

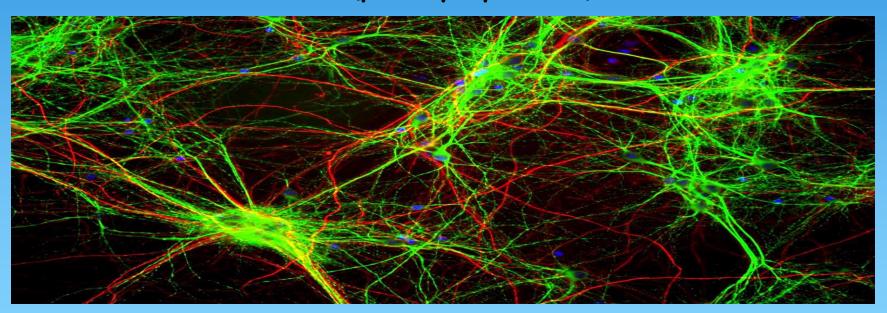
## Synapes and Synaptic Transmission

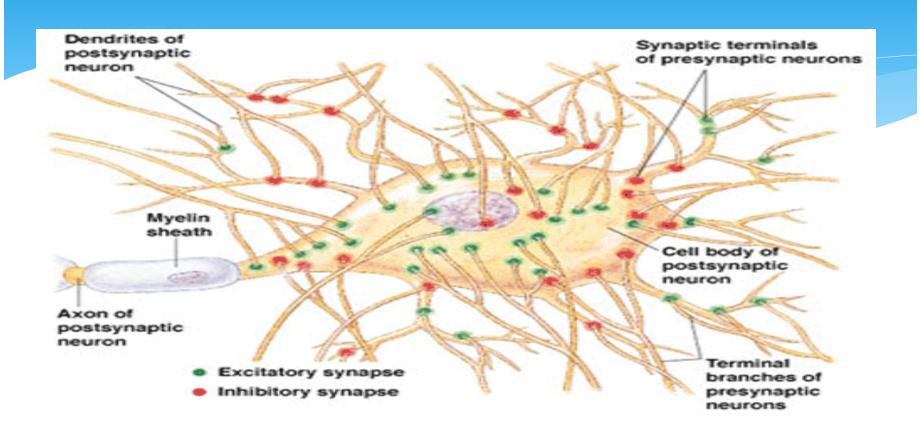
# Objectives

- Define synapses
- \* Functions of synapses.
- \* Structure of synapses
- \* Types of synapses: anatomical & functional.
- \* Synaptic transmission & neurotransmitters
- \* Fate of neurotransmitters.
- \* Electrical events at synapses (EPSPs & IPSPs).
- \* Properties of synaptic transmission
- \* Factors affecting synaptic transmission

# The CNS contains more than 100 billion neuron Synapse:

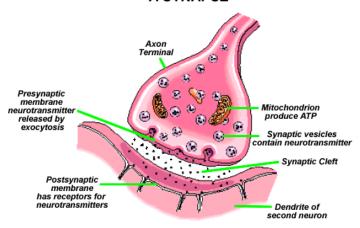
A junction where the axon or some other portion of one cell (= presynaptic cell) terminates on the dendrites, soma, or axon of another neuron (post synaptic cell).





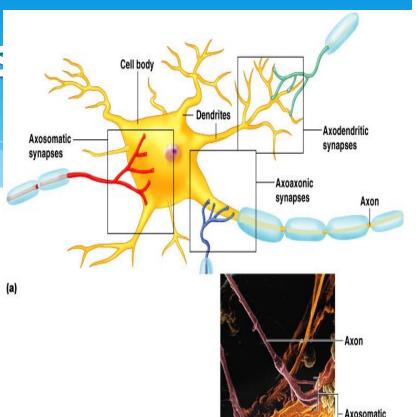
## **Functional Anatomy of a Synapse**

#### A SYNAPSE



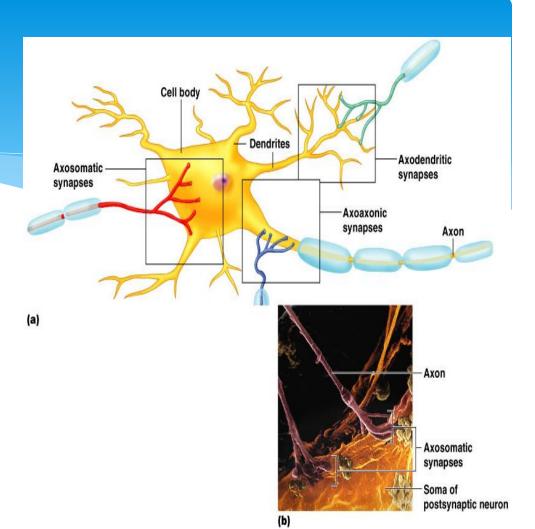
# Anatomical Types

- \* Axodendritic synapses between the axon of one neuron and the dendrite of another
- \* Axosomatic synapses between the axon of one neuron and the soma of another

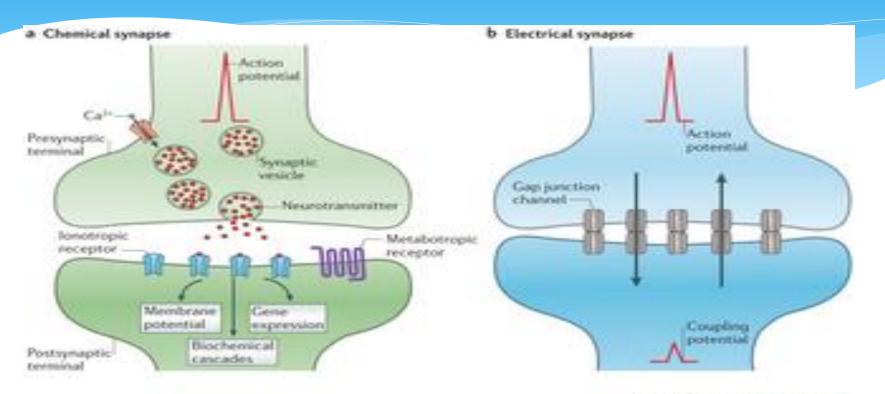


synapses

- Other types of synapses include:
- \* Axoaxonic (axon to axon)
- \* Dendrodendritic (dendrite to dendrite)
- Dendrosomatic(dendrites to soma)



# Functional types of synapses

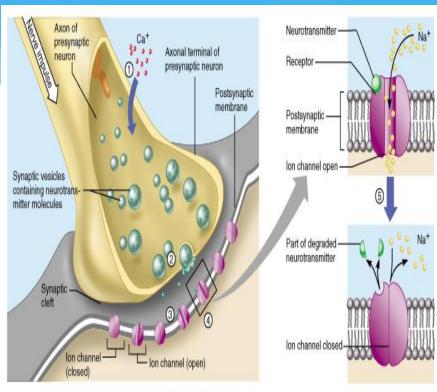


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## Functional types of synapses

#### \* A. Chemical synapse

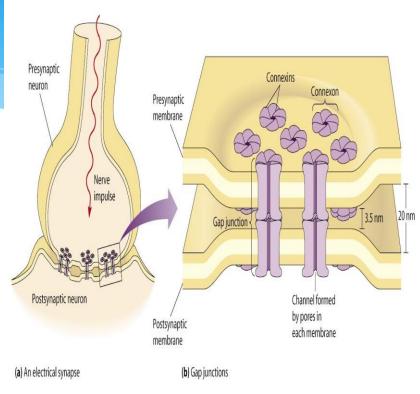
- \* Almost all synapses in the CNS
- \* ((i.e. first neuron secretes a chemical substance called neurotransmitter at the synapse to act on receptor on the next neuron to excite it, inhibit or modify its sensitivity))



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## B. Electrical Synapses

Membranes of the pre- and post-synaptic neurons come close together and gap junctions forms → no membrane borders which allow passage of ions.



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- Are less common than chemical synapses
- Correspond to gap junctions found in other cell types
- \* Are important in the CNS in:
  - \* Mental attention
  - \* Emotions and memory
  - \* Arousal from sleep

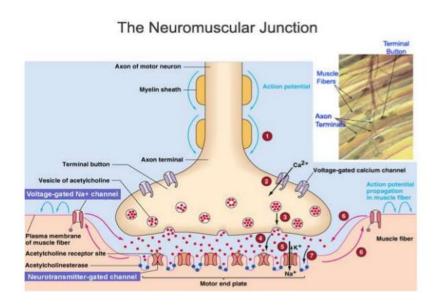
#### C. Conjoint synapse

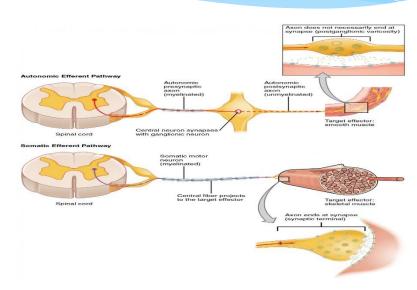
Both electrical and chemical.

Examples  $\rightarrow$  neurons in lateral vestibular nucleus.

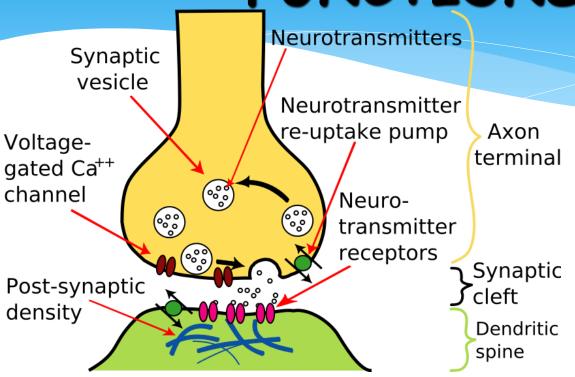
#### \* Examples of synapses outside CNS

- Neuromuscular junction
- Contact between: autonomic neurons and smooth, cardiac muscles, and other effector





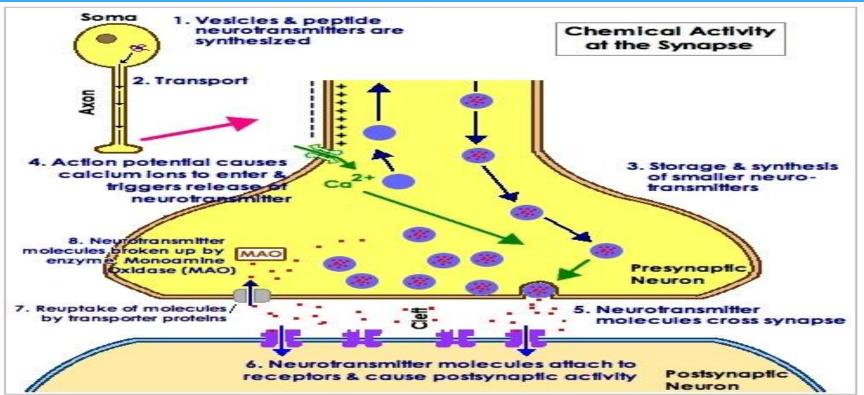
# SYNAPSE: STRUCTURE & FUNCTIONS



- \* Synaptic cleft: This the space between the axon terminal and sarcolemma. It has a width of 200-300 angstroms.
- \* Synaptic knobs (presynaptic terminal)



## Mechanism of Synaptic Transmission:



- AP >>>> NT relseas >>>> Binding to post synaptic receptors >>>> ((( inhibition or excitation))) of the post-synaptic membrane
- \* ((depending on the type of the neurotransmitter i.e. excitatory or inhibitory))



### These receptors have two components

1. Binding site that face the cleft to bind the neurotransmitter

2. Ionophore: It passes all the way through the membrane to the interior. It is of two types

Ion channels

2<sup>nd</sup> messenger system in the post-synaptic membrane.

#### **Cation channels**

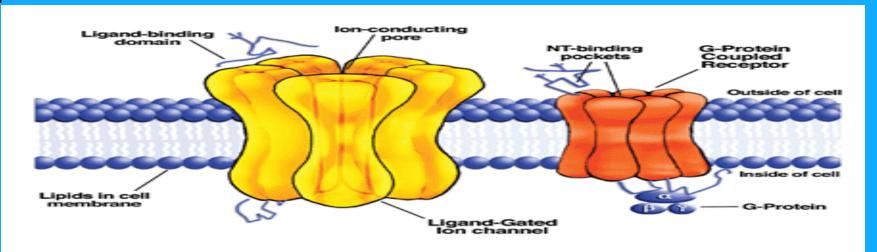
Na+ (most common)

K+

Ca++

#### **Anion channels**

Cl⁻ (mainly)



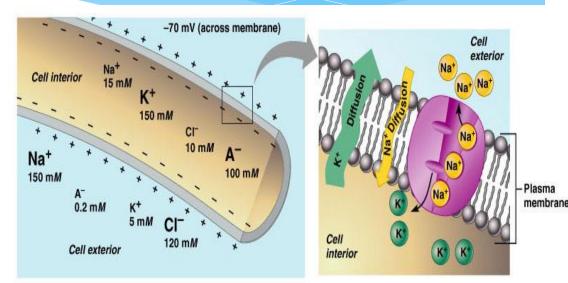
- \* Fate of a neurotransmitter
  - After a transmitter substance is released at a synapse, it must be removed by
- \* <u>Diffusion</u> out of synaptic cleft into surrounding fluid
- \* Enzymatic destruction e.g. Ach esterase for Ach
- \* Active transport back into pre-synaptic terminal itself e.g. norepinephrine

## Electrical events in post-synaptic neurons:

# 1. RMP of neuronal soma:

 $\simeq$  -65mV i.e. less than sk. ms. [-70 to -90mV]

 If the voltage is less negative → the neuron is excitable



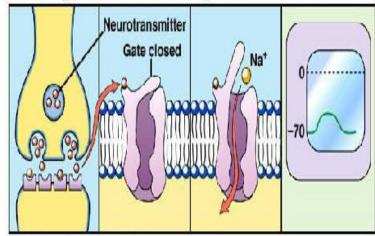
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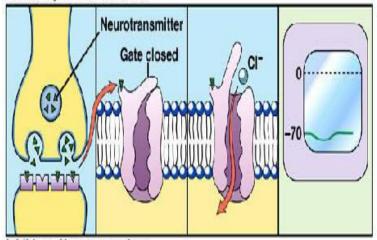
## 2. Excitatory postsynaptic potential [EPSPs]

When **excitatory** neurotransmitter binds to its receptor on postsynaptic membrane → partial depolarization [↑ Na influx] of post-synaptic cell membrane immediately under presynaptic ending, i.e. EPSPs If this potential rises enough to threshold level → AP will develop and excite the neuron

#### **Excitatory and Inhibitory Neurotransmitters**



**Excitatory Neurotransmitter** 



Inhibitory Neurotransmitter

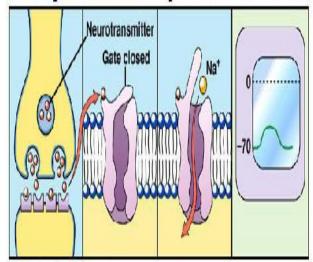
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# 3. Inhibitory post-synap (IPSPs):

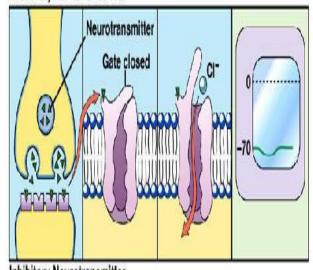
When an inhibitory NT binds receptor on post-synaptic membrane >>>> hyperpolarization of the post-synaptic membrane

1-Increasing membrane permeability to Cl of post-synaptic memb. (produced by inhibitory neurotransmitter)  $\rightarrow \downarrow$  excitability and m. potential (more negative)

### **Excitatory and Inhibitory Neurotransmitters**



**Excitatory Neurotransmitter** 



Inhibitory Neurotransmitter

## Synaptic properties

## 1. One-way conduction

Synapses generally permit conduction of impulses in one-way i.e. from pre-synaptic to post-synaptic neuron.

## 2. Synaptic delay

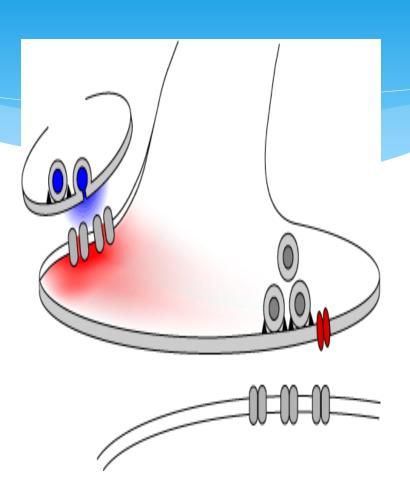
Is the minimum time required for transmission across the synapse. It is 0.5 ms for transmission across one synapse. This time is taken by

- Discharge of transmitter substance by pre-synaptic terminal
- Diffusion of transmitter to post-synaptic membrane
- Action of transmitter on its receptor
- Action of transmitter to ↑ membrane permeability
- Increased diffusion of Na+ to ↑ post-synaptic potential

# 3. Synaptic inhibition:

### **Types:**

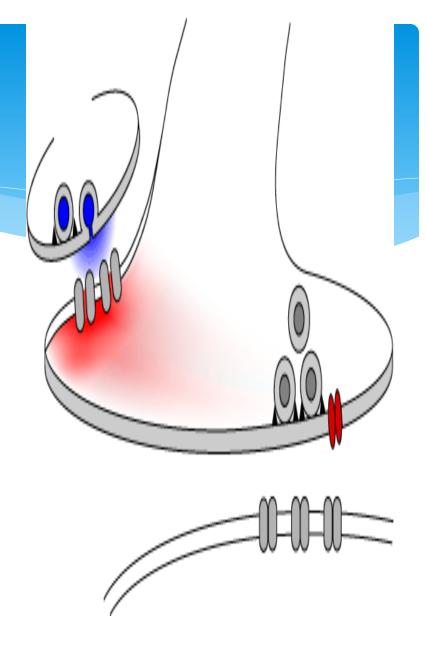
Direct inhibition
Indirect inhibition
Reciprocal inhibition
Inhibitory interneuron



#### A. Direct inhibition:

Occurs when an inhibitory neuron (releasing inhibitory substance) acts on a post-synaptic neuron leading to  $\rightarrow$  hyperpolarization due to opening of Cl [IPSPs] and/or K+ channels.

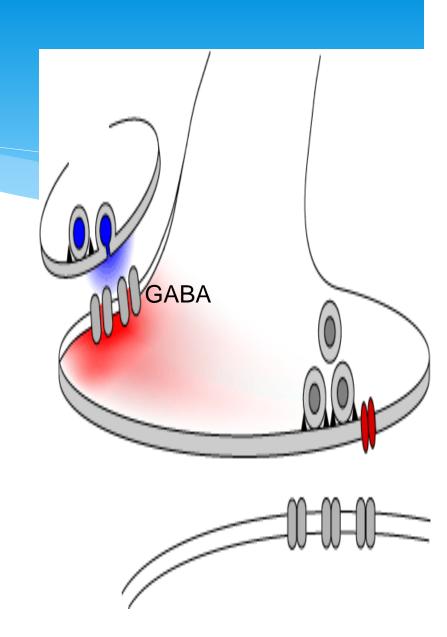
Example: Glycine at the level of the spinal cord to block pain impulses.



# B. Indirect Inhibition: (=Pre-synaptic inhibition).

This happens when an inhibitory synaptic knob lie directly on the termination of a pre-synaptic excitatory fiber. The inhibitory synaptic knob release a transmitter which inhibits the release of excitatory transmitter from the pre-synaptic fiber.

e. g. GABA (Pain modification)



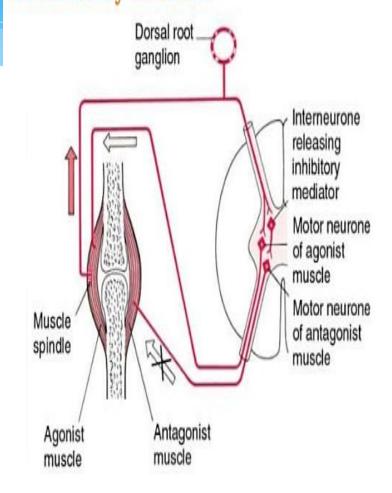
# C. Reciprocal inhibition

Inhibition of antagonist activity is initiated in the agonist muscle.

Impulses pass directly to the motor neurons supplying the same muscle and via branches to inhibitory interneurones that end on motor neurones of antagonist muscle.

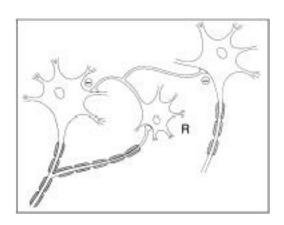
#### RECIPROCAL INHIBITION

When one muscle is contracted, its antagonist is automatically inhibited.



## D. Inhibitory interneuron (Renshaw cells)

Negative feedback inhibitory interneuron of a spinal motor neuron.



## 4. Summation:

#### **a.** Spatial summation.

When EPSP occurs in more than one synaptic knob at the same time

#### b. Temporal summation.

If EPSPs in a pre-synaptic knob are successively repeated without significant delay so the effect of the previous stimulus is summated to the next.

## 5. Convergence and divergence:

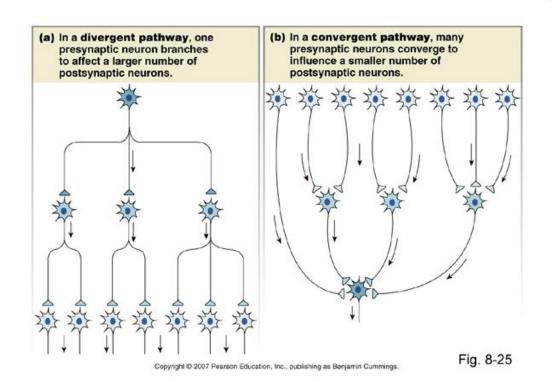
#### Convergence

When many pre-synaptic neurons converge on any single postsynaptic neuron.

#### **Divergence**

Axons of pre-synaptic neurons divide into many branches that diverge to end on many post-synaptic neurons.

# Divergence & convergence



## 6. Fatigue

It is due to exhaustion of neurotransmitter.

If the pre synaptic neurons are continuously stimulated there may be an exhaustion of the neurotransmitter. Resulting in stoppage of synaptic transmission.

## Factors affecting synaptic transmission:

#### **Alkalosis:**

#### **Increases neuronal excitability.**

Causes cerebral epileptic seizures (Increased excitability cerebral neurons)

e. g. overbreating in person with epilepsy

The over breathing blows off carbon dioxide and therefore elevates the pH of the blood momentarily

#### **Acidosis:**

Depresses neuronal activity; pH around 7.0 usually causes a coma (e. g. severe diabetic or uremic acidosis)

#### **Drugs:**

Caffeine found in coffee, tea, increases neuronal excitability, by reducing the threshold for excitation of neurons.



## Hypoxia

\* Depression of neurons

