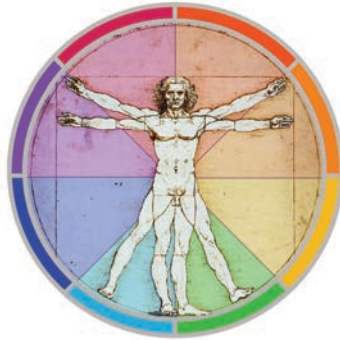


# HUBS191 Lecture Material

This pre-lecture material is to help you prepare for the lecture and to assist your note-taking within the lecture,  
it is NOT a substitute for the lecture !



Please note that although every effort is made to ensure this pre-lecture material corresponds to the live-lecture there may be differences / additions.

# Physiological Principles of Human Movement and Sensation



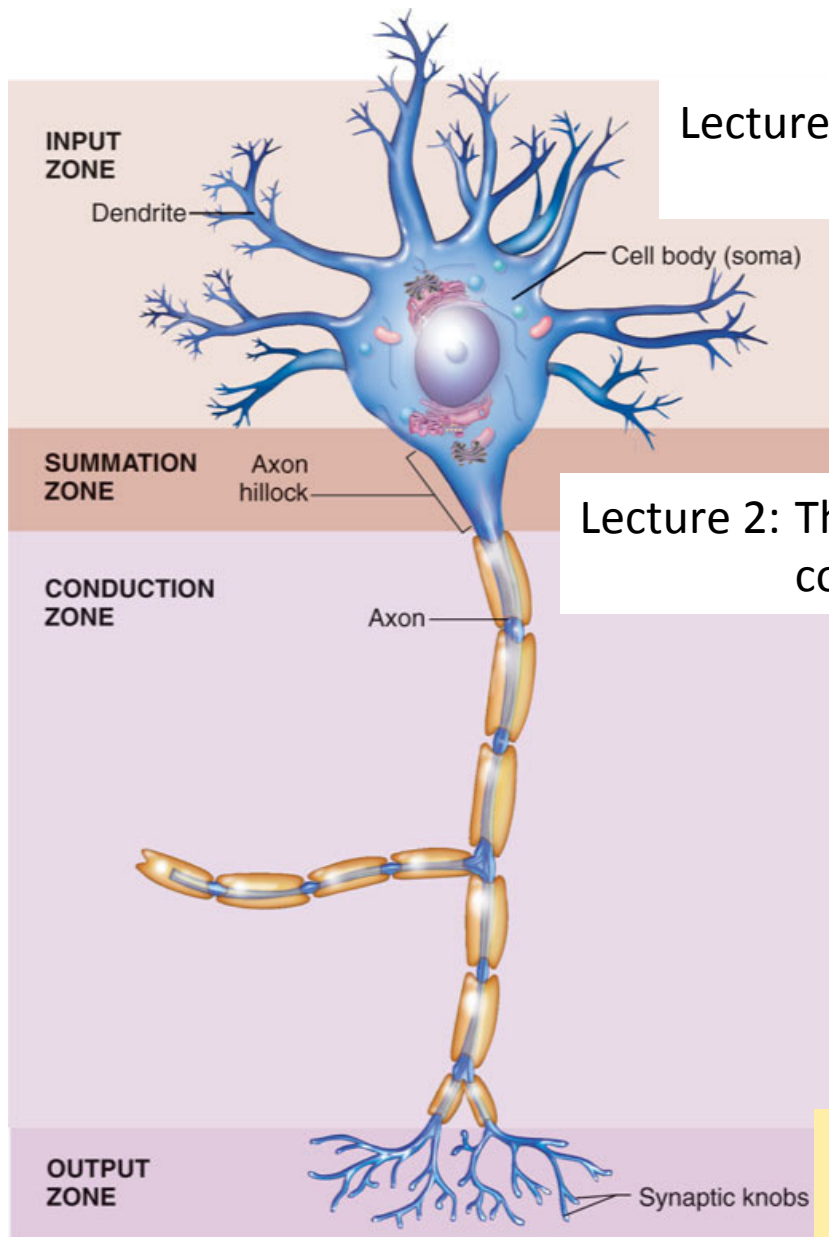
## **HUBS191**

### **Lecture 23**



Dr Martin Fronius  
Department of Physiology  
6. April 2017

# Lectures outline



Lecture 4: Synaptic Networks and Integration  
(input zone)

Lecture 2: The Action Potential and its  
conduction (axon hillock, axon)

Lecture 1: Bioelectricity and membrane potentials

Lecture 3: The Synapse – ‘connecting’ neurons  
(output zone)

# Aims of Today's Lecture

## Lecture 23 – The Synapse 'making connections'

### Outline

- Recap from lecture 2
- What is a synapse?
- Two types of Synapses: Electrical and chemical
- How does a chemical synapse work?
- How is a synapse switched off?

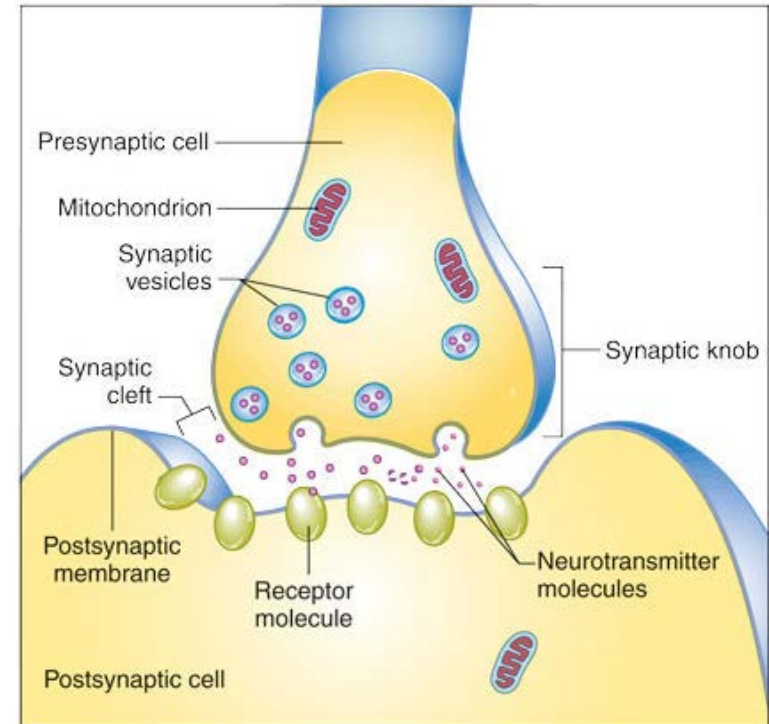


Fig. 13-23 Thibodeau and Patton, 8<sup>th</sup> Ed, p. 400

# To summarise the events during an AP...

## STEP DESCRIPTION

- 1 A stimulus triggers stimulus-gated  $\text{Na}^+$  channels to open and allow inward  $\text{Na}^+$  diffusion. This causes the membrane to depolarize.
- 2 As the threshold potential is reached, voltage-gated  $\text{Na}^+$  channels open.
- 3 As more  $\text{Na}^+$  enters the cell through voltage-gated  $\text{Na}^+$  channels, the membrane depolarizes even further.
- 4 The magnitude of the action potential peaks (at +30 mV) when voltage-gated  $\text{Na}^+$  channels close.
- 5 Repolarization begins when voltage-gated  $\text{K}^+$  channels open, allowing outward diffusion of  $\text{K}^+$ .
- 6 After a brief period of hyperpolarization, the resting potential is restored by the sodium-potassium pump and the return of ion channels to their resting state.

Table 13-2 from Thibodeau and Patton, 8<sup>th</sup> Ed p. 396

- **TIMING IS EVERYTHING !**
- Order of channel events  
→ opening/closing!
- Different types of channels  
(e.g. stimulus, voltage-gated)

Table 13-1 from Thibodeau and Patton  
8<sup>th</sup> Ed p. 395

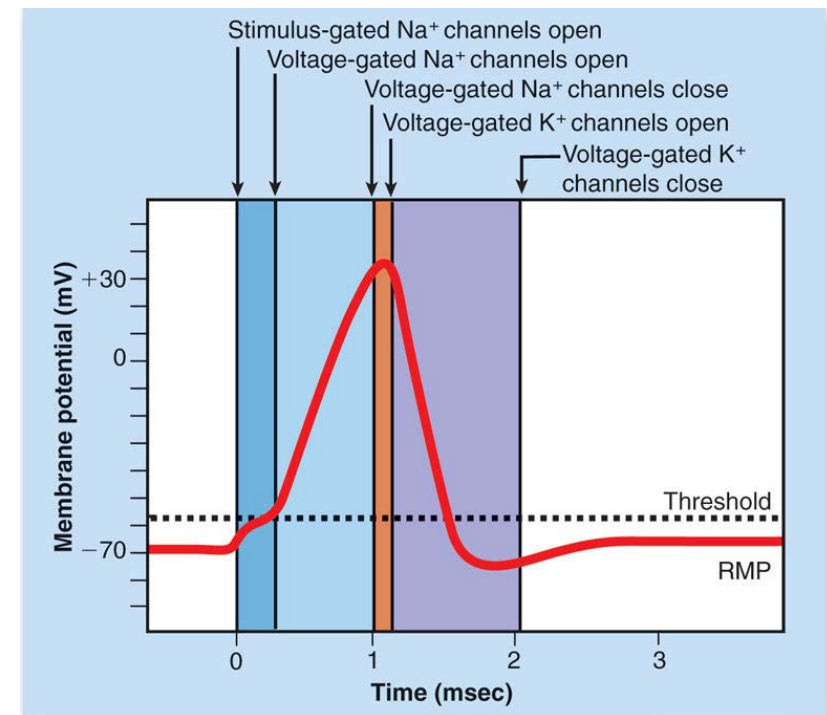
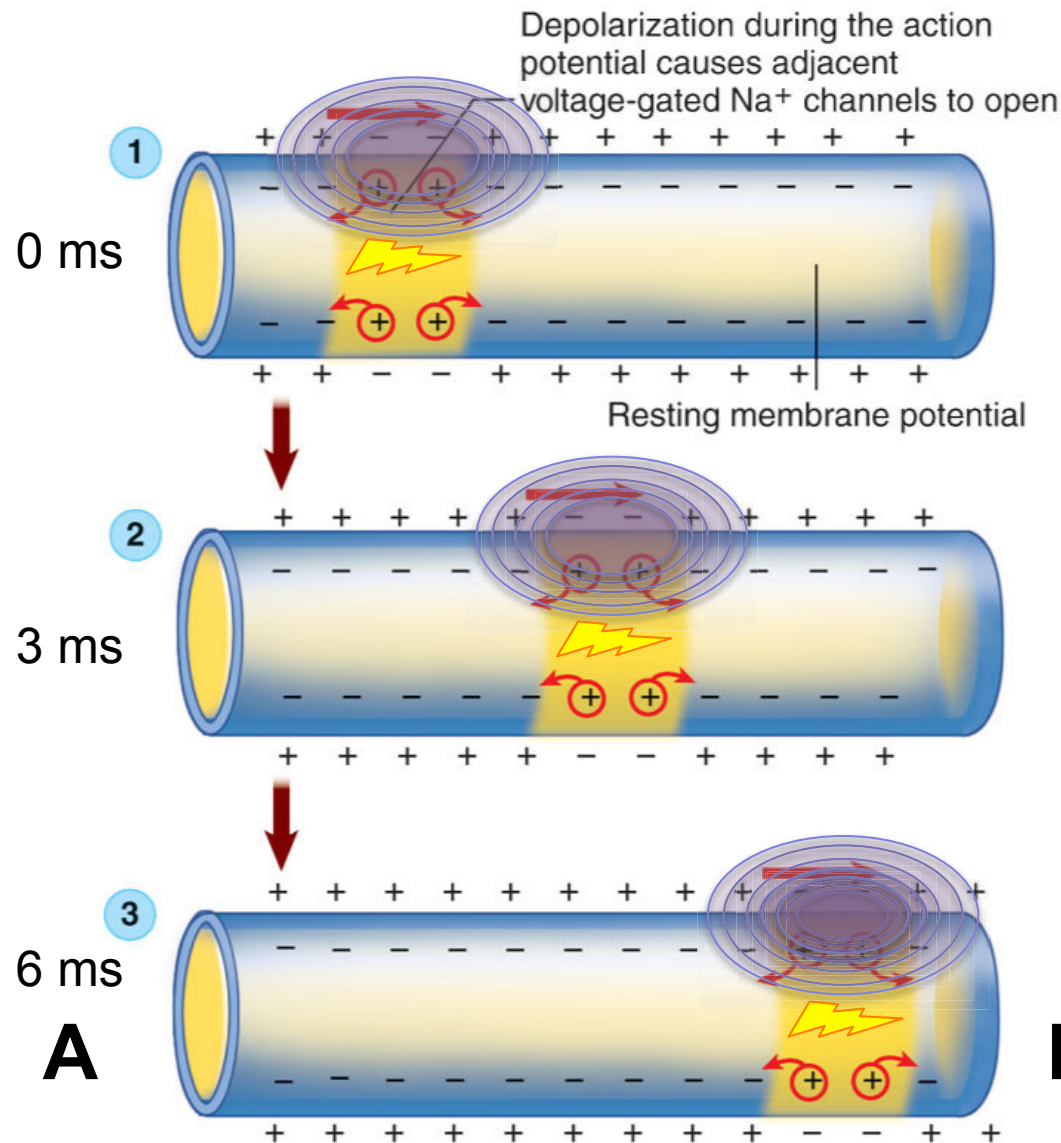


Fig. 13-19 Thibodeau and Patton 8<sup>th</sup> Ed p.396

# Conduction of the Action Potential down Axons



Relies on the spread of the DEPOLARISING electrical signal  
**Generation of an electric field**

ALONG

the axon to instantly activate the next set of voltage gated Na<sup>+</sup> channels



# What determines the propagation direction of an Action Potential

→ Refractory period

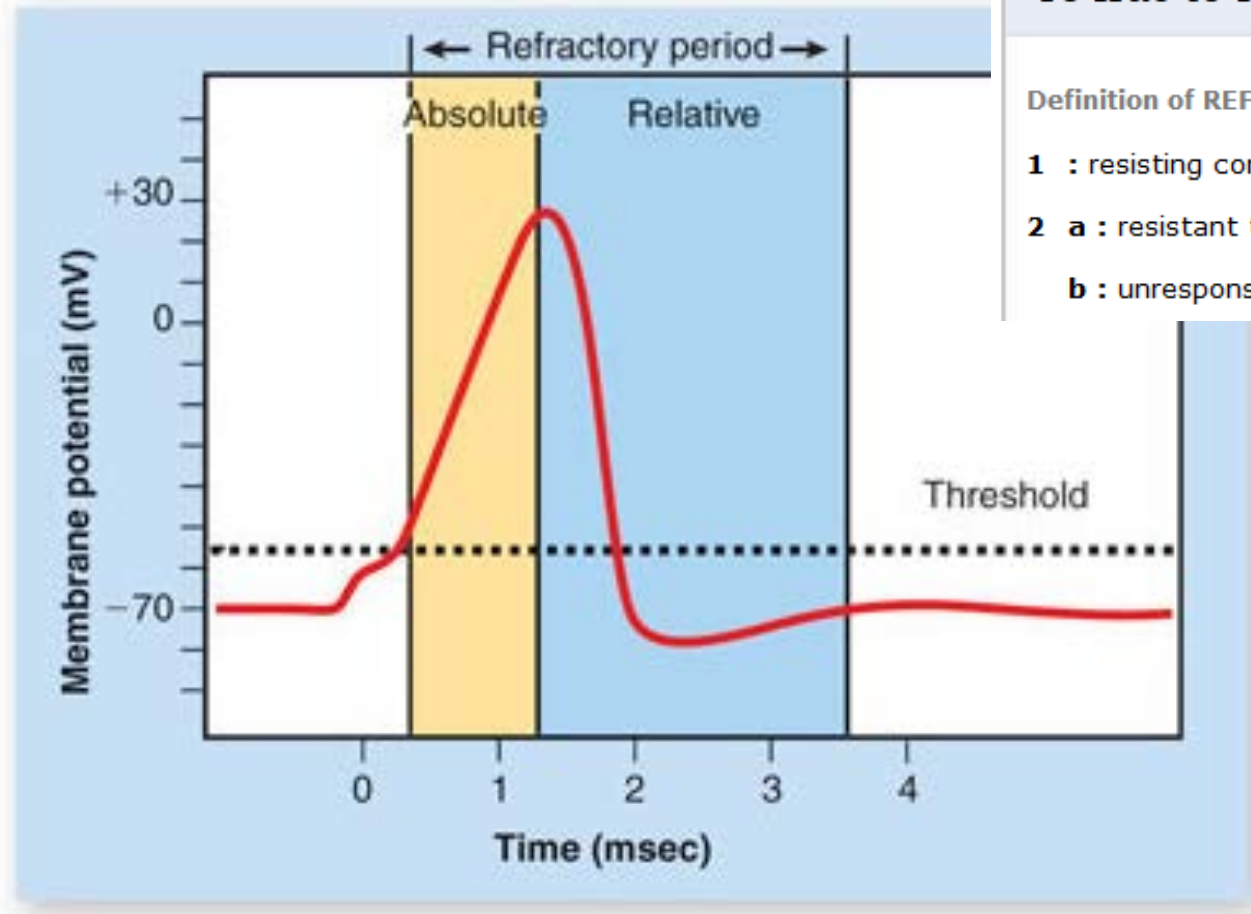


Fig. 13-20 Thibodeau and Patton 8<sup>th</sup> Ed p. 397

<sup>1</sup>re·frac·to·ry  *adj* \ri·'frak-t(ə-)rē\

Definition of REFRACTORY

- 1 : resisting control or authority : **STUBBORN**, **UNMANAGEABLE**
- 2 **a** : resistant to treatment or cure <a refractory lesion>
- b** : unresponsive to stimulus

From <http://www.merriam-webster.com>

## REFRACTORY PERIODS:

- **Absolute** → no response
- **Relative** → large stimulus required

## REFRACTORY PERIODS:

- Intrinsic feature of the Na<sup>+</sup> channels

→ It also limits the number of action potentials at a given time (frequency of action potentials)

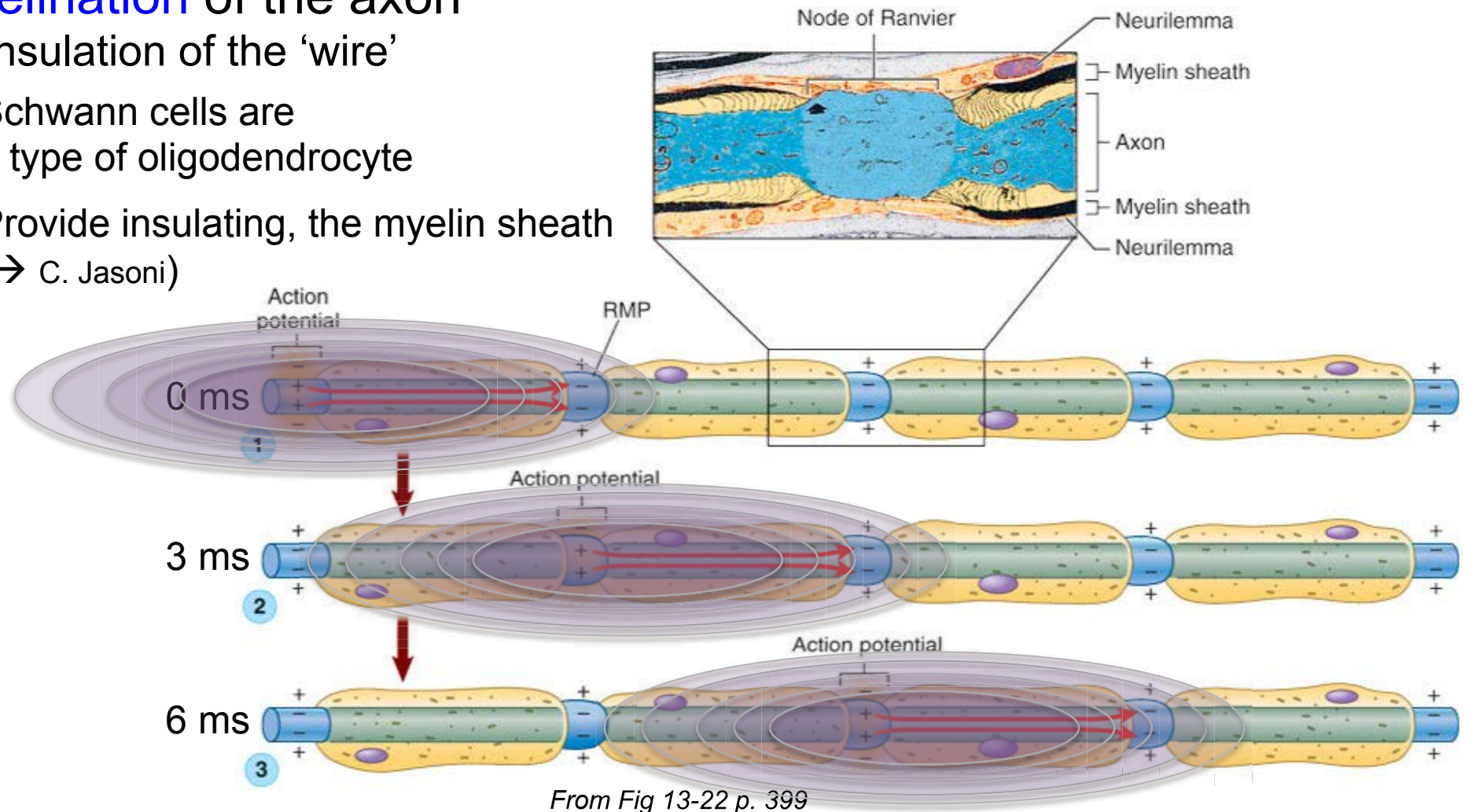
# IMPROVE CONDUCTION SPEED

## → MYELINATED AXONS

### Myelination of the axon

→ Insulation of the 'wire'

- Schwann cells are a type of oligodendrocyte
- Provide insulating, the myelin sheath (→ C. Jasoni)



**The action potential leaps between the nodes of Ranvier – saltatory conduction**



# Aims of Today's Lecture

## Lecture 23 – The Synapse 'making connections'

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- Two types of Synapses: Electrical and chemical
- How does a chemical synapse work?
- How is a synapse switched off?

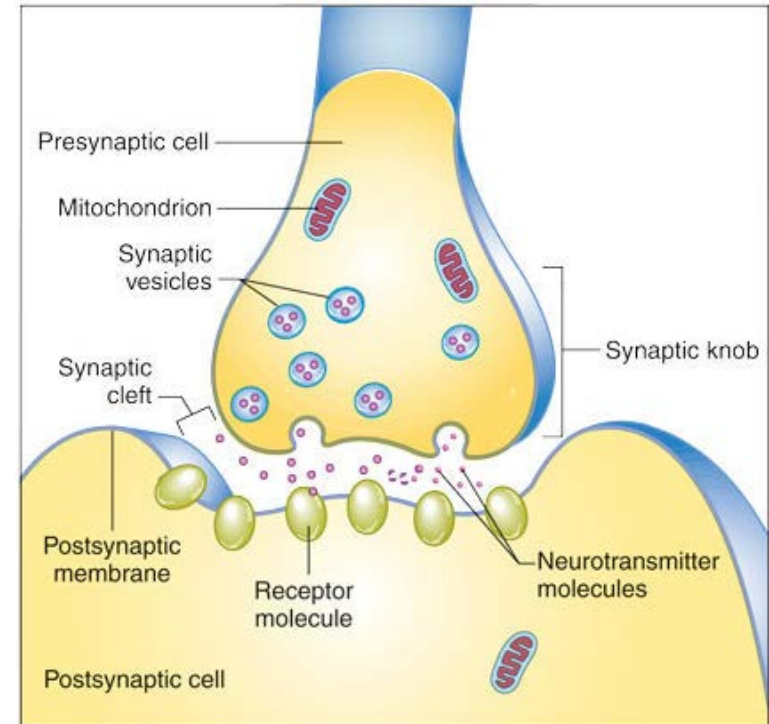


Fig. 13-23 Thibodeau and Patton, 8<sup>th</sup> Ed, p. 400

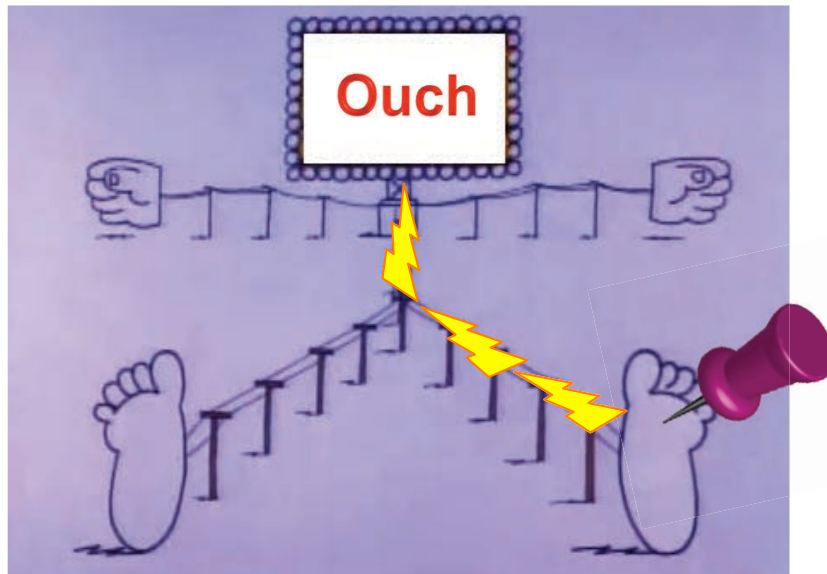
# Objectives

- Know the two types of synapse and how they differ
- List the structures that make up a chemical synapse
- Explain the steps that must occur for chemical synaptic transmission to occur
- List the mechanisms that terminate the chemical signal

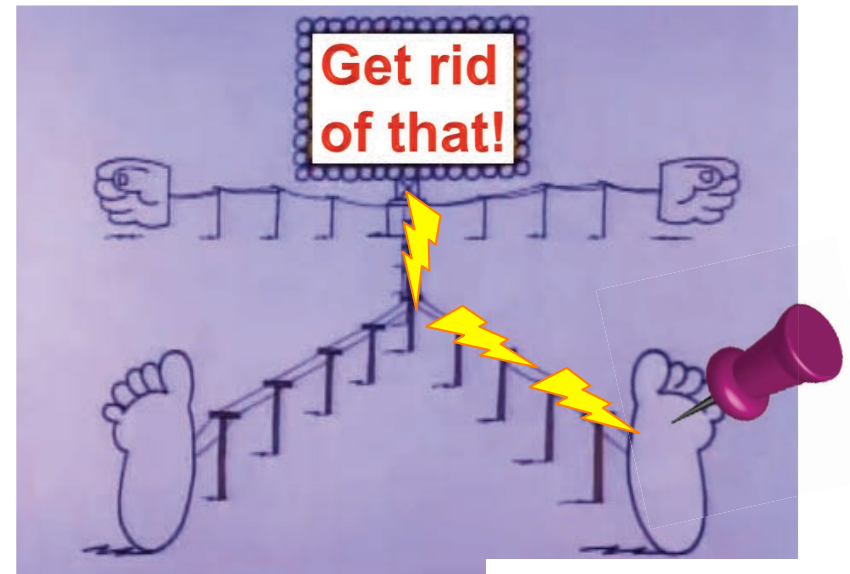
# Network Communication within our nervous system

REMEMBER --- From A/Prof Jasoni's Lecture

Integrating...



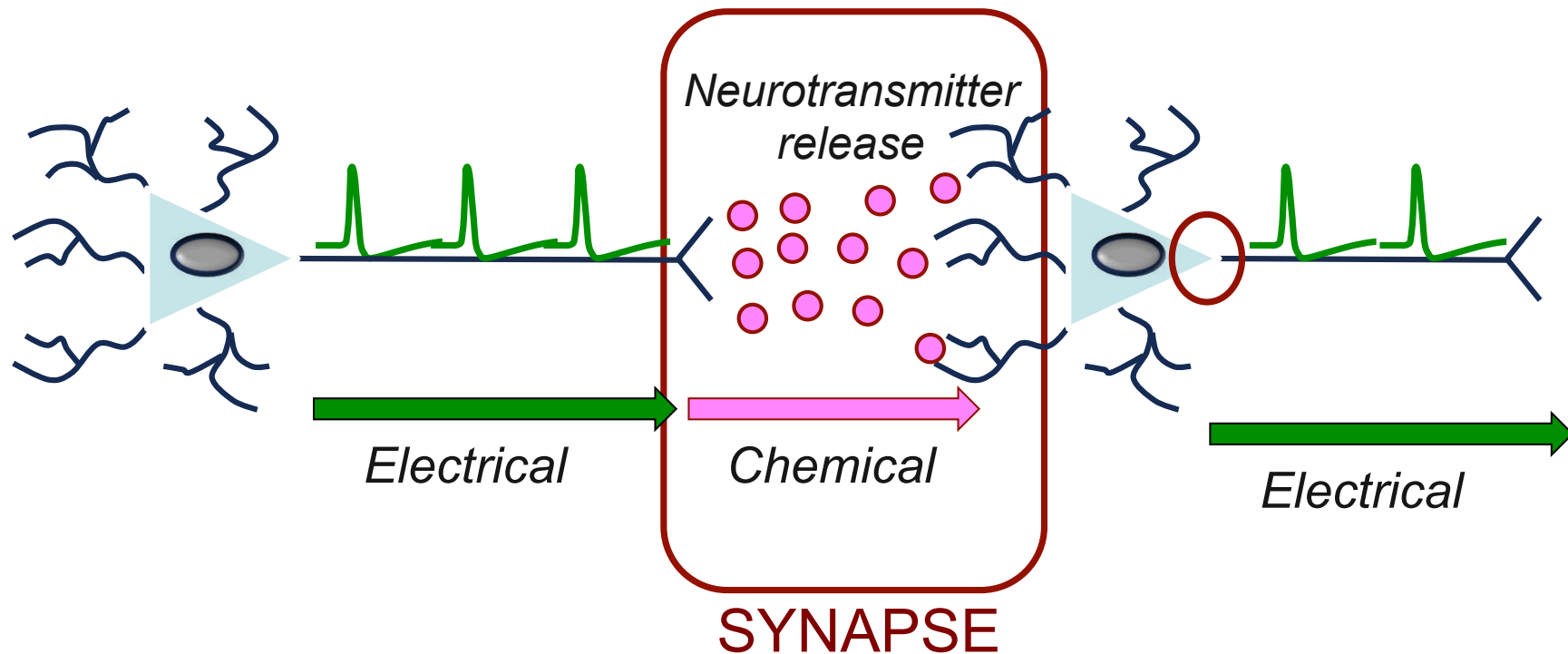
...and coordinating.



Shake foot!

- Action Potentials transmit information “up/down the wire”  
→ between neurons
- Within the NETWORK of the nervous system
- **SYNAPSES** join the network and allow control of information flow

*Recap: Cells of the nervous system:* Communication between neurons occurs through a junction called a *SYNAPSE*



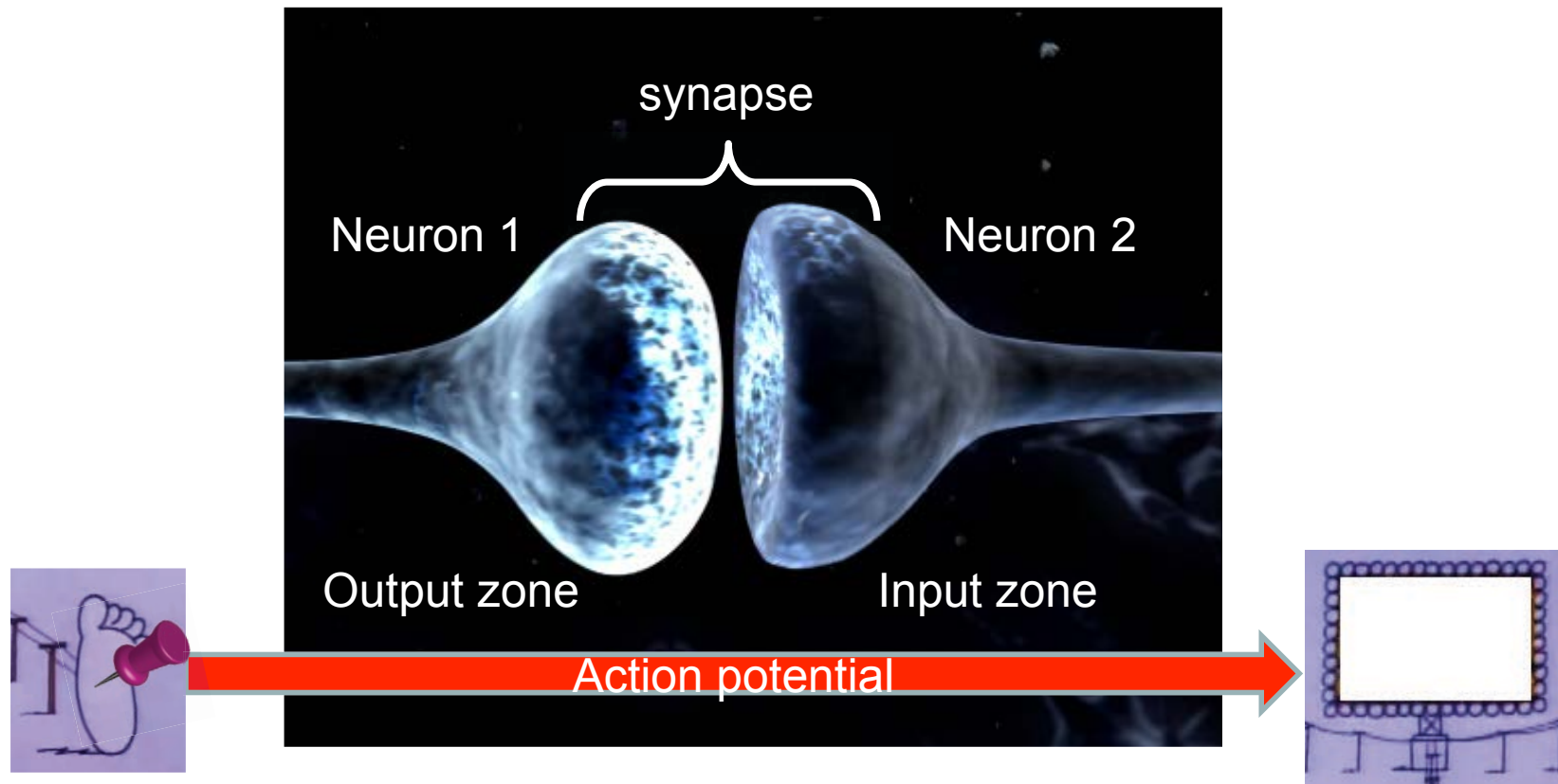
See also: Patton & Thibodeau 8<sup>th</sup> ed Fig 13-5 (7<sup>th</sup> ed Fig 12-5)

REMEMBER --- From A/Prof Jasoni's Lecture

# What is a Synapse ?

“The junction between nerve cells, where a nerve impulse is transferred from one neuron to another.”

<http://www.achenet.org/resources/glossary.php>





# Two different Types of Synapses – Electric

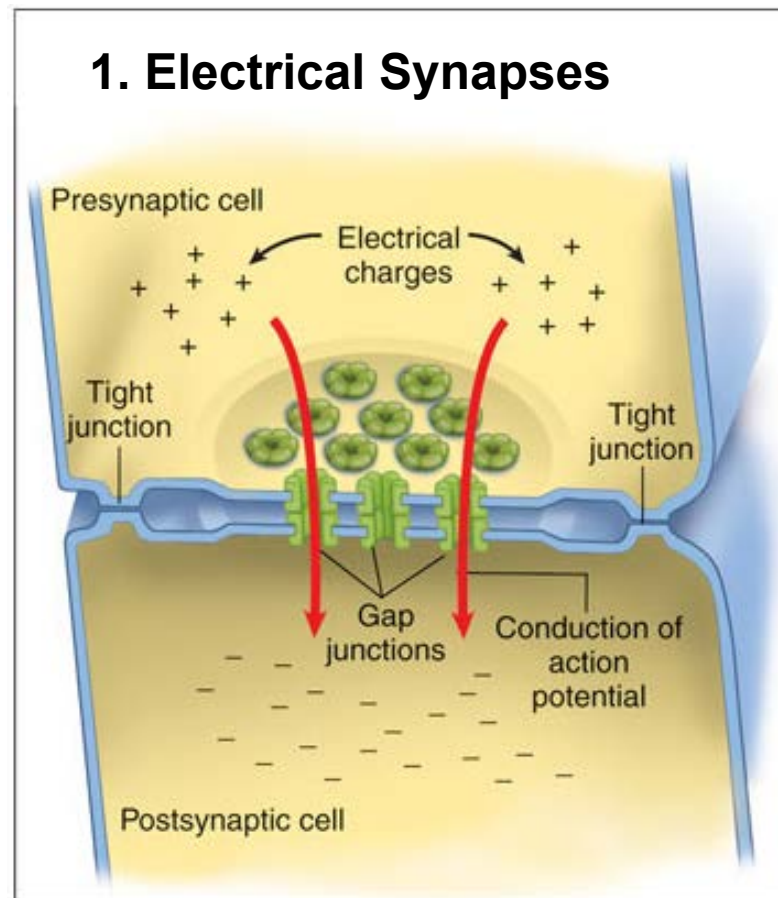


Fig. 13-23, Patton & Thibodeau, 8<sup>th</sup> Ed, p. 400

- Gap between neurons  
– linked by gap junctions
  - Gap junctions: specific type of membrane channels
  - ‘Tunnel’ connecting 2 different neurons
  - Physical link (direct connection)
- **Direct** propagation of **Action Potential**
- **VERY FAST** synapse

# Two different Types of Synapses – Electric

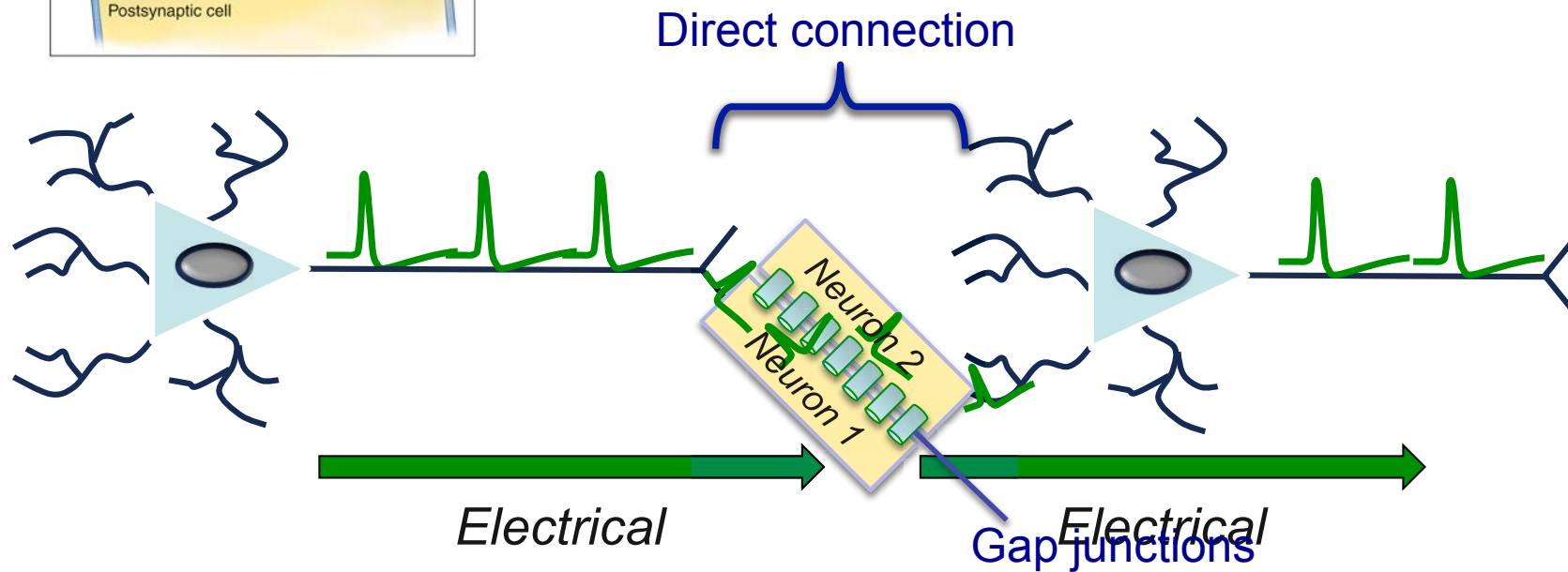
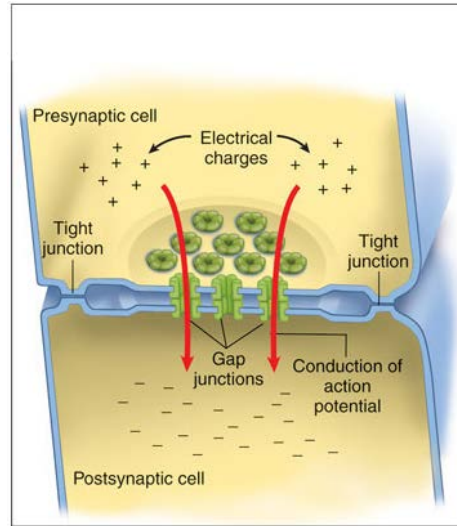


Fig. 13-23, Patton & Thibodeau, 8<sup>th</sup> Ed, p. 400

# Two different Types of Synapses – Chemical

## 2. Chemical Synapses

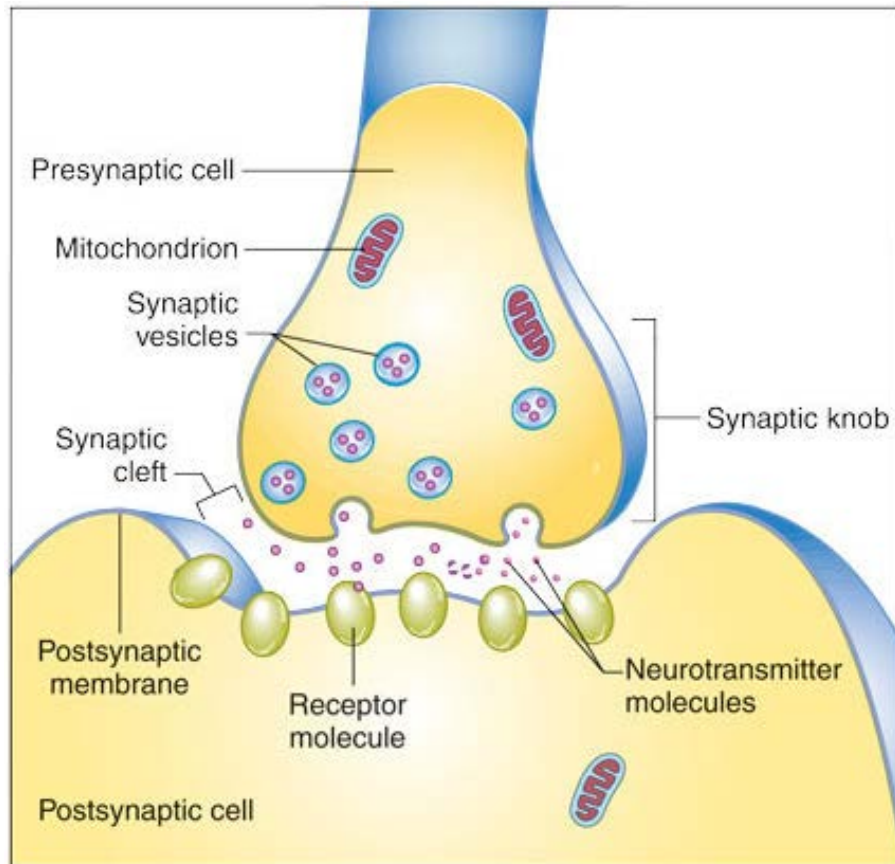
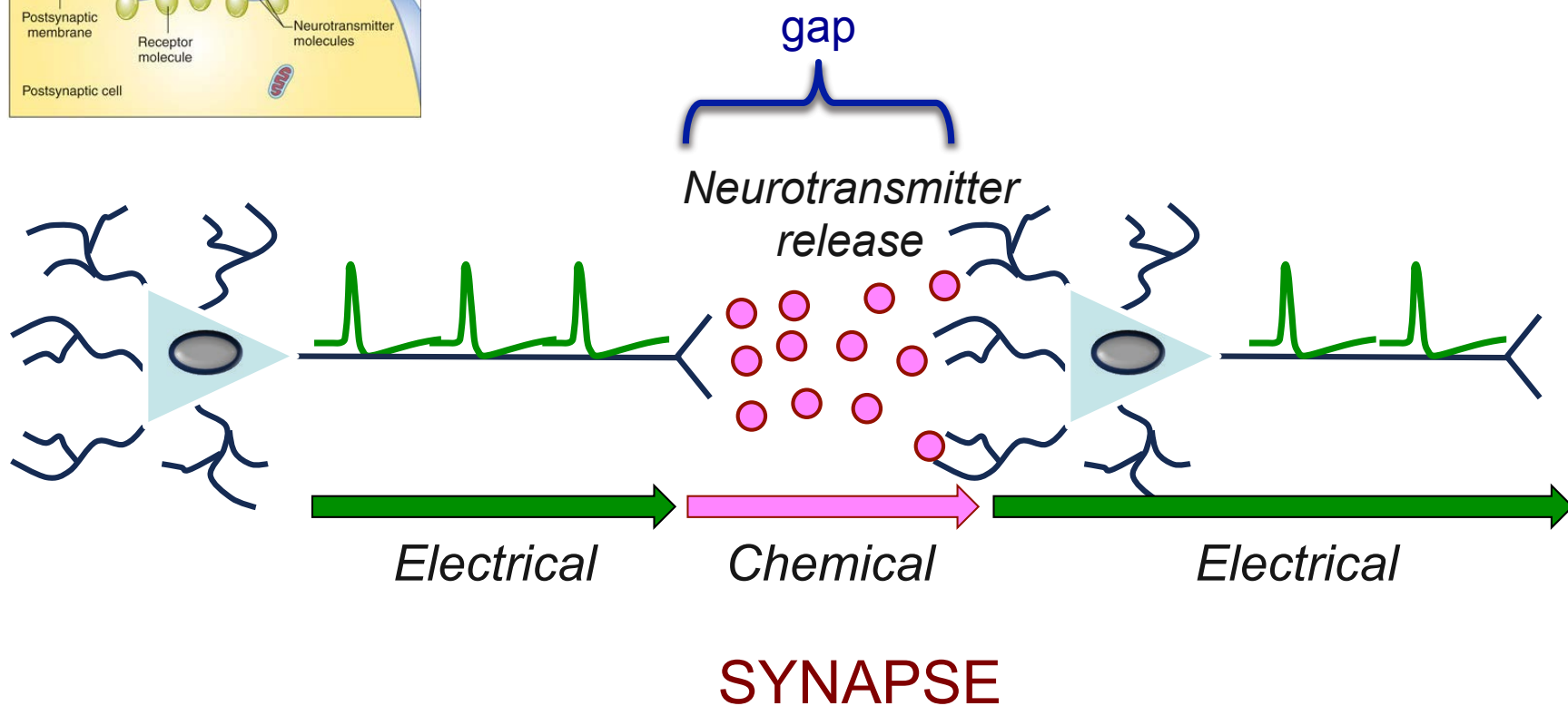
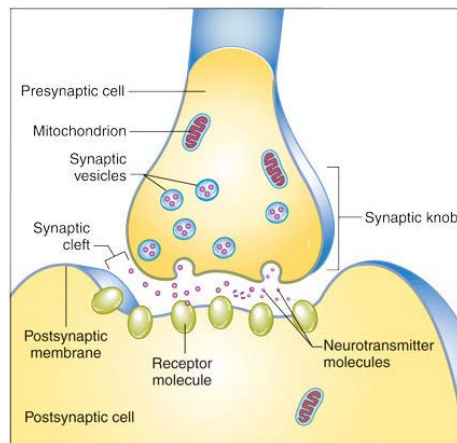


Fig. 13-23, Patton & Thibodeau, 8<sup>th</sup> Ed, p. 400

- Physical gap between neurons  
– linked by a chemical compound
  - Neurotransmitter! ‘messenger’
  - Released into the synaptic cleft
  - Bridges the gap
- **Indirect** propagation of **Action Potential**
- Slower than electrical synapse

# Two different Types of Synapses – Chemical



See also: Patton & Thibodeau 8<sup>th</sup> ed Fig 13-5 (7<sup>th</sup> ed Fig 12-5)

# Special type of Chemical Synapse

- Nerve to Muscle Synapse – the motor unit

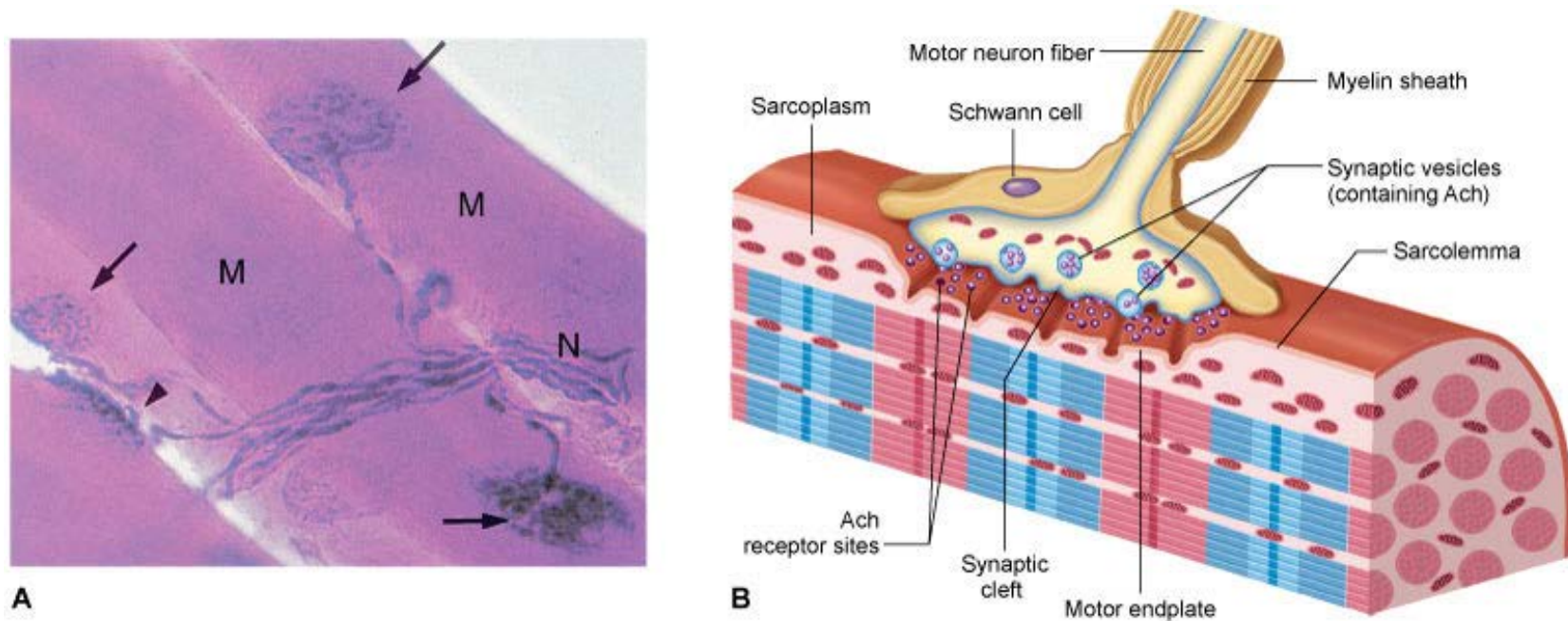


Fig. 13-23, Patton & Thibodeau, 8<sup>th</sup> Ed, p. 353

Also called the  
**Neuro-muscular junction**



# The chemical synapse – the key components

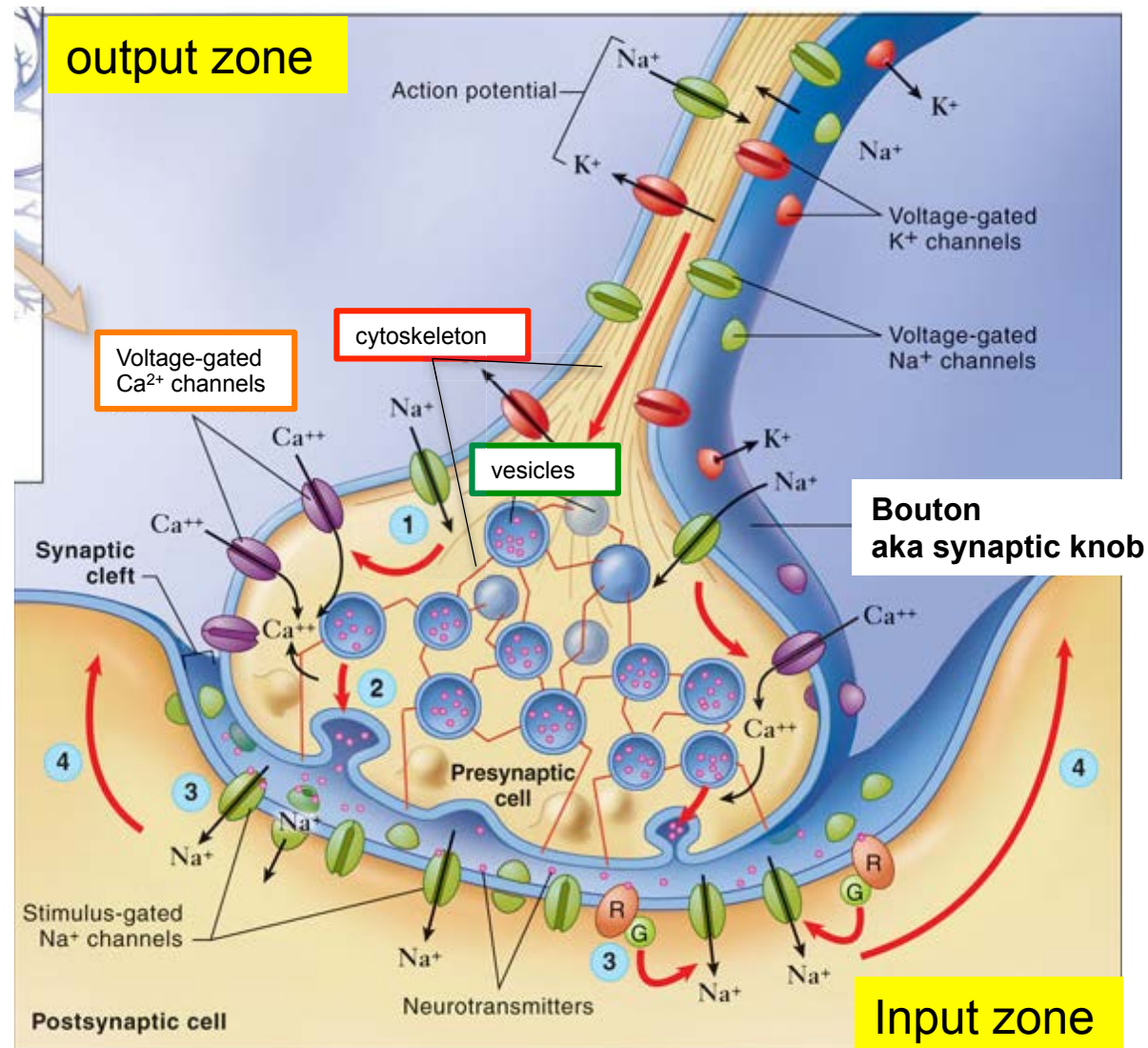
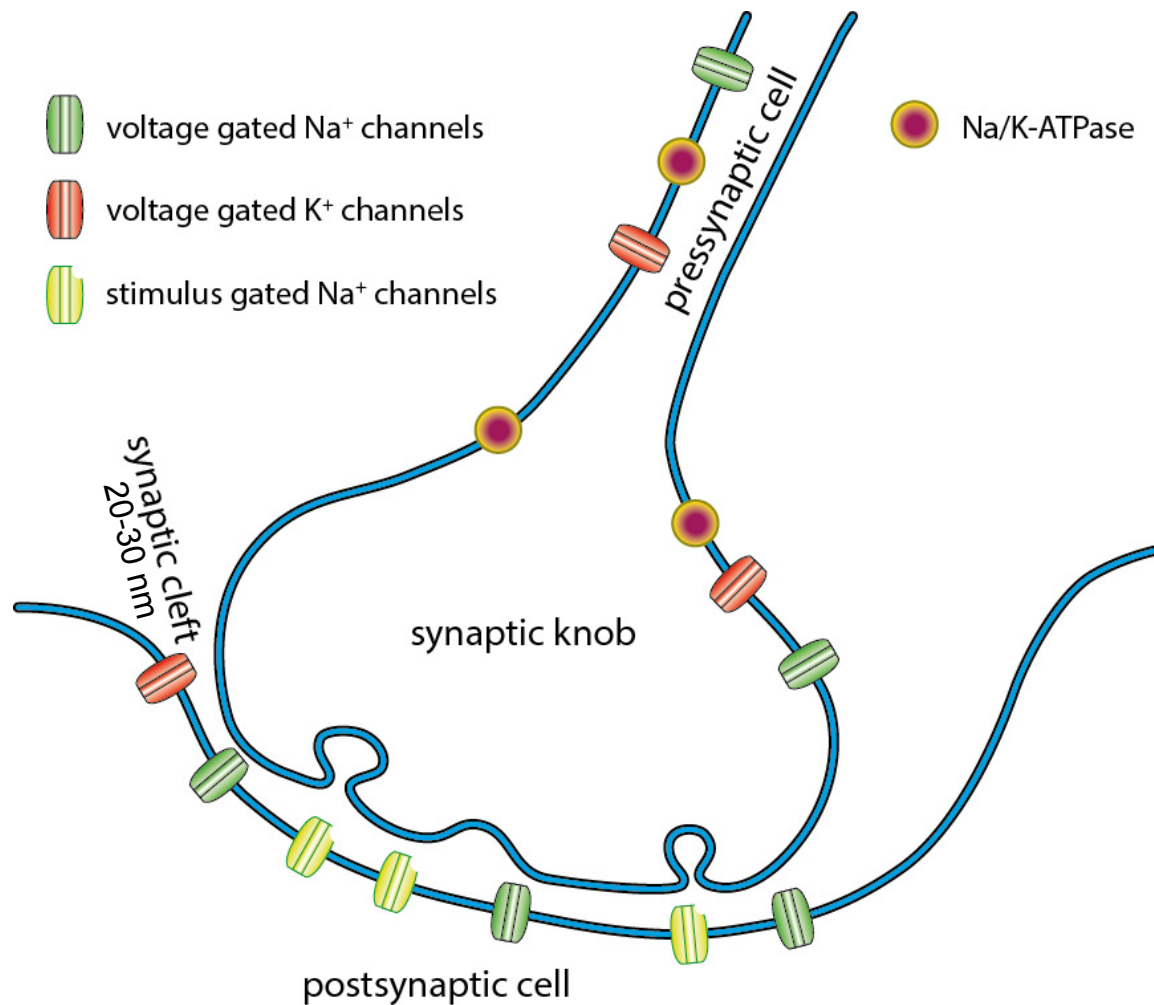


Fig. 13-25, Patton & Thibodeau, 8<sup>th</sup> Ed, p. 401

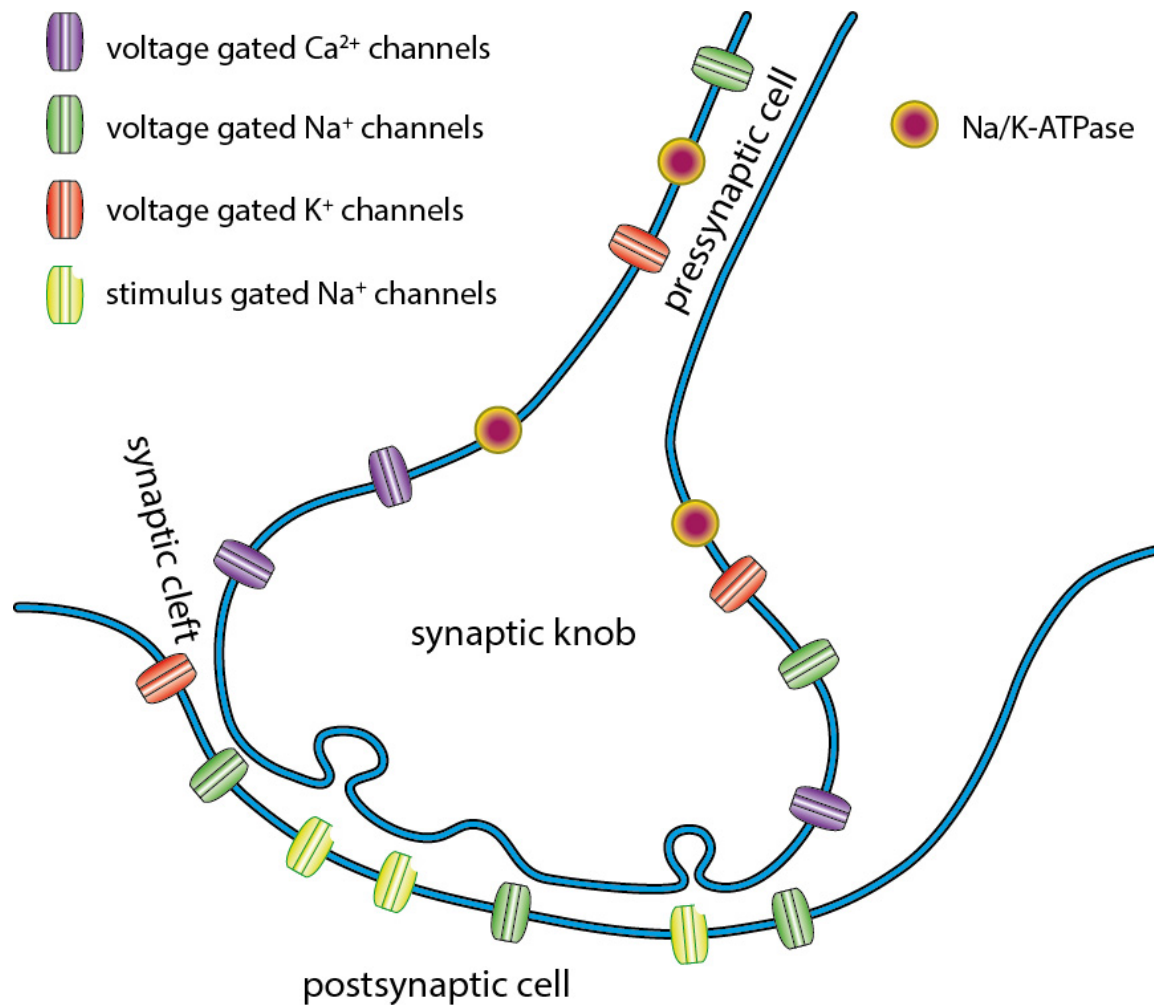
# The chemical synapse – the key components I



1. Presynaptic cell  
'output zone'
2. Synaptic knob
3. Synaptic cleft  
'the gap'
4. Postsynaptic cell  
'input zone'  
**with ion channels  
and receptors**

*Adapted from Fig. 13-25, Patton & Thibodeau, 8<sup>th</sup> Ed, p. 401*

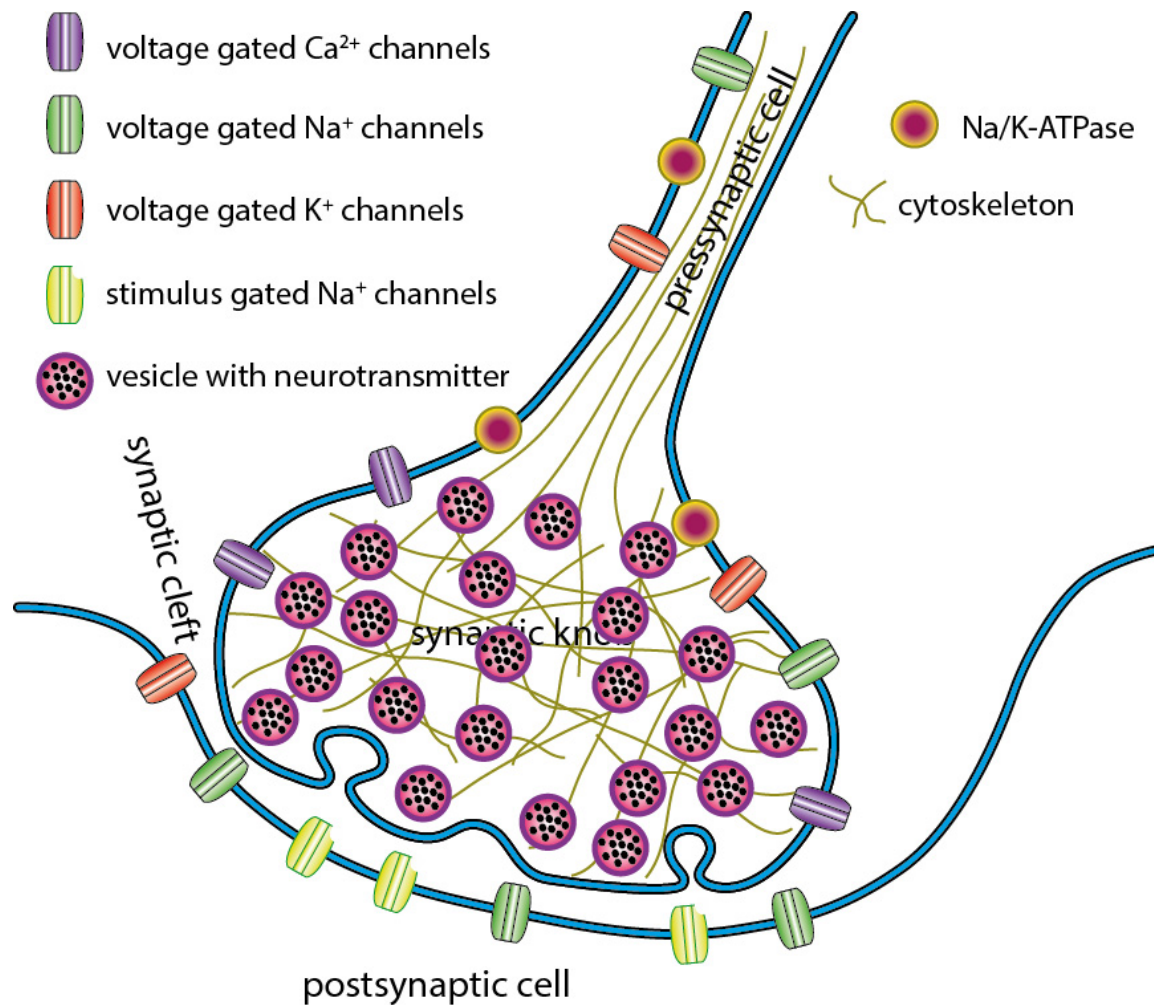
# The chemical synapse – the key components II



1. Presynaptic cell  
'output zone'
2. Synaptic knob
3. Synaptic cleft  
'the gap'
4. Postsynaptic cell  
'input zone'
5. Voltage-gated  
 $\text{Ca}^{2+}$  channels

Adapted from Fig. 13-25, Patton & Thibodeau, 8<sup>th</sup> Ed, p. 401

# The chemical synapse – the key components III

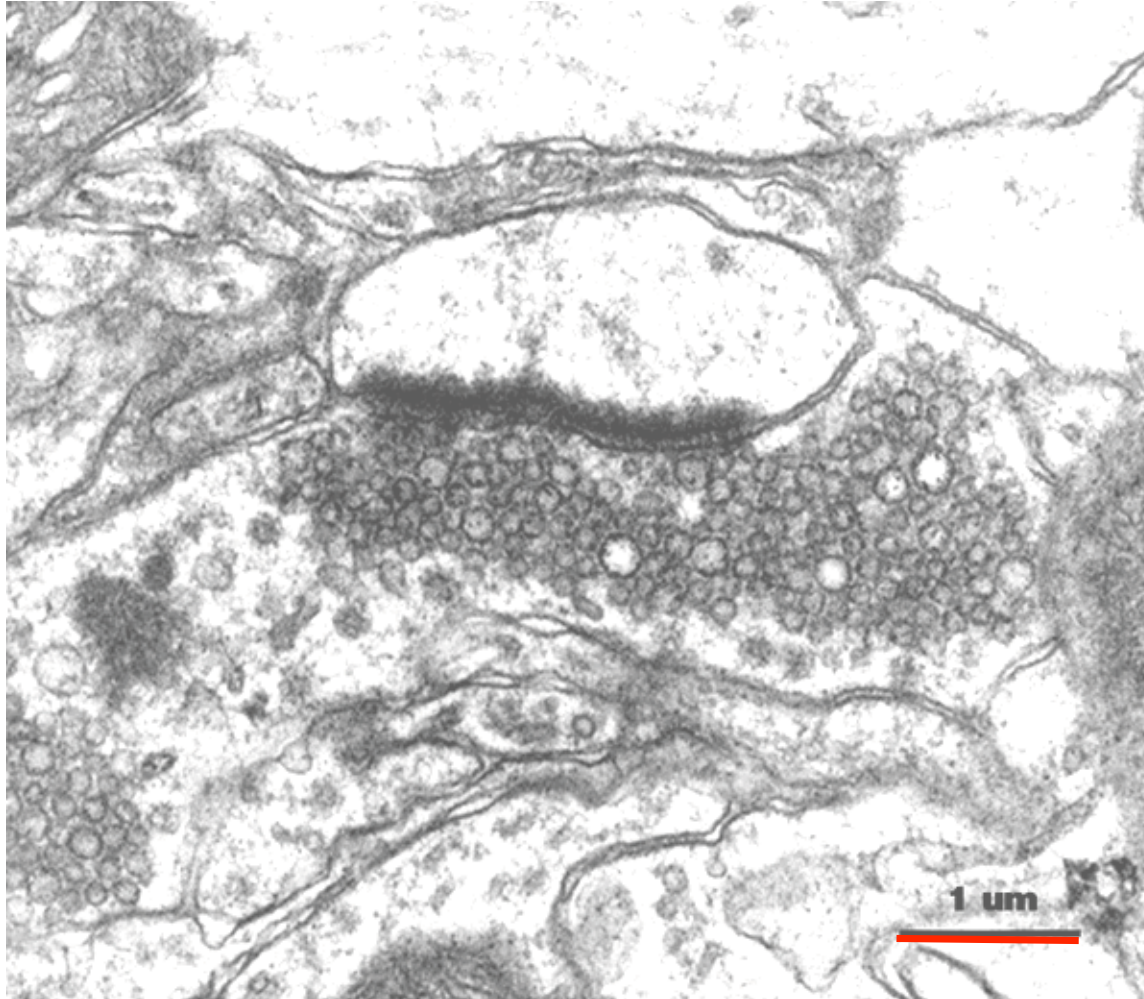


1. Presynaptic cell  
'output zone'
2. Synaptic knob
3. Synaptic cleft  
'the gap'
4. Postsynaptic cell  
'input zone'
5. Voltage-gated  $\text{Ca}^{2+}$  channels
6. Vesicles
7. Cytoskeleton
8. Na/K-ATPase
9. Mitochondria  
(not shown)

Adapted from Fig. 13-25, Patton & Thibodeau, 8<sup>th</sup> Ed, p. 401



# Electron micrograph of a synapse



Postsynaptic – why ?

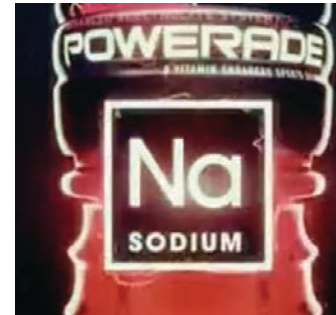
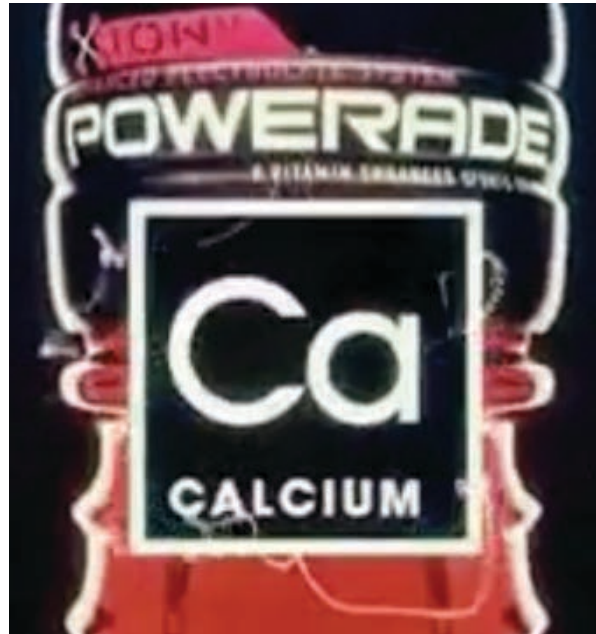
Presynaptic – why ?

**$1 \times 10^{-6} \text{ m}$**

From <http://www.itg.uiuc.edu/exhibits/gallery/pages/image-51.htm>

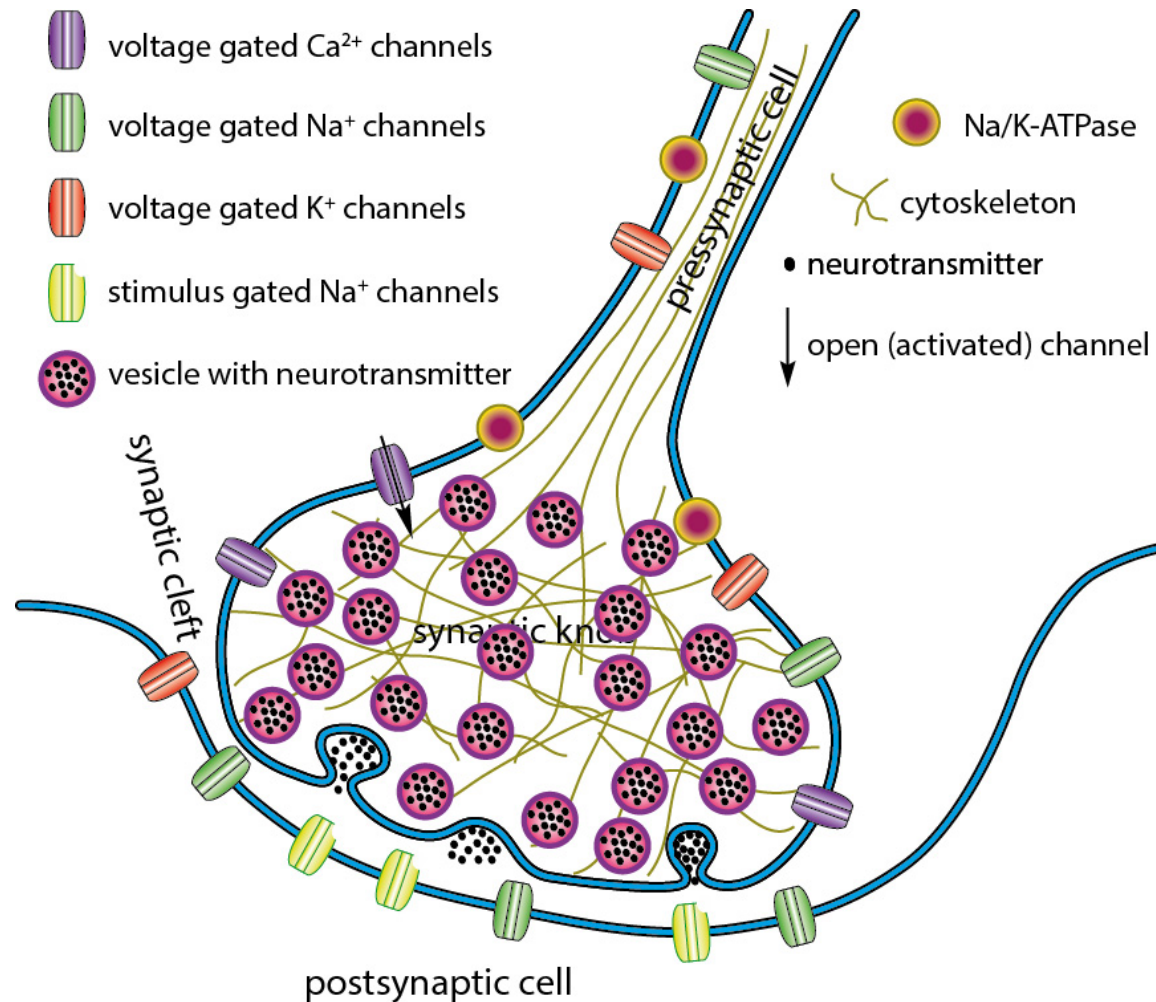


# Another CATION – Calcium ( $\text{Ca}^{2+}$ )



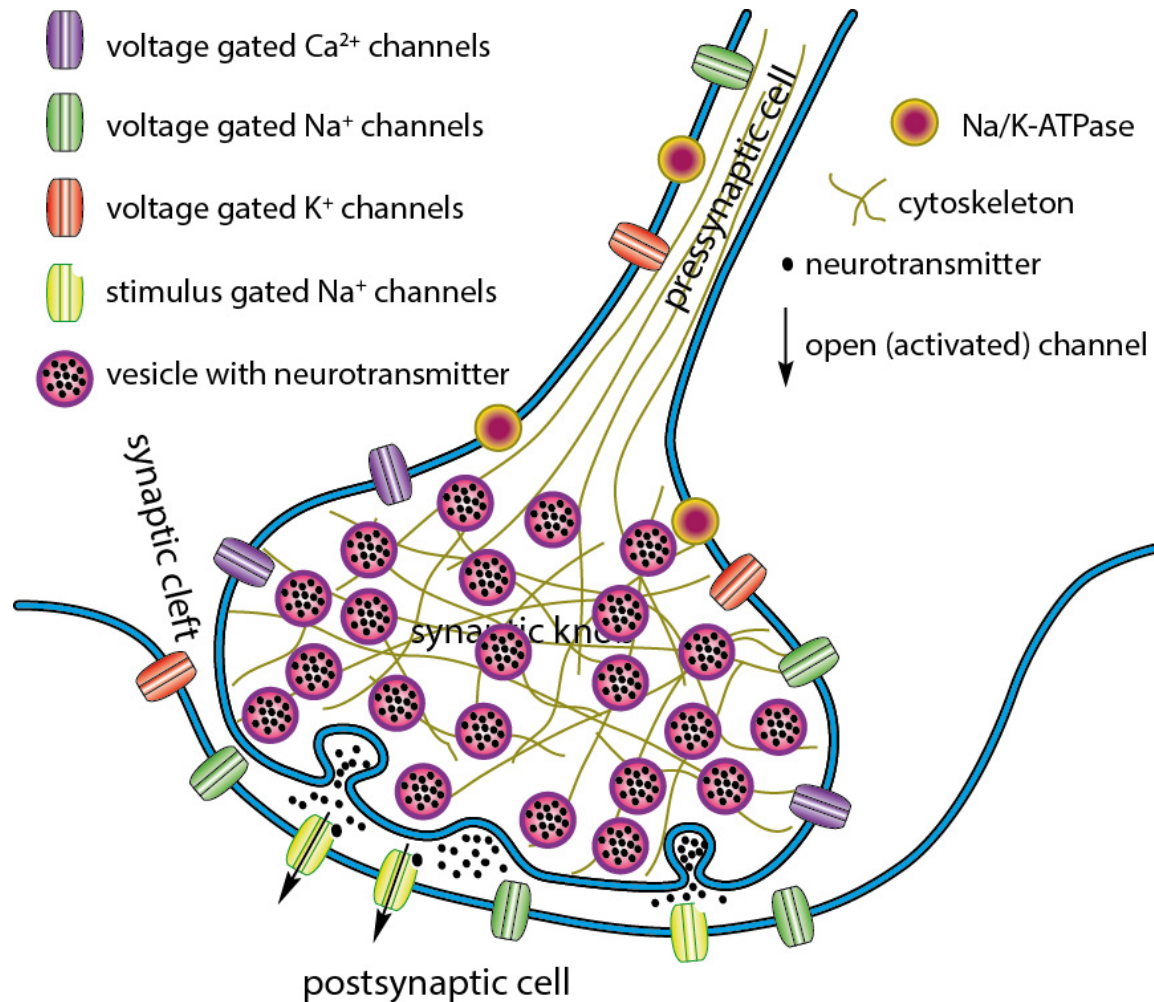
- 2 - 2.5 mM in extracellular fluid
- very low concentrations intracellularly ( $\mu\text{M}$  range)
- Important for bones...  
...and the **nervous system**
- → **triggers the release of vesicles**

# How is the signal transmitted?



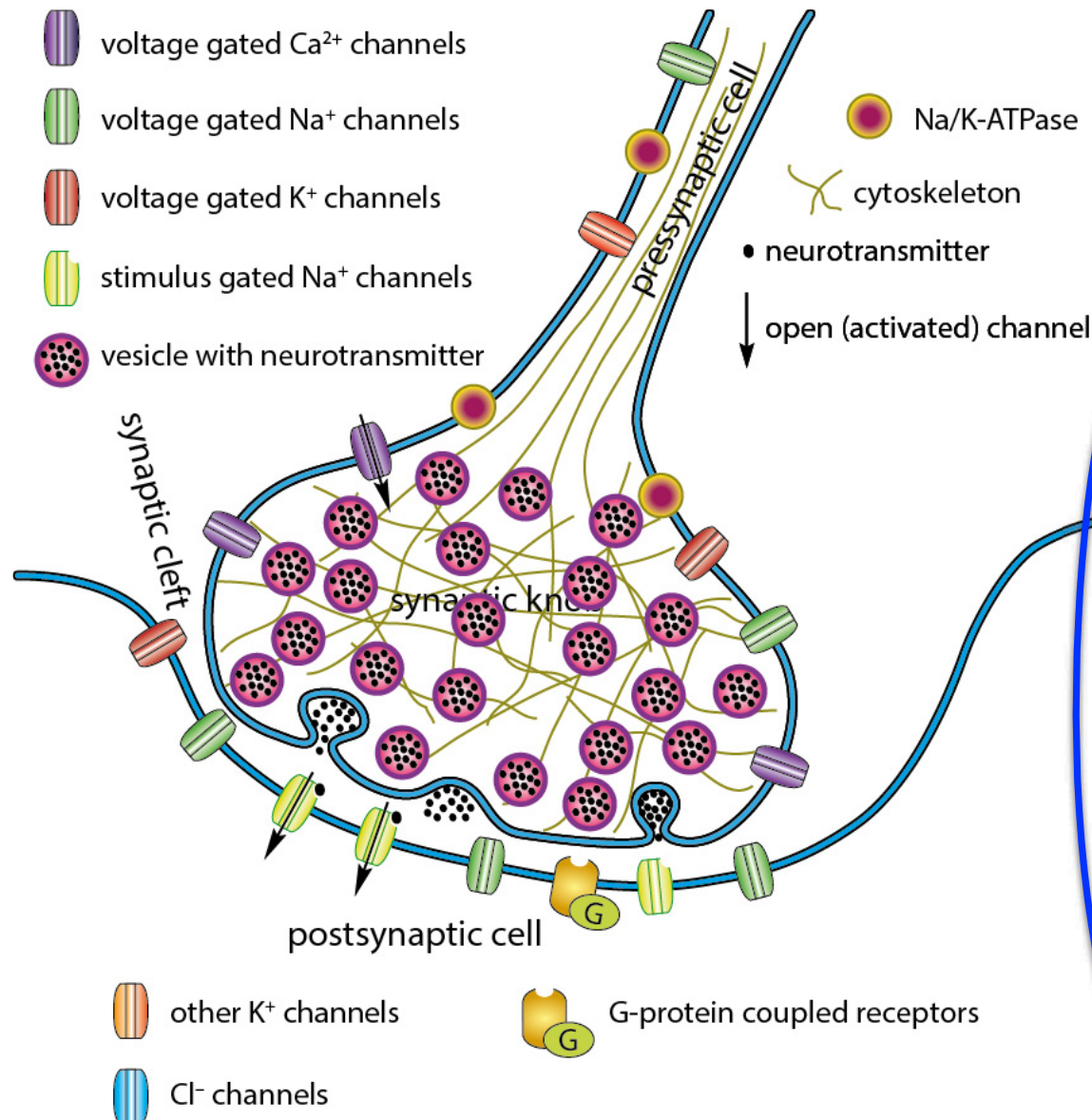
1. Action potential propagates down the axon – to the pre-synaptic knob
2. Pre-synaptic knob is depolarised – **voltage-gated  $\text{Ca}^{2+}$  channels open**
3.  $\text{Ca}^{2+}$  enters the cell
4.  **$\text{Ca}^{2+}$  ions** TRIGGER the fusion of vesicles with the presynaptic membrane.
5. This releases the neurotransmitter into the synaptic cleft

# How is the signal transmitted? II



- Neurotransmitter diffuses across the synaptic cleft to reach the postsynaptic membrane
- Neurotransmitter bind to specific receptors.  
aka 'stimulus gated channels'
- If 'stimulus gated channels' are  $\text{Na}^{+}$  channels open  
→ **depolarisation**
- Subsequent activation of  $\text{K}^{+}$  channels  
→ **repolarisation**

# How is the signal transmitted? III



6. Neurotransmitter diffuses across the synaptic cleft to reach the postsynaptic membrane

7. Neurotransmitter bind to specific receptors. aka 'stimulus gated channels'

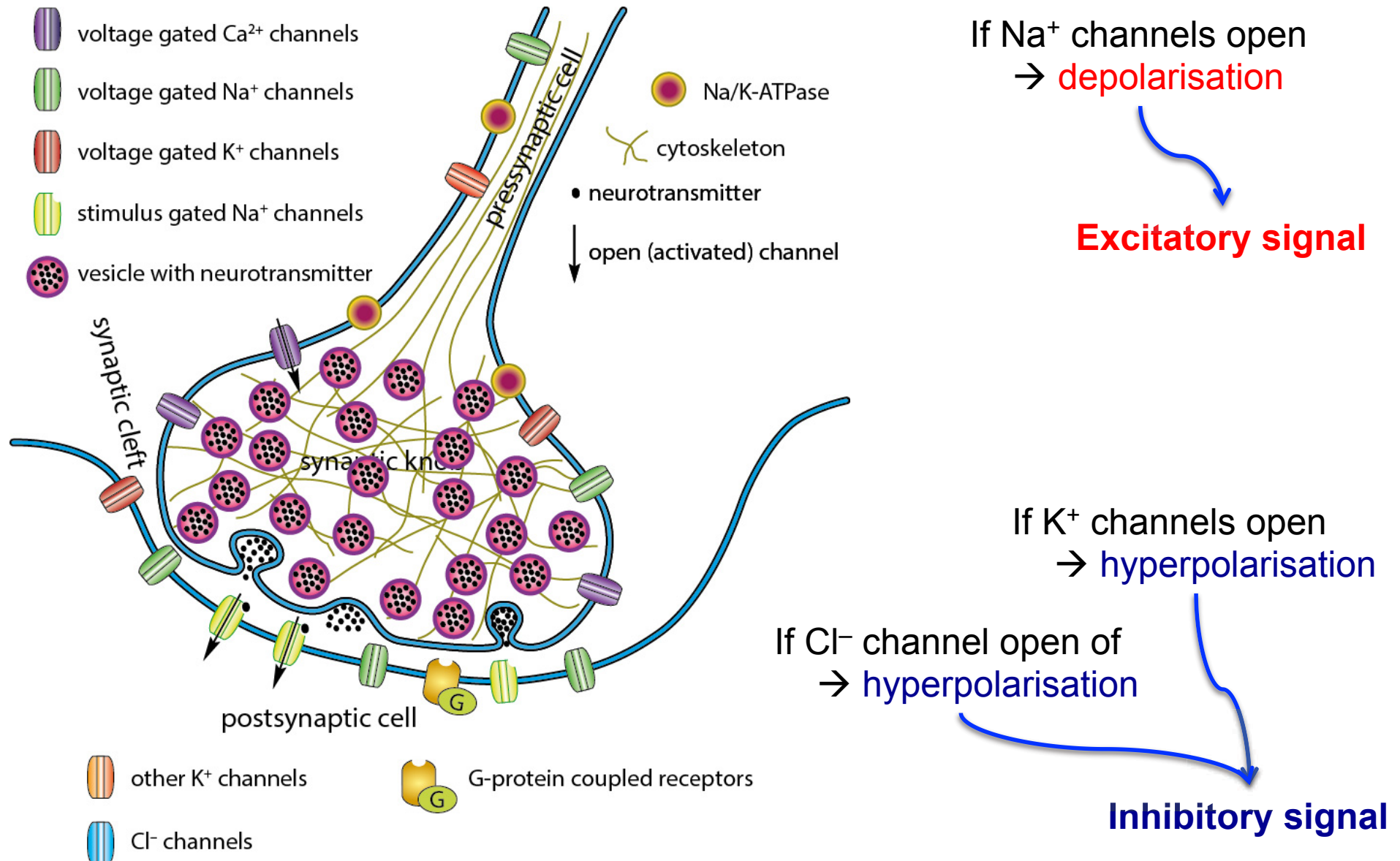
8. If 'stimulus gated channels' are  $\text{Na}^{+}$  channels open  
→ depolarisation

9. Subsequent activation of  $\text{K}^{+}$  channels  
→ repolarisation

8'. If stimulus gated channels are  $\text{Cl}^{-}$  or  $\text{K}^{+}$  channels  
→ hyperpolarisation

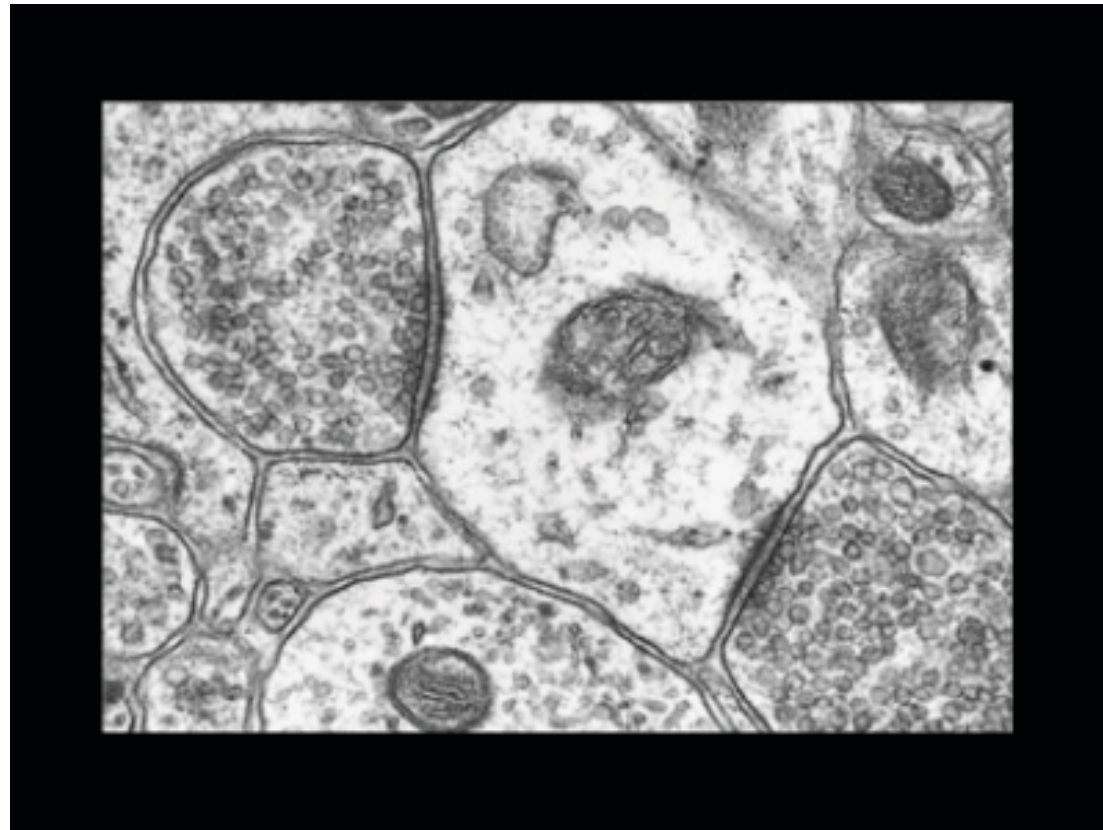


# Excitatory and inhibitory signals





# Synaptic transmission – chemical synapse



*Movie from: Molecular Biology of the Cell, 5<sup>th</sup> Edition, Alberts et al.*

# The process of synaptic transmission – form and function

1. Action potential propagates down the axon – to the pre-synaptic knob
2. Pre-synaptic knob is depolarised – voltage gated  $\text{Ca}^{2+}$  channels open
3.  $\text{Ca}^{2+}$  ions enter and TRIGGER the release of the neurotransmitter from the vesicles
4. Neurotransmitter is released INTO the synaptic cleft
5. Neurotransmitter diffuses across the cleft and binds to its SPECIFIC receptors (stimulus gated channels) on the POST SYNAPSE
6. If  $\text{Na}^+$  channels open – LOCAL depolarisation of post synaptic cell
7. Net depolarisation – called the

EXCITATORY POST SYNAPTIC POTENTIAL – or EPSP

- 7'. Net hyperpolarisation (opening of  $\text{Cl}^-$  or  $\text{K}^+$  channels) – called the

INHIBITORY POST SYNAPTIC POTENTIAL – or IPSP

# REMEMBER

- Depolarisation – Excitatory
- Hyperpolarisation – Inhibitory

.... Depending on neurotransmitter and targeted receptor  
(Na<sup>+</sup>, Cl<sup>-</sup> or K<sup>+</sup> channel)

# HOW IS A SYNAPSE SWITCHED OFF ?

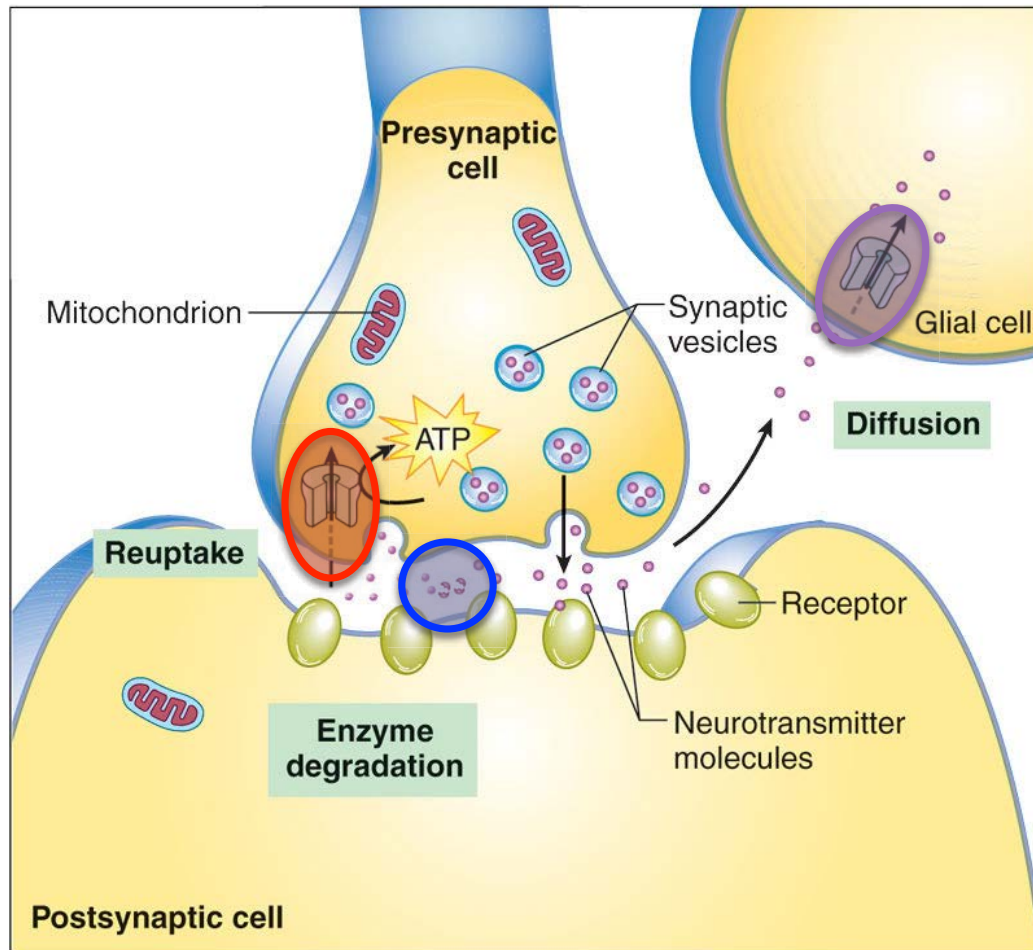


Fig. 13-26, Patton & Thibodeau, 8<sup>th</sup> Ed p. 402

- Excess transmitter released into the cleft.
- Excess transmitter must be removed
    - Degradation** – by enzymes
    - Reuptake** into the knob
    - Reuptake (diffusion) into **glia cells**  
(GLM3 – revisit)
  - Removal requires ATP – energy
  - Mitochondria in synaptic knob ...

Glia – the supporting cells – and revisit p. 384 T&P 8<sup>th</sup> Ed.

# HUBS191

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