B2.1 Membranes and Membrane Transport

Guiding Questions

How do molecules of lipid and protein assemble into biological membranes?

What determines whether a substance can pass through a biological membrane?

Linking Questions

What processes depend on active transport in biological systems?

What are the roles of cell membranes in the interaction of a cell with its environment?

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<u>Theme</u>: Form and Function <u>Level of Organization</u>: Cells

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https://www.artstation.com/ artwork/VgyEbR

SL Learning Content

| B2.1.1 | Lipid bilayers as the basis of cell membranes | Phospholipids and other amphipathic lipids naturally form continuous sheet-like bilayers in water. |
|---------|---|--|
| B2.1.2 | Lipid bilayers as barriers | Students should understand that the hydrophobic hydrocarbon chains that form the core of a membrane have low permeability to large molecules and hydrophilic particles, including ions and polar molecules, so membranes function as effective barriers between aqueous solutions. |
| B2.1.3 | Simple diffusion across membranes | Use movement of oxygen and carbon dioxide molecules between phospholipids as an example of simple diffusion across membranes. |
| B2.1.4 | Integral and peripheral proteins in membranes | Emphasize that membrane proteins have diverse structures, locations and functions. Integral proteins are embedded in one or both of the lipid layers of a membrane. Peripheral proteins are attached to one or other surface of the bilayer. |
| B2.1.5 | Movement of water molecules across membranes by osmosis and the role of aquaporins | Include an explanation in terms of random movement of particles, impermeability of membranes to solutes and differences in solute concentration. |
| B2.1.6 | Channel proteins for facilitated diffusion | Students should understand how the structure of channel proteins makes membranes selectively permeable by allowing specific ions to diffuse through when channels are open but not when they are closed. |
| B2.1.7 | Pump proteins for active transport | Students should appreciate that pumps use energy from adenosine triphosphate (ATP) to transfer specific particles across membranes and therefore that they can move particles against a concentration gradient. |
| B2.1.8 | Selectivity in membrane permeability | Facilitated diffusion and active transport allow selective permeability in membranes. Permeability by simple diffusion is not selective and depends only on the size and hydrophilic or hydrophobic properties of particles. |
| B2.1.9 | Structure and function of glycoproteins and glycolipids | Limit to carbohydrate structures linked to proteins or lipids in membranes, location of carbohydrates on the extracellular side of membranes, and roles in cell adhesion and cell recognition. |
| B2.1.10 | Fluid mosaic model of membrane structure | Students should be able to draw a two-dimensional representation of the model and include peripheral and integral proteins, glycoproteins, phospholipids and cholesterol. They should also be able to indicate hydrophobic and hydrophilic regions. |

HL Learning Content

| B2.1.11 | Relationships between fatty acid composition of lipid bilayers and their fluidity | Unsaturated fatty acids in lipid bilayers have lower melting points, so membranes are fluid and therefore flexible at temperatures experienced by a cell. Saturated fatty acids have higher melting points and make membranes stronger at higher temperatures. Students should be familiar with an example of adaptations in membrane composition in relation to habitat. |
|---------|---|---|
| B2.1.12 | Cholesterol and membrane fluidity in animal cells | Students should understand the position of cholesterol molecules in membranes and also that cholesterol acts as a modulator (adjustor) of membrane fluidity, stabilizing membranes at higher temperatures and preventing stiffening at lower temperatures. |
| B2.1.13 | Membrane fluidity and the fusion and formation of vesicles | Include the terms "endocytosis" and "exocytosis", and examples of each process. |
| B2.1.14 | Gated ion channels in neurons | Include nicotinic acetylcholine receptors as an example of a neurotransmitter-gated ion channel and sodium and potassium channels as examples of voltage-gated channels. |
| B2.1.15 | Sodium–potassium pumps as an example of exchange transporters | Include the importance of these pumps in generating membrane potentials. |
| B2.1.16 | Sodium-dependent glucose cotransporters as an example of indirect active transport | Include the importance of these cotransporters in glucose absorption by cells in the small intestine and glucose reabsorption by cells in the nephron. |
| B2.1.17 | Adhesion of cells to form tissues | Include the term "cell-adhesion molecules" (CAMs) and the understanding that different forms of CAM are used for different types of cell–cell junction. Students are not required to have detailed knowledge of the different CAMs or junctions. |



| B2.1.1-Lipid bilayers as the basis of cell membrar | nes. B2.1.2 – Lipid bilayers as barriers. |
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| 32.1.4—Integral and peripheral proteins in membrar | nes. B.2.1.9—Structure and function of |
| giycoproteins and giycolipids. B2.1.10—Fluid III | |
| of all cells: ① allows internal conditions to be different from surroundings ② c | ontrols entry and exit of substances 3 allows sensitivity and communication @ cell structure A2.2 |
| cell membrane structure proposed by Singer and Nicolson (1972): Oynamic, flexible str can move laterally + "mosaic": non-uniform, made of many different components | oucture made primarily of a phospholipid bilayer with proteins and carbohydrates. s (lipid, carbohydrate, and protein) in various configurations |
| by Davson and Danielli (1935) was falsified using: | Cholesterol: amphipathic lipid, allowing it to integrate within hileware help control membrane fluidity: ensuring they are not too fluid or too stiff |
| bilayer was split open showing proteins D integral transmembrane proteins | hydrophilic HO hydrophobic> expanded in HL |
| Glucoliaid chalesteed the | Phospholipids are amphipathic and spontaneously form bilayers in solutions @ lipids Bl.1 |
| | hydrophilic, polar head > semi-permeable barrier between aqueous solutions |
| | → phosphate group + glycerol i.e. inside and outside the cell: → hydrophobic molecules - high permeability H_C - C - CHZ + C - C - C - C |
| | phospholipid = c=0 very small non-polar molecules - high permeability bilayer ex: O2, CO2 |
| | ex: HzO, ethanol, urea, glycerol |
| | hydrophobic, non-polar tails ex: glucose, sucrose require transport |
| peripheral proteins | Statty acid hydrocarbon chains × charged molecules and ions proteins |
| | (saturated or unsaturated) ex: amino acids, Na ⁺ , K ⁺ , Cl., Ca ⁺ , H ⁺ |
| edded in the membrane Peripheral proteins proteins attached (often temp to the surface of the membrane | porarily) Glycoprotein: polypeptide bound rane to a carbohydrate space of cells |
| ane (both sides) or on an integral protein | (olycoliaid : liaid hand to a |
| ane proteins are amphipathic; Peripheral proteins are hydric in the interact | rophilic short carbohydrate chain |
| ng them to integrate within 000000000000000000000000000000000000 | Cell recognition & lipids BI.I Cell adhesion -> expanded in HL |
| d bilayer | unique shape of glycoproteins and glycocalyxes of adjacent cells fuse, holding |
| ex: transporter | glycolipids act as antigen or surface cells and tissues together |
| | from non-self through binding ex: N-CAM |
| | ex: red blood cells |
| receptor | have glycoprotein antigens (A, B, O) |
| | |

B2.1.3—Simple diffusion across membranes. B2.1.5—Movement of water molecules membranes by osmosis and the role of aquaporins. B2.1.6-Channel proteins for facilitate B2.1.7-Pump proteins for active transport. B2.1.8-Selectivity in membrane permea

Many of the processes of life (such as metabolism, homeostasis, nutrition, excretion) require transport of materials in and out of cells This can occur both passively or acti ↑[::] down concentration gradient Active transport : movement of particles from a · net movement of particles from an area Passive transport area of low concentration to of high concentration to low concentration Chooo **↑[**··] high concentration using AT without the use of ATP • particles are constantly in random motion (due to kinetic energy) and move independent of each other but as there are more particles in areas of high concentration that can randomly move to areas of low concentration than the other way around, a <u>net</u> movement from high to low concentration will occur passively. • ATP is the cell's 'energy currency' storing potential • rate can be increased by increasing concencentration gradient and membrane surface area and decreasing distance needed to move gas or liquid 🗲 energy from cellular respiration. When hydrolyzed Simple diffusion : passive net movement of particles from an area of high concentration to low concentration in fluids it releases this energy to power metabolic processes • Pump proteins : transmembrane proteins which use chemical energy - ATP in order to • <u>small</u> <u>non-polar</u> molecules can diffuse easily across membranes RRAMAN RECENTER RRAMMULU LARAM Sable to pass through hydrophobic core easily -> able to pass in between phospholipid molecules → ATTERN ATTERN A ex: oxygen (O_z) O=O and carbon dioxide (CO_z) O=C=O. [∴] ≫ [··] [::] > [*] = [*] ↑[..] X not selective: only depends on size and water-affinity Secilitated diffusion passive net movement of particles from an area of high concentration to low concentration using membrane proteins (such as channel or carrier proteins) ↓[] • Channel proteins : integral membrane proteins which have a hydrophilic **^ [N**a[†]] Nat Nat (00000 RITRIR RAUSS pore allowing a passage for charged (Nat, Kt, CI, Ca2+) or polar molecules (glucose, water) to cross hydrophilic core Choee ↓ [Nat] the membrane's hydrophobic core easily ↑[::] * channels are selectively permeable : shape / size / charge of pore ensures specificity ex: Nat channels only allow Nat some (like voltage-gated ion channels) can be open or closed, allowing control Somosis : passive net movement of water from an area of low solute concentration to high solute concentration ↓[] across a semi-permeable membrane & water potential D2.3 water entered cell • membranes allow the passage of water but restrict the movement Of larger or charged solutes, thus causing water to move more • greatly increases permeability • when water is interacting with and dissolving solutes its to water and osmotic rate movement is restricted **b** water in a dilute solution is located in collecting ducts ↑[solute] ↓ [solute] more free and likely to move to a more concentrated solution. in kidneys and plant roots

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• equilibrium when overall solute concentration is equal on both sides

cell is more concentrated : takes in water (net)

| across ed diffusion. ability | |
|--|-------------|
| ively | |
| o v[··] against concentration gradient > | ↑[∷] |
| | |

• particles in an area of low concentration are less likely to move to an area of high concentration then the other way around, thus for this to occur reliably an external energy input is required



transport particles across membranes against their concentration gradient

X: Pump proteins allow selective permeability: only transport specific ions or molecules in one direction

Pump protein in its stable conformation is open to one side of membrane. It has high affinity to particle (•) and binds to it.



Pump is open to other side and has low affinity for particle, releasing it against its concentration gradient. Pump reverts back to its original, stable conformation - ready for reuse

Aquaporins : channel proteins which selectively allow passage of water

for water absorption



Aquaporin has a very narrow pore which is positively charged only allowing water through (excluding H+)

B2.1.11 — Relationships between fatty acid composition of lipid bilayers and their fluidity.
 B2.1.12 — Cholesterol and membrane fluidity in animal cells. B2.1.13 — Membrane fluidity
 and the fusion and formation of vesicles. B2.1.17 — Adhesion of cells to form tissues

| Temperature impacts membrane <u>fluidity</u> i.e. the degree of random movement of the components, namely phospholipids which can move laterally, rotate and even switch positions ARRAR more movement greater fluidity greater permeability lower perm | nent The fluidity of the membrane allow dity and thus the formation (endocytosis eability |
|--|--|
| Impact of phospholipid fatty acid composition on membrane fluidity | Endocytosis: the formation of a ves of the plasma membrar Process requires ATP |
| > the fatty acid composition of phaspholipids can be saturated (no double bonds) or unsaturated (≥1 double bond) @ lipids Bl.1 | ex: |
| fatty acid tails → pack very tightly - high density 000000 less fluid /more viscous & stron saturated ∴ high intermolecular forces is less permeable ∴ high melting point is flexible & too s | Phagocytic white blood nger cells and paramecia engulf and destroy solid pathogens / food particles |
| fatty acid tails -> pack loosely - lower density fatty acid tails -> pack loosely - lower density forces fluid /less viscous forces | fluid |
| General trend: the relative percentage composition of saturated and I higher percentage of I higher percentage | of |
| Adaptation. Fish living in Antarctic waters examples: (such as <u>C.hamatus</u>) have membranes rich in unsaturated fatty acids The temperate freshwater fish <u>A. fulvescens</u> increases its membrane unsaturated content. when lake temperatures decrease The caribou <u>R. tarandus</u> has different membrane fatty acid composition in different tissues, as a trend: the further away tissues are from the body core, the more unsaturated fatty acids (hooves > upper leg > body | (y) (x) (x) (x) (x) (x) (x) (x) (x |
| Impact of cholesterol on membrane fluidity | ex: Neurotransmitters (such as Assilutation) |
| Scholesterol embeds itself in the plasma membrane as its amphipathic - OH 'head' is polar, body is non-polar K cholesterol is present in animal but not plant membrane fluid | are released by neurons during synaptic transmission in order to |
| Scholesterol octs as a membrane fluidity modulator: | |
| Cholesterol decreases fluidity by constraining phospholipid movement - stabilizing membrane | cell-adhesian molecules (CAMs) cell- cells erol |
| Cholesterol increases fluidity by disrupting Solid Solid | |
| 🕘 7 \ phospholipid packing - preventing stiffening low Temperature high | |

HL

us for phospholipid bilayers to change shape and fuse,) and fusion (exocytosis) of vesicles sicle from the invagination and pinching off of a piece ne - bringing <u>extracellular content</u> within the cell fluids (pinocytosis) <>>> solids (phagocytosis) Y Developing fetus takes maternal sid () in antibodies into its blood stream from the mother fetal sid via the placenta ATTATA PARTATA PARATTATA PARTATA PARTATA PARATTAT BIBLIDI NUSULISI UNUSULI DIVULU DU vesicle SEL AND STREET

with the plasma membrane - discharging vesicle contents ace (i.e. outside the cell). Process requires ATP

ncreas secretes pestive enzymes uch as lipase) to the duodenum and hormones uch as insulin.) into the blood ream for signaling

Paramecia osmoregulate by loading and expelling excess water using contractile vacuoles

surface proteins/glycoproteins which linK cells with other ; forming junctions - providing structure and tissue formation.

a) cells together, creating a barrier t as a bridge, allowing communication and material exchange inect cells to extracellular matrix, providing anchorage



B2.1.14—Gated ion channels in neurons. B2.1.15—Sodium-potassium pumps as an example of exchange transporters. B2.1.16—Sodium-dependent glucose cotransporters as an example of indirect active transport

| Channel proteins allow for facilitated diffusion, transporting specific materials that otherwise would only diffuse | Exchange transporters : Transmembrane protein which transports different substances in opposite directions across |
|--|--|
| slowly or not at all across a phospholipid bilayer (such as ions). While they do not control direction of flow | (antiporter) the membrane against their respective gradients, thus 'exchanging' them |
| (always a net flow high to low concentration) they can be opened or closed reversibly to control transport | Sodius Petersius auro i anche se la socia en 2414 - 1 aut - 1 11 12 - 1 |
| Name house it has a had in the male. Transmentance as here housed which allow the difference of a carefor in | (Net/Vt ATRice) exchange transporter moving 3 Na' and 2 K against their respective |
| 'hoted' as they are a class in cranics of a newatransmitter | (Na'/ K - AllPase) concentration gradients across the membrane using All |
| Sudd us they spen a close in response to binding of a neorod ansmitted | $\uparrow[N_{\star}^{\dagger}] \downarrow[\kappa^{\dagger}]$ Puma biads 3 Na ^t from subsolution |
| X neurotransmitters are signalling molecules which are secreted by a neuron at synapses to couse on effect | RIRAR RARA RARA RARA RARA RARA RARA RAR |
| ex: Acetylcholine, Glutamate, GABA, Dopamine, Scrotonin, Epinephrine | ATP is hydrolyzed to ADP, binding |
| | ↓[Nat] ↑[K+] ↓ phosphate to (phosphorylating) pump |
| Nicotinic acetylcholine receptors : gated ion channel allowing diffusion of sodium (Na+) into the cell | Coele Chosphorylation causes a change in pump |
| sl (when two acetylcholine (or nicotine) molecules bind to the receptor, | conformation, reducing its affinity for |
| causing the cell to depolarize and an action potential to be propagated | T[Na ⁺] +[K ⁺] RAPPROACHERSING 3 Na ⁺ outside cell RAPPROACHERSING 3 Na ⁺ outside cell |
| | $\left(\begin{array}{c} \left(\left(\begin{array}{c} \left(\begin{array}{c} \left(\left(\begin{array}{c} \left(\left(\begin{array}{c} \left(\left(\left(\begin{array}{c} \left($ |
| $N_{a}^{*} N_{a}^{*} N_{a}^{*} N_{a}^{*} N_{a}^{*} N_{a}^{*} N_{a}^{*} N_{a}^{*} Acetylcholine is secreted from the$ | 1 [wt] t[wt] t[wt] |
| $= \frac{1}{1000} \frac{1}{1000} \frac{1}{1000} \frac{1}{1000} \frac{1}{1000} \frac{1}{1000} \frac{1}{1000} \frac{1}{1000} \frac{1}{10000} \frac{1}{10000} \frac{1}{100000} \frac{1}{1000000} \frac{1}{10000000000000000000000000000000000$ | Give a court to aciainal |
| Synaptic citre and divises across | × as the sums undergoes unequal exchange of rations (lose 3, take in 2) ranformation and release 2, K+ into cell |
| s) ~ min (initial of the state | it maintains a relative negative membrane potential (-70mV) Pump ready for reuse. |
| Nat Nat Nat Nat Z Two Acetylcholine molecules bind | · · · · · · · · · · · · · · · · · · · |
| to the receptor, inducing a conformational | Indirect active transport . Uses the energy produced by the movement of one substance down its concentration gradient |
| Top view change, opening a pore and allowing | to transport another <u>against</u> its concentration gradient. Direct ATP use not required |
| Nat to diffuse through | |
| | Sodium-glucose cotransporter · cotransporter protein which transports glucose against its concentration gradient |
| Voltage-gated ion channel I ransmembrane protein channels which allow the diffusion of a specific ion | into the cell using the energy from simultaneously transporting Nat down its gradient |
| Gateo as they open or close in response to a specific membrane potential (voltage) | ↑[Nat] ↓[glucose]) |
| X potential difference or voltage is the amount of electric patential energy between two points (V or JC^{-1}) | RIRINGRANNING REAL AND SOLUTION OF CATRANSPORTER. |
| membrane potential is the difference in electric potential (V) between the inside and outside of the cell | booccoccoccoccoccoccoccoccoccoccoccoccoc |
| \rightarrow inside of the cell ~ -70 mV compared to the outside \rightarrow generated by the sodium-potassium pump | ↓[Nat] ↑[glucose] and move glucose into cell small intestines willi kidney |
| | |
| > voltage-gated sodium channel : gated ion channel allowing diffusion of sodium (Na+) into the cell | > voltage-gated potassium channel: gated ion channel allowing diffusion of potassium (K+) out of the cell |
| when there is an increase in membrane potential (-70mV to -55 mV, |) when the membrane potential has flipped (-70mV to 30mV) causing |
| cousing the cell to depolarize (negative to positive) | the cell to repolorize (positive to negative) |
| $N_{a}^{\dagger} N_{a}^{\dagger} N_{a$ | |
| $N_a^+ N_a^+ \times N_a^+ N_$ | K^{+} K^{+ |
| 1 KIIKI REAL OPENS, allowing | RIPRI REPAIR REPAIR 2 REPAIR REPAIR activation gate opens, allowing |
| - Bood - Bood - Bood - Bood - Bood - Bood - Nat to diffuse into cell (-55 mV) | about a bound a bound a bound a bound a bound a bound a kt to diffuse out of cell (30mV) |
| Nat Nat Nat V Nat a inactivate gate rapidly closes chann | iel, K+ K+ X K+ K+ K+ K+ K+ K+ V K+ K+ V K+ 2 inactivate gate rapidly closes channel, |
| activation gate Na ⁺ Na ⁺ Inactivation gate Na ⁺ Na ⁺ Inactivation gate Na ⁺ Na ⁺ Inactivation gate Na ⁺ Na ⁺ Inactivation gate Na ⁺ Na ⁺ Inactivation gate Na ⁺ Na ⁺ Inactivation gate < | κ ⁺ Ο κ ⁺ |
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BIBLIOGRAPHY





