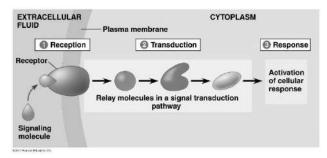
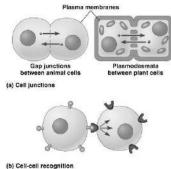
\*\*NOTE: This chapter is often considered difficult as you have not covered it in your general biology course. Plan on reading this chapter slowly!! I would suggest that you read the key concepts in bold first and then for each concept, look at the headings, then the figures and then READ for detail...and STUDY the figures and their captions.

- 1) What is a signal transduction pathway?
- 2) How do yeast cells communicate while mating?



3) How do intercellular connections function in cell to cell communication?

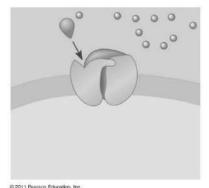


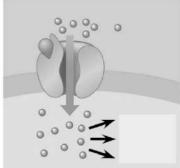
- 4) Explain the two types of local signaling:
  - A) Paracrine signaling
  - B) Synaptic signaling
- 5) How are long distance signals sent?
- 6) Explain Sutherland's investigations with epinephrine and the inferences that were derived from this work.
- 7) Define the three stages of cell communication:
  - A) Reception:
  - B) Transduction:
  - C) Response

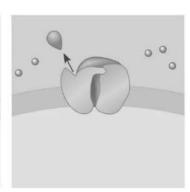
8) What is a LIGAND?	
9) What is special about <b>intracellular</b> receptors – hint think of the relates?	ne structure of the cell membrane and how this
10) Label this diagram of a steroid interacting with an intracellular receptor. (HINT: see fig. 11.9)	
11) Where would you expect most water soluble messengers to bind and why?	
12) What is a <b>G-protein-linked receptor</b> ? (see fig. 11.7)	200
13) (see fig. 11.7, p. 211 captions):	
(overview): A G-protein-coupled receptor is a	receptor that works with the help
of a <b>G protein</b> , a protein that binds the energy-rich molecule	<u> </u>
(1) When GDP is bound to the G protein, the G protein is	The receptor and G protein work
together with another protein, usually an	<u>_</u> :
(2) When the appropriate signaling molecule binds to the extract	ellular side of the receptor, the receptor is activated
and Its cytoplasmic side t	hen binds an inactaive G protein, causing a
to replace This	_ the G protein.
(3) The activated G protein leaves (dissociates from) the receptor	or, diffuses along the membrane, and then binds to
an, altering the enzyme's	Once
activated, the enzyme can trigger the next step, leading to a	
(4) The changes in the enzyme and G protein are only temporar	ry because the G pr otein also functions as a
enzyme – in other words, it then hydrol	lyzes its bound GTP to GDP. Now inactive again,
the G protein the enzyme, which returns to it	ts original state. The GTPase function of the G
protein allows the pathway to	rapidly when the
is no longer present.	

15) <b>(see fig. 11.7, p. 21</b>	2 captions):	
(overview): Receptor ty	rosine kinases belong to a major class of p	lasma membrane receptors characterized by
having	eceptor protein extending into the cytoplasm	
functions as a tyrosine I	fer of from ATP to the	
amino acid	on a substrate protein. One rece	ptor tyrosine kinase comlex may activate
differen	nt transduction pathways and cellular respo	onses. The ability of a single ligand-binding even
		and
(1) Before the signaling		referred to as monomers
Each monomer has an	extracellular	site, an $\alpha$ helix spanning the membrane,
	containing	
		(such as growth factor) causes 2 receptor
		, forming a complex known as a dimer
(dimerization).		
(3) Dimerization activate	es ther	egion of each monomer; each tyrosine kinase
	from an ATP molecule to a tyrosir	ne on the tail of the other monomer.
(4) Now that the receptor	or is,	it is recognized by specific relay proteins inside
		sine, undergoing a resulting structural change
that activates the bound	d protein. Each activated protein triggers a	,leading
to a		
16) <b>(see fig. 11.7, p. 21</b>	3 captions):	
(overview): What trigger	rs a ligand-gated ion channel to open/close	?
What then passes throu	ugh the channel once it is open?	
**study and read the ca	ptions for parts 1-3 of this diagram!	
(conclusion): How do no	erve cells make use of ligand-gated ion cha	nnels?
How is a voltage-gated	ion channel different?	
(label the diagram on th	ne next pagesee fig. 11.7,page 213)	

## **DIAGRAM: Ligand-gated ion channel**



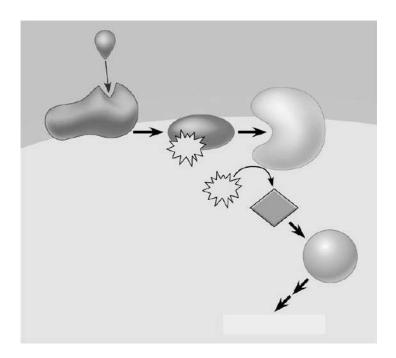




17) (see fig. 11.10, p. 215 captions):

, (555g, p. = 15 5p		
	a phosphorylation cascade:	
(2) Active protein kinase 1 transfers		to an inactive molecule of
protein kinase 2, thus	this 2 <sup>nd</sup> kianse.	
(3) Active protein kinase 2 then catalyze	es the phosphorylation (&	) of protein kinase 3.
(4) Finally, active protein kinase 3 phosp	phorylates a protein that brings about the	e cell's response to the signal.
(5) Enzymes called protein phosphatase	es catalyze the	
from the proteins, making them	&	for reuse.
18) What are <b>protein phosphatases</b> are	nd why are they so important?	
19) What are <b>second messengers</b> and characteristics significant)?	I what are two characteristics of a second	d messenger (& why are these
20) What did Sutherland find in his expe	eriments with regard to cyclic AMP and w	why is this important?
21) What is adenylyl cyclase?		

22) Complete the diagram here of cAMP as second messenger: (see fig. 11.12)

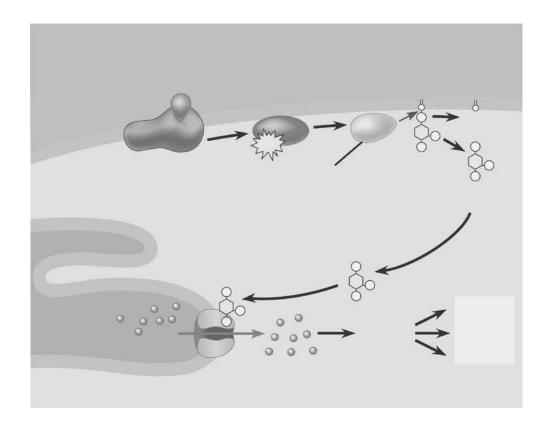


23) How does the cholera bacterium (& how this microbe causes disease) connect with the concepts of cell to cell communication?

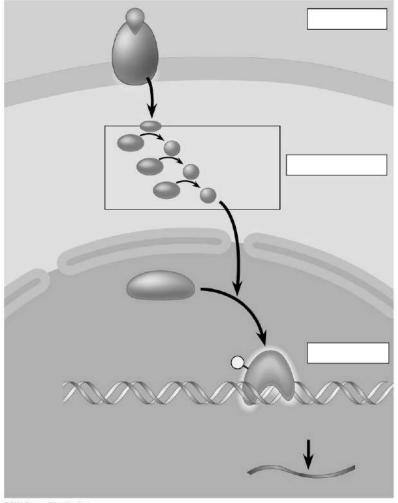
24) How does the drug known as Viagra work? Why was it originally prescribed for chest pain?

25) How and why are the calcium concentrations kept different and separate comparing the endoplasmic reticulum, mitochondria and cytoplasm?

26) Label the diagram below showing calcium and IP<sub>3</sub> in a cell. (see fig. 11.14)



27) Label the diagram below showing nuclear responses to a signal. (see fig. 11.15)

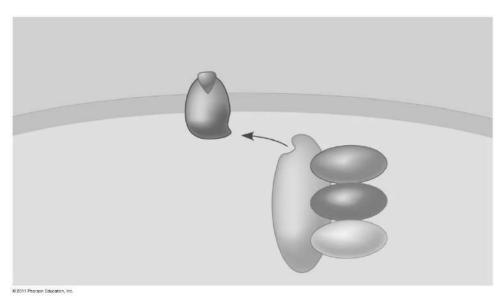


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28) How is <b>signal amplification</b> accomplished in the cell?	
29) How is <b>specificity</b> accomplished in cell signaling?	

30) What is a **scaffolding protein** and why is it important?

31) Label the diagram of a scaffolding protein shown here. (see fig. 11.19)



32) How is the termination of a signal accomplished and why is it so important that termination be accomplished?