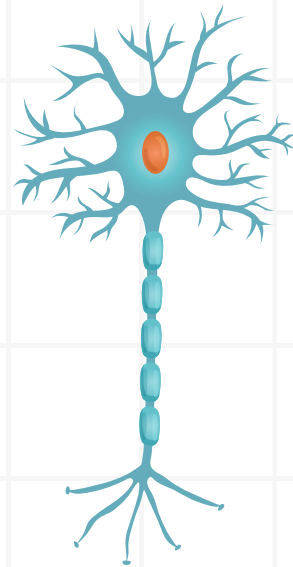


PHYSIOLOGY

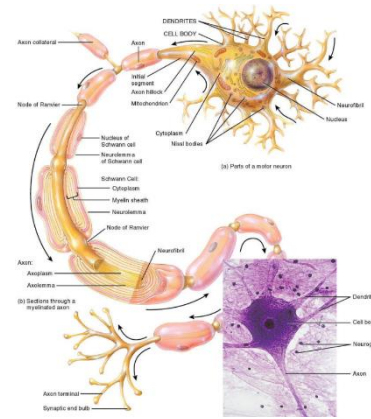
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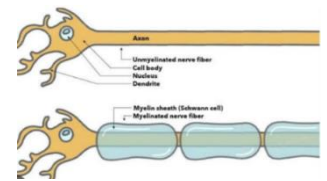
DR. AHMAD AL-QAWASMI

❖ Neural Structure

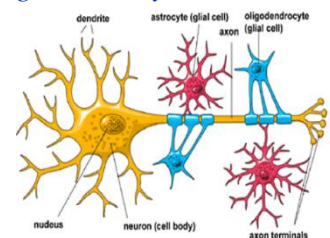
- The nervous system is formed of neurons and supportive cells
- The main parts of the Neuron (Neural cell):
 - **Cell body (Soma)** contains most organelles such as nucleus, ER, mitochondria, Nissl bodies (site of protein synthesis)
 - ✓ Neurons can't divide or regenerate so they *lack centriole*
 - **Dendrites** which are short projections from the cell body that *receive inputs* (collect signals) from neighboring neurons
 - ✓ Dendrites *can't generate* action potential, because they have a *very high threshold* due to the *lack of voltage gated Na⁺ channels* in addition to their *small diameter (high resistance)*



- **Axon (Nerve fiber)**
 - ✓ A long tubular like structure which projects from the cell body
 - ✓ Axons can be either myelinated or un-myelinated
 - **Myelin sheath** is composed mainly of glycolipids, and it appears white
 - **Nodes of Ranvier: gaps** in the myelin sheath, appear at intervals along the axon
 - ✓ At the end of the axon there are fine processes called **Axon terminals**
 - ✓ Some of these terminals end with bulb-shaped structures called **synaptic end bulb (Knob)**
 - At these end the neurotransmitters are stored in vesicles and ready for the release
- **Axon hillock (trigger zone)**
 - ✓ It is the junction between the cell body and axon, where **action potential is generated** because it has the *largest number* (density) of voltage gated sodium channels & the *lowest threshold*



- **Supportive cells (neuroglia)** include *microglia*, *astrocytes*, *Schwan* cells and *oligodendrocytes*
 - **Astrocytes** form the **blood brain barrier (BBB)** between the blood and the cerebrospinal fluid (in the brain)



- The functions of supportive cells:
 - **Maintenance of neural environment**, by the uptake of K⁺ and neurotransmitters from the interstitial fluid around the neurons
 - **Synthesize and release neurotrophic factors** to maintain the survival and protection of neurons
 - Other specialized supportive cells are responsible for **myelination of axons**
 - ✓ In the **CNS**, *oligodendroglia* are the responsible for the myelination process
 - ✓ In the **peripheral** nervous system, *Schwan cells* are the responsible for the myelination process

- Action potential propagation (conduction) along the axon (nerve fiber) occurs by:

1- Continuous conduction (Slow)

- Occurs in **unmyelinated** fibers
 - Local ionic currents flow between the adjacent regions (from + to -), causing activation of Na⁺ channels, which can bring potential to threshold and causing action potential generation in this region
 - This process is repeated **all along the nerve fiber** until the impulse has reached nerve terminals

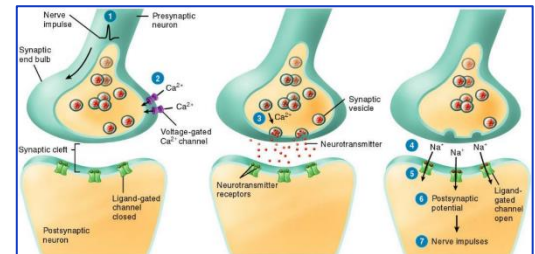
2- Salutory conduction (Fast)

- Occurs in **myelinated** fibers

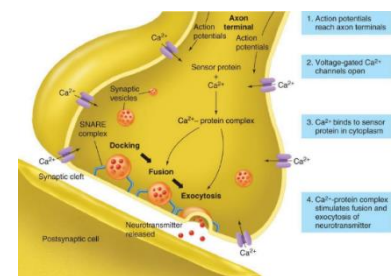
- The ionic current flow between two adjacent ***nodes of Ranvier***
- The impulse skips the myelinated regions in the axon and ***jumps*** from one node of Ranvier to the other
- It is 50 times faster than in unmyelinated fibers of the same size
- Velocity of action potential conduction depends on ***myelination*** and the ***diameter of nerve fibers***
- Larger fiber has lower resistance → higher velocity (faster)

❖ Transmission of action potential between neurons

- **Synapse:** The connection between a neuron and another cell
 - **Presynaptic membrane:** The membrane of the synaptic bulbs of the axon terminal of the first neuron
 - **Postsynaptic (subsynaptic) membrane:** The membrane of the dendrites & cell body of the second neuron
 - **Synaptic cleft:** The gap between the 2 cells



- When the impulse from the presynaptic neuron reaches the synaptic knob, it causes **activation** of ***voltage dependent Ca⁺⁺ channels***, causing diffuse of Ca⁺² into the synaptic knob, increasing its concentration inside the terminals which triggers the ***fusion of the neurotransmitter vesicles*** with the presynaptic membrane, ***releasing*** NTs into the cleft by ***exocytosis***
 - Ca⁺² triggers exocytosis by ***reducing (decreasing) the repulsion*** between the membranes of the vesicles and the plasma membrane



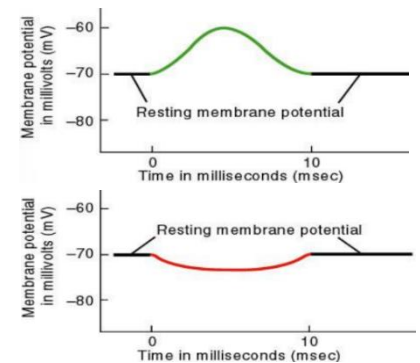
- **Bind of NTs to their receptors** on the postsynaptic membrane, inducing ***changes in the membrane potential*** (activity of ion channels), which can be excitatory or inhibitory

➤ EPSPs (*Excitatory Post Synaptic Potentials*)

- ✓ It ***decreases*** the membrane potential (become less negative, small depolarization), due to the activation of a few Na⁺ channels
- ✓ A single EPSP can ***not reach threshold***

➤ IPSPs (*Inhibitory Post Synaptic Potentials*)

- ✓ It ***increases*** the membrane potential (become more negative, hyperpolarization), due to the activation of ***K⁺ channels***
- ✓ Activating ***Cl⁻ channels*** is considered as inhibitory but it doesn't induce hyperpolarization where this inhibition is achieved by holding the membrane at its resting potential and preventing depolarization



- IPSP & EPSP are **graded potentials**

➤ The channels on the postsynaptic membrane are **chemical** gated channels

- **Acetylcholine** is a neurotransmitter causes **EPSP** by the activation of chemical gated Na⁺ channels
 - ✓ 2 molecules of Ach are required to bind & activate Na⁺ channels

- **Synaptic delay:** The period required to induce changes in membrane potential in the postsynaptic neuron

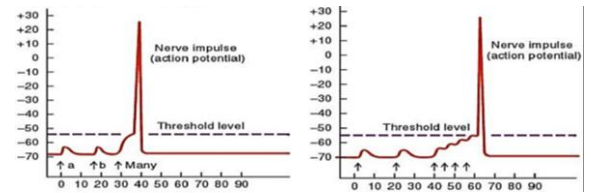
- After inducing the appropriate response at the postsynaptic membrane, the transmitter is ***inactivated*** (destroyed) or ***removed***, allowing the postsynaptic membrane to receive additional messages

➤ **Acetylcholine esterase** an enzyme that destroys acetylcholine (Ach) into acetyl and choline, which then transported back to the presynaptic knob, where they combine again to form new Ach

- **Blockers** are drugs can combine with receptor and **prevent** binding of transmitter to its receptor

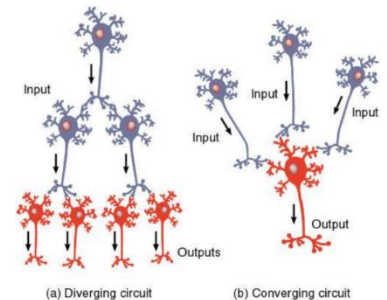
➤ **Hexamethonium** blocks the binding of acetylcholine (Ach) to its receptor

- **Summation:** is the addition and combination of many signals, producing a stronger signal which can reach threshold (in EPSPs) or cause more inhibition (in IPSPs)
- **Spatial summation:** 2 or more responses from *2 or more different presynaptic neurons* have appeared *simultaneously (at the same time)* at the same site of postsynaptic membrane
- **Temporal summation:** 2 or more postsynaptic potentials, which were elicited by *one presynaptic neuron (single input)* at *different times (repetitive excitation)*
- Summation can occur between EPSPs or IPSPs or between both (results in **cancellation**)



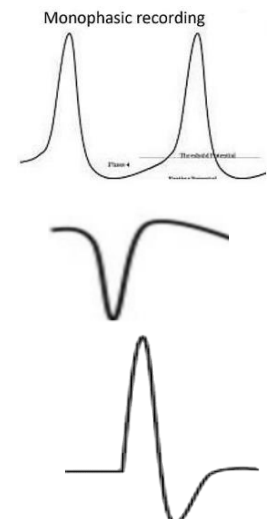
❖ Synaptic organizations

- Neurons are connected together forming neural networks, including:
 - **Convergence:** synapse of many axonal terminals from different neurons to one neural cell body
 - ✓ Many inputs (presynaptic) with 1 output (postsynaptic)
 - **Divergence:** It is the branching of 1 fiber (axon) to many terminals
 - ✓ One input (presynaptic) with many outputs (postsynaptic)



❖ Action Potential Recording

- There are 2 methods of recording action potential:
 - 1) **Monophasic action potential:**
 - We place one electrode **outside** and one electrode **inside** the cell
 - The recording would be either positive or negative but **not both**
 - It records potential in one region (one phase) [depolarization +, repolarization -]
 - 2) **Biphasic action potential:**
 - We place **both** electrodes **outside** but in 2 different regions
 - It records potential in 2 regions (2 different phases)



❖ Compound action potentials

- It is the sum of **all** recorded action potentials generated by all nerve fibers at a certain point on the nerve
 - It is done by placing one electrode at a source of zero voltage (high resistance source) and the other one at a point on the nerve, by sensing the voltage difference
 - It is used to check the **integrity** of the fibers
- **Aα** Fibers is the **fastest** in the conduction (transmission) of action potential
- **C** fibers are the **slowest**

