

AP Biology Unit 2

The Cell

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EXAM WEIGHT

10-13%

MCQs

6-8

FRQ APPEARANCE

Very Frequent

SPRINT TIME

~2 hours

Inside this pack

- **Quick Glance at every Unit topic with priority + format**
- **Topic-by-topic key traps, must-know rules, and exam frames**
- **Worked example questions on the highest-yield topics**

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AP Biology Unit 2 — The Cell

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Quick Glance — All Topics at a Glance

Topic	Priority	Exam Format	Key Trap
2.1 Cell Structure	★★★	MCQ, FRQ	Prokaryotes DO have ribosomes (70S); nucleolus is INSIDE nucleus
2.2 Cell Size / SA:V	★★★	MCQ, Calc	Larger cell = LOWER SA:V = LESS efficient (not more)
2.3 Plasma Membrane	★★★	MCQ, FRQ	Cholesterol is a BUFFER — increases fluidity in cold, decreases in heat
2.4 Permeability	★★	MCQ	Water crosses slowly (via aquaporins); ions CANNOT cross without channels
2.5–2.6 Passive Transport	★★★	MCQ, FRQ	Facilitated diffusion uses proteins but needs NO ATP — still passive!
2.7 Tonicity / Osmosis	★★★	MCQ, Calc, Data	Plant cells: hypo → turgid (NOT lysis); animal cells: hypo → lysis
2.8 Active Transport	★★★	MCQ, FRQ	Na ⁺ /K ⁺ pump: 3 Na ⁺ OUT, 2 K ⁺ IN — net charge moves out → electrochemical gradient
2.9–2.10 Compartments + Endosymbiosis	★★★	MCQ, FRQ	Must cite 3+ pieces of evidence for endosymbiosis; mitochondria in ALL eukaryotes

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2.1 Cell Structure & Function

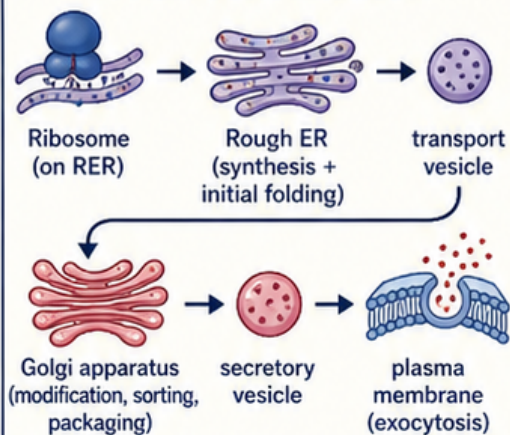
Prokaryotes, eukaryotes, organelles, and the secretory pathway

Prokaryote vs. Eukaryote — Must-Know Comparison

Feature	Prokaryote	Eukaryote
Nucleus	None — DNA in nucleoid region (no membrane)	Membrane-bound nucleus (double membrane)
DNA shape	Single circular chromosome + plasmids	Multiple linear chromosomes + histones
Membrane-bound organelles	None	ER, Golgi, mitochondria, etc.
Ribosomes	70S (smaller) — ALL cells have ribosomes!	80S cytoplasm; 70S in mitochondria/chloroplasts
Cell wall	Peptidoglycan (bacteria)	Cellulose (plant); chitin (fungi); none (animal)
Size	1–10 μm	10–100 μm
Examples	Bacteria, Archaea	Animals, Plants, Fungi, Protists

Secretory Pathway — High Frequency

Protein Secretion Route



- Ribosome (on RER) → Rough ER (synthesis + initial folding) → transport vesicle → Golgi apparatus (modification, sorting, packaging) → secretory vesicle → plasma membrane (exocytosis)
- Golgi is the “post office” — receives, modifies, and ships proteins
- Lysosomes bud from Golgi → contain hydrolytic enzymes for intracellular digestion

Energy Organelles

Mitochondria & Chloroplasts



- **Mitochondria:** double membrane; inner membrane highly folded (cristae = increased surface area for ATP synthesis); site of cellular respiration; matrix contains enzymes for Krebs cycle

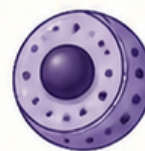


- **Chloroplasts:** double membrane + thylakoid membranes stacked into grana; site of photosynthesis; stroma = liquid between thylakoids

- Both have 70S ribosomes + circular DNA → endosymbiotic evidence (2.10)

Quick Reference

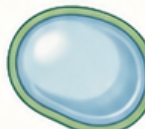
Other Key Organelles



- **Nucleus:** site of transcription; nuclear pores control traffic; nucleolus = rRNA synthesis (inside nucleus, NOT separate organelle)



- **Smooth ER:** lipid synthesis, detoxification (liver), calcium storage (muscle)



- **Vacuole:** central vacuole in plant cells → turgor pressure; contractile vacuole in protists (osmoregulation)



- **Cytoskeleton:** microfilaments (actin, cell shape/movement), microtubules (cilia/flagella, spindle fibers), intermediate filaments (structural)



Exam Sniper

- Classic FRQ: trace a secreted protein from synthesis to exit → Ribosome on RER → RER lumen → transport vesicle → Golgi → secretory vesicle → exocytosis
- A drug inhibits Golgi function → secretion of modified proteins is most affected
- Evidence for mitochondrial prokaryotic ancestry → double membrane, 70S ribosomes, circular DNA, binary fission
- Cells producing lots of lipids have abundant smooth ER



Trap Alert

- Prokaryotes **DO** have ribosomes — 70S, not 80S
- The nucleolus is **NOT** a separate organelle — it is inside the nucleus
- Archaea are prokaryotes but are more closely related to eukaryotes than to bacteria



2.2 Cell Size & Surface Area:Volume Ratio

Why cells stay small and how to calculate SA:V



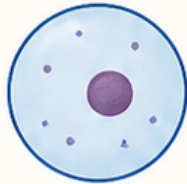
The Core Idea

Why SA:V Matters

- Surface area (membrane) = interface for exchanging nutrients, O₂, CO₂, waste
- Volume = metabolic demands of the interior
- As cell grows, volume increases faster than surface area
- Large cell = low SA:V = insufficient exchange = cell division triggered
- Small cells are **MORE** efficient at exchange



Small cell
High SA:V
Efficient



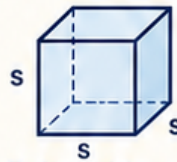
Large cell
Low SA:V
Inefficient



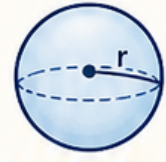
Formulas — Memorize

SA:V Calculations

- Cube (side = s): SA = 6s²; V = s³; SA:V = 6/s
- Sphere (radius = r): SA = 4πr²; V = 4/3 πr³
- Rule: Double the radius → SA ×4, V ×8 → SA:V halves
- Always show units; SA:V has units of 1/length (e.g., μm⁻¹)



Cube (side = s)
SA = 6s²
V = s³
SA:V = 6/s

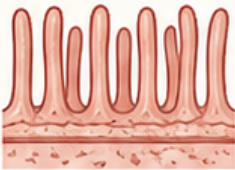


Sphere (radius = r)
SA = 4πr²
V = 4/3 πr³



Biological Applications

High SA:V Adaptations



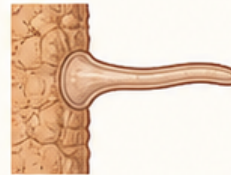
Microvilli in intestine → increased SA for nutrient absorption



Alveoli in lungs → small spheres, high SA for gas exchange



Cristae in mitochondria → increased inner membrane SA for ATP synthesis



Root hair cells → elongated projections increase SA for water/ion uptake



Red blood cells → flattened biconcave disc → maximizes SA relative to V



Exam Sniper

- Given two cells, calculate both SA:V values → smaller cell has higher SA:V and exchanges materials more efficiently
- Agar cube diffusion labs: smaller cubes equilibrate faster because of higher SA:V
- Cells divide because volume grows faster than surface area, so SA:V falls below what is needed for the interior



Calculation MCQ

Two cuboidal cells are compared: Cell A has sides of 1 μm and Cell B has sides of 3 μm. Which cell is more efficient at exchanging materials with its environment, and what is the SA:V ratio for each?

- (A) Cell B is more efficient; Cell A SA:V = 2, Cell B SA:V = 6
 (B) Cell A is more efficient; Cell A SA:V = 6, Cell B SA:V = 2
 (C) Both are equally efficient; SA:V = 1 for both
 (D) Cell B is more efficient; larger cells have more surface area



Answer: (B) — Cell A: SA = 6, V = 1, SA:V = 6.

Cell B: SA = 54, V = 27, SA:V = 2.

The smaller cell has proportionally more membrane per unit volume, so exchange is more efficient.



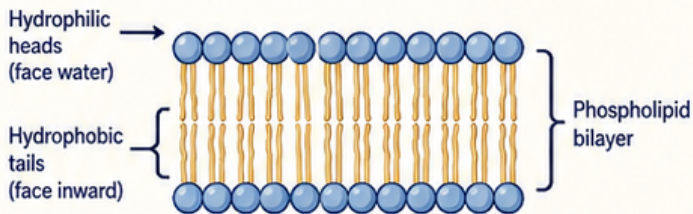
2.3–2.4 Plasma Membrane & Permeability

Fluid mosaic model, membrane proteins, and what can cross

Model Name

Fluid Mosaic Model

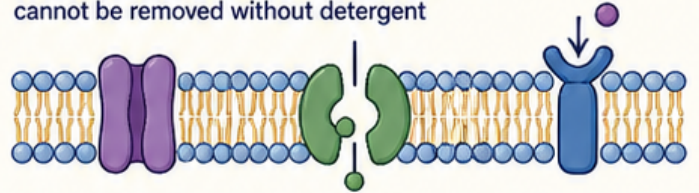
- “Fluid”: phospholipids and proteins can move laterally within each layer — not rigid
- “Mosaic”: proteins of many types embedded throughout
- Proposed by Singer & Nicolson, 1972
- Selectively permeable: controls what enters/exits
- Phospholipid bilayer: hydrophilic heads face water; hydrophobic tails face inward



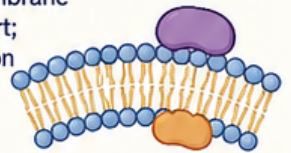
Components

Membrane Proteins

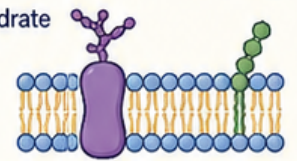
- **Integral proteins:** span the bilayer; channel proteins, carrier proteins, receptor proteins — cannot be removed without detergent



- **Peripheral proteins:** attached to membrane surface; signaling and structural support; removable by changing salt concentration



- **Glycoproteins/glycolipids:** carbohydrate chains on OUTER face only; cell-cell recognition, immune response, cell adhesion

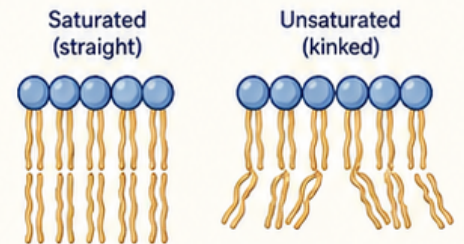


Fluidity Control — High Exam Yield

Factors Affecting Fluidity



- Higher temperature → higher kinetic energy → higher fluidity
- More unsaturated fatty acids (kinked tails) → cannot pack tightly → higher fluidity
- Cholesterol = fluidity buffer: at high temp restrains movement → lowers fluidity; at low temp prevents freezing → raises fluidity
- Shorter fatty acid chains → less van der Waals force → higher fluidity
- Cold-water organisms: more unsaturated lipids to keep membranes fluid



Membrane Permeability

Molecule Type	Example	Crosses Freely?	Why?
Small nonpolar	O ₂ , CO ₂ , N ₂ , steroids	Yes — simple diffusion	Dissolve in hydrophobic core; small enough to pass
Small uncharged polar	H ₂ O, urea, ethanol	Slowly (aquaporins speed up water)	Small helps, but polarity slows crossing
Ions (any charge)	Na ⁺ , K ⁺ , Cl ⁻ , Ca ²⁺	No — need ion channels	Hydration shells too large to enter hydrophobic core
Large polar molecules	Glucose, amino acids	No — need carrier proteins	Too large and polar to diffuse through lipid layer
Macromolecules	Proteins, RNA, polysaccharides	No — need vesicle transport	Far too large; require endo/exocytosis

Exam Sniper

- Fish moved from warm to cold water — increase unsaturated fatty acids to maintain membrane fluidity
- At body temperature, cholesterol stabilizes the membrane by reducing fluidity
- Glycoproteins are on the outer face because they were glycosylated in the ER/Golgi lumen and remain facing outward after vesicle fusion
- Glucose cannot cross by simple diffusion because it is both large and polar
- Aquaporins show that water can cross slowly without them, but much faster with them

Trap Alert

- Cholesterol does NOT simply increase or decrease fluidity — it is a buffer, and the answer depends on temperature
- Glycoproteins are on the outer face only
- Phospholipids move laterally easily, but flip-flop between leaflets is rare



2.5–2.6 Passive Transport

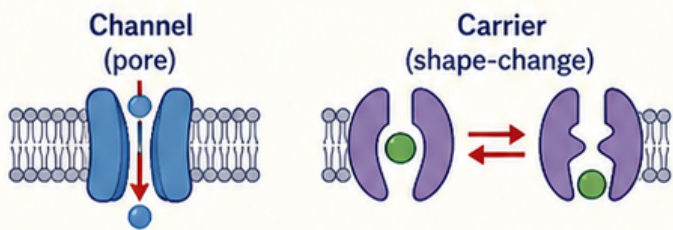
Simple diffusion, osmosis, and facilitated diffusion

All Passive Transport = No ATP, Down Gradient

Type	What Moves	Protein Required?	ATP Required?	Direction
Simple Diffusion 	Small nonpolar gases (O ₂ , CO ₂), steroids	No	No	High → Low concentration
Osmosis 	Water (H ₂ O)	Via aquaporins (speeds up)	No	Low solute → High solute (High Ψ → Low Ψ)
Facilitated Diffusion 	Ions, glucose, amino acids	Yes (channel or carrier)	No	High → Low concentration

Facilitated Diffusion – Key Details

Channel vs. Carrier Proteins



- **Channel proteins:** form a pore; allow ions to flow through; can be gated by voltage, ligand, or mechanical signals
- **Carrier proteins:** bind a specific molecule, change shape, release it on the other side; slower; shows saturation
- Both are specific, require no ATP, and move substances down the gradient
- **Saturation:** when all carriers are occupied, rate maxes out — unlike simple diffusion

Diffusion Rate Factors

What Affects Diffusion Speed

- Higher concentration gradient → higher diffusion rate
- Higher temperature → higher kinetic energy → higher rate
- Higher surface area → higher rate
- Shorter distance → higher rate (Fick's Law: rate \propto SA \times Δ C / distance)
- More fluid membrane → faster diffusion
- Smaller molecule size → faster crossing



Exam Sniper

- Carrier protein + no ATP + down gradient = facilitated diffusion
- A rate curve that plateaus suggests carrier-mediated transport (saturation), not simple diffusion
- Alveoli are efficient for gas exchange because they have high surface area, short diffusion distance, and thin membranes



Trap Alert

- Uses a protein does NOT automatically mean active transport
- Facilitated diffusion requires NO ATP
- Simple diffusion has no maximum rate; carrier-mediated transport does



Question:

Glucose enters a cell using a carrier protein, and no ATP is used. What type of transport is this?

Answer:







Facilitated diffusion — because glucose moves through a carrier protein down its concentration gradient with no ATP required.



2.7 Tonicity, Osmosis & Water Potential

The highest-frequency calculation and reasoning topic

Osmosis = diffusion of water across a selectively permeable membrane, from low solute → high solute (equivalently: from high water potential → low water potential).

Hypotonic	Isotonic	Hypertonic
 <ul style="list-style-type: none"> Water moves INTO cell Animal cell → swells → LYSES 	 <ul style="list-style-type: none"> No NET water movement Animal cell → normal 	 <ul style="list-style-type: none"> Water moves OUT of cell Animal cell → crenates (shrivels)
 <ul style="list-style-type: none"> Plant cell → TURGID (normal healthy state) RBC in distilled water → lysis 	 <ul style="list-style-type: none"> Plant cell → flaccid 0.9% NaCl is isotonic to human blood 	 <ul style="list-style-type: none"> Plant cell → plasmolysis (membrane pulls away from wall) Examples: saltwater fish, salted meat preservation



Ψ_s Calculation

Solute Potential Formula

$$\Psi_s = -iCRT$$

- i = ionization constant (1 for glucose/sucrose; 2 for NaCl; 3 for CaCl_2)
- C = molar concentration (mol/L)
- R = 0.0831 L·bar/mol·K
- T = temperature in Kelvin ($^{\circ}\text{C} + 273$)
- Result is always negative — solutes lower Ψ



Plant Cell Water Potential

Turgor Pressure in Plants

- $\Psi_p = 0$ in open systems (like a beaker); positive in turgid plant cells
- At equilibrium: Ψ of cell = Ψ of solution
- Turgor pressure prevents wilting; drives stomatal opening/closing
- Guard cells: gain water → become turgid → stoma opens
- Wilted plant: low turgor pressure → $\Psi_p \approx 0$ → gains water when watered



Exam Sniper

- Given concentrations and temperature, calculate Ψ_s and determine water movement
- If a plant cell is -4 bar and the solution is -2 bar, water moves from -2 to -4 → into the cell
- In mass-vs-sucrose graphs, the isotonic point is where mass change is zero
- Plant cells in hypotonic solution become turgid, **NOT** lysed



Trap Alert

- Water moves from low solute to high solute
- More negative Ψ = lower Ψ ; water moves there
- Isotonic means no NET movement, not no movement at all
- Plant cells in hypotonic solution become turgid, not lysed



A student places red blood cells and plant cells into three solutions: distilled water, 0.9% NaCl, and 3% NaCl. In which solution would the plant cell become turgid AND the red blood cell lyse?

- (A) 3% NaCl (C) 0.9% NaCl
- (B) Distilled water (D) Both 3% NaCl and distilled water



Answer:

(B) — Distilled water is hypotonic to both cells. Water enters both; the plant cell becomes turgid because its wall prevents lysis, while the red blood cell bursts.



2.8 Active Transport, Bulk Transport & Co-transport

Against the gradient, ATP, pumps, vesicles, and indirect transport



Definition

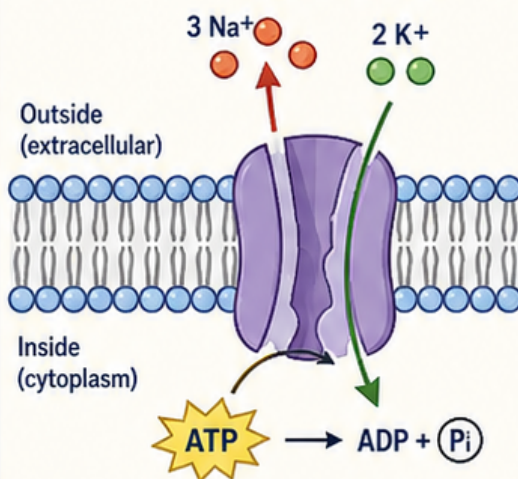
Active Transport

- Moves substances against concentration gradient (low → high)
- Requires ATP
- Uses pump proteins
- Maintains gradients needed for nerve impulses, nutrient uptake, and pH regulation



Most-Tested Pump

Na⁺/K⁺-ATPase Pump



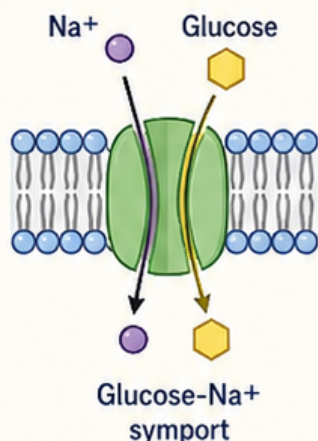
- Pumps 3 Na⁺ OUT and 2 K⁺ IN per ATP hydrolyzed
- Both move against their gradients
- Net positive charge moves out → inside becomes more negative → resting membrane potential (~-70 mV)
- Critical for nerve impulses, muscle contraction, and secondary active transport
- Memory: **NaKe 3-2 rule**



Indirect Active Transport

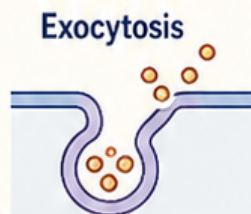
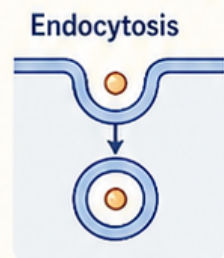
Co-transport (Secondary Active)

- Uses an existing ion gradient (usually Na⁺) to drive another molecule
- Example: glucose-Na⁺ symport in intestinal epithelium
- The Na⁺ gradient was created by the Na⁺/K⁺ pump, so ATP is used indirectly
- Symport = same direction; Antiport = opposite directions



Bulk Transport

Endo & Exocytosis



- Endocytosis: cell engulfs material using membrane vesicles; requires ATP
- Phagocytosis = solids; pinocytosis = fluids/small molecules; receptor-mediated endocytosis = specific ligands
- Exocytosis: vesicles fuse with plasma membrane and release contents outside
- Both require ATP and involve membrane fusion



Exam Sniper

- Blocking ATP production most directly affects active transport, co-transport indirectly, and endocytosis/exocytosis; passive transport continues
- The Na⁺/K⁺ pump creates resting membrane potential by exporting 3 Na⁺ and importing 2 K⁺
- Intestinal glucose uptake via Na⁺/glucose co-transporter is secondary active transport
- Vesicle required = endocytosis or exocytosis



Trap Alert

- Na⁺/K⁺ pump = 3 Na⁺ OUT, 2 K⁺ IN — never reverse it
- Co-transport is not passive transport; it uses an ion gradient created by ATP
- Phagocytosis and pinocytosis are both types of endocytosis



2.9–2.10 Compartmentalization & Endosymbiotic Theory

Why membrane compartments matter and how mitochondria/chloroplasts evolved



Why Compartmentalize?

Advantages of Membrane Compartments

- Allows simultaneously incompatible reactions to occur
- Concentrates reactants → higher reaction efficiency
- Creates H⁺ gradients for ATP synthesis in mitochondria and chloroplasts
- Isolates digestive enzymes in lysosomes to prevent self-digestion
- Nucleus separates transcription from translation, allowing post-transcriptional processing



FRQ Must-Know

Endosymbiotic Theory – Evidence

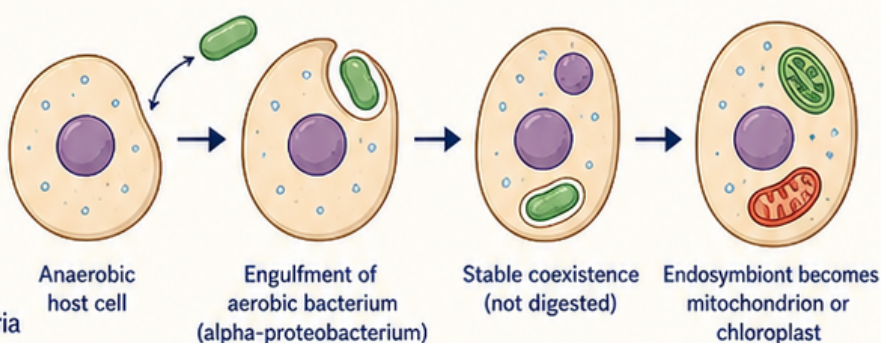
- ★ Double membrane
- ★ 70S ribosomes
- ★ Circular DNA
- ★ Reproduce by binary fission
- ★ Size similar to bacteria
- ★ Antibiotic sensitivity: some prokaryote-targeting antibiotics also affect mitochondria



Theory Details

Who Was Engulfed?

- Proposed by Lynn Margulis
- Mitochondria descended from aerobic alpha-proteobacteria; found in ALL eukaryotes
- Chloroplasts descended from cyanobacteria; found only in photosynthetic eukaryotes
- Host cell was an anaerobic archaean-like cell that phagocytosed but did not digest these bacteria
- Over time, most bacterial genes transferred to the host nucleus



Exam Sniper

- To support endosymbiosis, list multiple independent pieces of evidence: 70S ribosomes, circular DNA, double membrane, binary fission
- An antibiotic that targets 70S ribosomes may also affect mitochondria and chloroplasts
- Compartmentalization helps cells run incompatible reactions and create specialized microenvironments
- Plant cells have chloroplasts, central vacuoles, and cell walls; animals do not



Common Confusions

- Mitochondria are in ALL eukaryotes, not just plants
- Chloroplasts are only in photosynthetic eukaryotes
- The evidence must show prokaryotic ancestry, not just organelle function



Sprint Practice — Mixed Questions

Cross-topic AP-style reasoning

1



MCQ Cross-Topic: Transport

A researcher treats intestinal epithelial cells with ouabain, a drug that specifically inhibits the Na^+/K^+ -ATPase pump. Which of the following would be the MOST DIRECT effect on glucose absorption in these cells?

- (A) Glucose diffusion would increase because the concentration gradient would steepen
- (B) Glucose uptake via the $\text{Na}^+/\text{glucose}$ co-transporter would decrease because the Na^+ gradient would dissipate
- (C) Glucose would be exported from the cell by exocytosis instead
- (D) Passive diffusion of glucose would compensate for the blocked pump



Answer:

(B) — The $\text{Na}^+/\text{glucose}$ co-transporter depends on the Na^+ gradient maintained by the Na^+/K^+ pump. Block the pump → Na^+ builds up inside → the gradient collapses → co-transport decreases → glucose absorption falls.

2



Data Analysis MCQ: Tonicity

A student places identical pieces of potato tissue (each 5 grams) into sucrose solutions of different concentrations and records mass after 2 hours. Results: 0.0 M → 5.6 g; 0.2 M → 5.2 g; 0.4 M → 4.9 g; 0.6 M → 4.8 g; 0.8 M → 4.5 g. What is the approximate solute concentration of the potato cells?

- (A) 0.0 M
- (B) 0.8 M
- (C) Approximately 0.3–0.4 M
- (D) 0.6 M



Answer:

(C) — The isotonic point is where mass change is closest to zero. The sample gains mass at 0.2 M and loses a little at 0.4 M, so the internal concentration is about 0.3–0.4 M.

3



FRQ-Style Endosymbiosis

A scientist discovers a new eukaryotic organism with organelles resembling mitochondria. She claims these organelles evolved via endosymbiosis. Describe THREE independent pieces of evidence she could present to support this claim.

- (A) 80S ribosomes, double membranes, and ATP production
- (B) 70S ribosomes, circular DNA without histones, and reproduction by binary fission
- (C) A single membrane, ATP synthase, and maternal inheritance
- (D) Presence in all eukaryotes, oxygen use, and cristae



Answer:

(B) — The strongest evidence is 70S ribosomes, circular DNA without histones, and binary fission. These are prokaryotic features and support endosymbiotic ancestry.



Tip: For Unit 2 questions, always identify the gradient, the transport mechanism, and whether ATP is used.



Final Review — Exam Traps + Checklist

Use this page the night before and the morning of the exam

1. Unit 2 High-Frequency Exam Traps

- Facilitated diffusion uses proteins but NO ATP
- Plant cells in hypotonic solution become TURGID, not lysed
- Water moves from low solute to high solute; from high Ψ to low Ψ
- Larger cells have LOWER SA:V and are LESS efficient
- Prokaryotes DO have ribosomes — 70S
- Cholesterol is a fluidity BUFFER — answer depends on temperature
- Na⁺/K⁺ pump = 3 Na⁺ OUT, 2 K⁺ IN
- The nucleolus is inside the nucleus

2. Pre-Exam 10-Minute Checklist

A. Cell Structure (2.1)

- Prokaryotes: no nucleus, circular DNA, 70S ribosomes, peptidoglycan wall
- Secretory pathway: ribosome on RER → RER → vesicle → Golgi → vesicle → plasma membrane/lysosome
- Nucleolus is inside the nucleus
- Smooth ER = lipids/detox; Rough ER = proteins

B. Cell Size & Membrane (2.2–2.4)

- SA:V for cube = 6/s; larger cell = lower SA:V
- Can calculate SA:V for cube and sphere
- Fluid mosaic model = fluid + mosaic
- Cholesterol reduces fluidity in heat, prevents rigidity in cold
- Glycoproteins are on the outer face only
- More unsaturated fatty acids = more fluid membrane

C. Transport (2.5–2.8)

- Simple diffusion: small nonpolar molecules
- Facilitated diffusion: ions + glucose + amino acids; no ATP
- Osmosis: water from low solute to high solute
- Tonicity outcomes: plant turgid/flaccid/plasmolyzed; animal lyse/normal/crenate
- Water potential: $\Psi = \Psi_s + \Psi_p$; pure water $\Psi = 0$
- Na⁺/K⁺ pump = 3 out / 2 in
- Co-transport uses Na⁺ gradient; no direct ATP

D. Compartmentalization & Endosymbiosis (2.9–2.10)

- Compartments allow incompatible reactions and create proton gradients
- Endosymbiotic evidence: 70S ribosomes, circular DNA, double membrane, binary fission
- Mitochondria in all eukaryotes; chloroplasts only in photosynthetic eukaryotes

3. Final Sprint Strategy for Unit 2

- Biggest point-earners: tonicity outcomes, water potential calculations, SA:V calculations, Na⁺/K⁺ pump, endosymbiosis evidence
- FRQ danger zone: always state direction of movement, mechanism, ATP use, and protein involved if any
- Data questions: isotonic point = where mass does not change
- Connections forward: membrane fluidity → Unit 1; chloroplasts/mitochondria → Unit 3; Na⁺ gradients → Unit 4; endosymbiosis → Unit 7



If you can explain every checked item out loud, you're ready.

Done with the sheet? Now get marked.

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See

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