



AP[®] Biology 2007 Scoring Guidelines

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Question 1

Membranes are essential components of all cells.

- (a) **Identify** THREE macromolecules that are components of the plasma membrane in a eukaryotic cell and **discuss** the structure and function of each. **(6 points maximum; 1 point for each macromolecule + structure, 1 point for each macromolecule + function)**

NOTE: Only first three molecules mentioned will be scored.

Macromolecule	Structure	Function (must match selected macromolecule)
Phospholipids OR Lipid with phosphate	<ul style="list-style-type: none"> • Glycerol, two fatty acids, and polar head group w/phosphate • Amphipathic • Hydrophilic or polar (head) and hydrophobic or nonpolar (tails) • Forms a lipid bilayer 	<ul style="list-style-type: none"> • Selectively permeable • Fluidity • Creates compartment/ separates cell from environment; barrier • Signals, inositol pathway (IP3) diacylglycerol (DAG)
Cholesterol	<ul style="list-style-type: none"> • Ring structure • Steroid • Amphipathic • Embedded in bilayer 	<ul style="list-style-type: none"> • Moderates fluidity • Stabilizes membrane
Proteins OR <u>The following specific types must indicate that they are proteins</u> Integral Peripheral Pump Receptor Transport Recognition Tight junction Desmosomes Gap junctions Integrins Enzyme Channel	<p style="text-align: center;"><u>General Structure</u></p> <ul style="list-style-type: none"> • Polypeptides; amino acids • 2°, 3°, 4° structure description <p style="text-align: center;"><u>Specific Structure</u></p> <ul style="list-style-type: none"> • Integral, transmembrane, embedded; forms a channel • Peripheral, on surface • Structure fit to substrate or ligand 	<ul style="list-style-type: none"> • Transport • Enzyme, catalysis • Signal transduction • Attachment: extracellular matrix (ECM)-cytoskeleton • Recognition • Cell junction
Glycolipid/Glycoprotein	<ul style="list-style-type: none"> • Carbohydrate (chains) linked to lipid/protein 	<ul style="list-style-type: none"> • Cell recognition • Attachment to external molecule or another cell

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Question 1 (continued)

- (b) **Explain** how membranes participate in THREE of the following biological processes:
(6 points maximum; 2 points maximum per section)

Muscle contraction

- Motor neuron or axon terminal releases neurotransmitter or acetylcholine (ACh)
- ACh binds to receptors
- Depolarization or Na^+ moves in through membrane channels or membrane depolarizes
- Action potential propagates along cell membrane (sarcolemma) or T tubules
- Depolarization changes permeability of sarcoplasmic reticulum (SR) or Ca^{2+} released from SR
- Ca^{2+} active transport into SR (reuptake of Ca^{2+})
- Repolarization or maintenance of membrane potential (Na^+/K^+ pump)
- Smooth or cardiac muscle gap junctions directly transfer membrane potential between cells

Fertilization of an egg

- Part of the acrosomal reaction or sperm acrosome releases hydrolytic enzymes (by exocytosis)
- Sperm binds to receptors on egg
- Fusion of sperm and egg plasma membranes
- Change in membrane electrical charge or fast block (depolarization) to prevent further fertilization (polyspermy)
- Cortical reaction or slow block by exocytosis (prevents polyspermy) or “hardening” of membrane
- Separation of fertilization membrane (envelope)
- Fusion of egg and sperm nuclear membranes or nuclei

Chemiosmotic production of ATP

- Electron transport chain (ETC) in membrane pumps H^+ across membrane
- H^+ gradient established across membrane
- H^+ move through ATP synthase embedded in membrane to produce ATP
- Membrane infolding increases surface area

Intercellular signaling

- Release of chemical signals by exocytosis
- Receptors in membrane bind ligands or chemical signals or chemical signals pass through the membrane (examples: neurotransmitters, hormones, pheromones)
- Ligand-gated ion channels opening/closing
- Cascade of cellular events, including enzymatic reactions and second messengers (examples: G-proteins, cAMP, IP_3 , Ca^{2+})
- Antibodies activate immune function
- Descriptions of gap junctions, plasmodesmata (communicating junctions)

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Question 2

Cephalization and the development of a brain were important steps in animal evolution.

(a) **Discuss** the evolutionary origin and adaptive significance of cephalization in animal phyla. **(3 points)**

- **Cephalization (1 point)**

Defined: The concentration of the nervous system toward the anterior end of the organism

OR

Association: Cephalization tied to bilateral symmetry development

- **Origin (1 point)**

Origin identification: (Platyhelminthes/flatworms)

OR

Evolutionary progression of development

- **Adaptive Significance/Advantage (1 point)**

Efficient response to a stimulus (e.g., protection, predation, avoidance, movement toward or away)

During movement sensory organs encounter the environment first

(b) **Describe** the development of the nervous system in the vertebrate embryo. **(4 points maximum)**

- **Tissue of origin (1 point)**

- Ectoderm gives rise to the nervous system.

- **Processes of development (2 points)**

- Neurulation described (neural tube formation) Note: The notochord does not become the nerve cord.
- Other nerve development processes
 - Neural crest cells migrate to form the peripheral nervous system
 - Anterior portion of the neural tube/cord bulges to become the brain or brain regions

- **Endpoints with structures described at the end of a process step of development (1 point)**

- The ectoderm folds into the neural crest/tube or dorsal nerve/spinal cord
- Neural tube expands or develops into developmental brain region (e.g., fore-mid-hind brain, prosen-mesen-rhombencephalon)
- Spinal column/vertebrae/cranium that protects the CNS

- **Signaling (1 point)**

- Notochord (mesodermal in origin) signals or directs development of neural tube (ectodermal in origin)
- *Hox* genes, morphogens (diffusible developmental signal)

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Question 2 (continued)

(c) At the sound of shattering glass, people quickly turn their heads. **Discuss** how the human nervous system functions to produce this type of response to an external stimulus. **(5 points)**

- **Stimulus/Intermediating Structure of Receptor Action (1 point)**
Stimulus (sound waves, pressure, heat, etc.) producing an appropriate receptor action (eardrum vibrating, cochlear hairs vibrating or bending, pressure receptors firing, heat receptors firing, etc.)
- **Input/Sensory/Afferent (1 point)**
Signal direction toward the central nervous system
- **Integration (1 point)**
Processing/Interpretation by CNS
Interneurons/Association/Communicating/Internuncial
- **Output/Motor/Efferent Response (1 point)**
Signal direction toward effectors (peripheral NS) **or** description of the response or autonomic nervous response (e.g., increase in blood pressure or heart rate, muscle contraction **but not just** turning of head)
- **Possible Elaboration (1 point)**
Neural electrophysiology (e.g., action potential, neurotransmitters, synapse)
Neuron structure and impulse pathway
Sensory physiology

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Question 3

Compared with other terrestrial biomes, deserts have extremely low productivity.

- (a) **Discuss** how temperature, soil composition, and annual precipitation limit productivity in deserts. **(3 points maximum)**

Abiotic factor (description)	How abiotic factor limits productivity (must be linked) (1 point per factor)
Temperature Increase in transpiration/evaporation Desiccation Loss of water from tissues/guard cells Not optimal temperatures	Lowers photosynthetic rate Lowers plant growth Lowers biomass production PS/metabolic enzymes/proteins hindered
Soil composition Low organic content/nutrients Low water retention Sandy Compacted soil	Lowers photosynthetic rate/plant growth Lowers photosynthetic rate/plant growth Poor root anchorage limits plant growth Root limitations decrease photosynthesis
Annual precipitation Low rainfall Seasonal rainfall	Little water available for photosynthesis Lowers plant growth Period of high productivity/wildflowers

Clear definition/discussion of productivity: e.g., a measure of the amount of biomass produced by autotrophs/photosynthetic organism/plants...amount of light energy converted to chemical energy by autotrophs per unit time...reduced community productivity **(1 point)**

- (b) **Describe** a four-organism food chain that might characterize a desert community, and **identify** the trophic level of each organism. **(2 points)**

- **Written description** of a minimum of 4 organisms (must include a producer/plant) **(1 point)**
- **Clear identification** of 4 distinct trophic levels of the organisms discussed **(1 point)**
 (producer → primary consumer → secondary consumer → tertiary consumer
 or top carnivore or decomposer or scavenger)

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Question 3 (continued)

- (c) **Describe** the results depicted in the graph. **Explain** one anatomical difference and one physiological difference between species *A* and *B* that account for the CO₂ uptake patterns shown. **Discuss** the evolutionary significance of each difference. **(6 points maximum)**

Graph interpretation (3 points)

- Describe graph (plant *A* takes up CO₂ during day AND plant *B* takes up CO₂ at night) **(1 point)**
- Species *B* as CAM **(1 point)**
- Species *A* as C₃ or species *A* as C₄ **(1 point)**

Anatomical difference (1 point)

- Species *A* is C₄ with bundle sheath/wreath/Kranz anatomy
- Stomata location (pits/crypts, underside stems) linked to CO₂ uptake
- Stomata density linked to CO₂ uptake
- In species *B*/CAM vacuole/mesophyll of organic acids (malate)

Physiological difference (1 point)

- Species *A* stomata open during day
- CAM/species *B* stomata open at night/closed during day
- Species *A* uses C₃ pathway; CAM/ species *B* uses C₄ pathway
- C₃ uses Rubisco/C₄ uses PEP Carboxylase
- Organic acids synthesis for CO₂ storage
- Carbon fixation during day vs. night

Evolutionary significance (2 points)

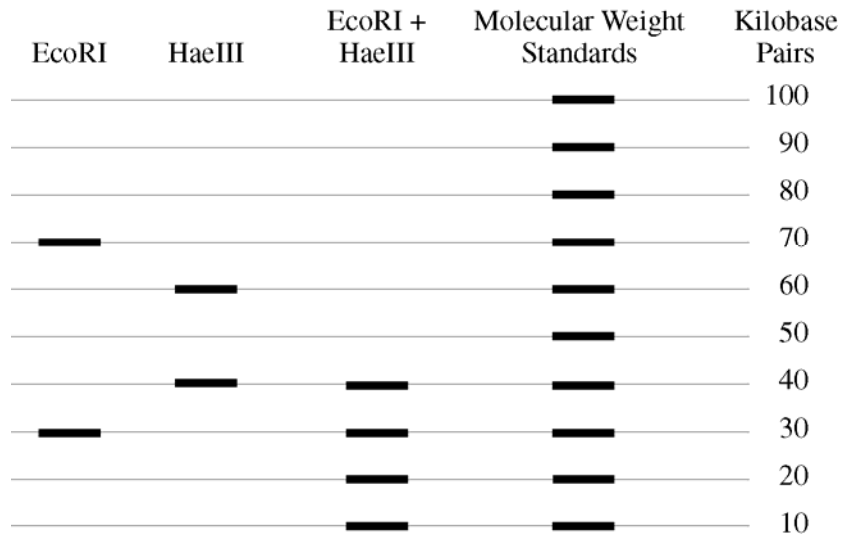
Discuss the evolutionary significance linked to each difference **(2 points, 1 point per difference)**
e.g., increased evolutionary success due to decrease in water loss in the desert environment
e.g., C₄ pathway circumvents the problem of photorespiration

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Question 4

A bacterial plasmid is 100 kb in length. The plasmid DNA was digested to completion with two restriction enzymes in three separate treatments: EcoRI, HaeIII, and EcoRI + HaeIII (double digest). The fragments were then separated with electrophoresis, as shown.

RESULTS OF GEL ELECTROPHORESIS

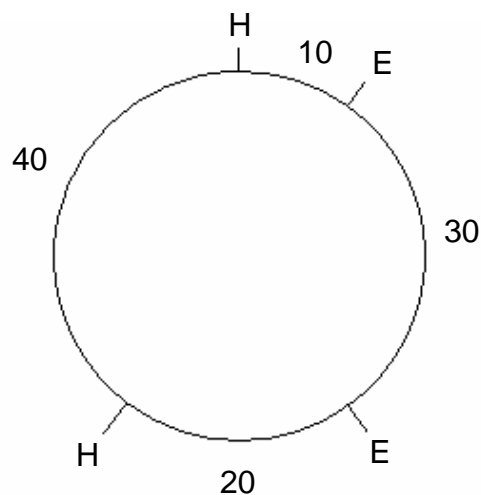
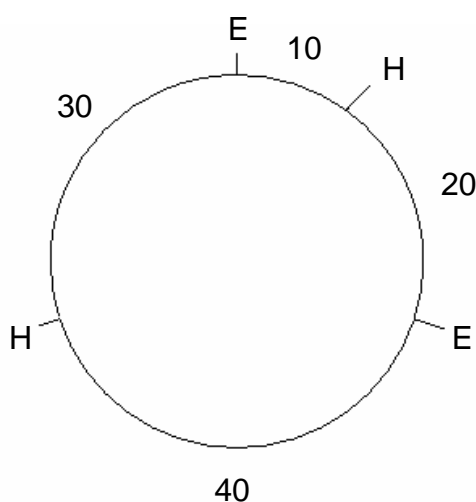


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Question 4 (continued)

- (a) Using the circle provided, **construct** a labeled diagram of the restriction map of the plasmid. **Explain** how you developed your map.

Construct a labeled map and **explain (3 points maximum)**



E = EcoRI Restriction Point H = HaeIII Restriction Point

- Restriction sites correctly placed and kilobase sizes shown (**2 points**)
- Explanation (**1 point**)
(NO POINTS for explanation with incorrect or missing map OR for interpreting gel only)
 - trial and error discussion
 - restriction site within larger fragment

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Question 4 (continued)

(b) **Describe** how:

- Recombinant DNA technology could be used to insert a gene of interest into a bacterium
- Recombinant bacteria could be identified
- Expression of the gene of interest could be ensured

Describe how to: (6 points maximum)

(1) Insert gene of interest (4 points maximum)

- Cut gene of interest from source and/or cut plasmid with restriction enzyme
- Use SAME restriction enzyme on both
- Anneal/ligate/mix/combine gene of interest with vector (plasmid/virus/phage)
- “Sticky ends”/bp matches/complementarity
- Treatment for competent cells (CaCl₂/heat shock); incubate together
- Chemical modification can prevent restriction enzyme activity (e.g., methylation)
- Gene = cDNA (without introns) to fit into plasmid

(2) Identify recombinant bacteria (1 point)

- Phenotypic selection (antibiotic resistance/blue-white colony selection/“glo” gene, product produced [e.g., insulin])
- Radioactively/fluorescently labeled probe (tag/dye) / mRNA
- Electrophoresis of cut recombinant vs. original (gene/plasmid) **OR** with sequence comparison of recombinant vs. original (gene/plasmid) **(Not bacterial genome)**

(3) Ensure expression of gene of interest (1 point)

- Promoter [for prokaryote]
- cDNA/removal of introns for prokaryotic expression
- Operon (e.g., nutrient/arabinose induced)

(c) **Discuss** how a specific genetically modified organism might provide a benefit for humans and at the same time pose a threat to a population or ecosystem. **(3 points maximum)**

Discuss GM, benefit to humans, and threat to population/ecosystem

- Nonhuman organism with specific, heritable GM trait
- Plausible benefit to humans related to the GM trait
- Plausible or unknown threat to population/ecosystem related to GM trait/modified organism