

---

---

# COGS 17 Week 3

— SPRING 2024, A03 —

---

---

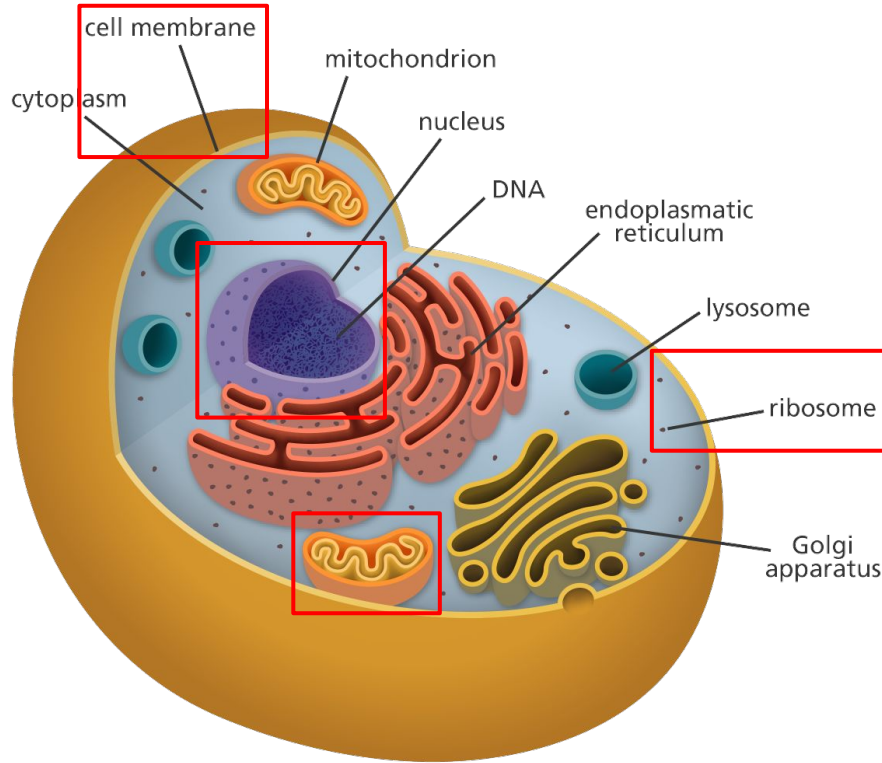
# Problem Set for Today

- Link:

<https://docs.google.com/document/d/1HYoP3TxzcKxuYK0lqclrwgEDAaeffJb91cn3GmwXvQ4/edit?usp=sharing>

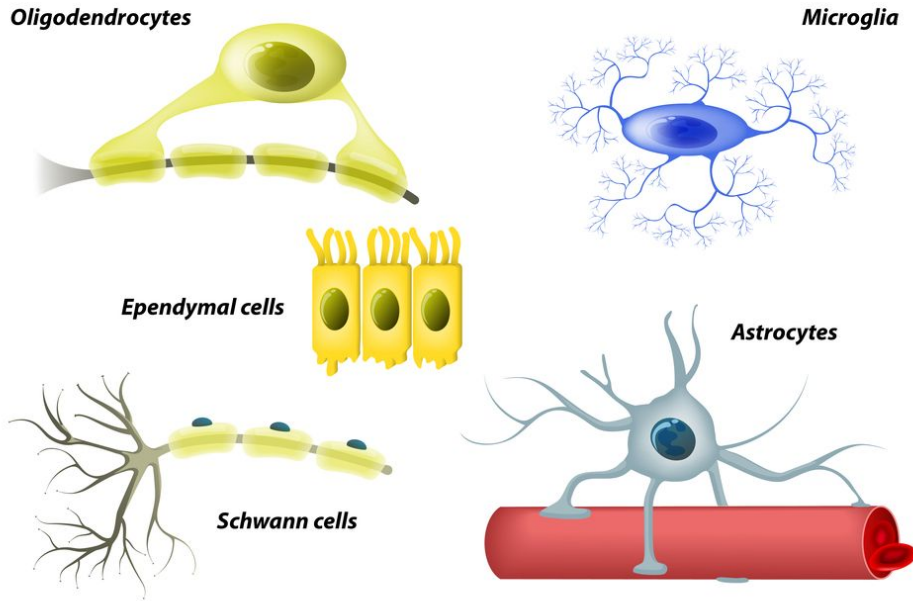


# Cells



- Nucleus -- where DNA stored
- Ribosome -- where protein synthesis occurs
- Mitochondria -- Produce ATP, the “powerhouse” of the cell
- Cell membrane -- lipid, **BILAYER**, semi permeable membrane

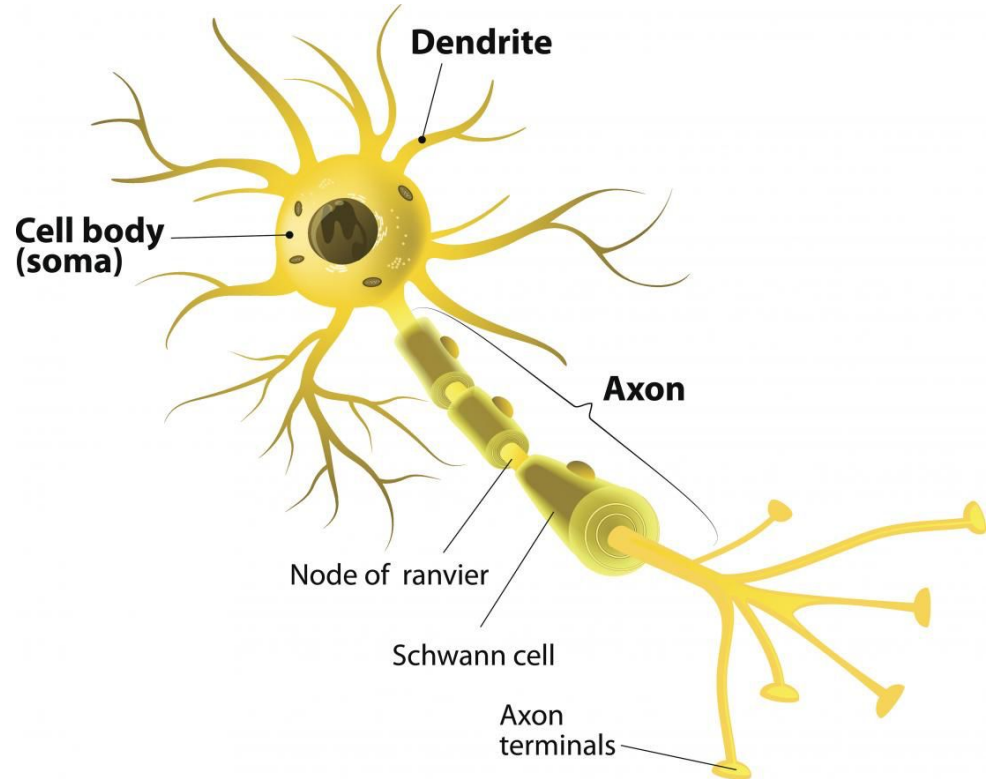
# Glial Cells



- Non-neural cells of the nervous system with multiple functions
- Do **NOT** participate in information transfer
- Much smaller than neurons, but much more numerous
- ~50% of brain by weight

# Neurons

- Cells that are specialized for **INFORMATION TRANSFER**
- Modified processes (i.e. dendrites & axon)
- Dendrites -- reception of **INCOMING** message via receptors
- Axon -- Site of release of **OUTGOING** message
- Modified membrane -- ion channels

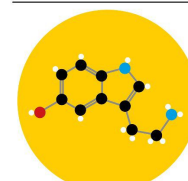
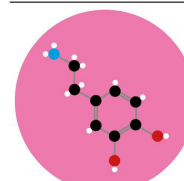
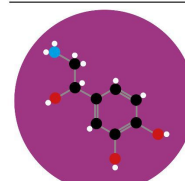
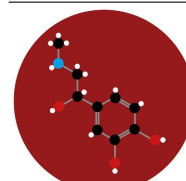


# Neurotransmitters (NT)

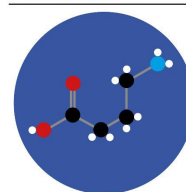
- Endogenous chemicals that allow neurons to **communicate** with each other throughout the body
- E.g. Acetylcholine (ACh), GABA, Glutamate, Serotonin (5-HT), Norepinephrine, etc.
- May have different effects in different parts of body

## CHEMICAL STRUCTURES OF NEUROTRANSMITTERS

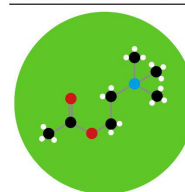
ADRENALINE  $C_9H_{13}NO_3$  THE FIGHT OR FLIGHT NEUROTRANSMITTER  
 NORADRENALINE  $C_8H_{11}NO_3$  THE CONCENTRATION NEUROTRANSMITTER  
 DOPAMINE  $C_8H_{11}NO_2$  THE PLEASURE NEUROTRANSMITTER  
 SEROTONIN  $C_{10}H_{12}N_2O$  THE MOOD NEUROTRANSMITTER



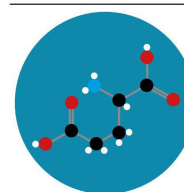
$\gamma$ -AMINOBUTYRIC ACID  $C_4H_9NO_2$   
THE CALMING NEUROTRANSMITTER



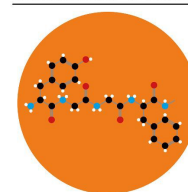
ACETYLCHOLINE  $C_7H_{16}NO_2^+$   
THE MOVING NEUROTRANSMITTER



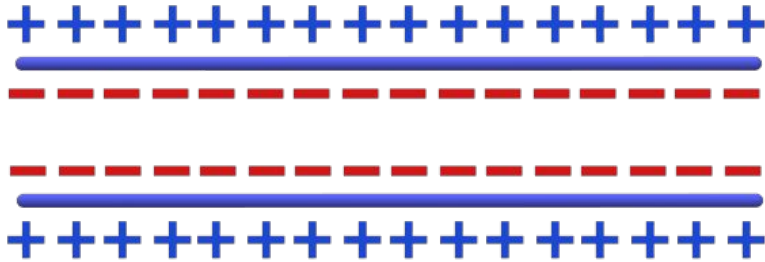
GLUTAMATE  $C_5H_9NO_4$   
THE REMOXY NEUROTRANSMITTER



ENDORPHINS 20+ TYPES IN THE HUMAN BODY  
THE PAINKILLER NEUROTRANSMITTERS



# Resting Potential

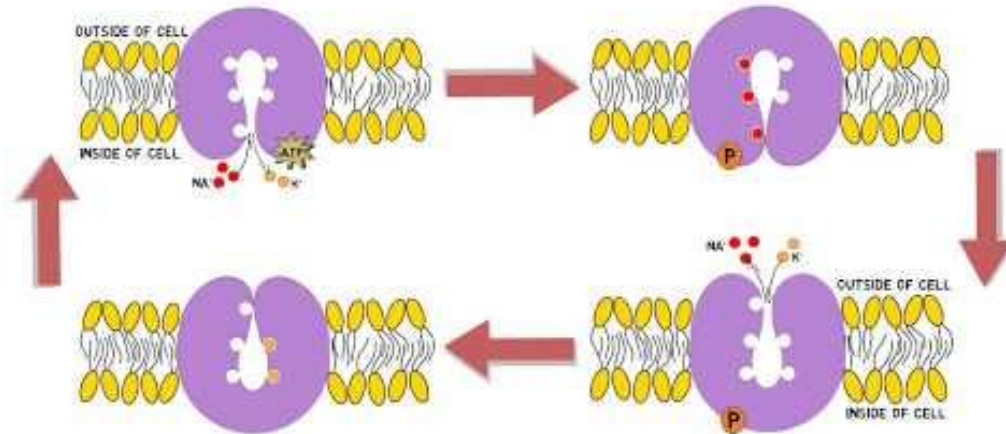


- Generally -70 mV
- More negative inside & more positive outside
- Established in part by Na-K pump, which actively transport 3 Na<sup>+</sup> out and 2 K<sup>+</sup> in
- Highly **POLARIZED** -- ready to “fire”

# Sodium-Potassium Pump

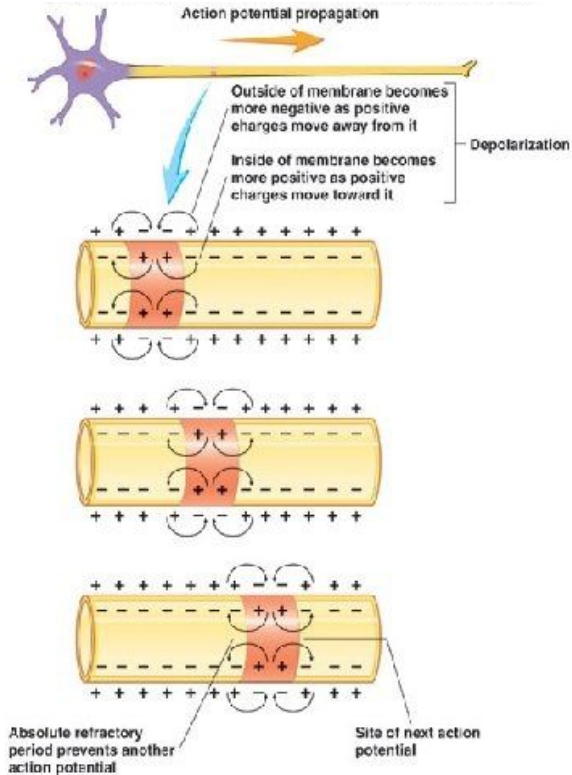
## SODIUM-POTASSIUM PUMP CYCLE

-USES ATP TO TRANSPORT SODIUM AND POTASSIUM IONS ACROSS THE CELL MEMBRANE



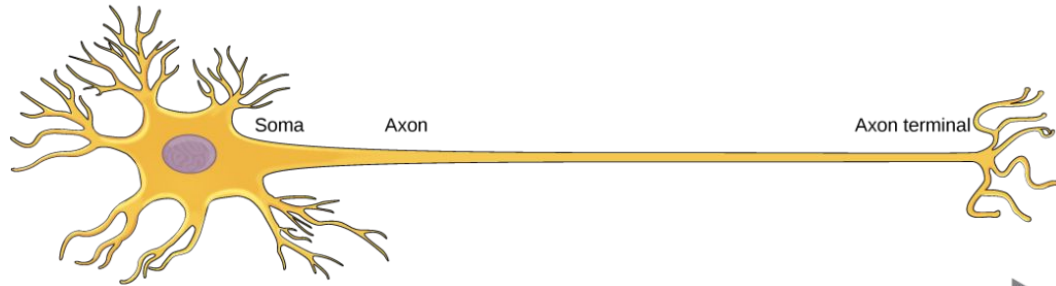


# Action Potential

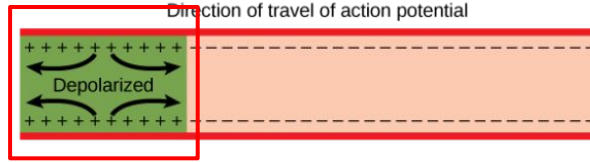


- **DEPOLARIZATION** of neuron
- Triggered by neurotransmitters from other neuron, electrical stimulation, or other stimuli
- Starting at Axon Hillock

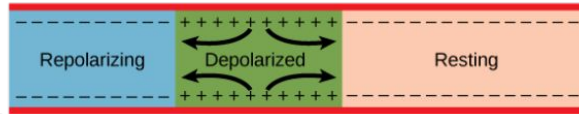
# Action Potential



a. In response to a signal, the soma end of the axon becomes depolarized.



b. The depolarization spreads down the axon. Meanwhile, the first part of the membrane repolarizes. Because  $\text{Na}^+$  channels are inactivated and additional  $\text{K}^+$  channels have opened, the membrane cannot depolarize again.

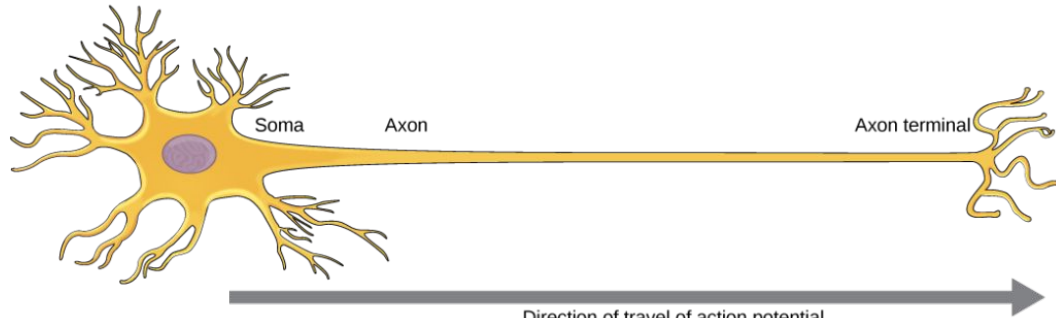


c. The action potential continues to travel down the axon.



- Voltage-activated  $\text{Na}^+$  gates open --  $\text{Na}^+$  in
- Reverse local polarization to  $\sim +50 \text{ mV}$
- $\text{Na}^+$  influx causes adjacent  $\text{Na}^+$  voltage-activated gates to open & previously opened gates close
- Propagation of the action potential to the terminal

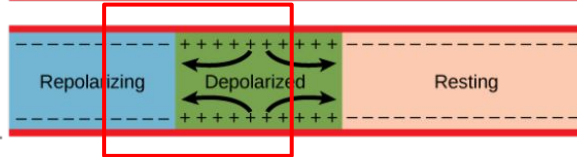
# Action Potential



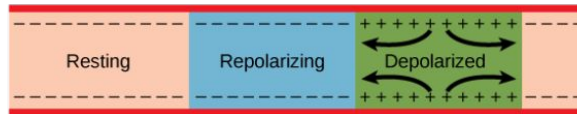
a. In response to a signal, the soma end of the axon becomes depolarized.



b. The depolarization spreads down the axon. Meanwhile, the first part of the membrane repolarizes. Because  $\text{Na}^+$  channels are inactivated and additional  $\text{K}^+$  channels have opened, the membrane cannot depolarize again.

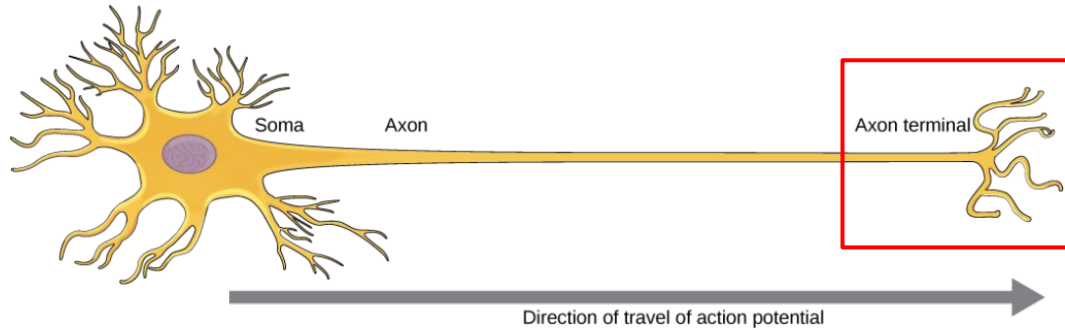


c. The action potential continues to travel down the axon.



- As previous  $\text{Na}^+$  gates close, local  $\text{K}^+$  gates open --  $\text{K}^+$  leaves the cell
- Because of opened  $\text{K}^+$  channels & closed  $\text{Na}^+$  channels, the membrane **cannot** depolarize again
- Refractory period -- **prevents backflow** of action potential

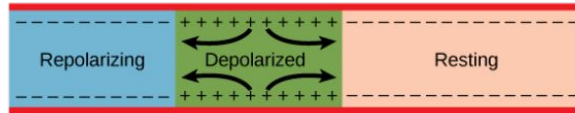
# Action Potential



a. In response to a signal, the soma end of the axon becomes depolarized.



b. The depolarization spreads down the axon. Meanwhile, the first part of the membrane repolarizes. Because  $\text{Na}^+$  channels are inactivated and additional  $\text{K}^+$  channels have opened, the membrane cannot depolarize again.

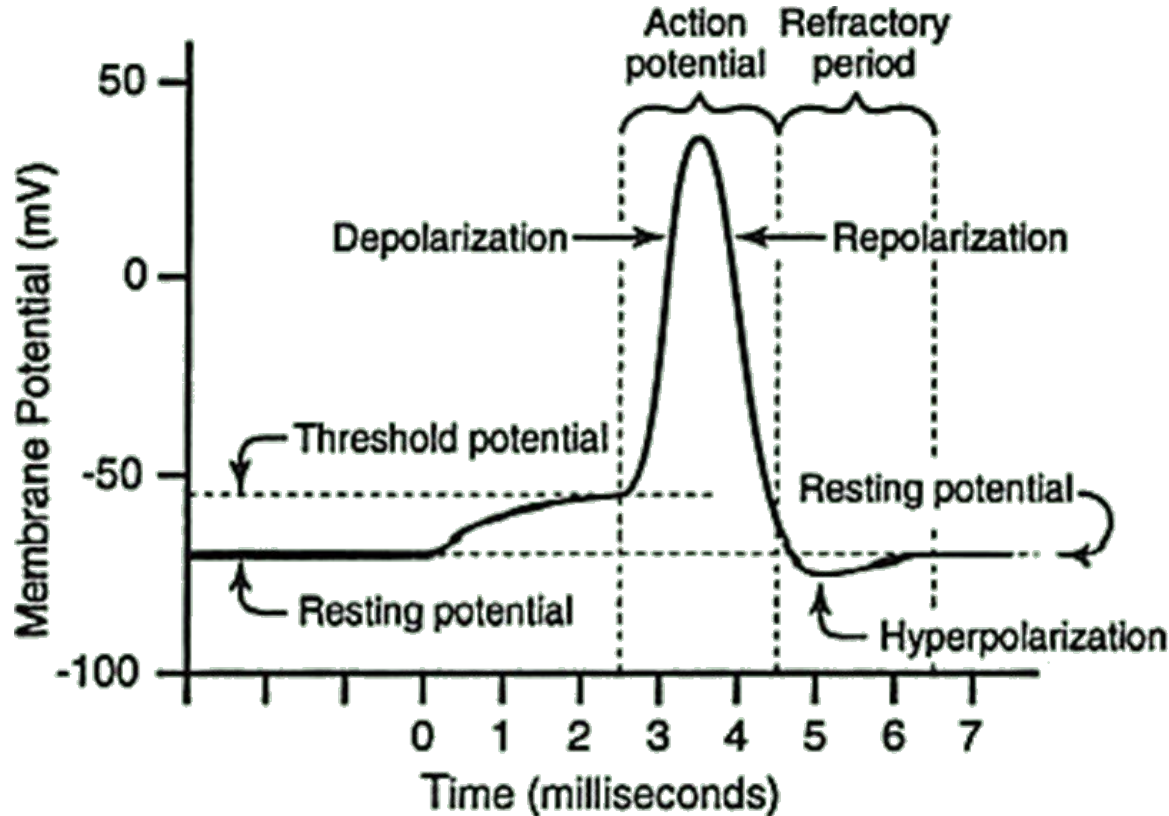


c. The action potential continues to travel down the axon.



- When reaches terminal,  $\text{Ca}^{2+}$  enters cell & Neurotransmitter released
- When outside becomes more positive again,  $\text{K}^+$  channels close
- Na-K pumps restore resting potential to -70 mV
- Ca pump reject  $\text{Ca}^{2+}$  from terminal
- Pumps **require energy**

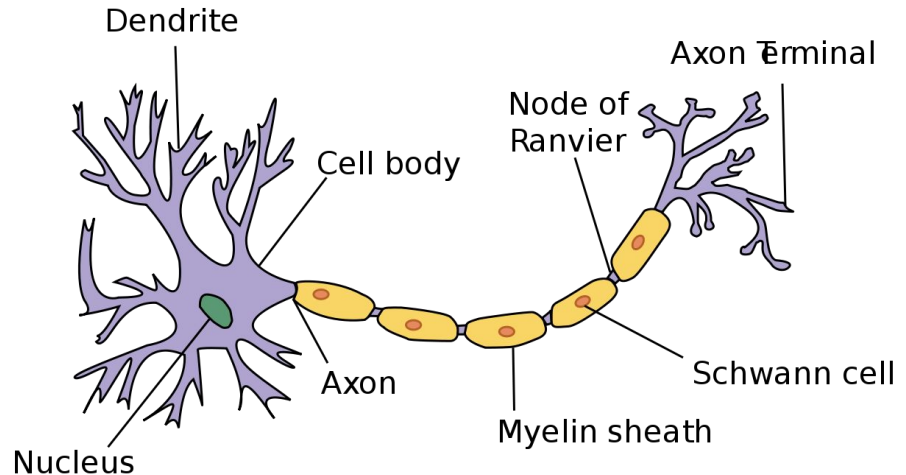
# All-or-none Law



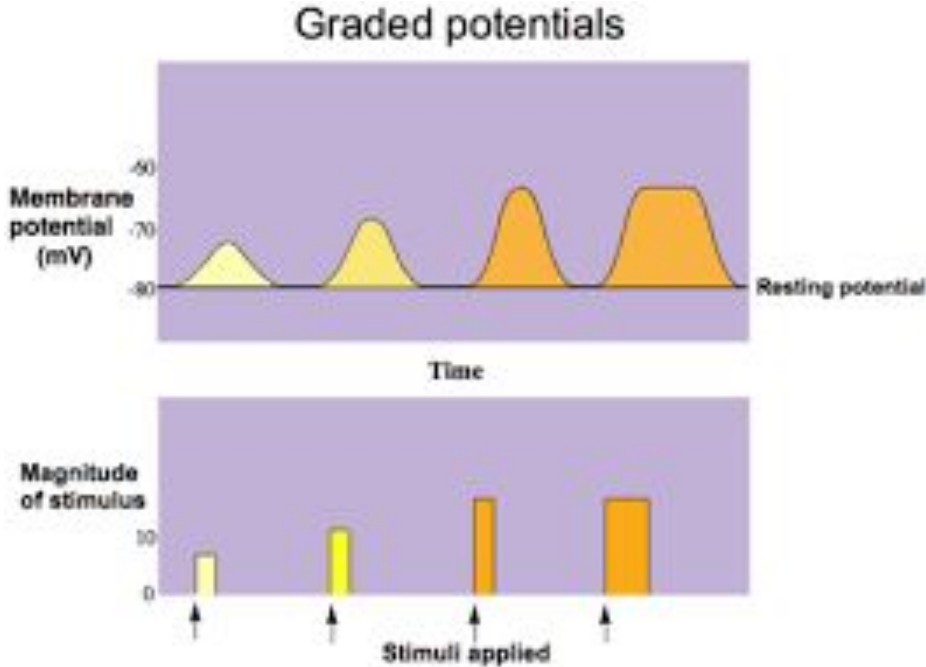
- An Action Potential always has the same amplitude and velocity, **regardless** of the intensity of the stimulus that triggered it
- Either full response or no response at all

# Myelination and Saltatory Conduction

- Increase the speed of propagation
- Glial cells form **insulating sheaths** around axons, with small gaps in between sheaths
- **Electrical** conduction -- fast, but degrades as it moves
- Node of Ranvier -- **ionic** conduction, **reboost** the signal to original strength and pass it to next sheath
- Saltatory Conduction -- **"Jumping"** of the signal from node to node along axon

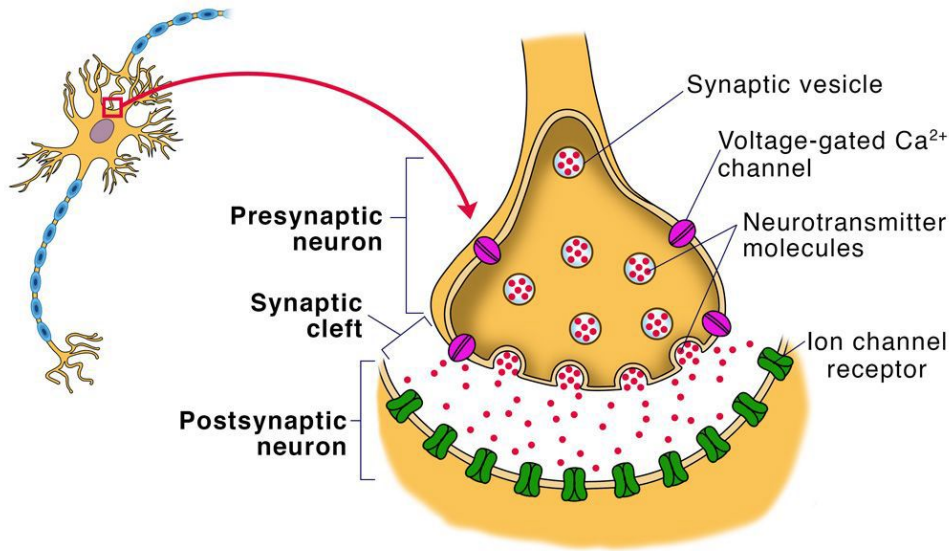


# Graded potentials



- Releasing NT from a Neuron does **NOT** always require an Action Potential
- Can vary in amplitude
- i.e. Strong response due to strong stimulus; weak response due to weak stimulus
- Examples: some receptor cells (e.g. retina), lateral inhibitors, local Neurons, etc.

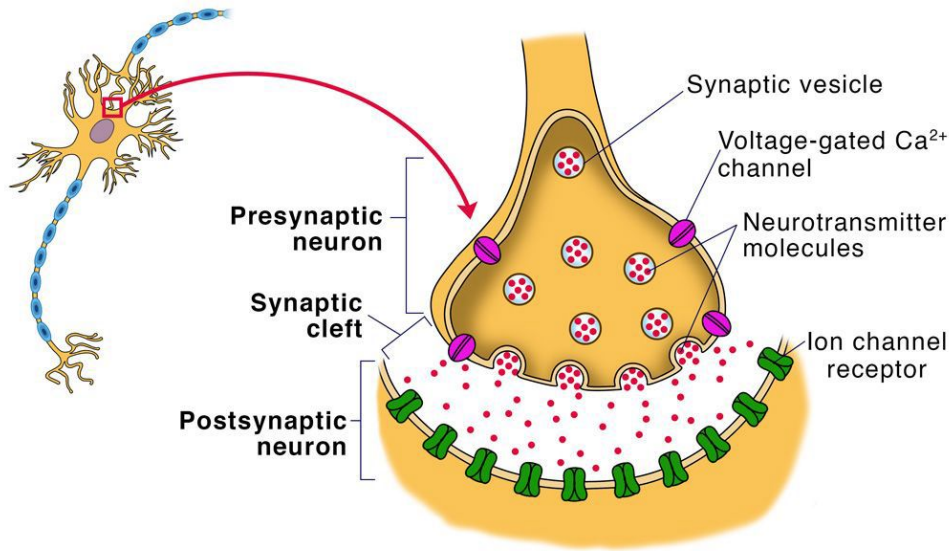
# The synapse



- Presynaptic cell releases NT into synaptic cleft, affects postsynaptic cell via **Exocytosis** (release of vesicles of NT).
- When membrane depolarization reaches the axon terminal, voltage-gated  $\text{Ca}^{++}$  channels open, causes influx of  $\text{Ca}^{++}$

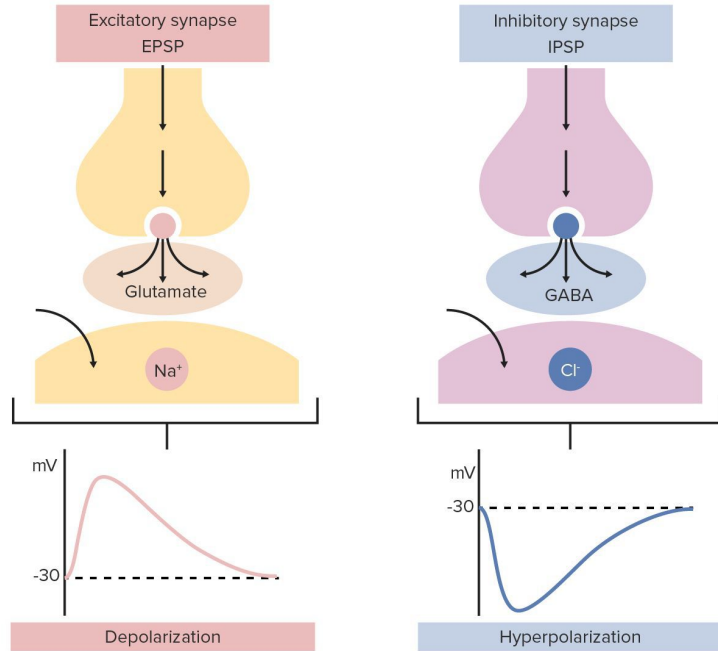


# The synapse



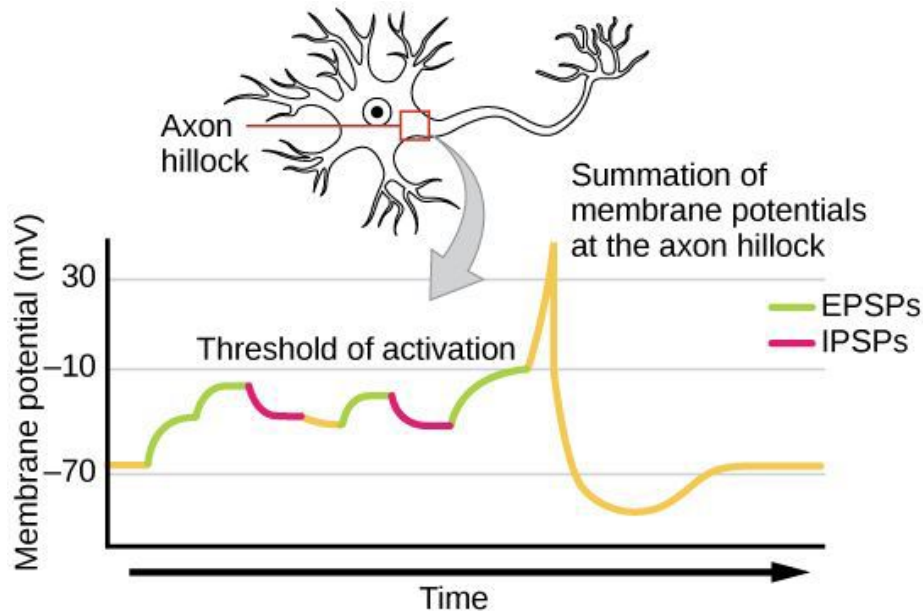
- NT **passively** diffuses across synaptic cleft
- NT binds to **NT-specific** receptor site on (usually) dendrites of postsynaptic cell
- Postsynaptic cell may fire its NT, etc.
- NT soon **detaches** from receptor
- NT is often then **deactivated** by enzymes or Glia cells

# EPSP VS. IPSP



- EPSP: **Excitatory** Postsynaptic Potential - postsynaptic cells become hypopolarized.
- IPSP: **Inhibitory** Postsynaptic Potential - postsynaptic cells become hyperpolarized.

# Summation

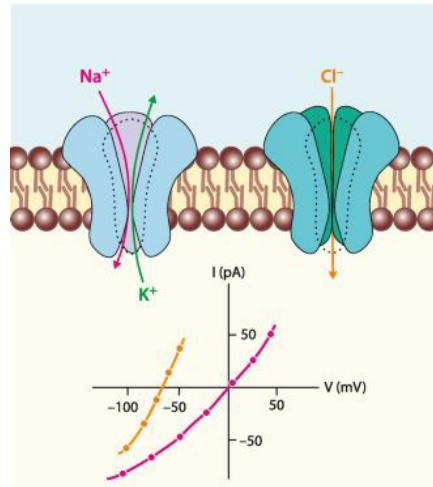


- The response of a given neuron is nearly always the product of a **summation of EPSPs and/or IPSPs**
- Two types of summation: **Spatial** summation and **Temporal** summation

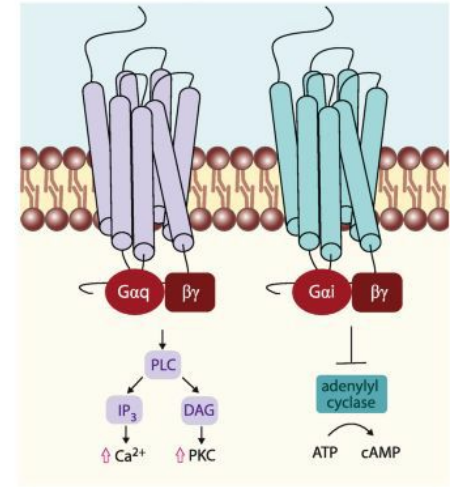
# Ionotropic Vs. Metabotropic

- Ionotropic: NT **directly** affects ion gates
  - Effects are **rapid, short-lived**
- Metabotropic: Triggers **metabolic changes** in Postsynaptic cell
  - Activates/triggers production of **Secondary Messenger**, which binds w/**G-Protein** to open separate ion gate, **requires energy, slower and long-lasting**

a ionotropic receptors

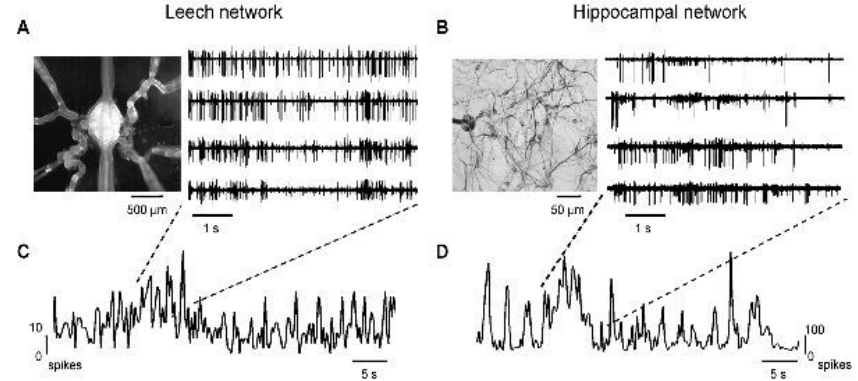


b metabotropic receptors



# Spontaneous Activity

- Some Neurons show spontaneous Firing, **in absence of incoming NT**
- Usually graded potentials
- Converging NTs increase or decrease such cells' activity, modifying how much NT they will then release



# Agonism Vs. Antagonism

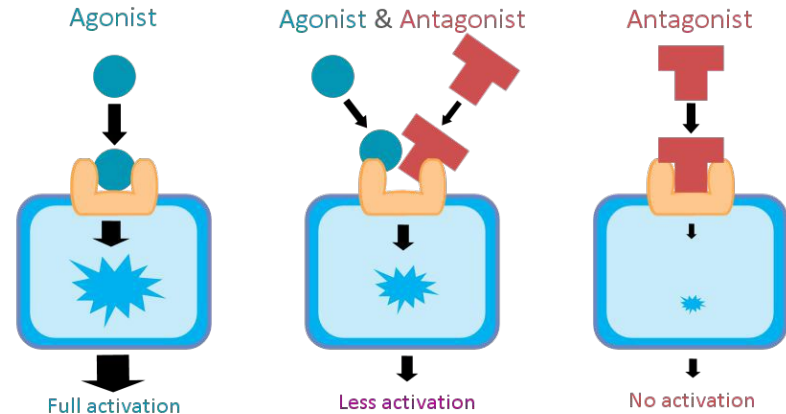
- Agonist: chemicals that **increase** the effect of a NT
- Antagonist: chemicals that **decrease** the effect of a NT

## Agonists and Antagonists

Agonists - Drugs that occupy receptors and activate them.

Antagonists - Drugs that occupy receptors but do not activate them.

Antagonists block receptor activation by agonists.



# Questions?

Office Hours: Mon 5-6 pm

To get the section slides: <https://github.com/JasonC1217/COGS17-A03-Sp24>

OR:

