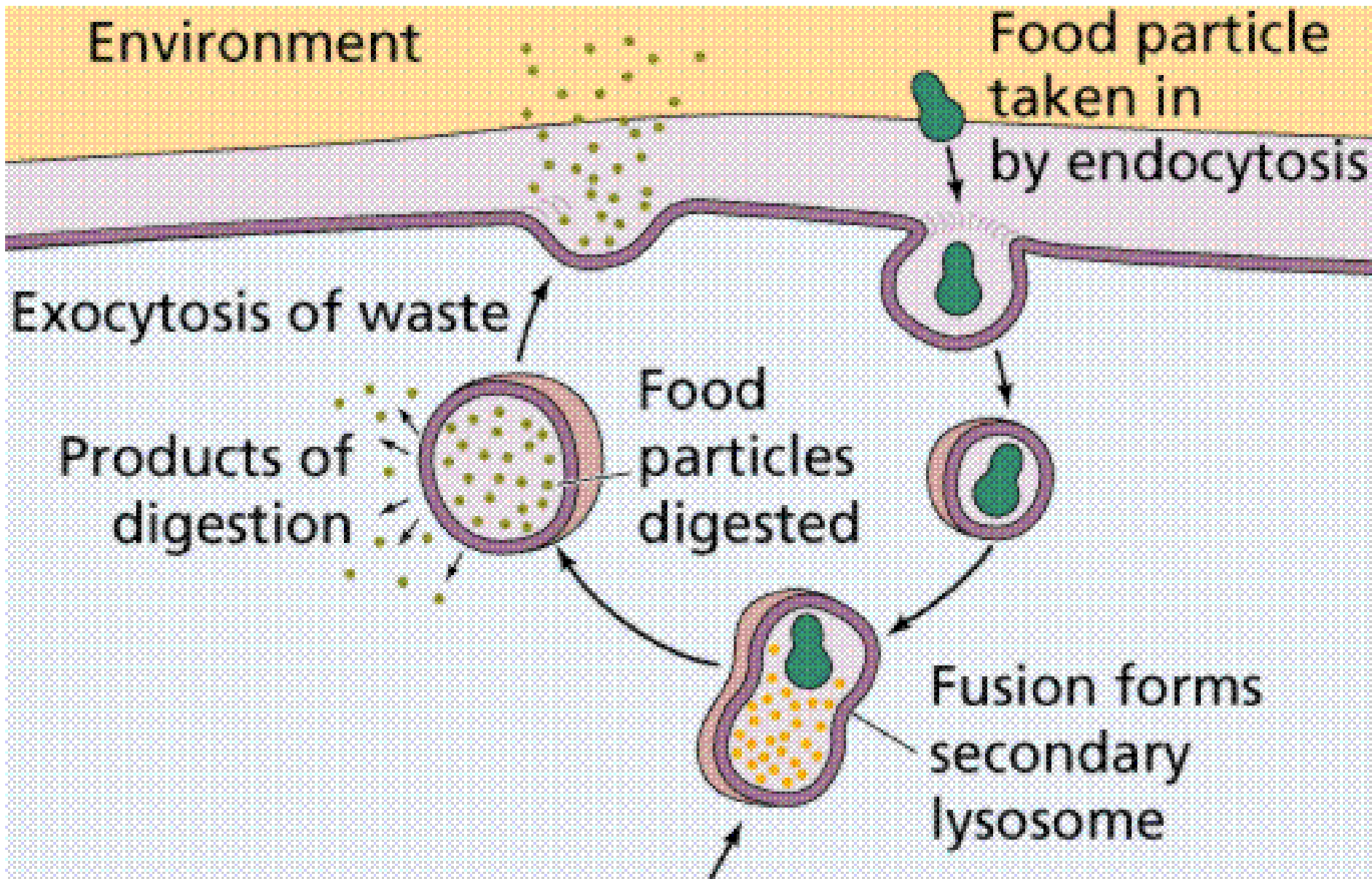
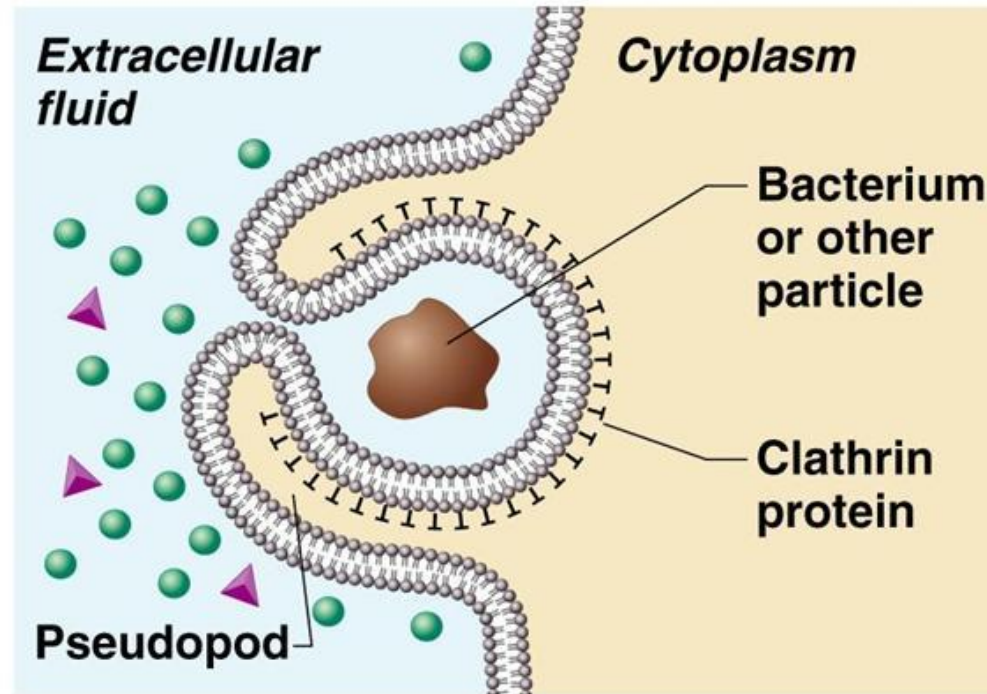


Figure 3.19

# Endocytosis



# Phagocytosis

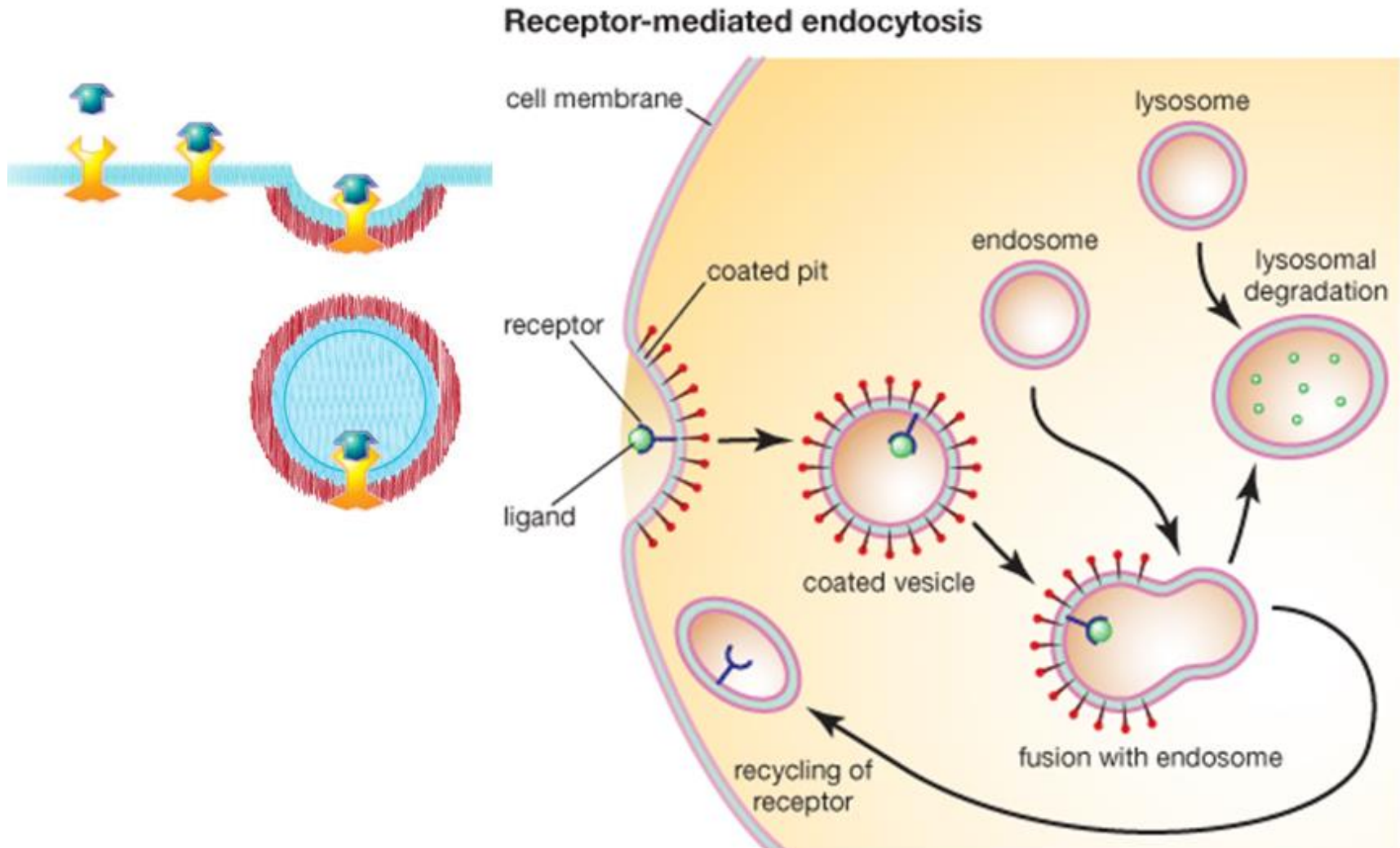


**(b) Phagocytosis**

Copyright © 2006 Pearson Education, Inc., publishing as Benjamin Cummings.

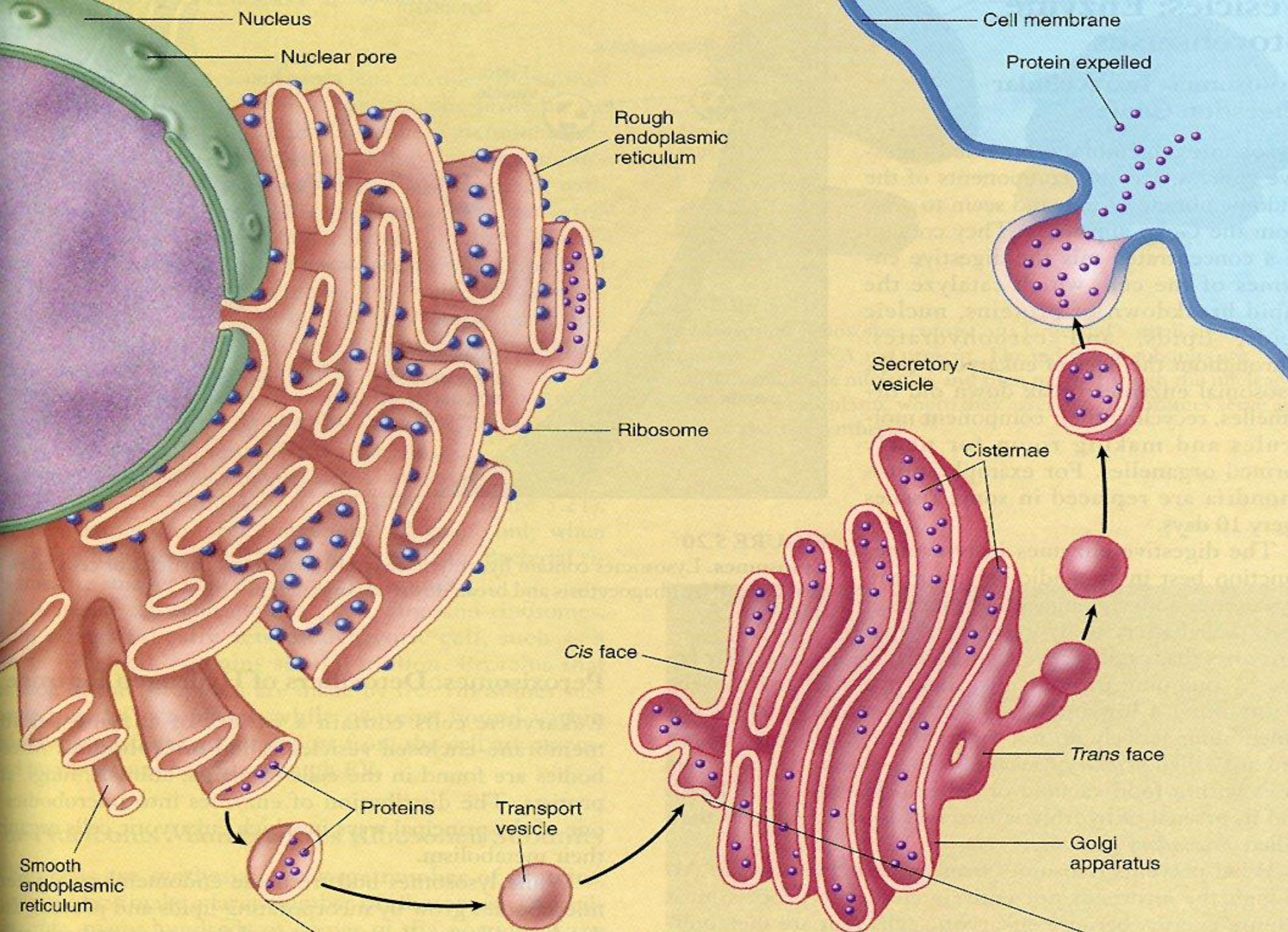
Figure 3.13b

# Receptor mediated endocytosis



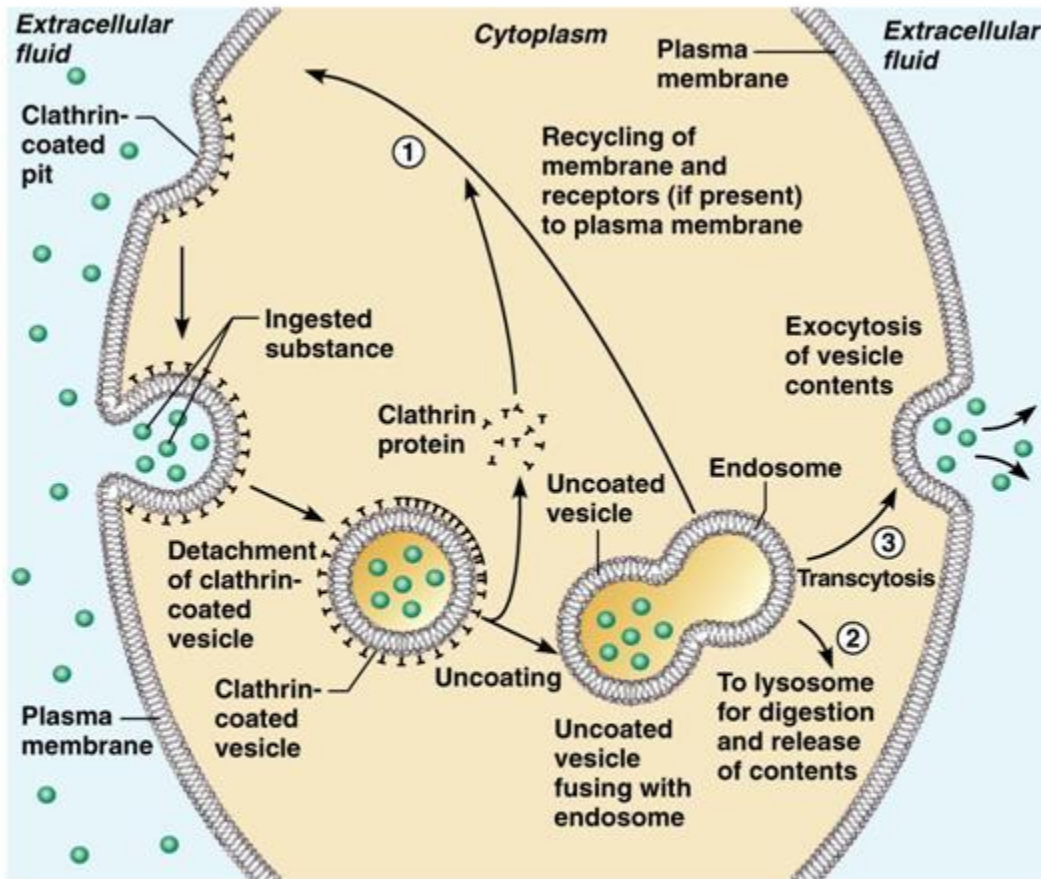


# Exocytosis





# Transcytosis



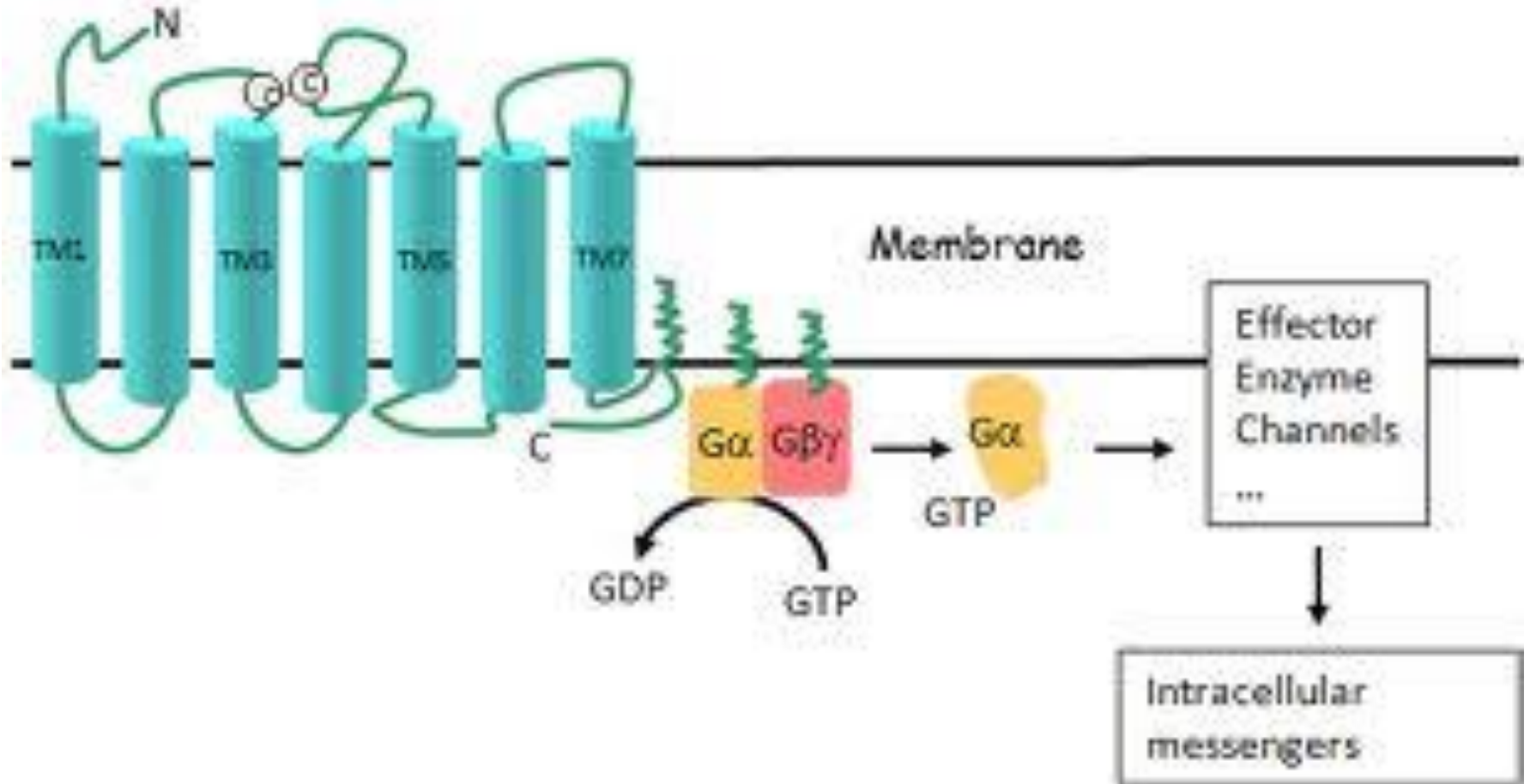
(a) Clathrin-mediated endocytosis

**Table 3.1 Transport of Materials Into and Out of Cells**

<b>Transport Process</b>	<b>Description</b>	<b>Substances Transported</b>
<b>Osmosis</b>	Movement of water molecules across a selectively permeable membrane from an area of higher water concentration to an area of lower water concentration.	Solvent: water in living systems.
<b>Diffusion</b>	Random mixing of molecules or ions due to their kinetic energy. A substance diffuses down a concentration gradient until it reaches equilibrium.	
<b>Diffusion through the lipid bilayer</b>	Passive diffusion of a substance through the lipid bilayer of the plasma membrane.	Nonpolar, hydrophobic solutes: oxygen, carbon dioxide, and nitrogen; fatty acids, steroids, and fat-soluble vitamins; glycerol, small alcohols; ammonia. Polar molecules: water and urea.
<b>Diffusion through membrane channels</b>	Passive diffusion of a substance down its electrochemical gradient through channels that span a lipid bilayer; some channels are gated.	Small inorganic solutes, mainly ions: $K^+$ , $Cl^-$ , $Na^+$ , and $Ca^{2+}$ . Water.
<b>Facilitated Diffusion</b>	Passive movement of a substance down its concentration gradient via transmembrane proteins that act as transporters; maximum diffusion rate is limited by number of available transporters.	Polar or charged solutes: glucose, fructose, galactose, and some vitamins.
<b>Active Transport</b>	Transport in which cell expends energy to move a substance across the membrane against its concentration gradient through transmembrane proteins that act as transporters; maximum transport rate is limited by number of available transporters.	Polar or charged solutes.
<b>Primary active transport</b>	Transport of a substance across the membrane against its concentration gradient by pumps; transmembrane proteins that use energy supplied by hydrolysis of ATP.	$Na^+$ , $K^+$ , $Ca^{2+}$ , $H^+$ , $I^-$ , $Cl^-$ , and other ions.
<b>Secondary active transport</b>	Coupled transport of two substances across the membrane using energy supplied by a $Na^+$ or $H^+$ concentration gradient maintained by primary active transport pumps. Antiporters move $Na^+$ (or $H^+$ ) and another substance in opposite directions across the membrane; symporters move $Na^+$ (or $H^+$ ) and another substance in the same direction across the membrane.	Antiport: $Ca^{2+}$ , $H^+$ out of cells. Symport: glucose, amino acids into cells.
<b>Transport In Vesicles</b>	Movement of substances into or out of a cell in vesicles that bud from the plasma membrane; requires energy supplied by ATP.	
<b>Endocytosis</b>	Movement of substances into a cell in vesicles.	
<b>Receptor-mediated endocytosis</b>	Ligand-receptor complexes trigger infolding of a clathrin-coated pit that forms a vesicle containing ligands.	Ligands: transferrin, low-density lipoproteins (LDLs), some vitamins, certain hormones, and antibodies.
<b>Phagocytosis</b>	"Cell eating"; movement of a solid particle into a cell after pseudopods engulf it to form a phagosome.	Bacteria, viruses, and aged or dead cells.
<b>Pinocytosis</b>	"Cell drinking"; movement of extracellular fluid into a cell by infolding of plasma membrane to form a pinocytic vesicle.	Solutes in extracellular fluid.
<b>Exocytosis</b>	Movement of substances out of a cell in secretory vesicles that fuse with the plasma membrane and release their contents into the extracellular fluid.	Neurotransmitters, hormones, and digestive enzymes.

# Control of Transport and activity of Enzymes

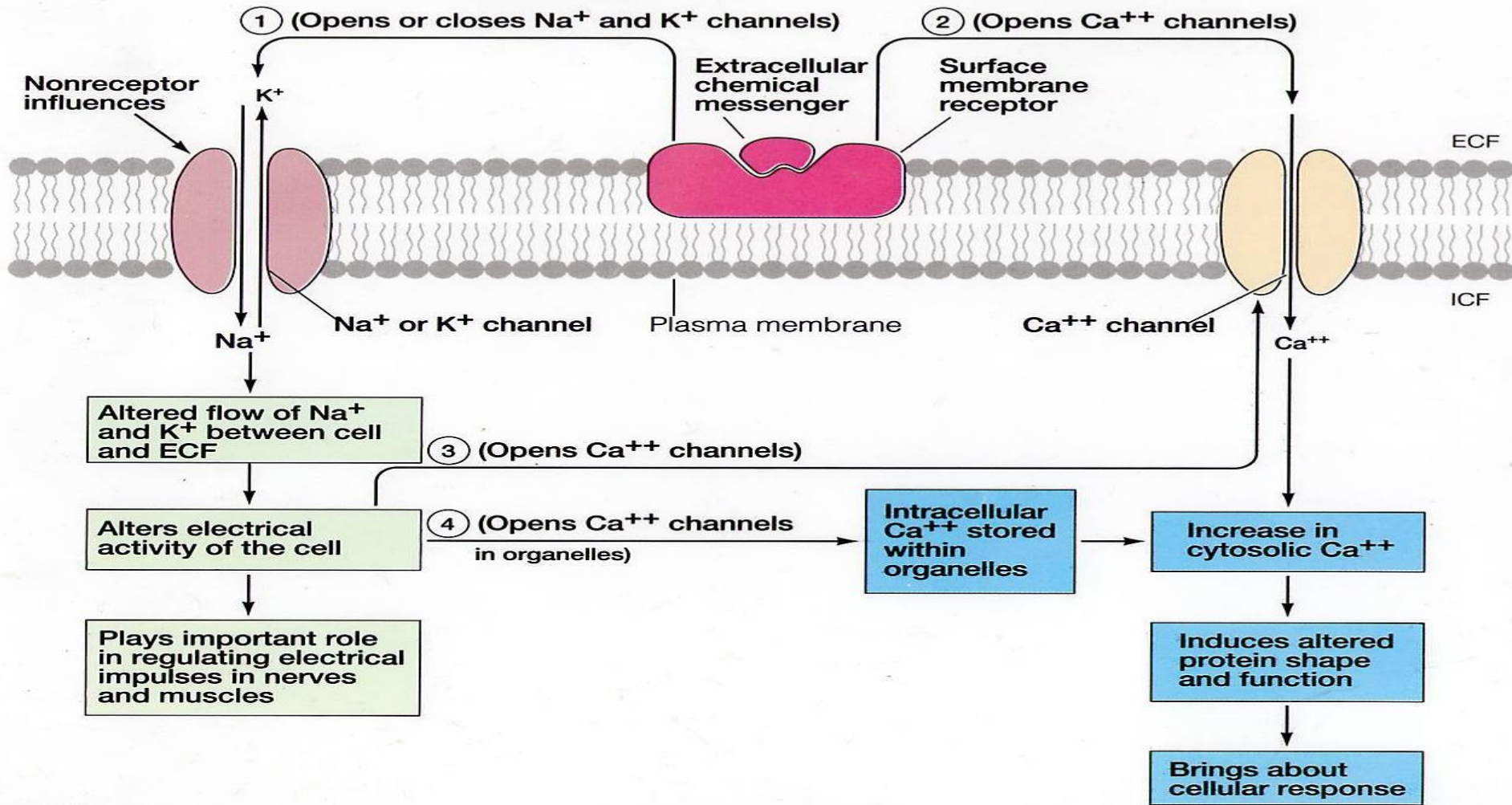
# Receptors & Enzymes



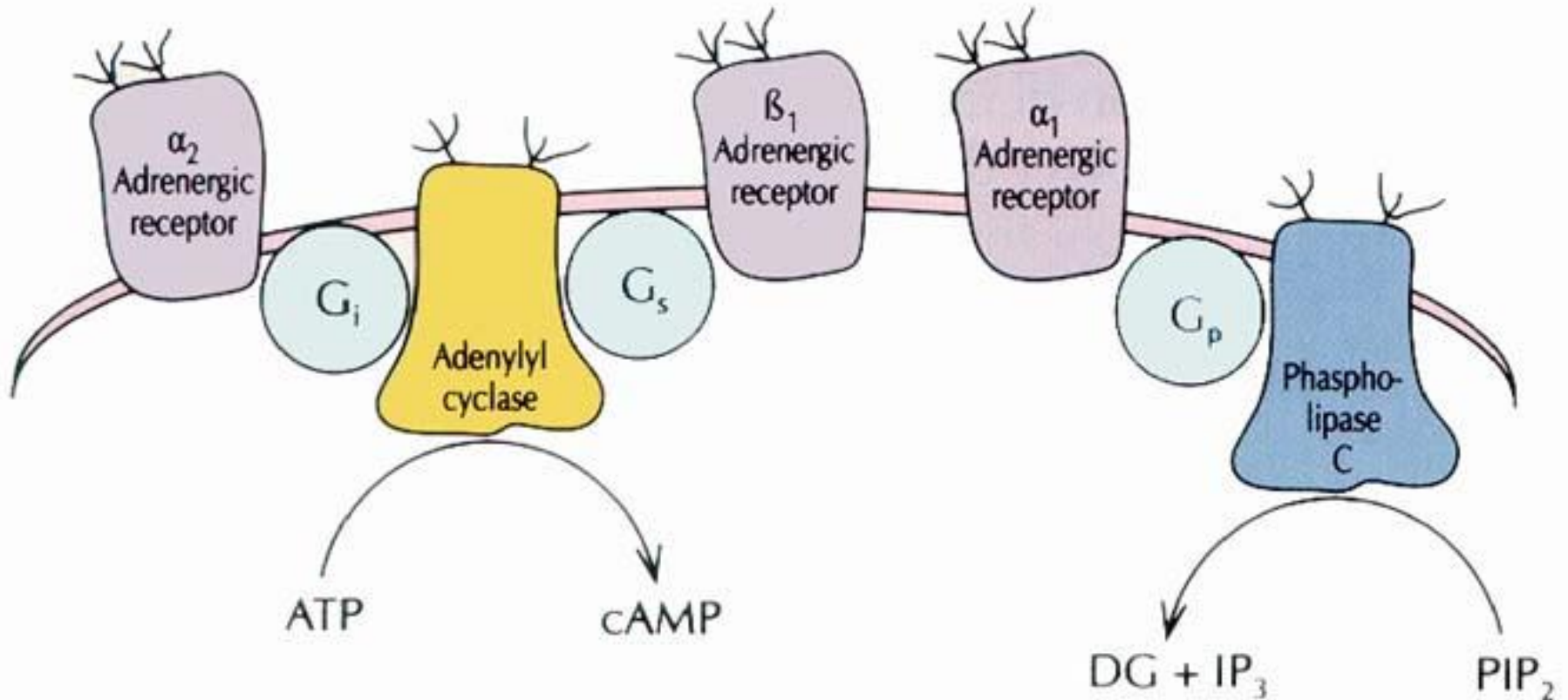


# Receptors & Channels

## Postreceptor Event: Channel Regulation

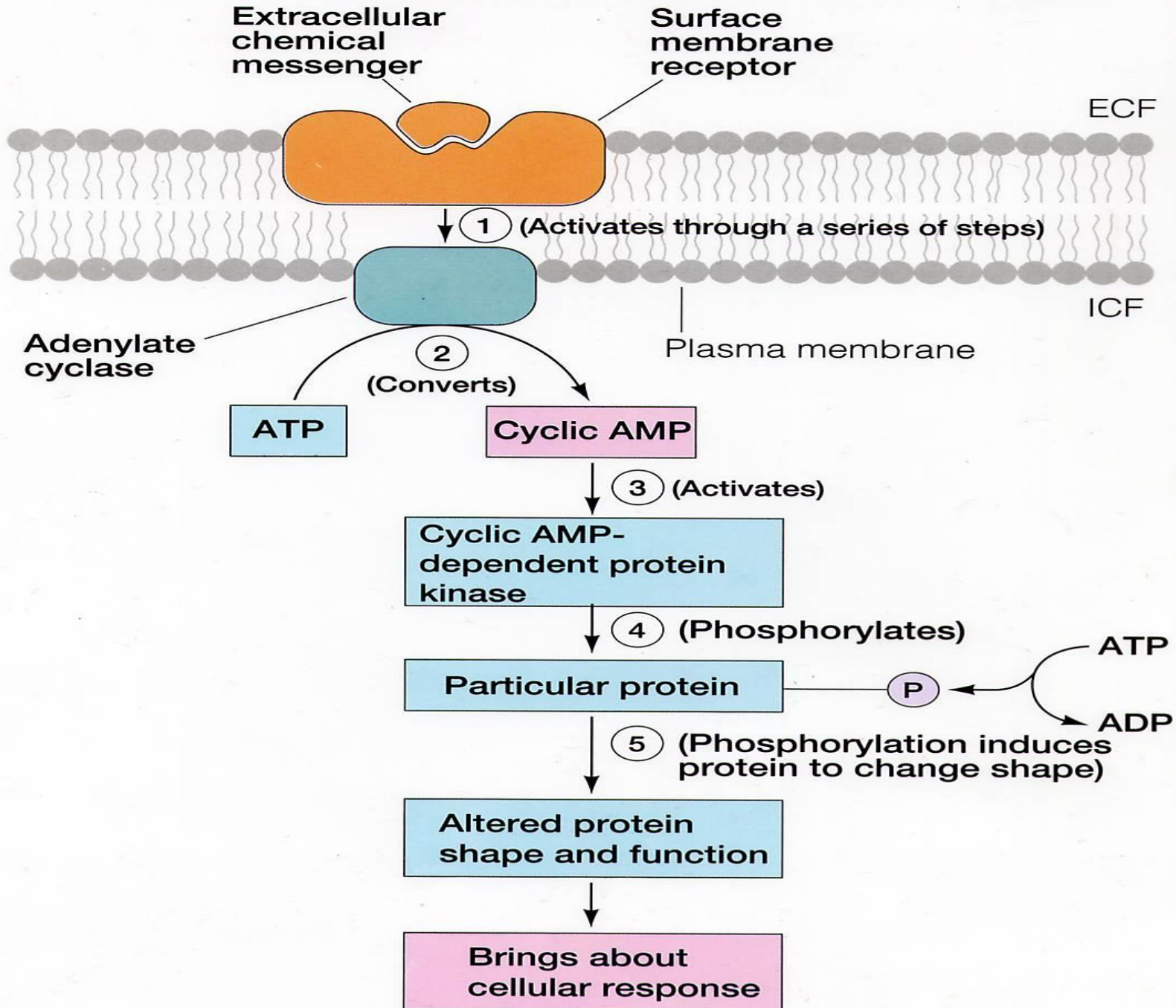


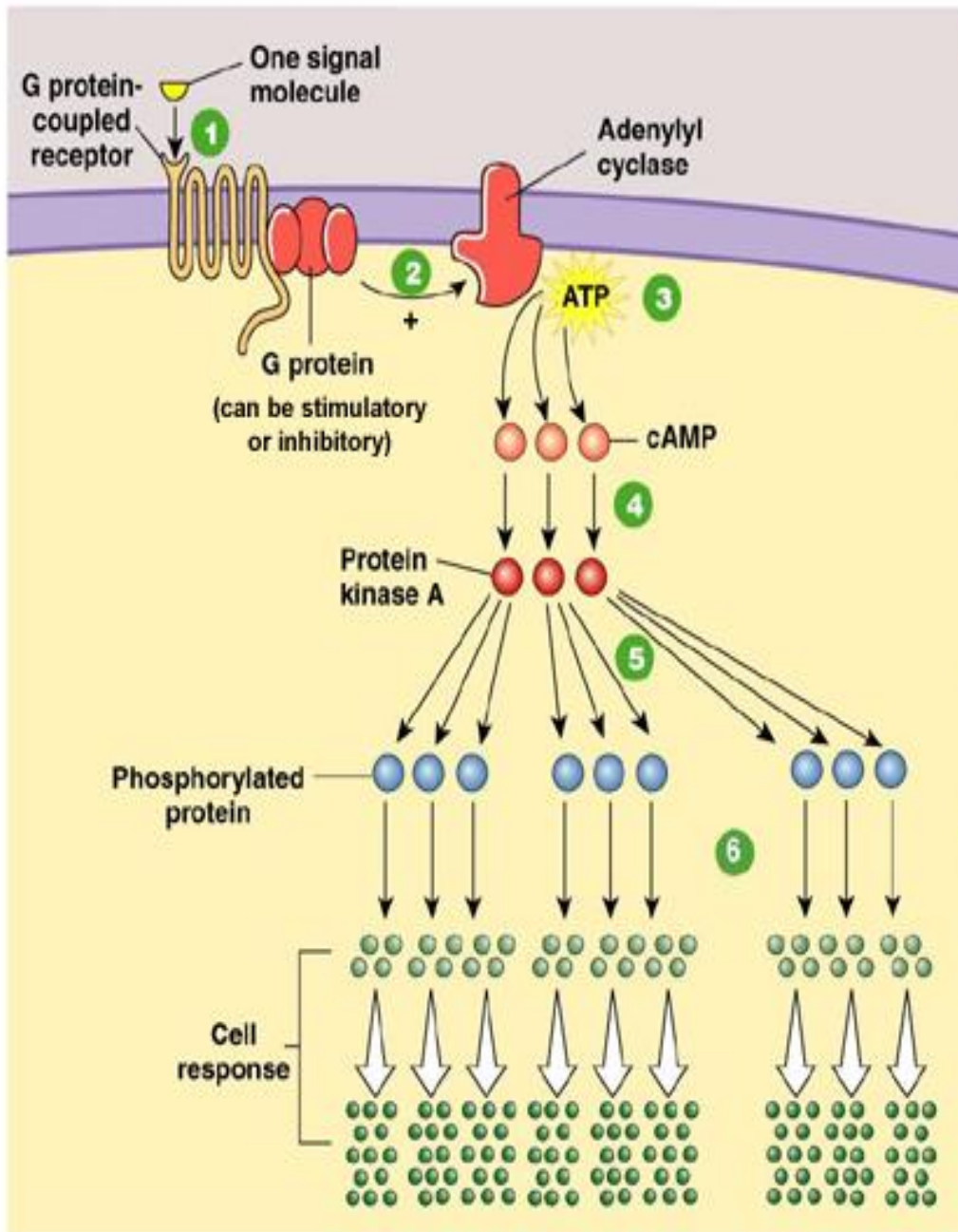
# Receptors & G proteins



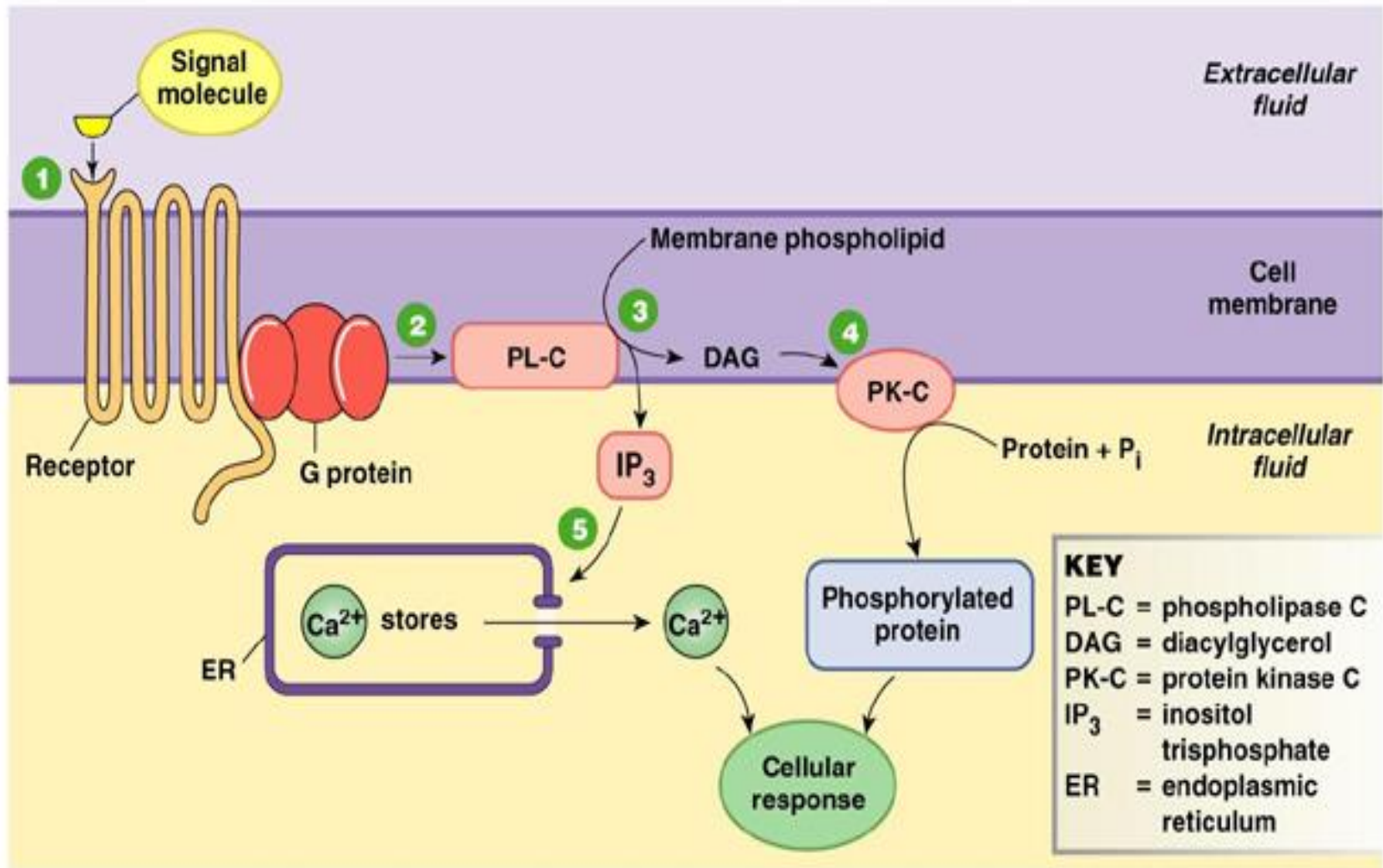


# Postreceptor Event: Cyclic AMP Second Messenger System





- 1** Signal molecule binds to G protein-linked receptor, which activates the G protein.
- 2** G protein turns on adenylyl cyclase, an amplifier enzyme.
- 3** Adenylyl cyclase converts ATP to cyclic AMP.
- 4** cAMP activates protein kinase A.
- 5** Protein kinase A phosphorylates other proteins, leading ultimately to a cellular response.
- 6** Note how the initial signal is amplified.

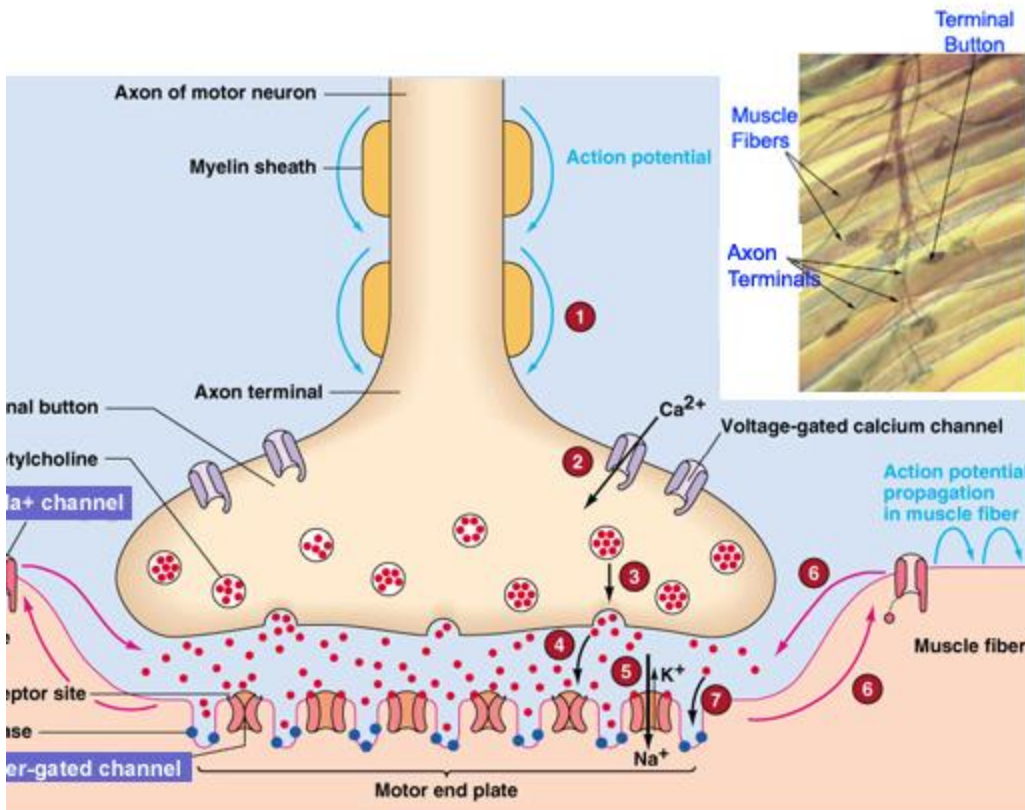


- 1** Signal molecule activates receptor and associated G protein.
- 2** G protein activates phospholipase C (PL-C), an amplifier enzyme.
- 3** PL-C converts membrane phospholipids into diacylglycerol (DAG), which remains in the membrane, and IP<sub>3</sub>, which diffuses into the cytoplasm.
- 4** DAG activates protein kinase C (PK-C), which phosphorylates proteins.
- 5** IP<sub>3</sub> causes release of Ca<sup>2+</sup> from organelles, creating a Ca<sup>2+</sup> signal.



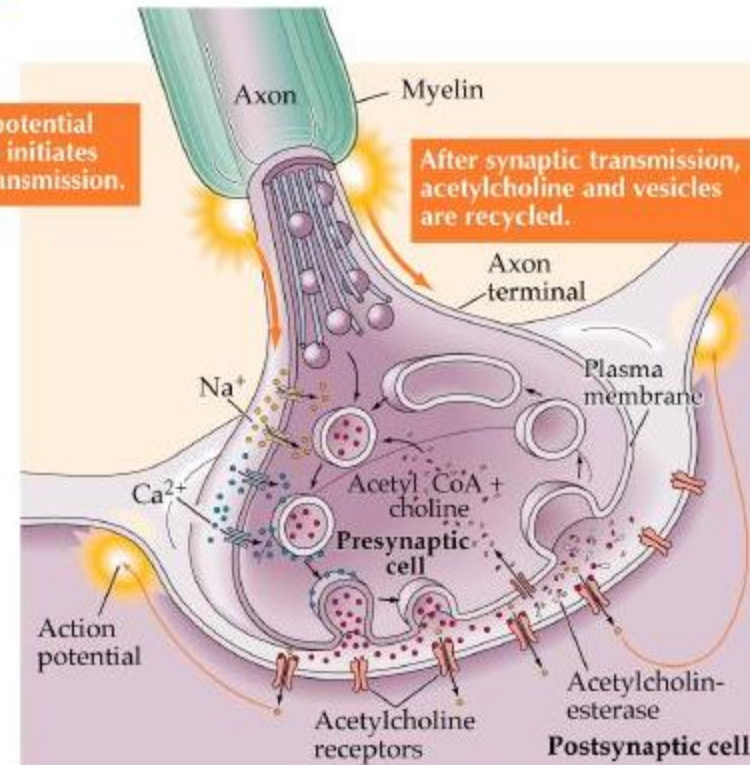
# Control of Exocytosis

## The Neuromuscular Junction



## Terminal

An action potential arrives and initiates synaptic transmission.



# Summary of Lectures

**-Homeostasis**

**-Control of body functions** by feed-back mechanisms to keep homeostasis

**-Functions of Cell organells** (ER, Golgi complex, Mitochondria, Lisosomes)

**-Cytoskeletal structures** and Functions:

# Plasma membrane

- **Membrane structure and function of lipids of plasma membrane** including phospholipids, cholesterol and PIP2)
- **Functions of Protein structures of plasma membranes** (Channels, Carriers, Receptors, Pumps, enzymes, cell markers, G proteins, adhesion molecules)
- **Activation of chemical gated channels**

# Transport through plasma membranes

## Passive transport modalities

**-Simple diffusion:** transport through lipid bilayer, transport through channels, Ficks law of diffusion.

**-Facilitated diffusion:** by carriers

Differences in diffusion Kinetics between the previous modalities

Equivalent Concentration of particles.

# Transport through plasma membranes

## Passive transport modalities

-**Osmosis**: concept of osmotic pressure (Van't Hoff's law), Oncotic (Colloid-osmotic) pressure. Osmolarity, osmolality  
Hydrostatic pressure



# Transport through plasma membranes

## Active transport modalities

- **Primary active transport** (ATP-ase carriers or Pumps) (functions of pumps:  $\text{Na}^+/\text{K}^+$  pump,  $\text{Ca}^{++}$  pump,  $\text{H}^+$  pump,  $\text{H}^+/\text{K}^+$  pump).
- **Secondary active transport** ( $\text{Na}^+$  dependent carriers) examples

# Transport through plasma membranes

## Active transport modalities

- **Vesicular transport:** endocytosis, phagocytosis, transcytosis, pinocytosis and exocytosis and its control in secretory cells.-