

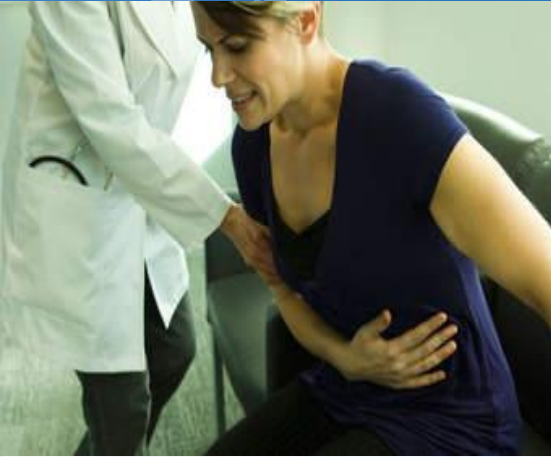


بِسْمِ اللَّهِ الرَّحْمَنِ الرَّحِيمِ

اللهم صل على محمد وآل محمد

السلام عليكم ورحمة الله وبركاته

Physiology of



*Dr. Hayam Gad
Associate Professor of
Physiology
College of Medicine, K S U*

Learning Objectives

What is pain and its significance

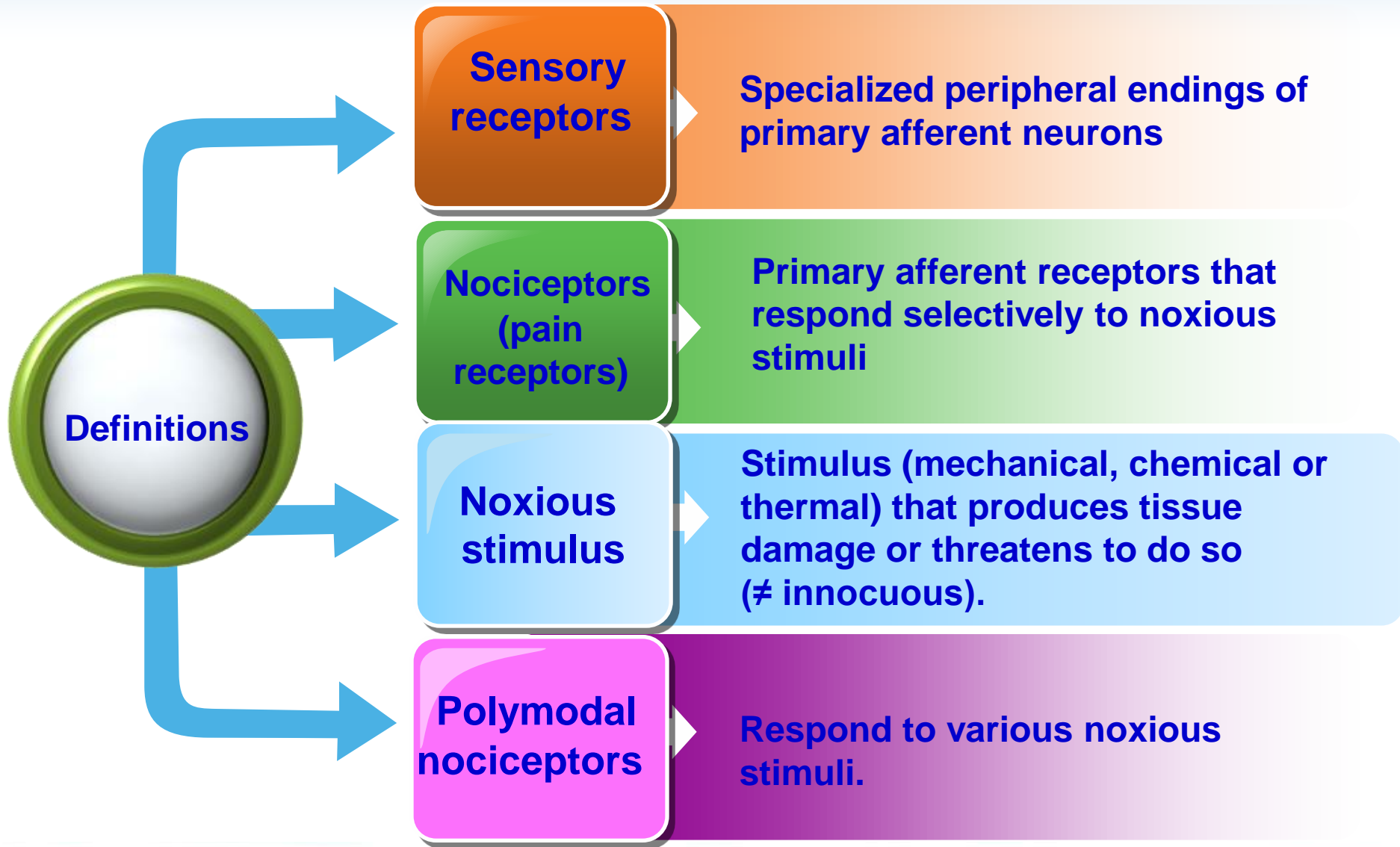
Pain reception and perception

Pain receptors & mechanism of stimulation

Qualities & types of pain

Pathway of pain

Role of cerebral cortex in pain perception

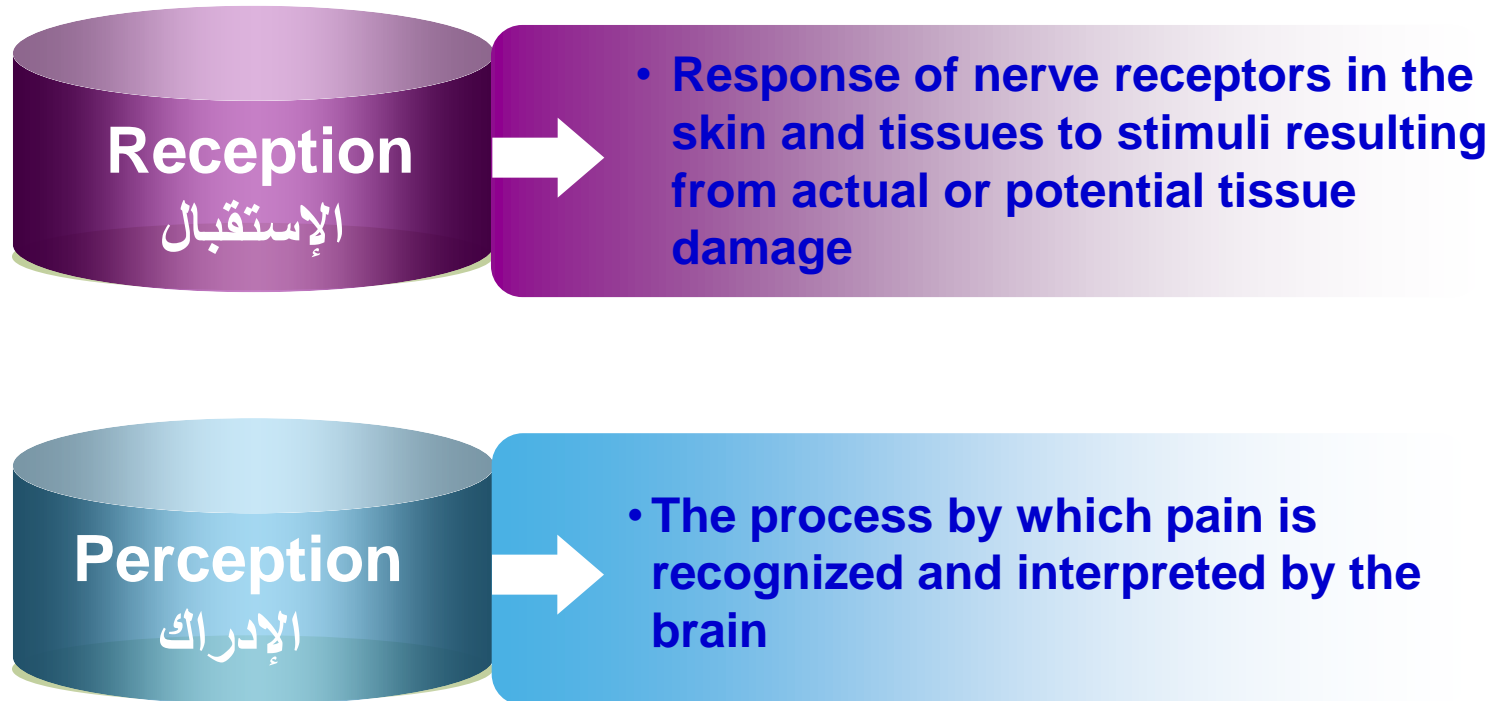


Significance of Pain: Why do we feel pain?

- It is a protective mechanism meant to make us aware that tissue damage is occurring or is about to occur:-
 - Avoid noxious stimuli
 - Remove body parts from danger
 - Promote healing by preventing further damage
 - Storage of painful experiences in memory to avoid potentially harmful event in the future
- The sensation of pain may be accompanied by behavioral responses (withdrawal, defense) as well as emotional responses (crying, anxiety or fear).
- Pain is perceived at both the cortical & thalamic levels.



Pain Reception and Perception



Pain & Nociception

Nociception

Pain

What is?

Transmission of signals evoked by activation of nociceptors from periphery to the CNS.

Perception of unpleasant sensation that originates from a specific body region

Pain is

an unpleasant sensory and emotional experience associated with actual or potential tissue damage.

Pain Receptors 'Nociceptors'?

“ Are special receptors that respond only to noxious stimuli and generate nerve impulses which the brain interprets as "pain".

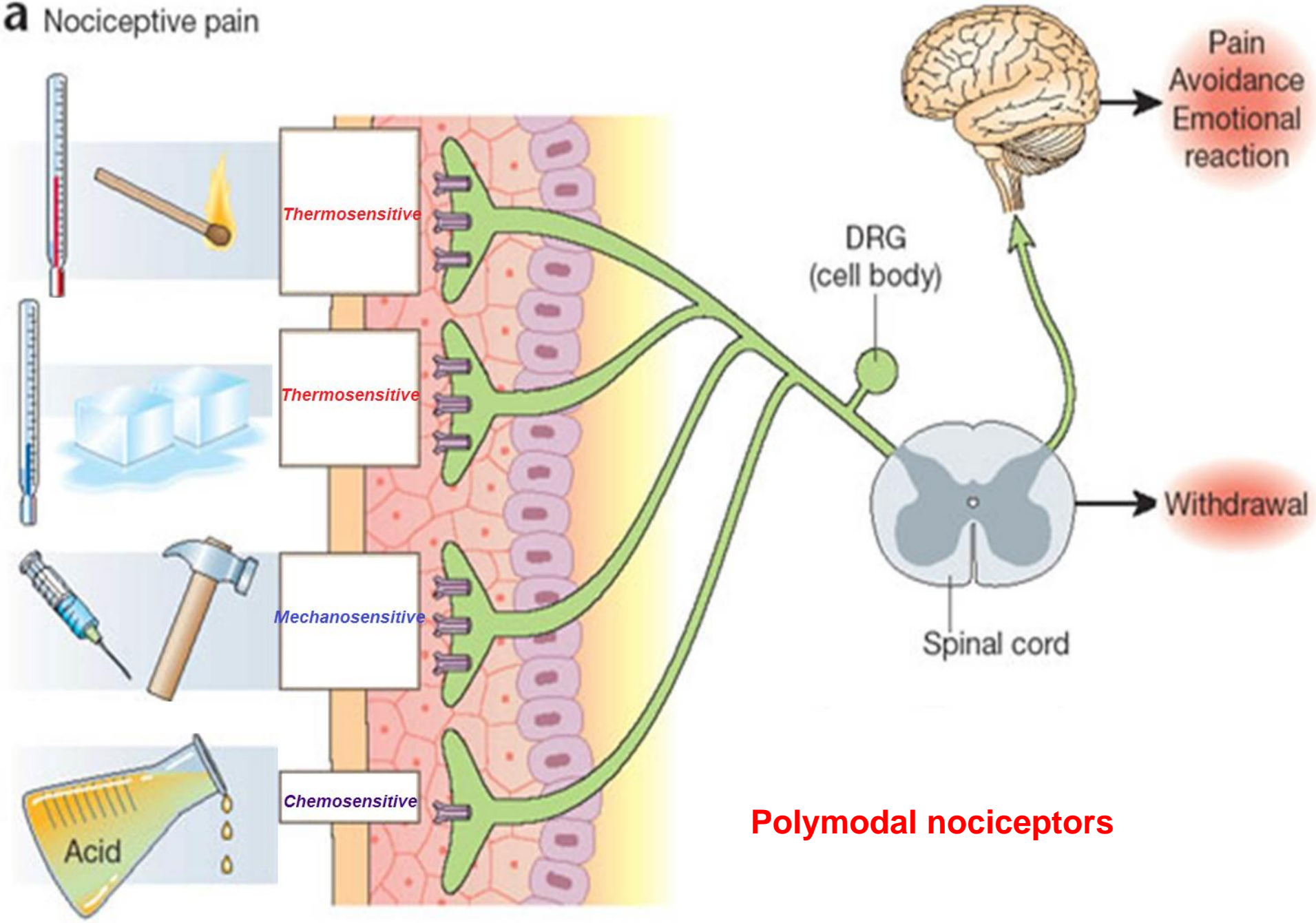
(Sherrington 1906)



Sir Charles Scott
Sherrington

- *They are the most widely distributed.*
- *They are specific (have adequate stimulus) in that pain is not produced by overstimulation of other receptors.*
- *They are high threshold receptors i.e. painful stimuli must be strong & noxious to produce tissue damage.*
- *Do not adapt (or very little) to repetitive stimulation (it allows the pain to keep the person apprised of a tissue-damaging stimulus as long as it persists.)*

a Nociceptive pain



Thermosensitive

Thermosensitive

Mechanosensitive

Chemosensitive

DRG
(cell body)

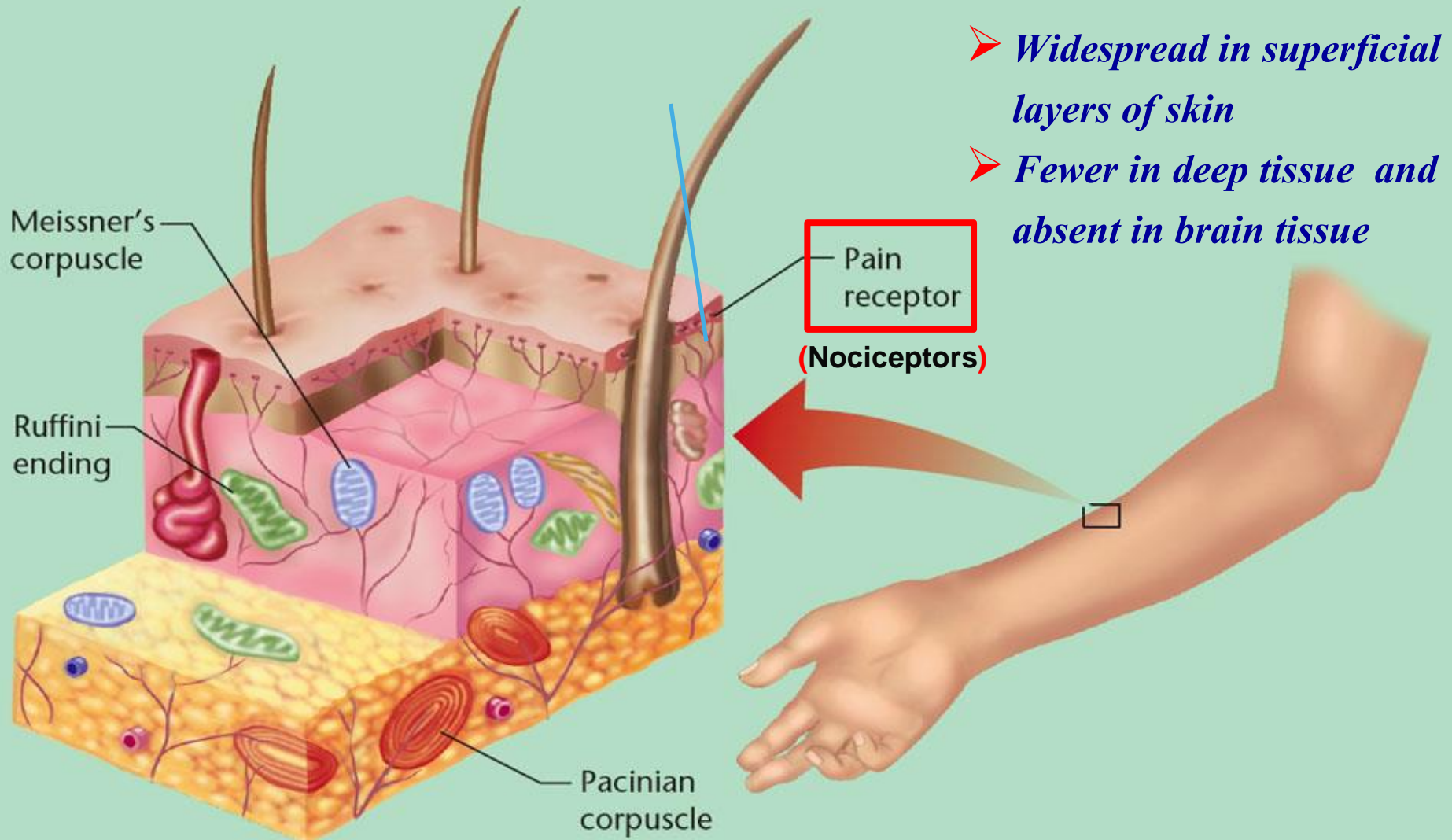
Spinal cord

Pain
Avoidance
Emotional
reaction

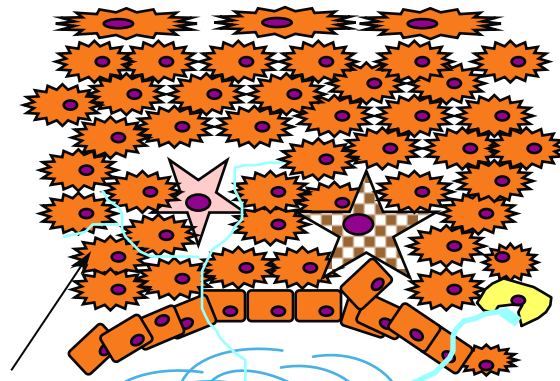
Withdrawal

Polymodal nociceptors

Distribution of Pain Receptor(Nociceptors)



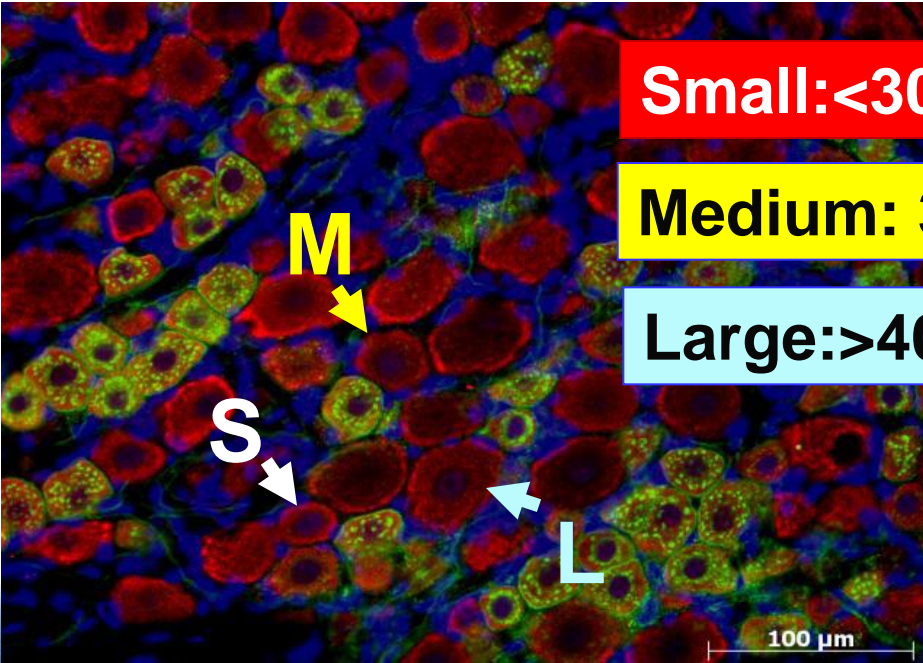
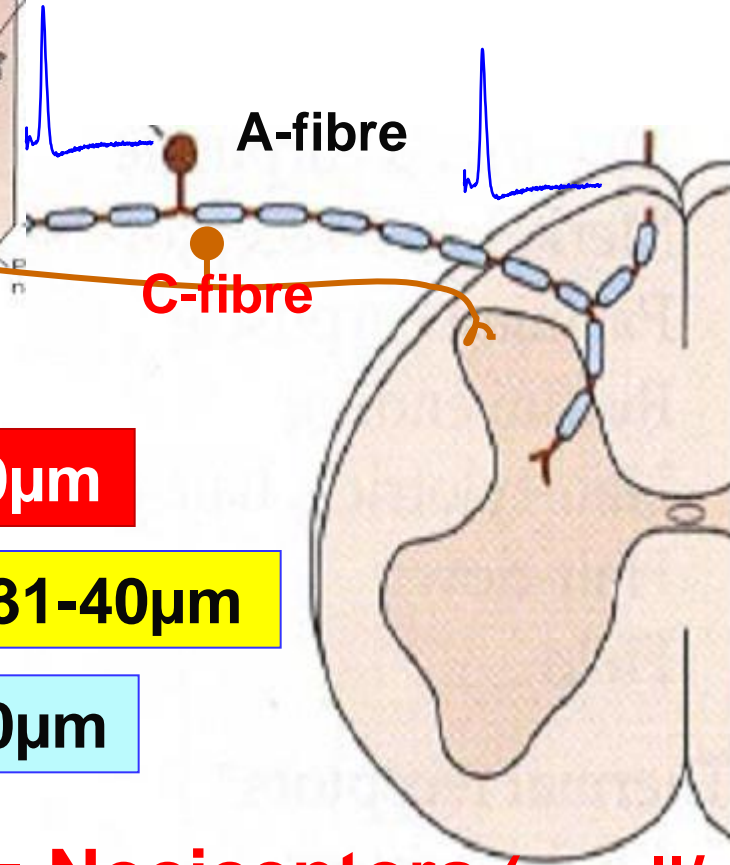
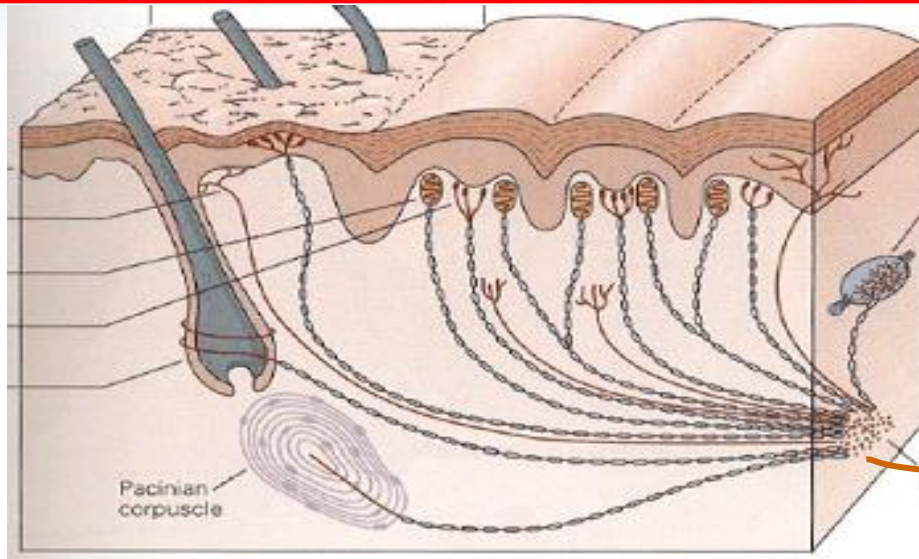
○ *All pain receptors are free nerve endings of unmyelinated **C fibers** & small diameter myelinated **A δ fibers**.*



Free endings

Pain receptors

Type-A & Type-C Fibers



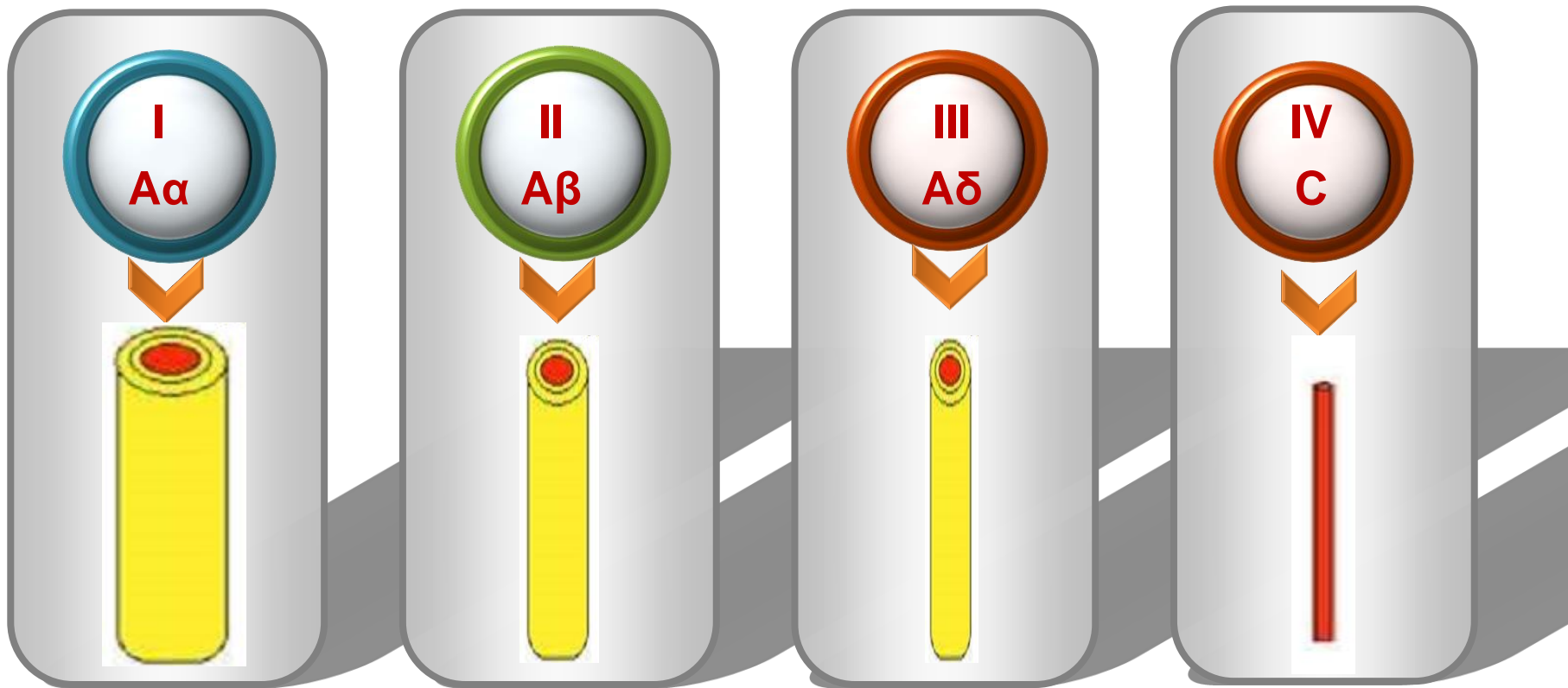
Small: <math><30\mu\text{m}</math>

Medium: 31-40μm

Large: >40μm

- Nociceptors (small/medium)
- Non-nociceptors (large)

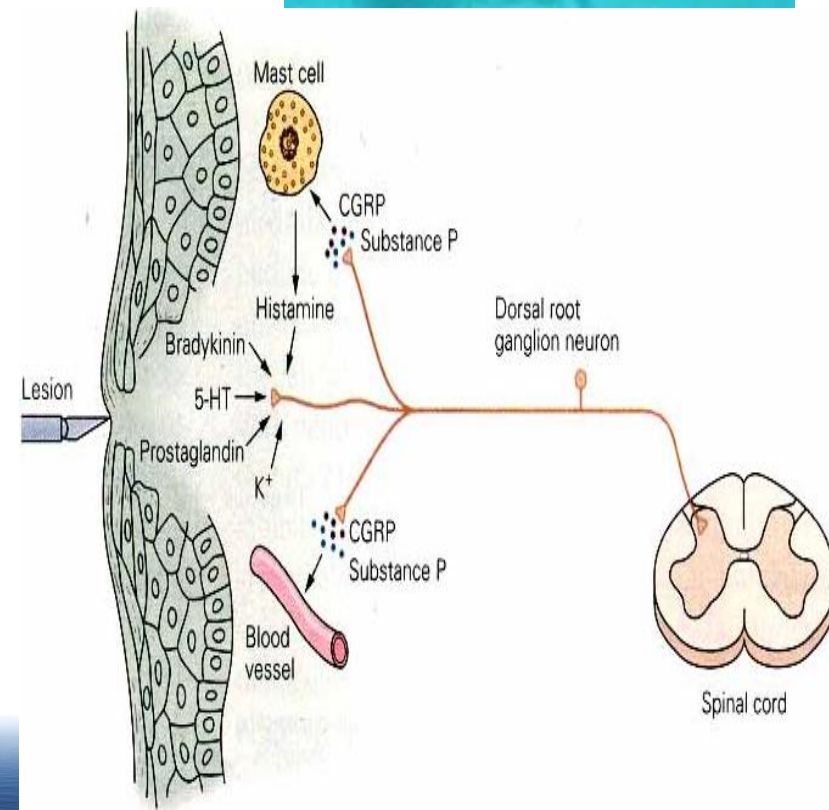
Classification of Nerve fibres



Diameter (μm)	10-20	5-10	2-5	0.5-2
Conduction Velocity (m/s)	70-120	30-70	5-30	0.5-2

Mechanism of stimulation of nociceptors

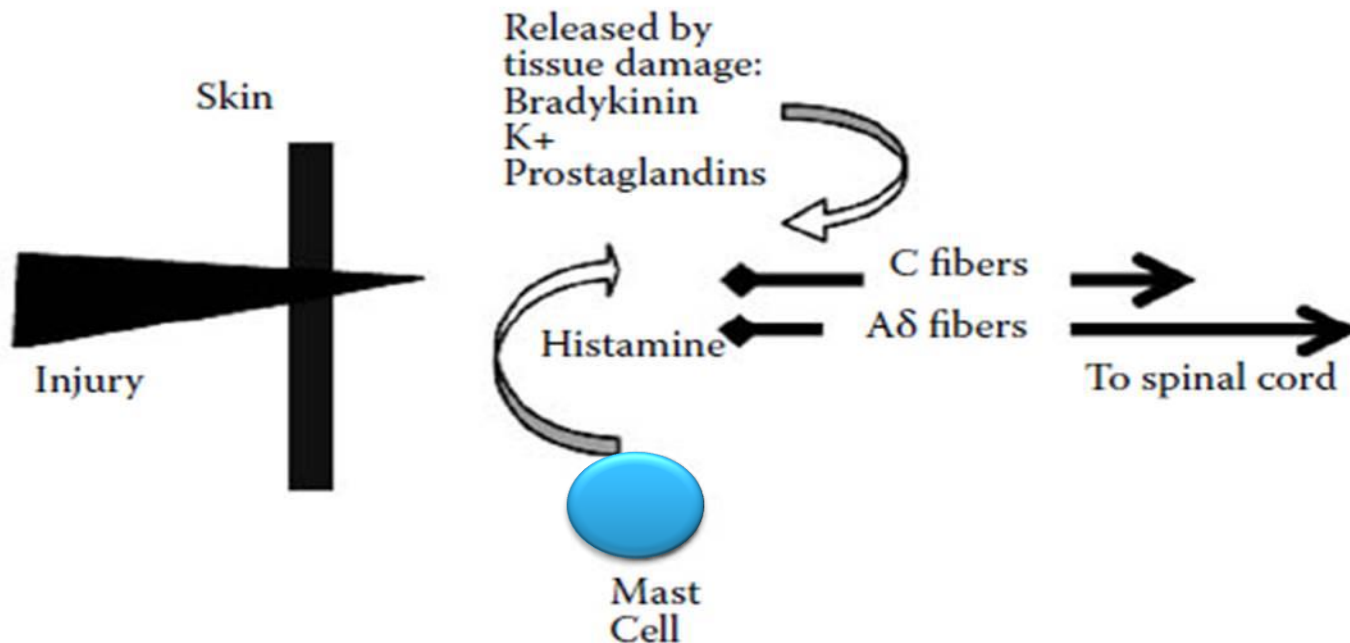
- ❖ Pain receptors are depolarized either directly or through the production of pain producing substances (inflammatory mediators) from damaged tissues
- ❖ e.g. bradykinin, histamine, substance P, calcitonin gene-related peptide (CGRP), interleukins, prostaglandins, K^+ , Ach, proteolytic enzymes.



Chemical substances released during tissue damage

<i>Substance</i>	<i>Source</i>
<i>Potassium</i>	<i>Damaged cells</i>
<i>Serotonin</i>	<i>Platelets</i>
<i>Bradykinin</i>	<i>Plasma</i>
<i>Histamine</i>	<i>Mast cells</i>
<i>Prostaglandins</i>	<i>Damaged cells</i>
<i>Leukotrienes</i>	<i>Damaged cells</i>
<i>Substance P</i>	<i>Primary nerve afferents</i>

Pain Mechanism



Some chemicals released by tissue damage that stimulates nociceptors. In addition release of substance-P, along with histamine, produce vasodilation and swelling.

Pain Mechanism

Damage and inflammation release chemical mediators as cytokines, bradykinin, prostaglandin, Substance P

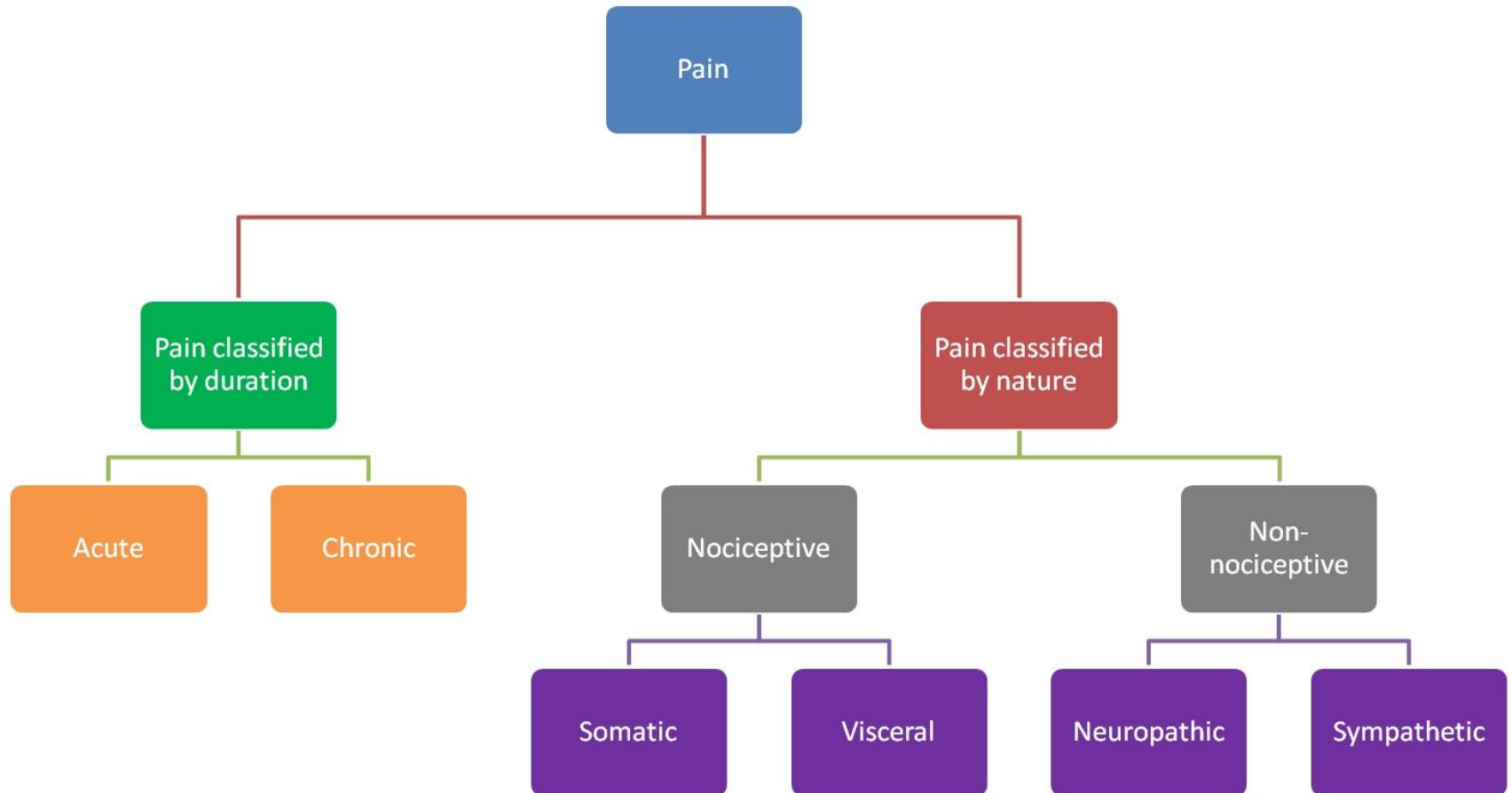


Activate or sensitize the receptor endings

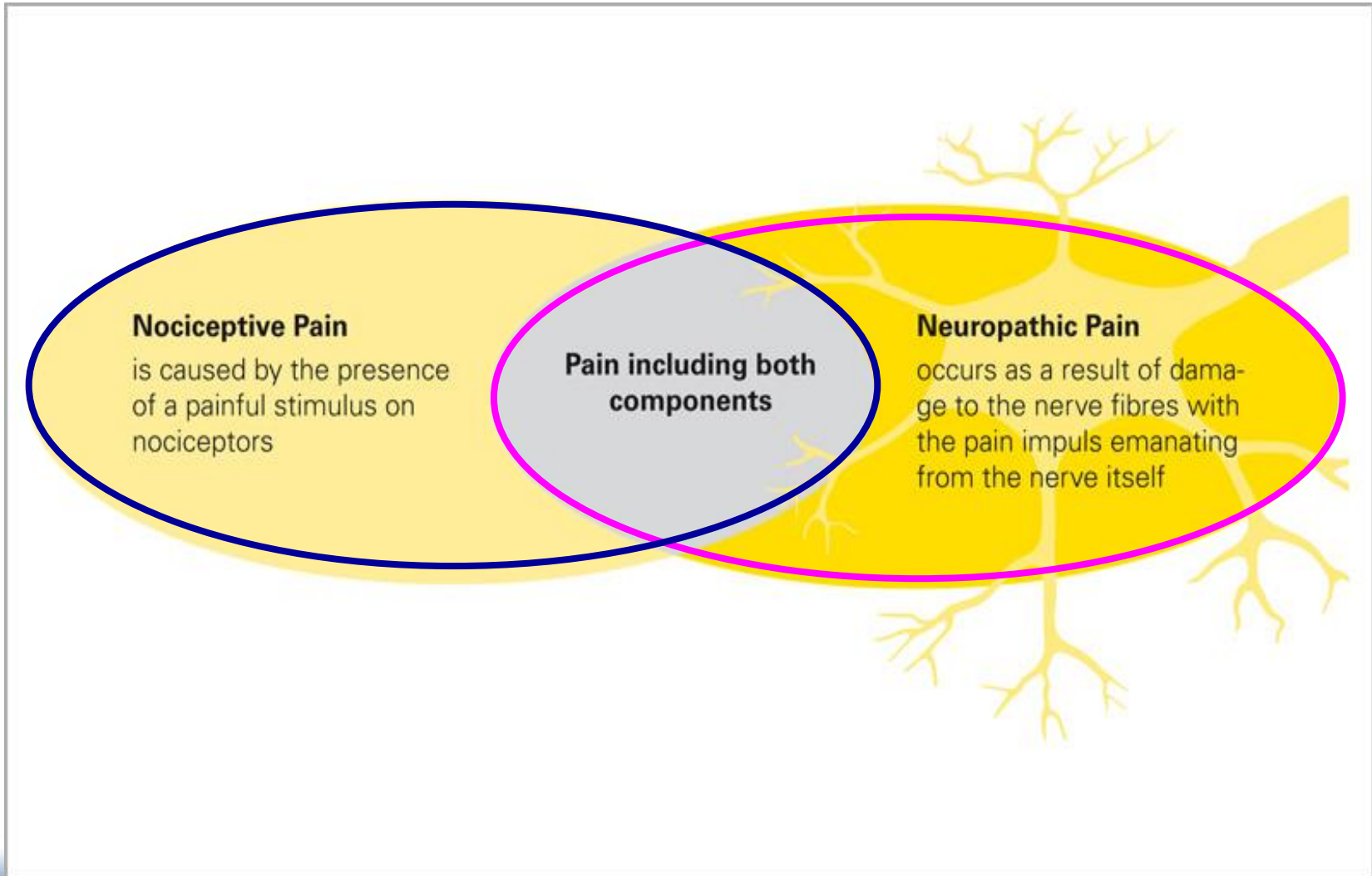


Transduction and conduction of nerve impulse

Classification of Pain



Nociceptive & Neuropathic Pain



Qualities of pain

(Phenomenon of double-pain)

Fast/immediate (1st) pain vs slow/delayed (2nd) pain

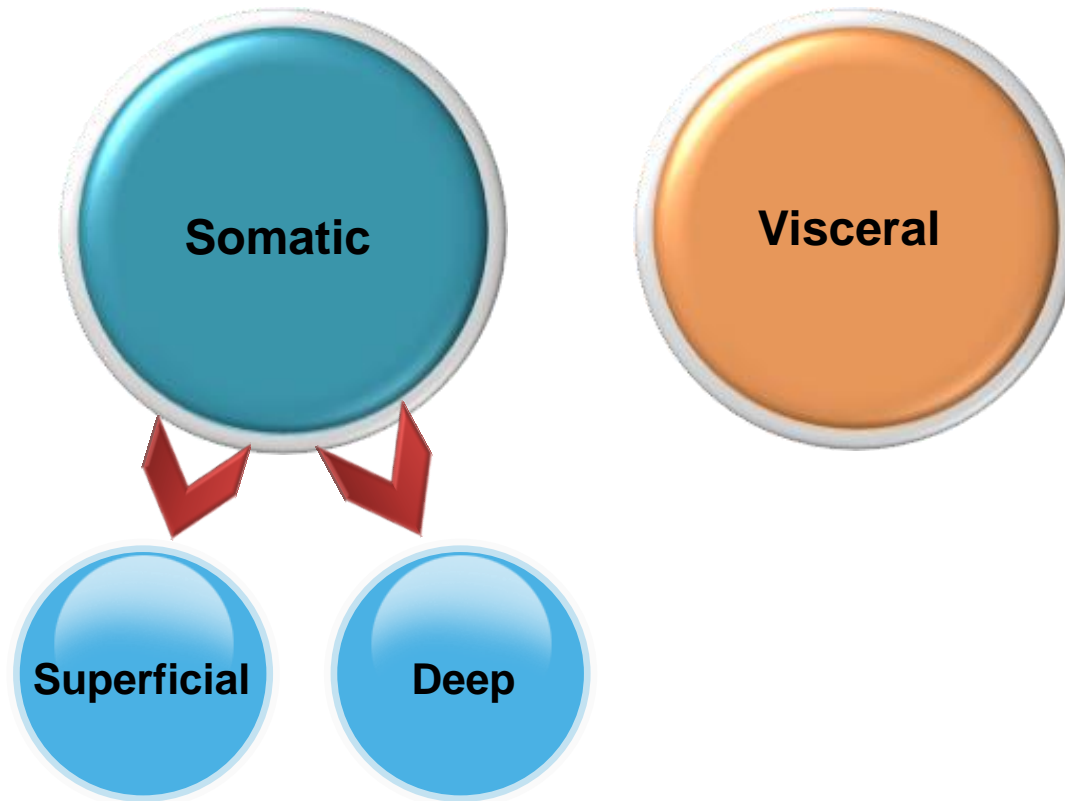
Fast (immediate) pain

Slow (second) pain

- | <i>Fast (immediate) pain</i> | <i>Slow (second) pain</i> |
|---|---|
| <ul style="list-style-type: none">■ <i>Sharp, intense, pricking</i>■ <i>Felt within 0.1 sec</i>■ <i>Associated with reflex withdrawal</i>■ <i>Usually somatic not visceral</i>■ <i>Well localized and is mediated by $A\delta$-fiber nociceptors</i>■ <i>Terminate at I and V laminae</i>■ <i>Neurotransmitter – glutamate</i> | <ul style="list-style-type: none">■ <i>Burning, aching, throbbing “unbearable” diffuse, dull, or chronic pain</i>■ <i>Felt after 1 sec or more</i>■ <i>Associated with destruction of tissue</i>■ <i>Can occur in skin or any internal organ/tissue</i>■ <i>Poorly localized and is mediated by C-fiber nociceptors: → misery (responsible for emotional aspect of pain)</i>■ <i>Terminate at II and III laminae</i>■ <i>Neurotransmitter – Substance- P</i> |

Pain

- Pain can be classified according to the site of stimulation into:-



Superficial Pain

Arises from skin or other superficial structures

Occurs in 2 phase (fast pricking , slow burning pain)

Can be well localized

Associated with motor, autonomic, emotional reactions

Characteristics

Deep Pain

*Arises from muscles,
joints, periosteum,
tendons & ligaments*

*Diffuse, slow
prolonged
conducted by
type C fibers*

*May be referred,
initiate reflex
contraction of
nearby muscles.*

*Caused by: trauma,
bone fracture,
inflammation,
arthritis, muscle
spasm & ischemia*

Characteristics

Visceral pain

- *There are few pain receptors in most viscera*
- *Some viscera are pain insensitive e.g. **liver**
parenchyma, **lung alveoli**, **brain tissue**,
visceral layer of peritoneum, **pleura** and
pericardium.*

Visceral Pain

*Slow, diffuse,
poorly localized,
conducted by C
fibers*

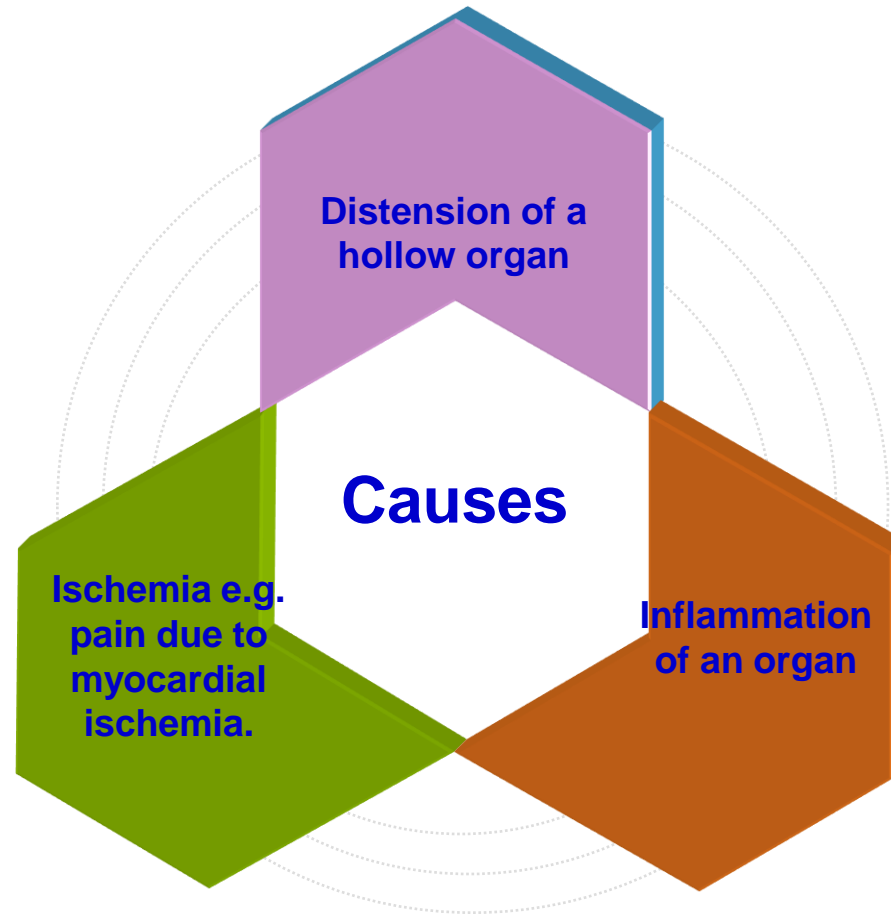
*Pain arising from
parietal peritoneum,
pleura & pericardium
is sharp, pricking type*

*Often referred
associated with
rigidity of nearby
muscles and
autonomic reactions*

*caused by:
distension,
inflammation or
ischemia*

Characteristics

Causes of visceral pain



N.B: Cutting, crushing are not painful when applied to viscera

Referred pain

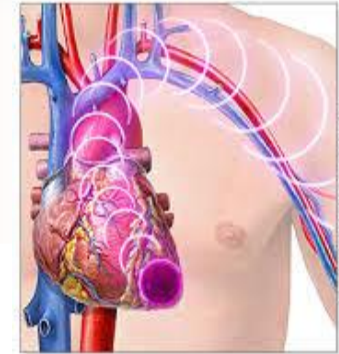
- *This is pain that is felt away from its original site.*
- *It is most frequent with visceral pain & deep somatic pain but **cutaneous pain is not referred.***
- *Pain is referred according to dermatomal rule.*

Examples of referred pain



Cardiac pain

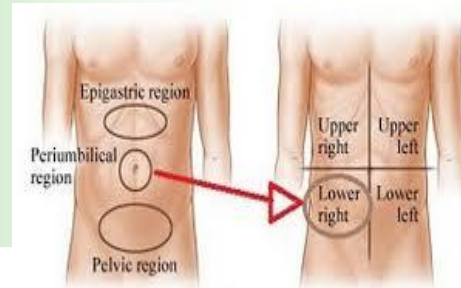
Is referred to the jaw, left shoulder & inner side of left arm



Pain in the chest radiating up to the jaw or down the left (or, less often, right) arm might signal a heart attack

Pain of appendicitis

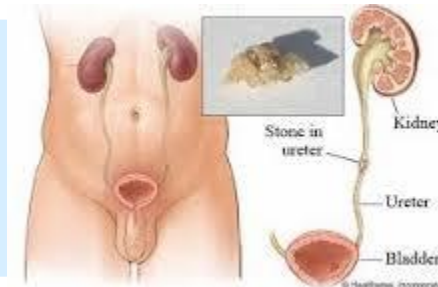
Is referred to periumbilical region



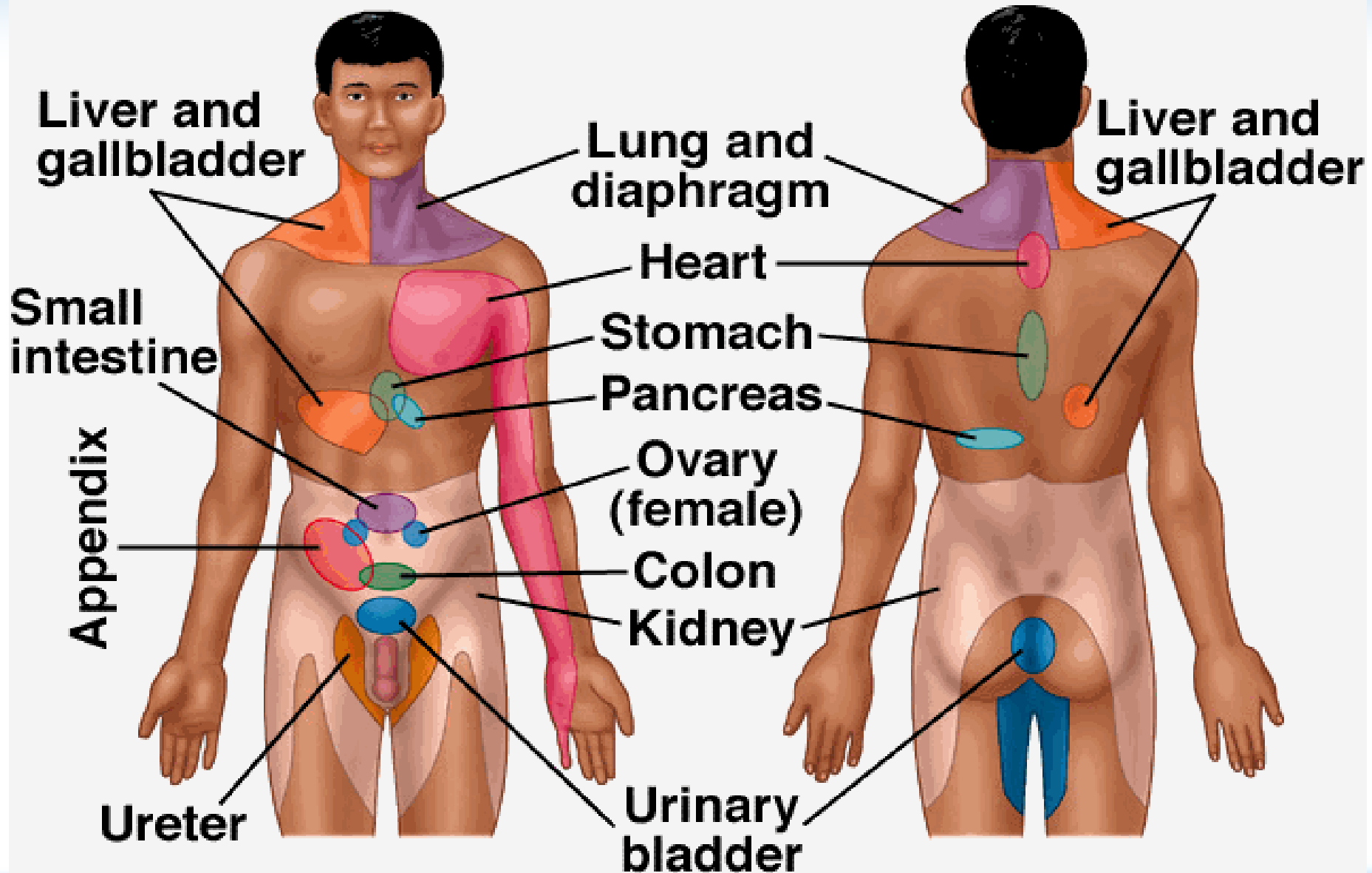
Progression of Pain in Appendicitis

Pain from ureter

Is referred to testicular region



Referred Pain Regions

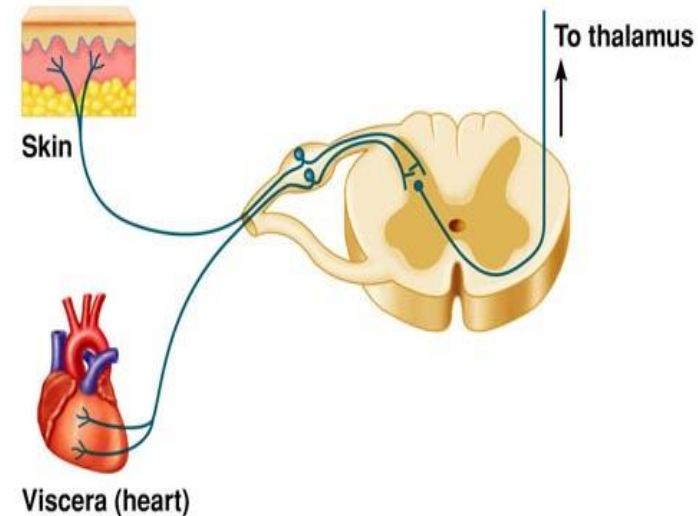


<i>Organ</i>	<i>Site of referred pain</i>
<i>Meninges</i>	<i>Back of head & neck</i>
<i>Heart</i>	<i>Central chest, left arm</i>
<i>Diaphragm</i>	<i>Shoulder tip</i>
<i>Esophagus</i>	<i>Behind sternum</i>
<i>Stomach, duodenum</i>	<i>Epigastrium</i>
<i>Small bowel, pancreas</i>	<i>Around umbilicus</i>
<i>Large bowel, bladder</i>	<i>Lower abdomen</i>
<i>Kidney</i>	<i>Loin</i>
<i>Ureter</i>	<i>Testicles</i>
<i>Trigon of bladder</i>	<i>Tip of penis</i>
<i>Hip</i>	<i>Knee</i>
<i>Appendix</i>	<i>Umbilicus</i>
<i>Uterus</i>	<i>Low back</i>

Mechanism of referred pain

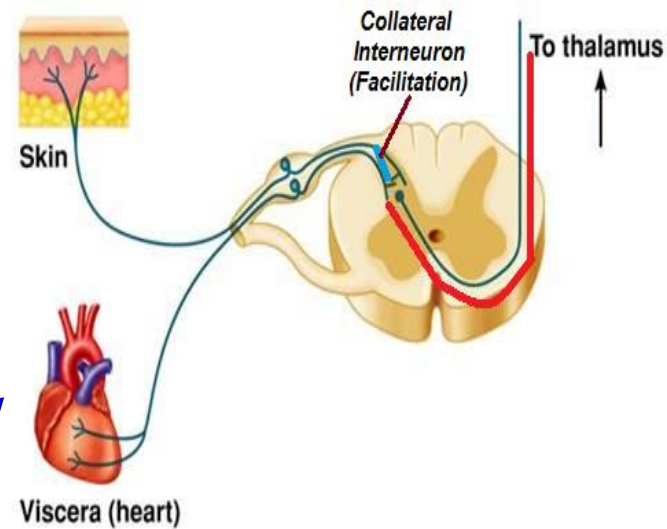
Convergence theory

- *Afferent pain fibers from skin area & diseased viscera that develop from same embryonic segment converge on same 2nd order neuron and finally stimulate the same cortical neuron.*
- *The brain interprets the information coming from visceral nociceptors as having arisen from cutaneous nociceptors, because this is where nociceptive stimuli originate more frequently*



Facilitation theory

- *Pain fibers from skin are always carrying impulses, not enough to produce pain.*
- *Impulses from diseased viscus pass through afferents which give collaterals to ST neurons receiving pain fibers from skin.*
- *As a result, ST neurons' excitability is raised (they are facilitated) to reach a threshold level.*
- *The signals reaching the brain are projected to skin area and pain is felt in skin dermatome*



Pathway of Pain

Pain sensation is carried by lateral spinothalamic tracts which includes 2 separate pathways:-

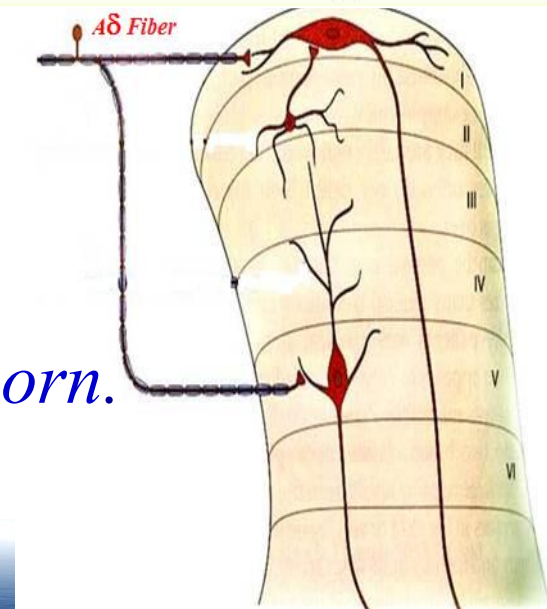
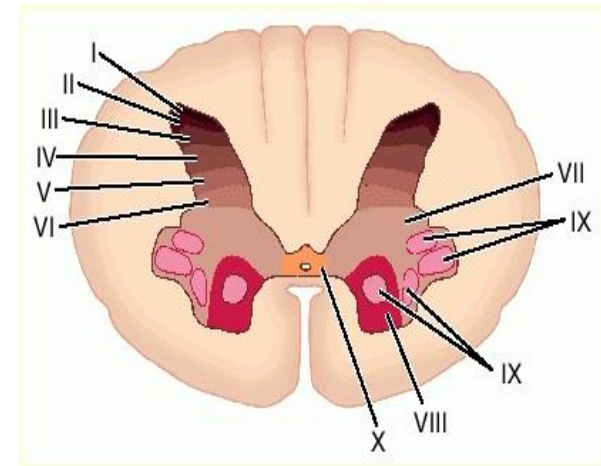
A) The neospinothalamic pathway:

This transmits fast pain.

○ First order neurons

Are mainly $A\delta$ afferent nerves.

They terminate at lamina I & V of dorsal horn.

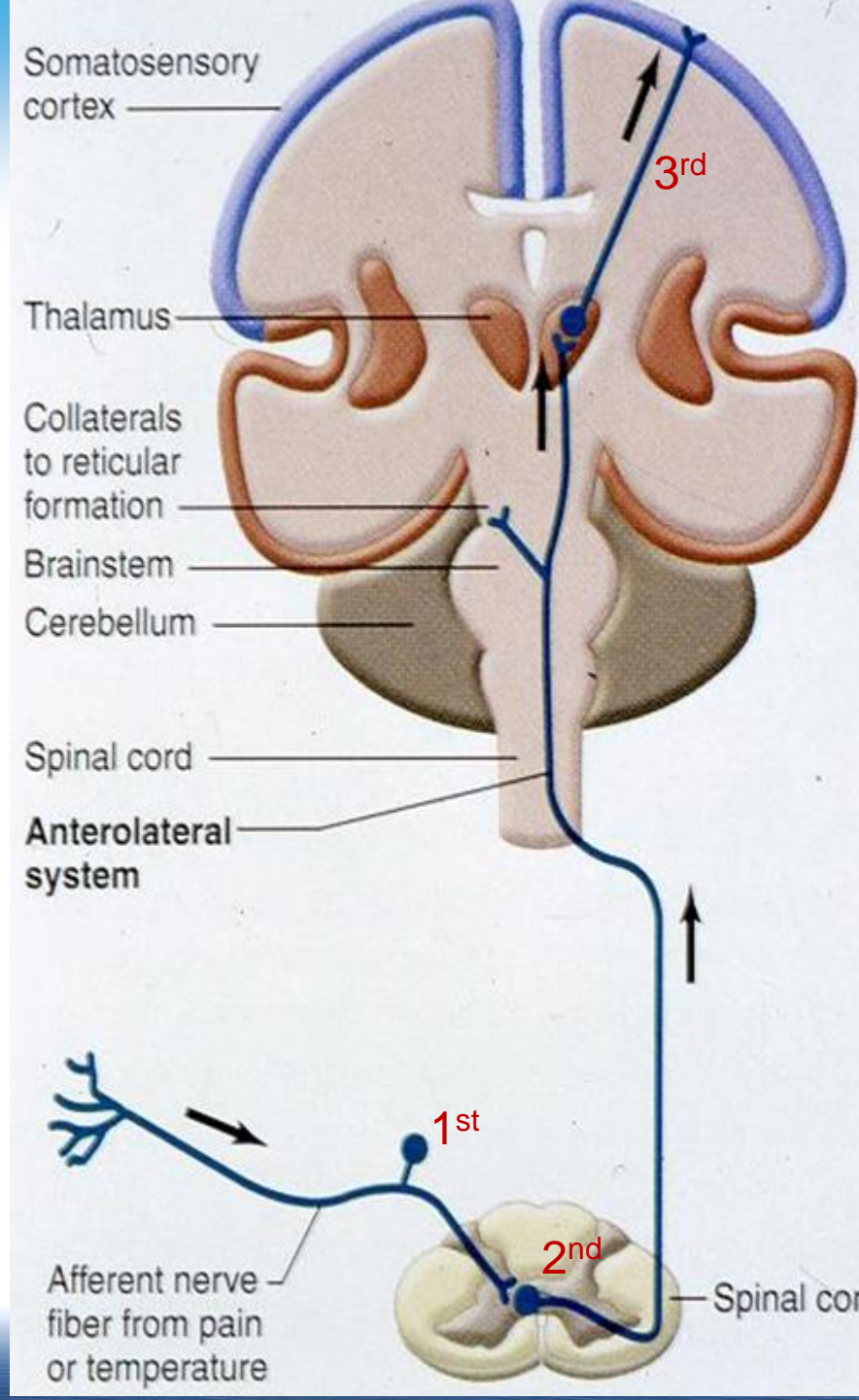


○ Second order neurons

These constitute the tract. They start at dorsal horn, cross to opposite side and ascend in lateral column of spinal cord. The fibers ascend in brain stem to terminate in ventrobasal complex of thalamus.

○ Third order neurons

These start at thalamus & most fibers project to somatosensory cortex.



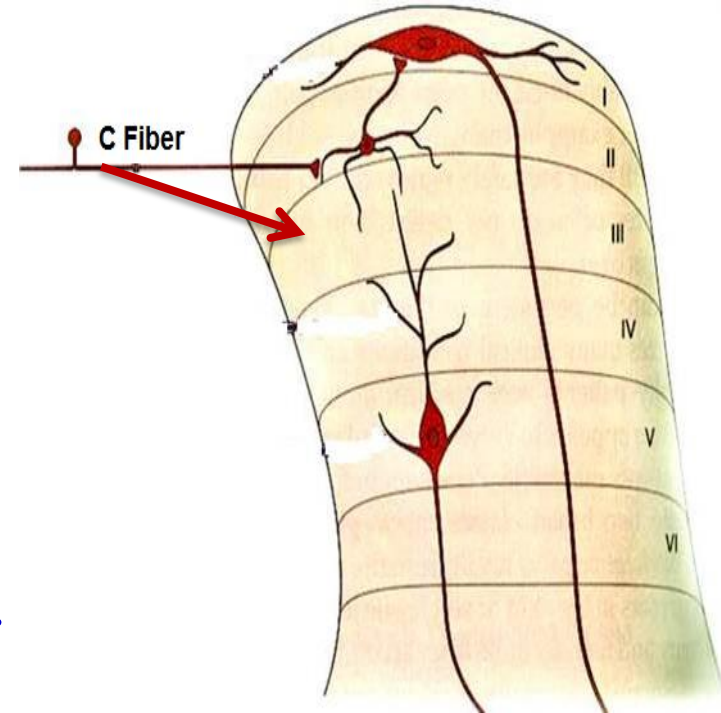
B) The paleospinothalamic pathway:

This transmit slow pain sensation.

○ First order neurons

*They are mainly **type C fibers**.*

They enter spinal cord via dorsal roots, terminate at substantia gelatinosa in laminae II & III of dorsal horn(substantia gelatinosa).



○ Second order neurons

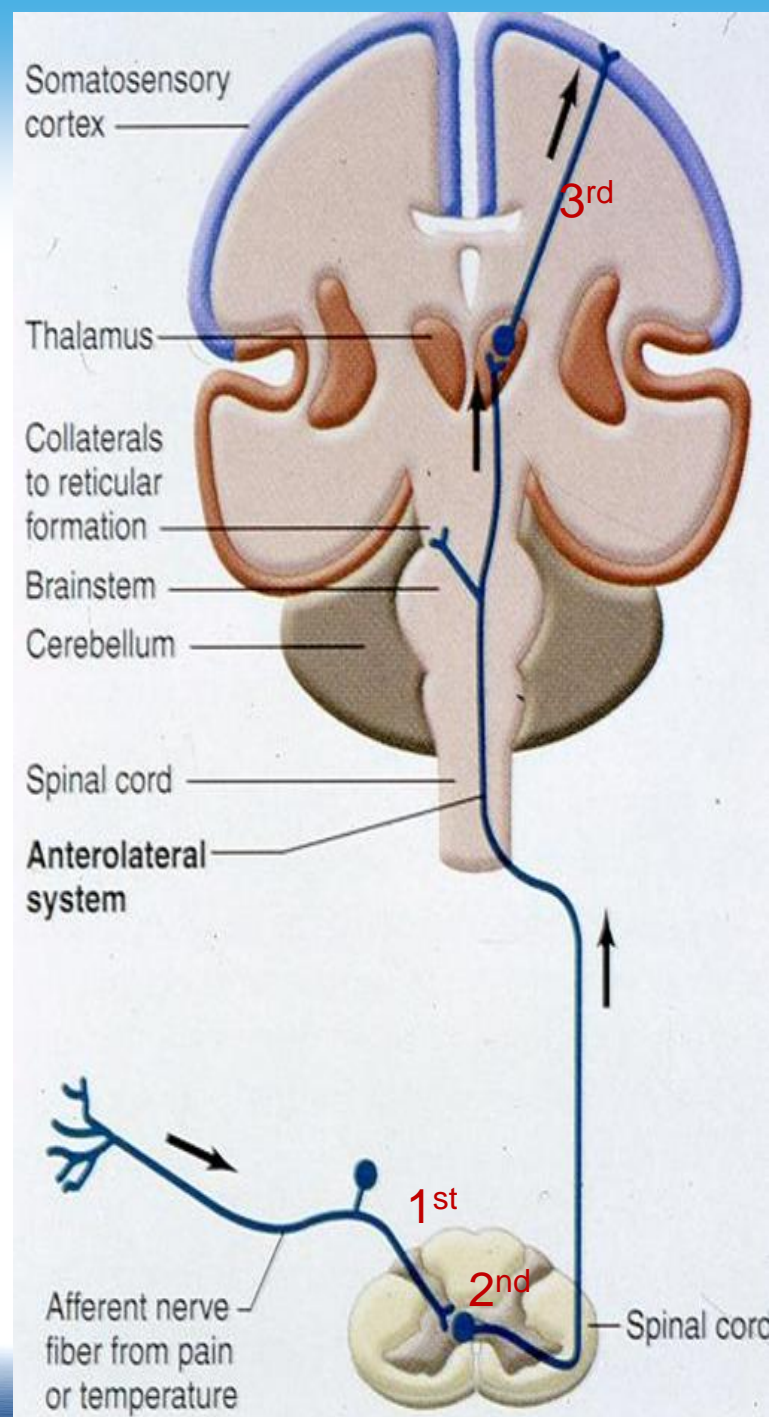
They start at SGR, cross to opposite side in front of central canal, ascend in lateral column of SC & terminate at:-

- *Reticular formation of brain stem.*
- *Intralaminar nuclei of thalamus.*
- *Hypothalamus & adjacent region of basal brain.*

Impulses arriving these regions have strong arousal effects and can be perceived.

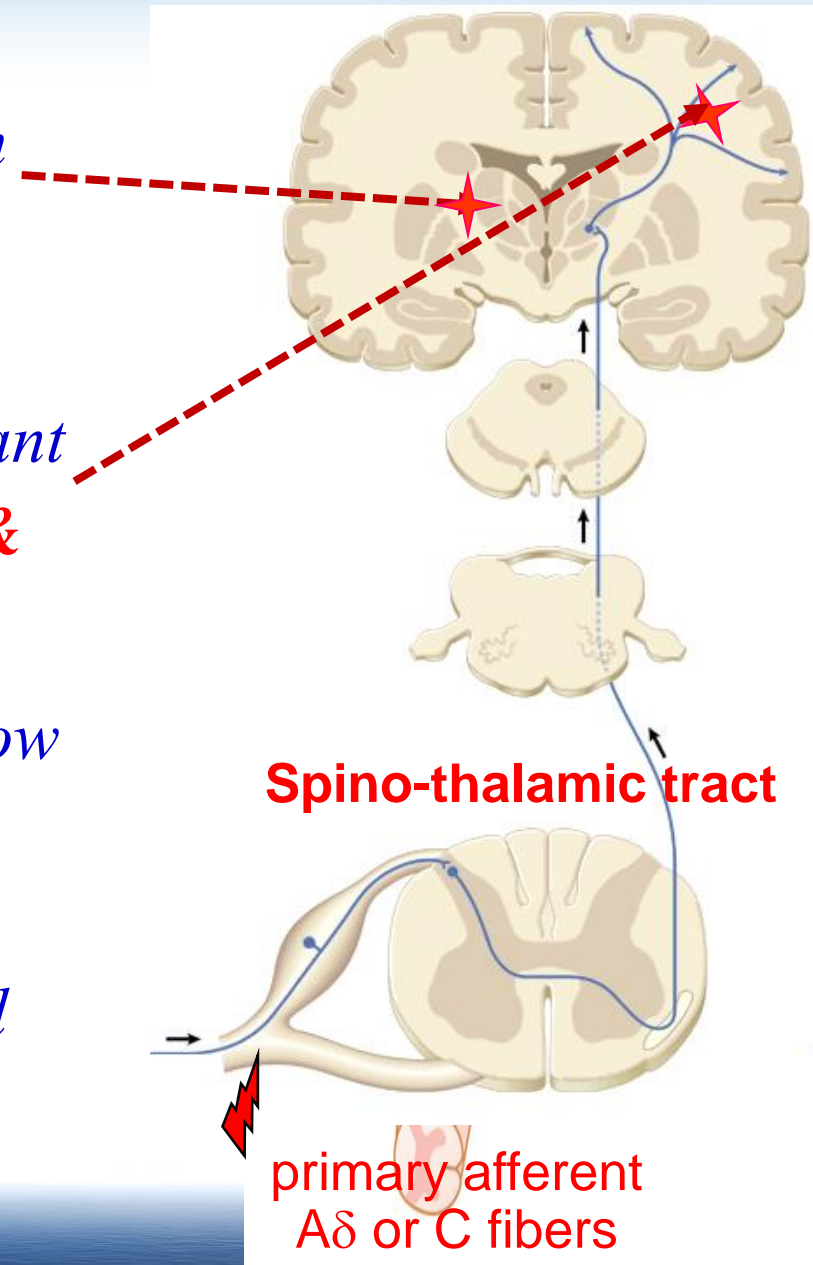
○ Third order neurons

- *These start at thalamus,*
- *Few fibers project to cerebral cortex.*



Role of cerebral cortex in pain perception

- *Full perception of pain occurs when signals enter RF of brain stem, thalamus & basal regions.*
- *Somatosensory cortex plays important role in topognosis i.e. **localization & interpretation of pain quality.***
- *Fast pain is localized better than slow pain because signals carried in neospinothalamic tract reach somatosensory cortex, while a small proportion of paleospinothalamic pathway reach there.*



Thank You

The text "Thank You" is rendered in a highly decorative, cursive script. The letters are filled with a deep red color and outlined with a bright gold border, giving them a three-dimensional appearance. The text is embellished with clusters of vibrant red roses and green foliage. Two white doves are depicted in flight, one positioned above the letter 'T' and the other above the letter 'Y'. The entire graphic is set against a white background with a soft blue gradient at the top and bottom.