



Enlightism
Spreading Inspiration

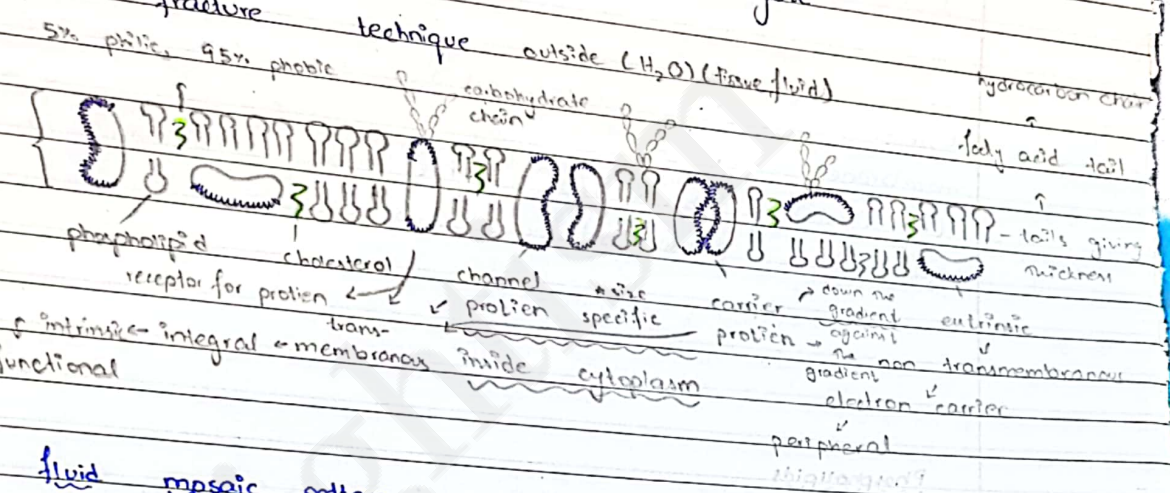
AS Biology

Unit:
Cell membrane

Contributed by Saima

Cell Membrane

- thickness : 7nm
- observed with electron microscope
- common feature in prokaryote and eukaryote
- fluid mosaic model
- freeze fracture technique



philic → β pleated
phobic → α helix

fluid mosaic pattern

cell membrane contains 2 monolayers of phospholipids which diffuse into each other. Between them proteins are scattered (in diff: pattern / mosaics). This is fluid mosaic model.

phospholipids ↓ movable
protein ↓ 99% movable
*increasing temp → increases kinetic energy → mobility increase
↓
denaturation of memb: since it does not remain intact

Sajid

Sajid

Day / Date

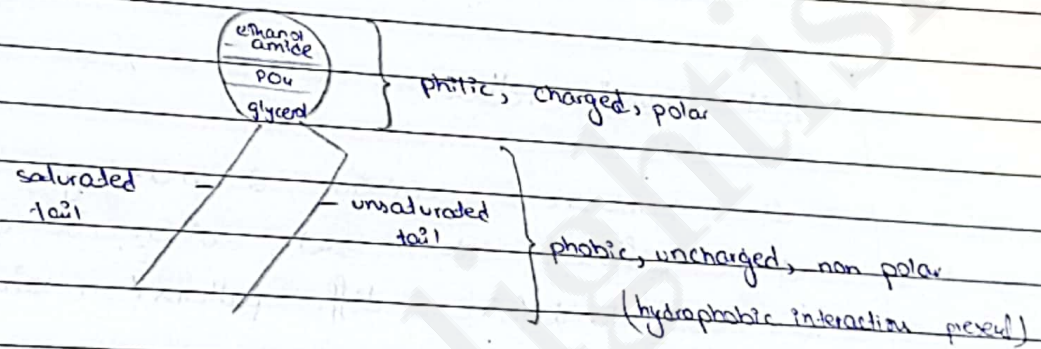
freeze fracture technique

Cell membrane is fluid therefore it is difficult to cut it's cross sections. For taking the cross sections, we keep the cell in a freezer. Membrane becomes hard, easy to cut the cross section. This is called freeze fracture technique.

⇒ Composition varies. Mitochondria membrane and Thylakoid membrane contain 80% protein and 20% other things (cholesterol & phospholipid). myelin sheath contains 20% protein & 80% phospholipids and cholesterol.

↑ protein = metabolically active
↑ phospholipid & cholesterol = metabolically less active

Phospholipids



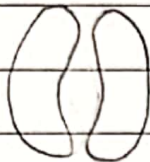
proportion of the tails keeps on changing membrane

↓	↓
rigid	fluid

- | | |
|--|---|
| <ul style="list-style-type: none"> → increase saturated tails → less cholesterol → longer tail (more phobic interactions) | <ul style="list-style-type: none"> → increase unsaturated tails → increase cholesterol → shorter tails |
|--|---|

Proteins

vary dep: on the size



of mol: that

will pass channels

→ single specific

passive movements → no energy reqs

→ no ATPase

→ always down the gradient

→ movement of mol: faster

→ contains no carbohydrate chains

→ keeps on changing its position while transporting

→ fixed shape

→ allows H₂O soluble things



substrate specific

(lock & key)

carriers

enzyme ATPase

→ having receptors

passive / active

→ enzyme ATPase

→ down / against depends on need

→ slower

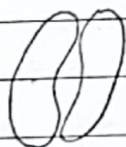
→ contains carbohydrate chains

→ does not keep on changing its location

→ keeps on changing its shape

→ H₂O soluble as well as H₂O insoluble

} while things are passing through otherwise both do



aqua porins



only permeable to water



glycoproteins



receptor for protein (signaling)
↓
antigen (attached on cell membrane)
↓
when falls in plasma antibody

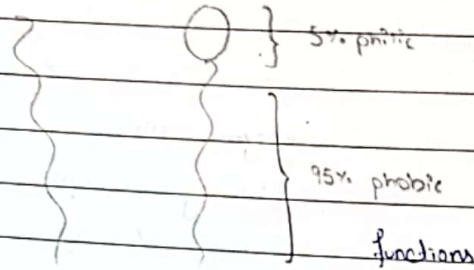


electron carriers

Day / Date

O₂ & CO₂
pass through
phospholipid
bilayer

Cholesterol



increasing fluidity

↓
plugs entry of
ionic solution via
phospholipid bilayer
which is totally non-polar

→ chains are
recognition
site

Carbohydrates (chains)

always on outside of cell membrane

→ binding
(lock & key)

occurs on the
protein

develops hydrogen
bonding with H₂O
molecules around

helps in anchoring of cell memb;
with H₂O mol: around

↓
giving mechanical stability
to cell memb

length of chain keeps
on changing



mechanical stability
anchoring

glycoprotein

↓
receptor

(fallen out in
plasma) antibody

(B lymphocytes)

↓
protein receptor

(in memb) antigen

↳ glycoprotein

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glycolipid

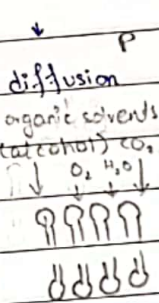
- receptor for steroid (align)
- helps in joining cells to make tissues

(the carbohydrate chains in glycolipid binds (similar to protein))

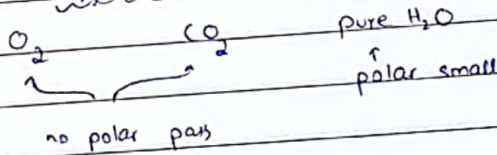
→ plant cells are joined via plasmodesmata

exchange processes

→ water is polar but passes through phospholipids due to its small size



diffusion via phospholipids



through non polar tail

↓ P facilitated diffusion

↓ P osmosis

↓ A active transport

↓ A bulk transport

diffusion depends upon

SA of molecule

size of molecule

thickness of membrane

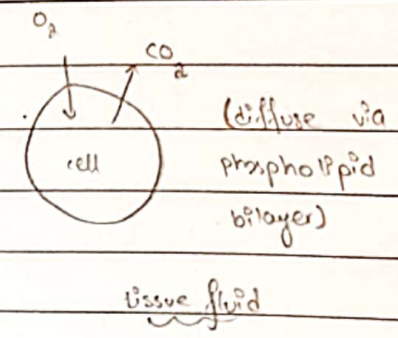
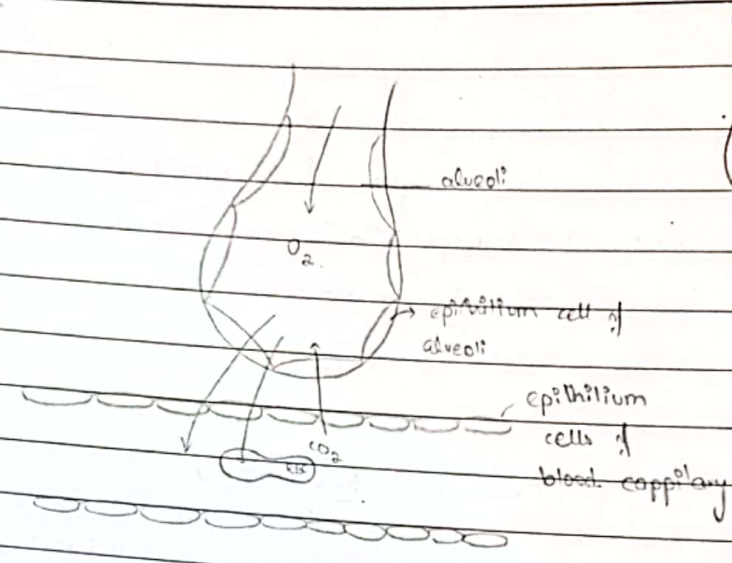
nature of molecule

Conc: gradient

temp:

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* alcohol causes more damage to animal cells compared to plant cells since cholesterol also dissolves in alcohol

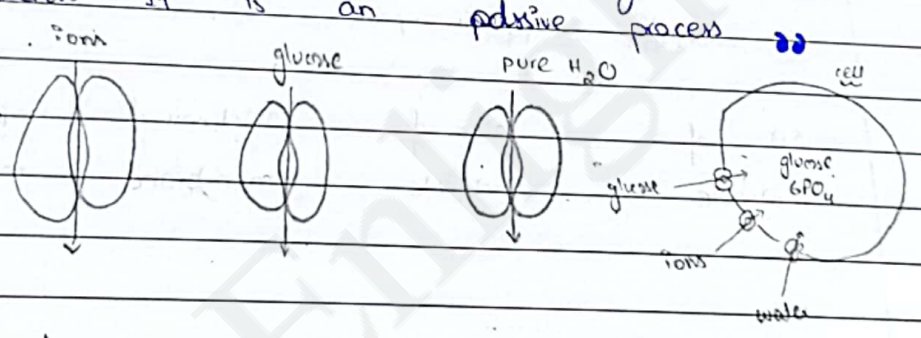


of membranes crossed
 O₂ → CO₂

of monolayers of phospholipids
 in plasma ← 4, 5 → in RBC ← 8, 10 →
 in plasma ← 4, 5 → in RBC ← 8, 10 →

facilitated diffusion

“ diffusion via channel (protein) always down the gradients. It is an passive process ”



O₂ →
 to prevent backflow once ions inside increased

Why is glucose converted in glucose 6 P O₆?
 + no channel / carrier for glucose 6 P O₆
 bigger molecule is trapped in cytoplasm of cell and cannot leave via the same channel. It does not move via phospholipid bilayer as it has become charged.

diff & fac: diff
sim

diff

- movement down the conc: gradient
- passive process
- channels
- pore substances

Q If cell membrane of mitochondria (inner) is less permeable to ions, water & salts, what should be its composition. [3]

→ channels are preferred over carriers for movement however channels enhance uptake

- more cholesterol
- less channels & carriers
- unsaturated phospholipid tails
- longer phospholipid tails

active transport

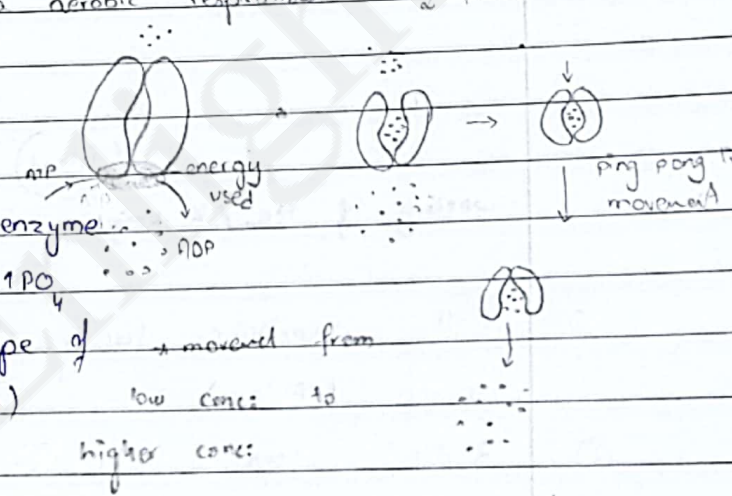
Q [4]

“ movement of molecules from an area of lower conc: to an area of higher conc: against the conc: grad: using energy it is an active process. ”
from mitochondria
via aerobic respiration → O₂ present

- 1) always via Carriers
- 2) against the gradient
- 3) enzyme present on carrier.

ATPase we provide ATP enzyme. will break ATP → ADP + P_o

is used for changing shape of carrier (configurational changes) of molecules against gradient. → moved from low conc: to higher conc:



Day / Date

carriers work depending on need of cell

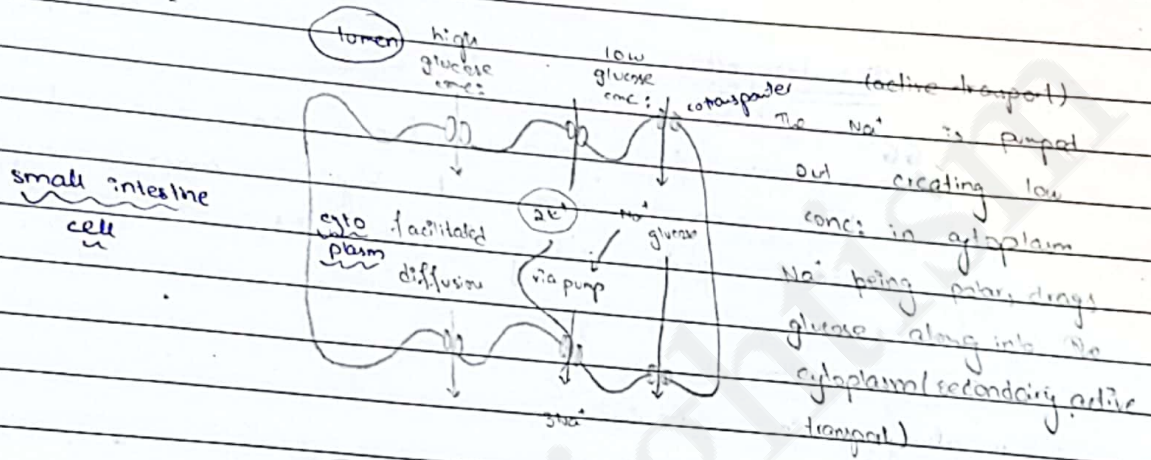
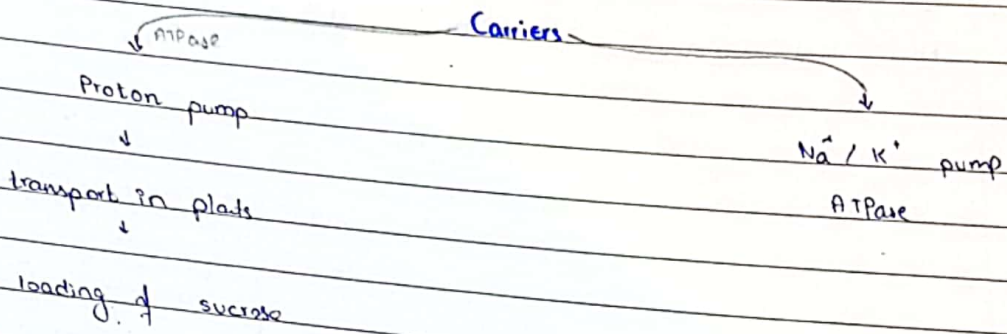
if

glucose more outside

glucose more inside

enter via fac diff

enter via active transport



working of Na^+ / K^+ pumps

- ① Cell membrane facing blood capillary contains Na^+ / K^+ pump (ATPases)
- ② 3 Na^+ ions enter the blood, in return 2 K^+ ions enter the cytoplasm. K^+ ions enter the lumen via facilitated

diffusion.

(3) due to the movement of Na^+ ion from the cytoplasm, there is deficiency of Na ion in the cytoplasm of cell.

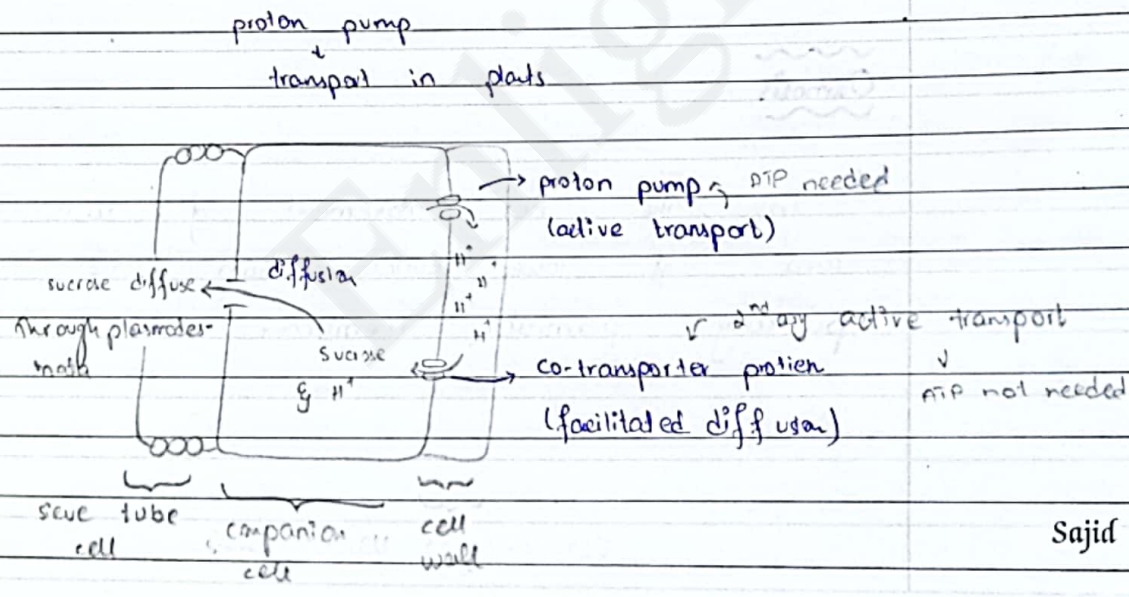
(4) To maintain the concs of Na^+ ion. in the cytoplasm, Na^+ ion with glucose enters the cytoplasm via co-transporter protein or symporter (a channel from which 2 things pass at a time)

(5) Na^+ ions are charged ions. They produce a dragging force or driving force which takes glucose with it inside cytoplasm.

(6) from cytoplasm, glucose enters the blood via facilitated diffusion.

(7) Secondary active transport depends for its working on active transport. No active transport means no secondary active transport.

(8) When Na^+ ions enter into the cytoplasm, water potential becomes lower or more negative. To maintain it, water rushes from lumen into the cytoplasm, back into the blood via aquaporin. Reabsorption of water also occurs.



Day / Date

Q What is passive movement?

Q What is active process?

Q Give diff: b/w diffu: & fae: diffu:

Q Give diff: b/w fae: diffu: & active transport.

Osmosis

“ In the net movement of water molecules from area of lower conc: down the gradient through partially permeable membrane. ”

wilting \rightarrow plasmolysis



carrier works according to need
whilst channel does not

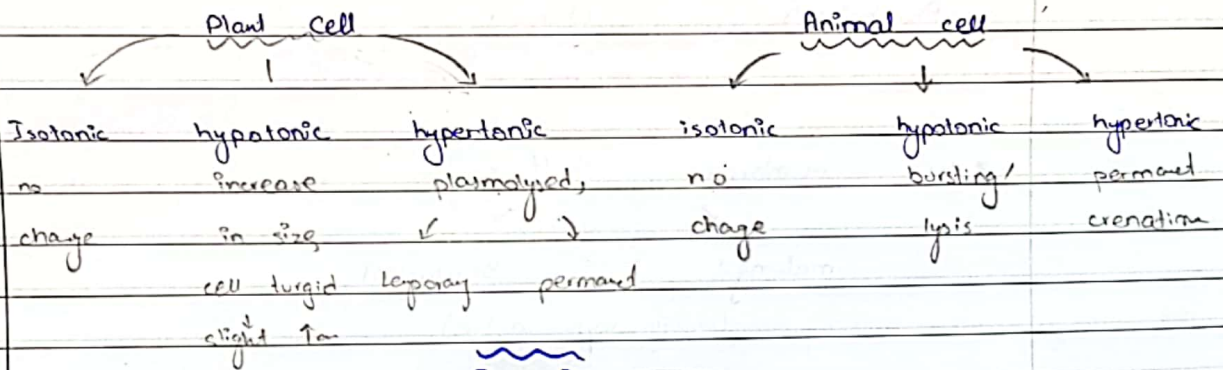
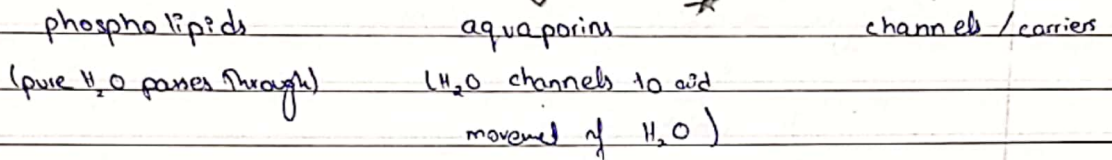
Day / Date

24th Oct 19

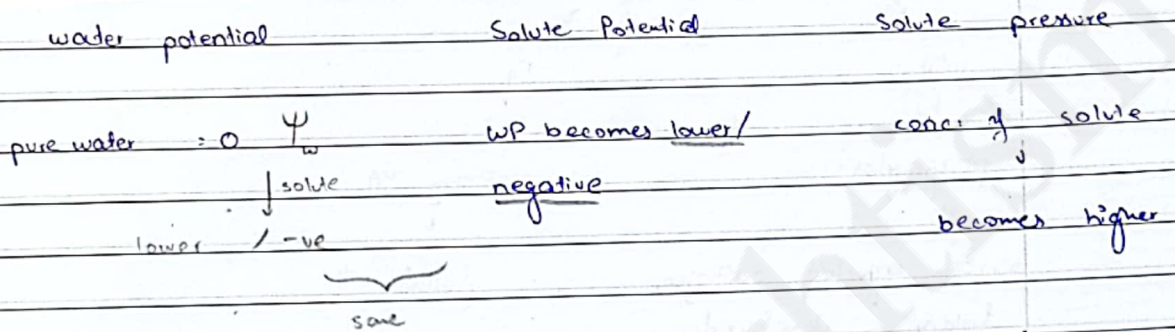
Thursday

* inward
plasmolysis
 \downarrow
start of
plasmolysis

Osmosis / H₂O



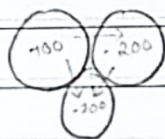
Osmosis



* when solute is added, both Ψ_w & sp become lower / -ve
however the solute pressure becomes higher / +ve.
concn of solute

exam
ans

When sucrose is added in sieve tube cell, Ψ_w or sp becomes lower / -ve, solute pressure becomes higher
turgid pressure



water movement

Sajid

Bulk transport :-

Bigger molecules → Proteins

↳ which

Phospholipids

Channels

Carriers

* cell

membrane

? itself

making it

breaking it

(energy req. active)

bulk transport

solid

phagocytosis

lig

ovary egg

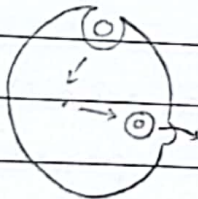
follicle cell

(pinocytosis)



memb: engulfed pathogen, foldings developed around pathogen

endo cytosis



no memb combining waste throw out it's contents, ?) itself becomes a

part of the csm and therefore

we can say the membrane grows

ADH, glucagon, insulin, FSH, LH → protein in nature

Quin
Eis
Toment

Date

28th Oct '19

Monday

Cell Signalling:

It is communication b/w cells



① Insulin dumped in plasma, produced by pancreas, binds with CSM of liver cells.

→ glucose uptake increases

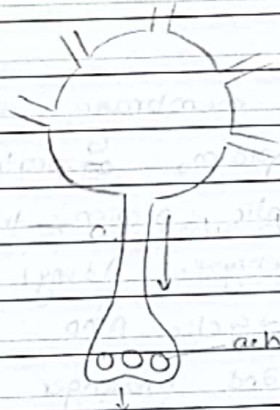
→ CSM's permeability increases

→ glucose converted into glycogen

→ rate of resp: ↑



②



acetylcholine

protein in nature

never enters

ion channels

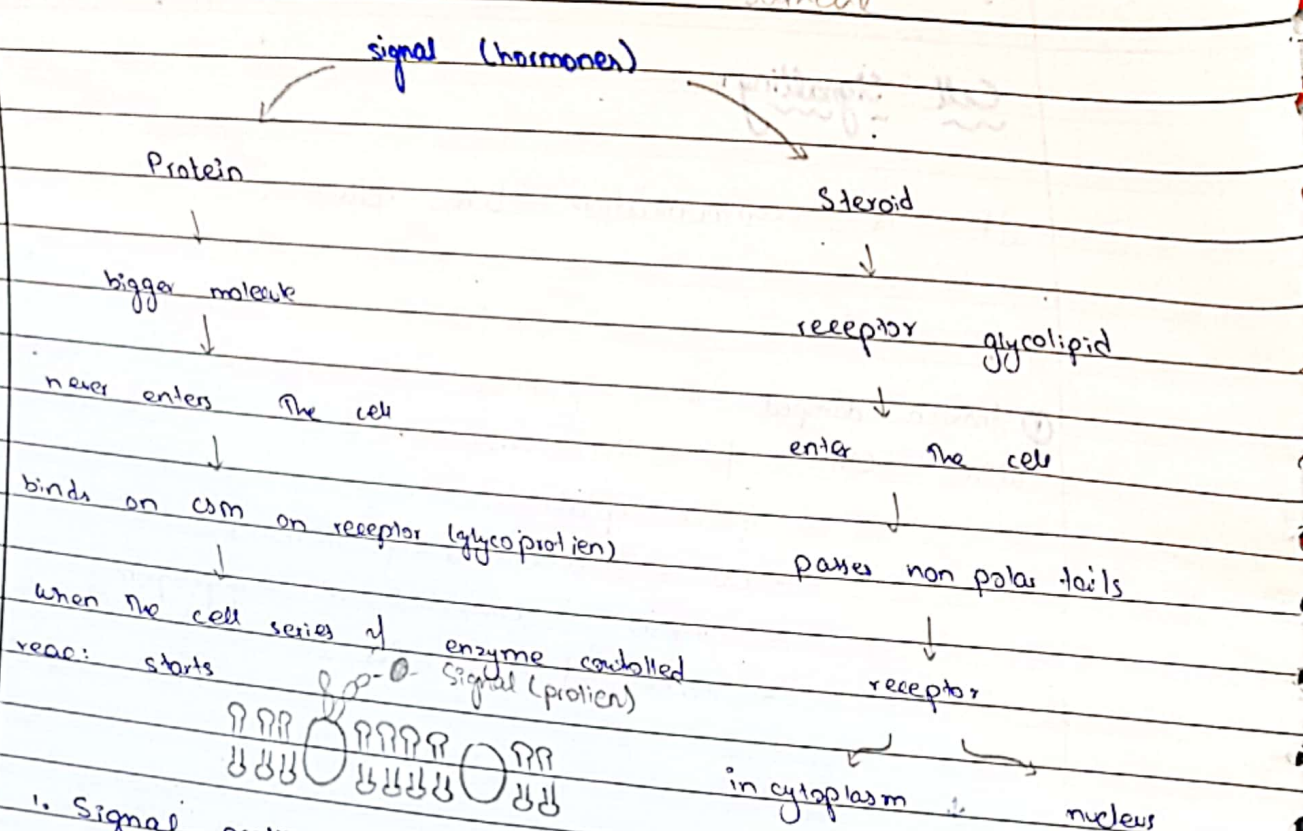
open

signal transferred

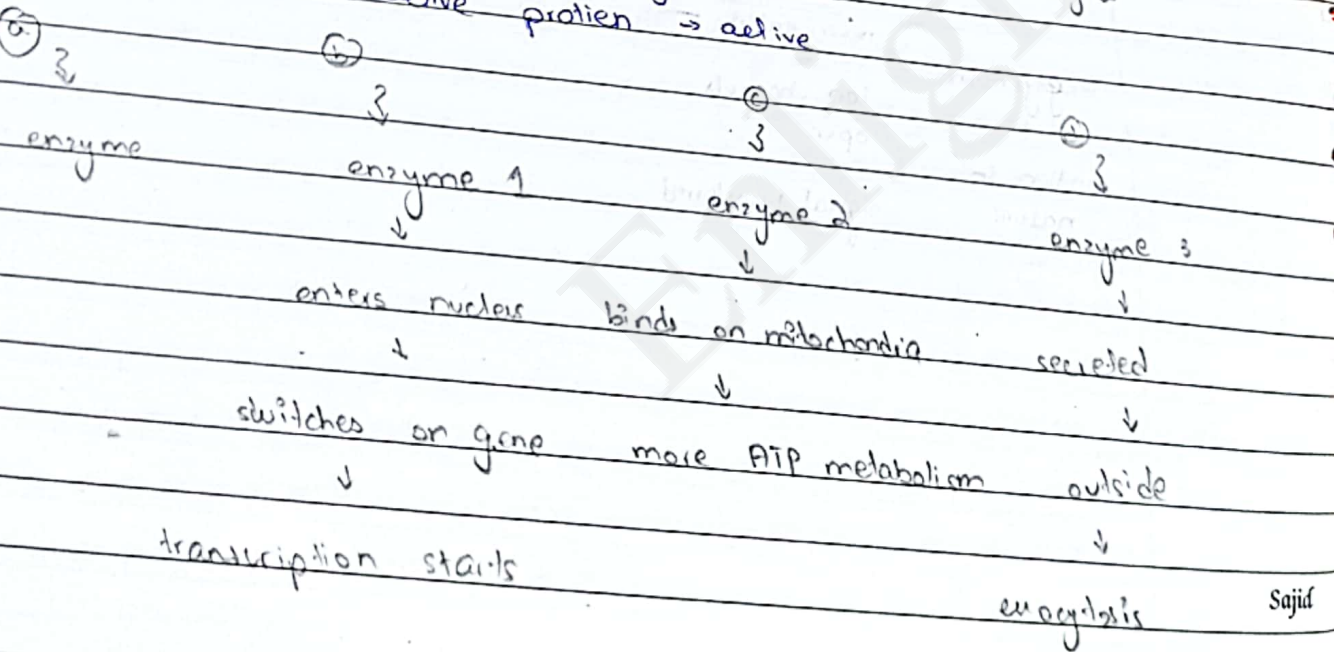
ach neurotransmitter

falls on CSM of another neurone

Ep's same cell

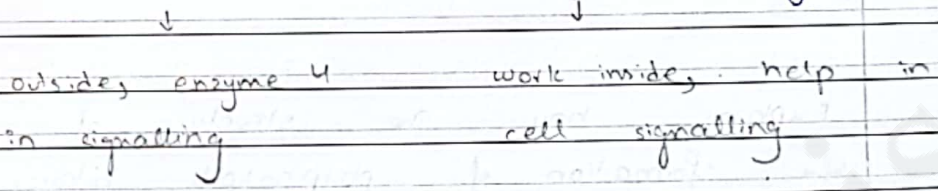


1. Signal protein
2. Binds on cell membrane receptor
3. Inside in cytoplasm, G protein activated
4. Activates enzymatic protein, transmembranous enzyme
5. Activates on enzyme. Adeny cyclase
6. Converts ATP → cyclic AMP
7. Cyclic AMP and messenger
8. Converts inactive protein → active



Cell membrane Q's

Q, what are extracellular and intracellular enzymes?



Q there is a protein (toxin) produced by bacteria, works on RBC.

Can dissolve / damage

- a) phospholipids bilayer
- b) channels ions
- c) carriers down & against grad

what will be effect on working of RBC if in tissue fluid

- a) uncontrolled water enters, lysis happens (haemolysis)
- b) glucose & ions will have an inhibitor, wp will drop so water will also rush in, lysis occurs

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↳ Water enters, bursting

steroids: cell wall of a plant cell damaged

→ cell will not burst.

Q1. How mitochondria destroyed by cell signalling and why is it destroyed? [3]

*signal is never
either protein
or steroid
always
inter

signal within the cell binds on mitochondria, series of enzyme controlled reactions, mitochondria will be denature. OR

signal sent to lysosome to destroy worn out mitochondria. It will engulf.

reason: Mitochondria working inefficiently

Q2. Explain how the structure of phospholipids allows the formation of phospholipid bilayer of cell membrane. [3]

head phobic towards water, phobic tails come close, phobic interactions created, bilayer made

Q3. RBC placed in NaCl solution is crenated because it has an abnormal shrivelled appearance. Explain how this RBC has shrivelled. [3]

placed in hypertonic, water moved out, cell crenates, end osmosis

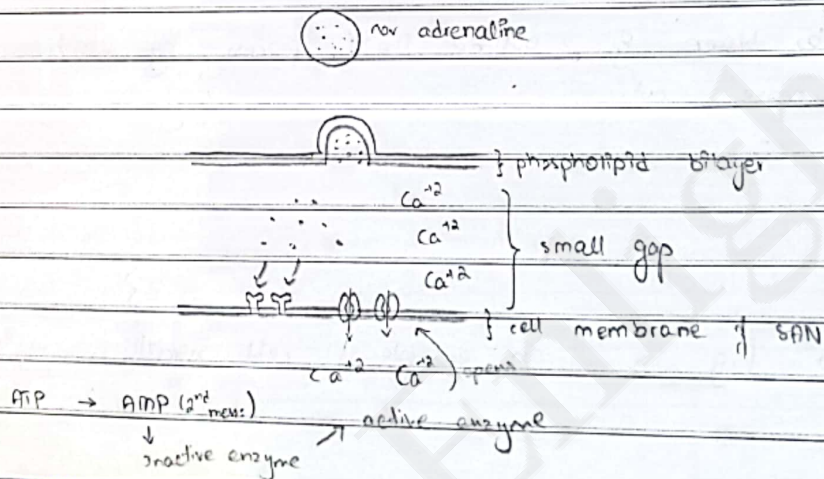
* partitioning of a cell into 2 is cytokinesis.

Q1 Following liver tissue damage chemicals are produced and released into circulation. These chemicals are able to stimulate liver cells to help tissue repair. Explain how this is an example of cell signalling. [37]

signal from blood, plasma. Binds on liver cells (mitotic gene activated (rapid repair is mitotic)). receptor is specific. lock and key series of enzyme controlled reaction. cell divides

lock & key • specificity

Q2 The activity of SAN (Sino adrial node) on heart controlled by nervous system nerve cell in SAN

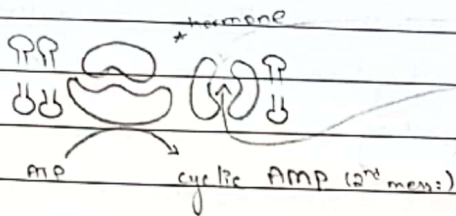


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a) with reference to figure outline the process of cell signalling

- non-adrenergic secreted (exocytosis)
- bind on receptor
- lock & key
- within cell $ATP \rightarrow cAMP$ (2nd mess.)
- calcium channel is opened

a₂)



Inactivation → active enzyme → active enzyme ATGL adipose triglyceride

Lipase

Adipose contains fats

fats → fatty acid + glycerol

a) Name the bond broken by active ATGL to break fats.

ester

b) Name & outline the process by which fatty acids move out.

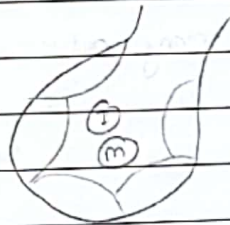
facilitated diff: / active transport
(any one)

c) fig is an example of cell signalling. within the

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adipocyte. with ref: to fig: outline process of cell signalling.

Q) All ^{has fat deposits} adipocyte use fats as source of energy why not RBC.
↓
does not have enzymes, nucleus,



Q) As some alveolar cell produces a surfactant that helps to prevent collapse of alveoli on exhalation. Too much surfactant decreases efficiency of gas exchange. In alveoli a glycoprotein known as GM-CSF is released by some cell of immune system which there is too much surfactant in alveoli. Excess surfactant is then broken down by alveolar macrophage. Receptors GM-CSF are on cell surface membrane of alveolar macrophage. Explain how maintaining correct quantity of surfactant in alveoli is result of cell signalling process.

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