

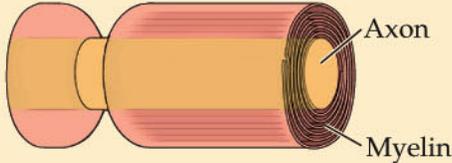
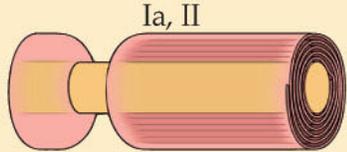
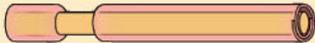
Somatosensory system

(touch and pain)

General, historical

- compelling "Physiology" (vs psychology)
- submodalities (e.g. pain vs hot)
- von Frey (around 1900) punctate sensitivity -
- correlation receptor with sensory experience.
- Muller's (mid-1800's) "doctrine of specific nerve energies"
- the quality comes from the nervous system not the physics of the stimulus.

TABLE 9.1 Somatic Sensory Afferents that Link Receptors to the Central Nervous System

Sensory function	Receptor type	Afferent axon type ^a	Axon diameter	Conduction velocity
Proprioception	Muscle spindle		13–20 μm	80–120 m/s
Touch	Merkel, Meissner, Pacinian, and Ruffini cells		6–12 μm	35–75 m/s
Pain, temperature	Free nerve endings		1–5 μm	5–30 m/s
Pain, temperature, itch	Free nerve endings		0.2–1.5 μm	0.5–2 m/s

^aDuring the 1920s and 1930s, there was a virtual cottage industry classifying axons according to their conduction velocity. Three main categories were discerned, called A, B, and C. A comprises the largest and fastest axons, C the smallest and slowest. Mechanoreceptor axons generally fall into category A. The A group is further broken down into subgroups designated α (the fastest), β , and δ (the slowest). To make matters even more confusing, muscle afferent axons are usually classified into four additional groups—I (the fastest), II, III, and IV (the slowest)—with subgroups designated by lowercase roman letters!

(After Rosenzweig et al., 2005.)

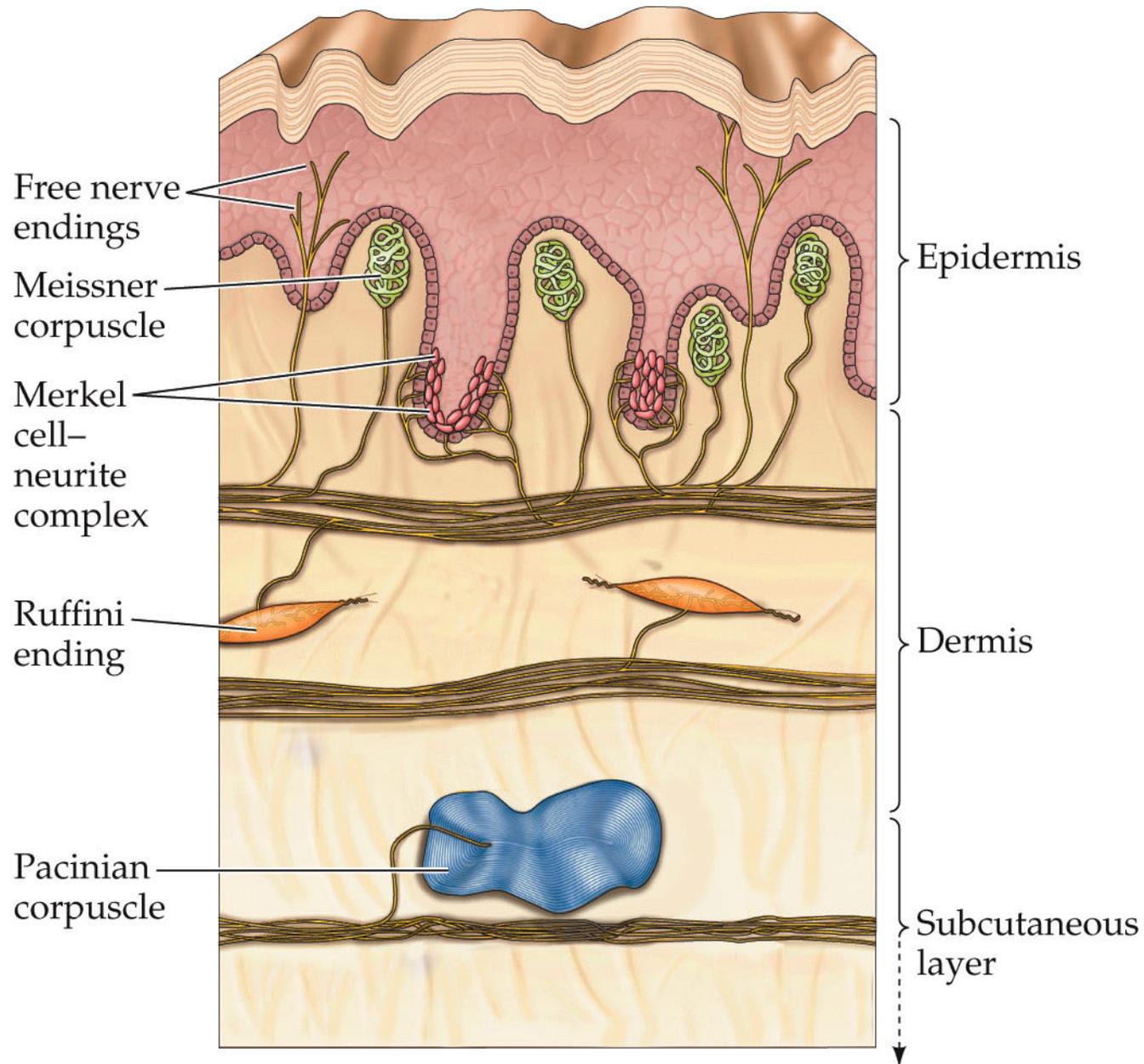
TABLE 9.2 Afferent Systems and Their Properties

	Small receptor field		Large receptor field	
	Merkel	Meissner	Pacinian	Ruffini
Location	Tip of epidermal sweat ridges	Dermal papillae (close to skin surface)	Dermis and deeper tissues	Dermis
Axon diameter	7–11 μm	6–12 μm	6–12 μm	6–12 μm
Conduction velocity	40–65 m/s	35–70 m/s	35–70 m/s	35–70 m/s
Sensory function	Form and texture perception	Motion detection; grip control	Perception of distant events through transmitted vibrations; tool use	Tangential force; hand shape; motion direction
Effective stimuli	Edges, points, corners, curvature	Skin motion	Vibration	Skin stretch
Receptive field area ^a	9 mm ²	22 mm ²	Entire finger or hand	60 mm ²
Innervation density (finger pad)	100/cm ²	150/cm ²	20/cm ²	10/cm ²
Spatial acuity	0.5 mm	3 mm	10+ mm	7+ mm
Response to sustained indentation	Sustained (slow adaptation)	None (rapid adaptation)	None (rapid adaptation)	Sustained (slow adaptation)
Frequency range	0–100 Hz	1–300 Hz	5–1000 Hz	0–? Hz
Peak sensitivity	5 Hz	50 Hz	200 Hz	0.5 Hz
Threshold for rapid indentation or vibration:				
Best	8 μm	2 μm	0.01 μm	40 μm
Mean	30 μm	6 μm	0.08 μm	300 μm

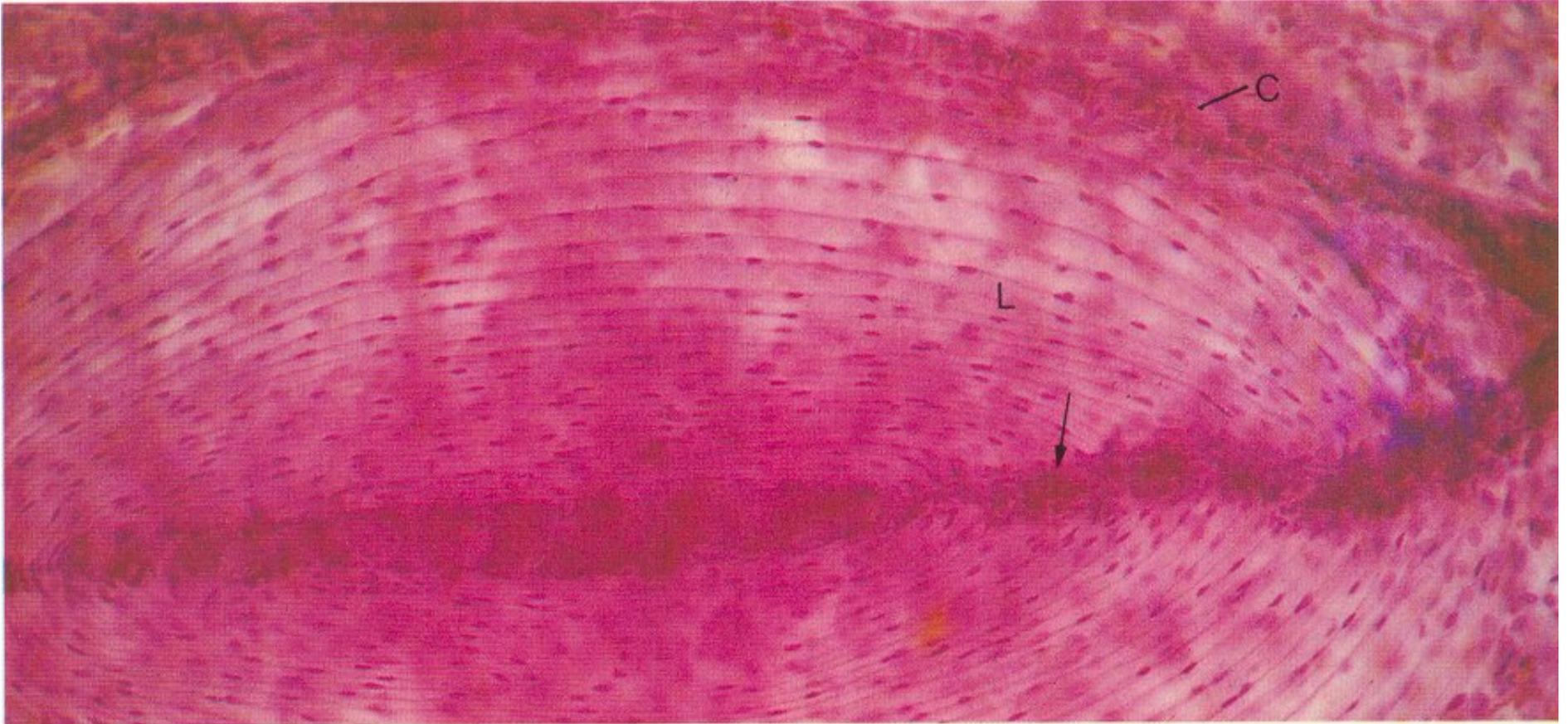
^aReceptive field areas as measured with rapid 0.5-mm indentation.
(After K. O. Johnson, 2002.)

Current thinking

- conduction velocities
- different sizes of myelinated (A) axons, alpha biggest and delta is smallest,
- unmyelinated (C) axons.
- Skin (glabrous, there is also hairy)
- The different types of receptors
- free nerve endings and encapsulated
- Free nerve endings for pain, temperature and crude touch the axons are C fibers (unmyelinated) and A delta, also slow



NEUROSCIENCE, Fourth Edition, Figure 9.5



Vater-Pacini corpuscle H&E Med

C - peripheral capsule; L - surrounding lamellae; arrow - nerve containing core

Loewenstein

- Pacinian corpuscle is rapidly adapting
- because of the layers surrounding the nerve ending
- dissecting off these layers
- Also, there is an electrical adaptation preventing continued spikes after stimulus onset.

Pacinian corpuscle

- rapid adaptation
- A beta
- very sensitive,
- very large receptive field (area which, if stimulated, will affect the receptor [or higher order sensory nerve])
- vibration - 250 - 300 Hz

Meisner's corpuscles

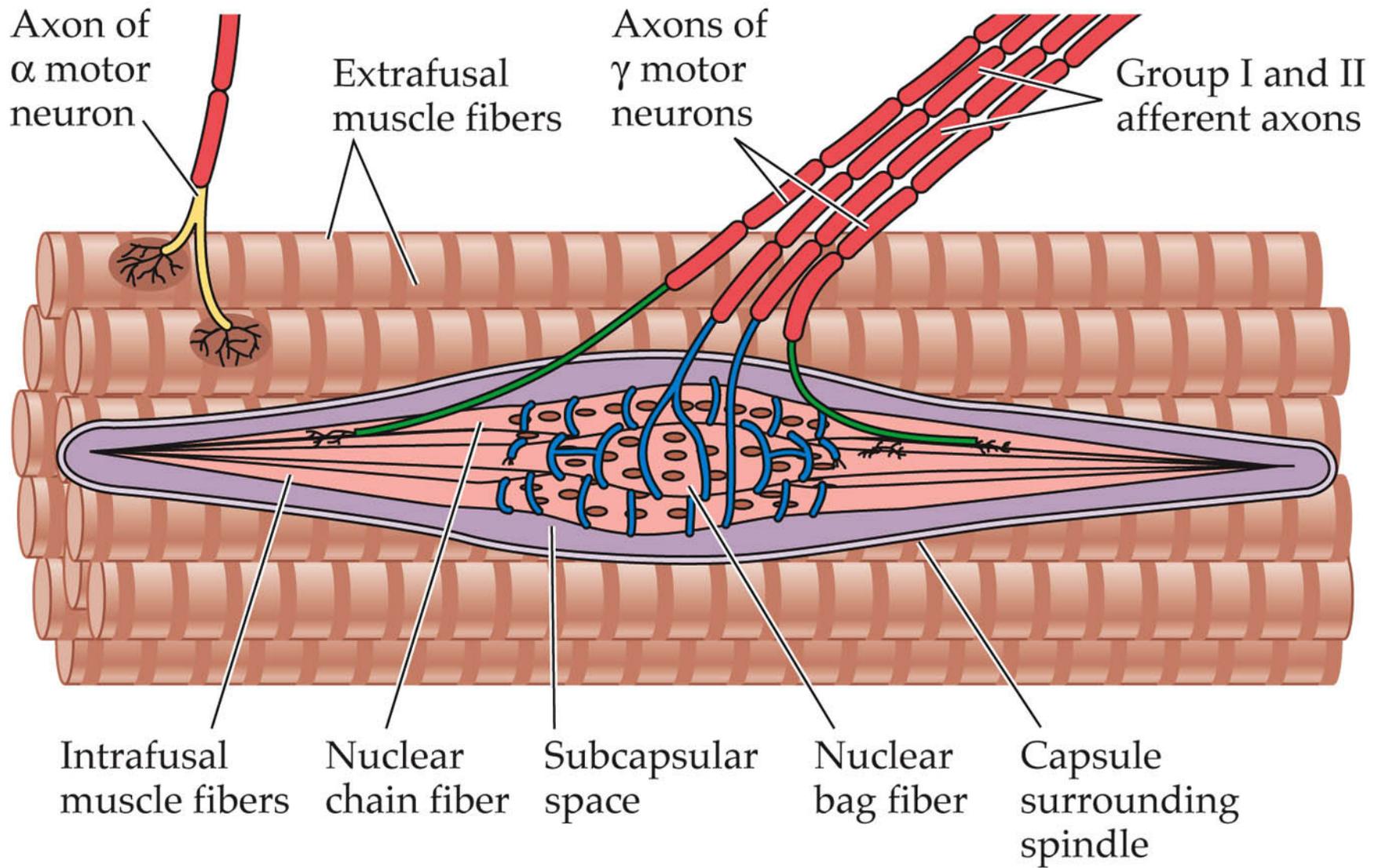
- not as fast as Pacinian
- encapsulation with Schwann cell layers
- most common receptors of fingers, palms and soles
- A beta axons
- smaller receptive field
- "feeling" - active touch - would use fast as finger moves across textured surface

Merkel's disks

- Slow, small receptive field, light touch
- finger tips, lips and genitals
- A beta axons
- static discrimination of shape

Others

- Ruffini slow - large receptive field - sensitive to stretching in deep skin, ligaments and tendons
- A beta axons
- Krauss in lips and genitals (dry vs mucous skin)



Proprioceptors

- muscle spindles (nuclear bag fibers)
- muscle spindle tension presets readiness for reflex,
- gamma motor neurons to intrafusal fibers
- Ia sensory axon
- also Golgi tendon organs Ib afferents

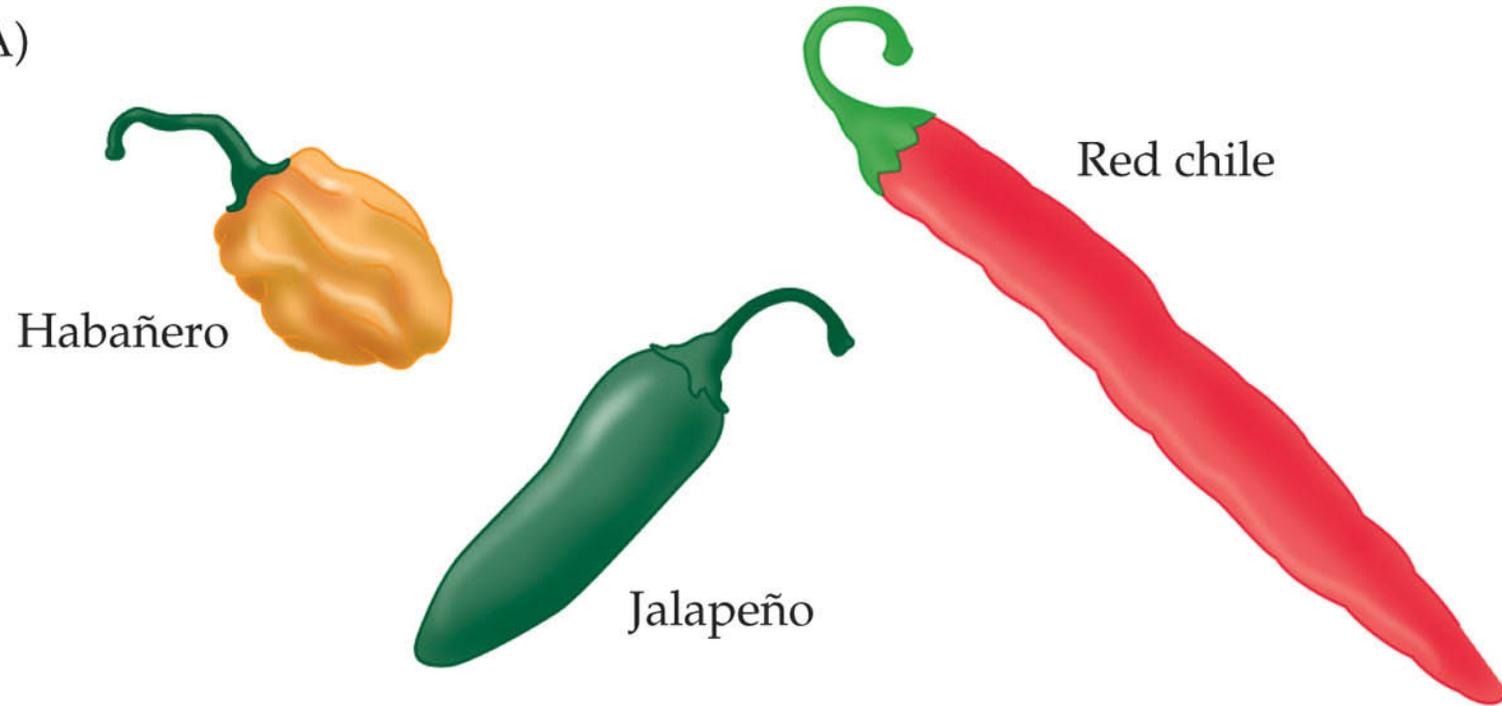
warm and cold

- person can feel a difference of 0.01°C
- relation to body temperature
- (cold have additional peak at high temp
- paradoxical cold - "pins and needles"

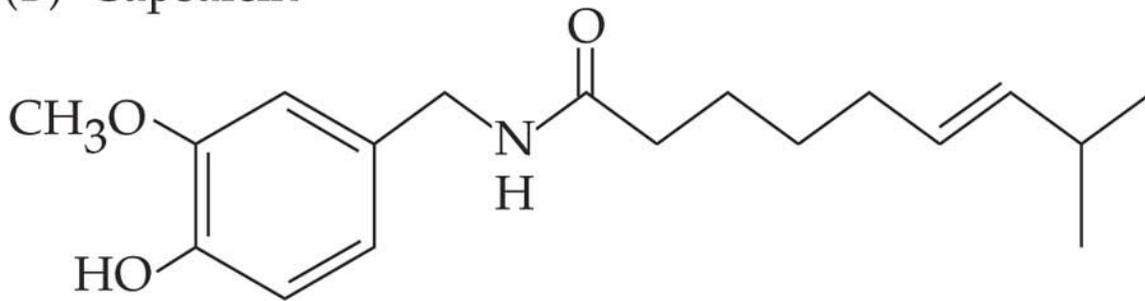
Recent progress

- channel properties
- How neurons know that it's cold outside,
- Hot and cold trp ion channels,
- cold related to menthol
- hot related to capsaicin

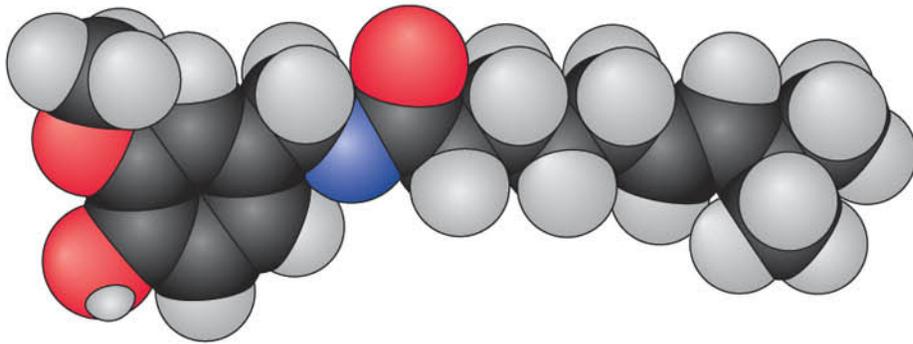
(A)



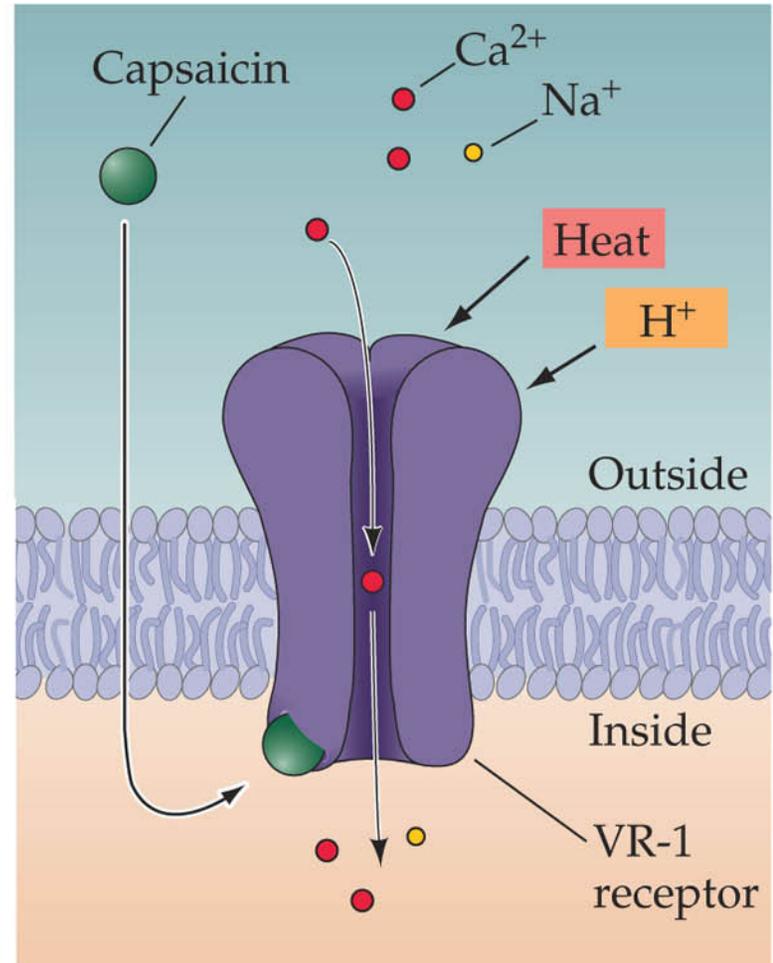
(B) Capsaicin



(C)

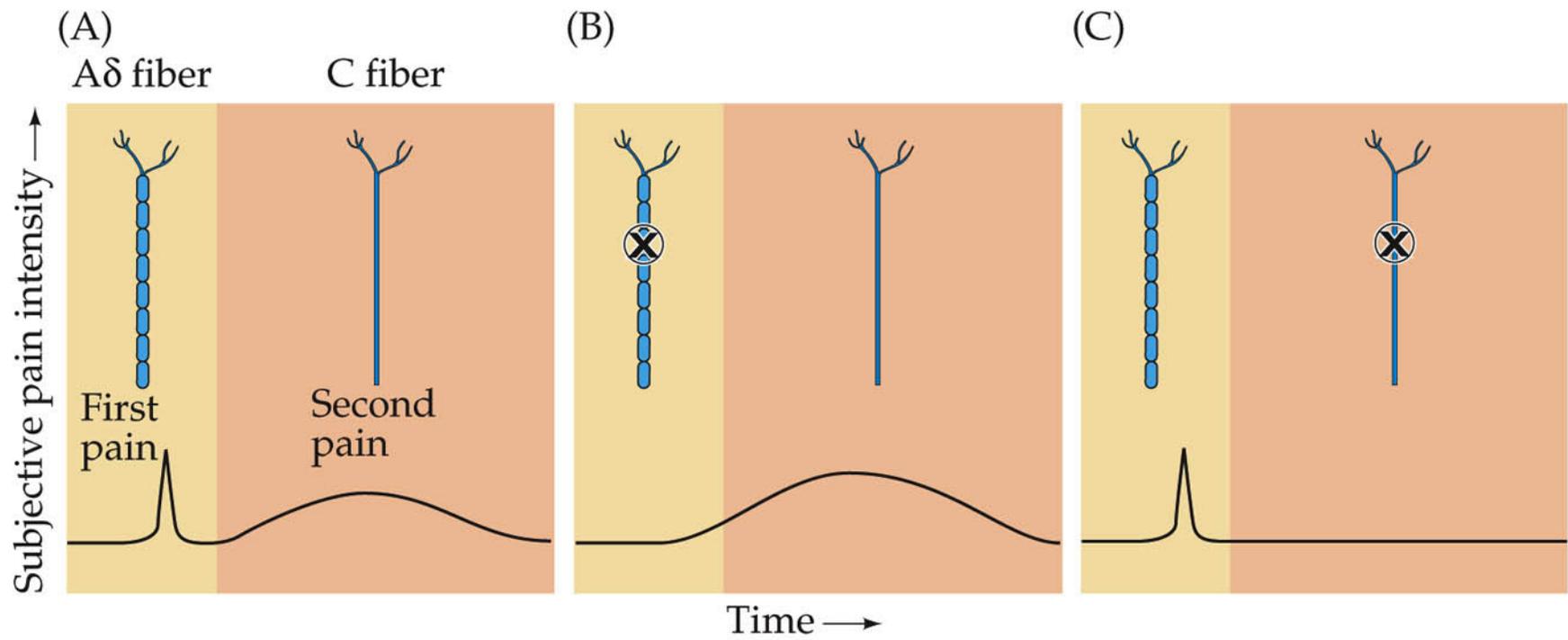


(D)



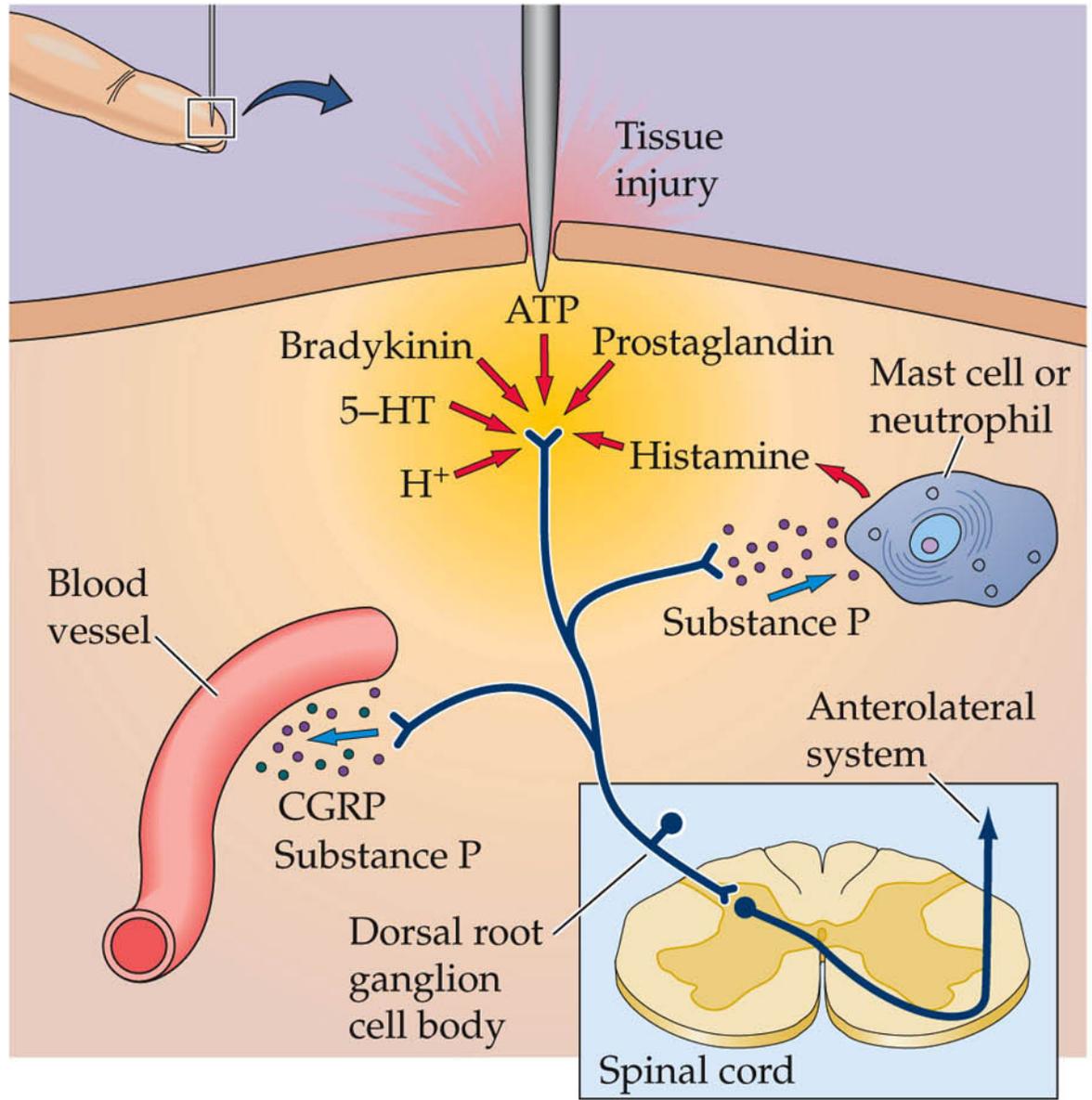
Personal reflection

- Cosens & Manning *Drosophila* mutant
- Cosens - abnormal mating.
- Baruch Minke,
- trp (transient receptor potential)



Pain axons

- Pain is faster in A delta fibers than in C fibers
- Nociceptors
- A delta mechano and mechano-thermal,
- C fiber polymodal



NEUROSCIENCE, Third Edition, Figure 9.6 © 2004 Sinauer Associates, Inc.

From outline

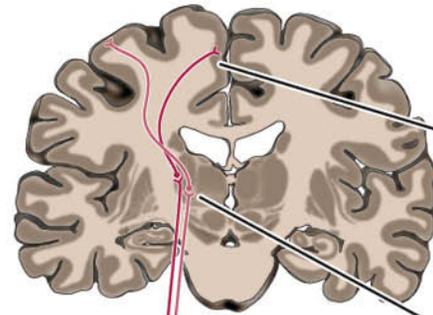
- sting venoms (serotonin, histamine, acetylcholine).
- Also tissue damage substances
- serotonin (platelets), prostaglandins, leukotrienes, Histamine from mast cells, substance P, Bradykinin from blood borne precursor - enzyme from injury

In summary

- nociceptor is really a chemoreceptor
- Nociceptors are in many places,
- but not in brain,
- hence brain surgery under local anesthesia
- used in mapping studies in humans by Penfield.

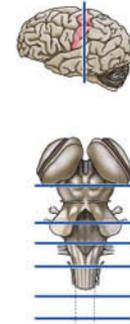
(A)

Cerebrum



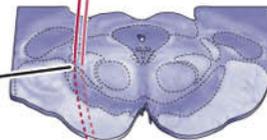
Primary somatic sensory cortex

Ventral posterior lateral nucleus of the thalamus



Midbrain

Medial lemniscus



Mid-pons



Medial lemniscus

Rostral medulla

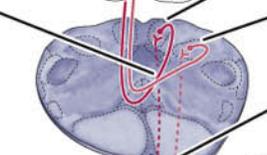
Internal arcuate fibers



Gracile nucleus

Cuneate nucleus

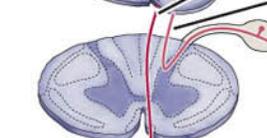
Caudal medulla



Gracile tract

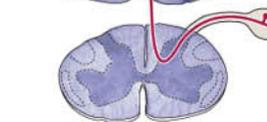
Cuneate tract

Cervical spinal cord



Mechanosensory receptors from upper body

Lumbar spinal cord



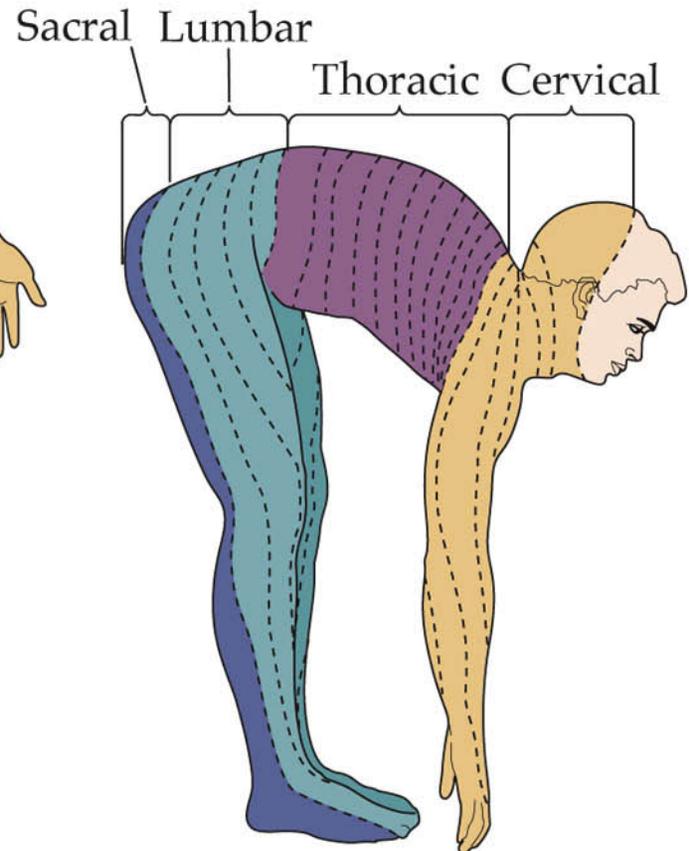
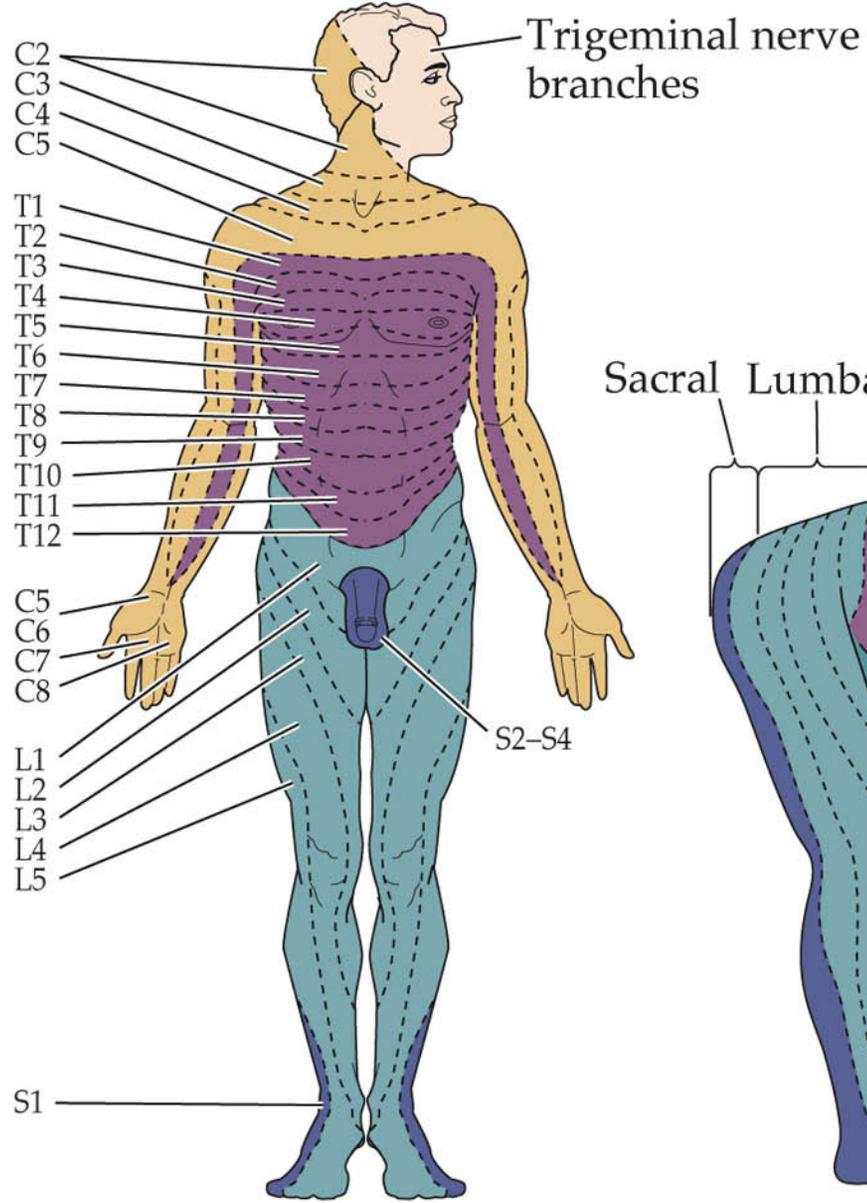
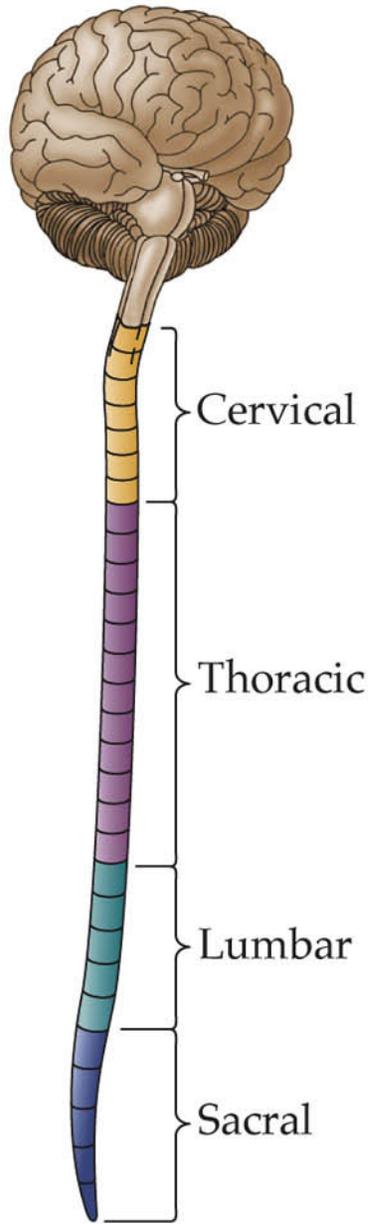
Mechanosensory receptors from lower body

From outline

- Lower limbs medial in gracile tract.
- Upper limbs lateral in cuneate tract.
- ipsilateral projection
- First nucleus is in lower medulla
- There is a cross-over, and then the next nucleus is in the thalamus.
- This lemniscal system is evolutionarily "new" (reptiles and above) and is for localized touch.

More

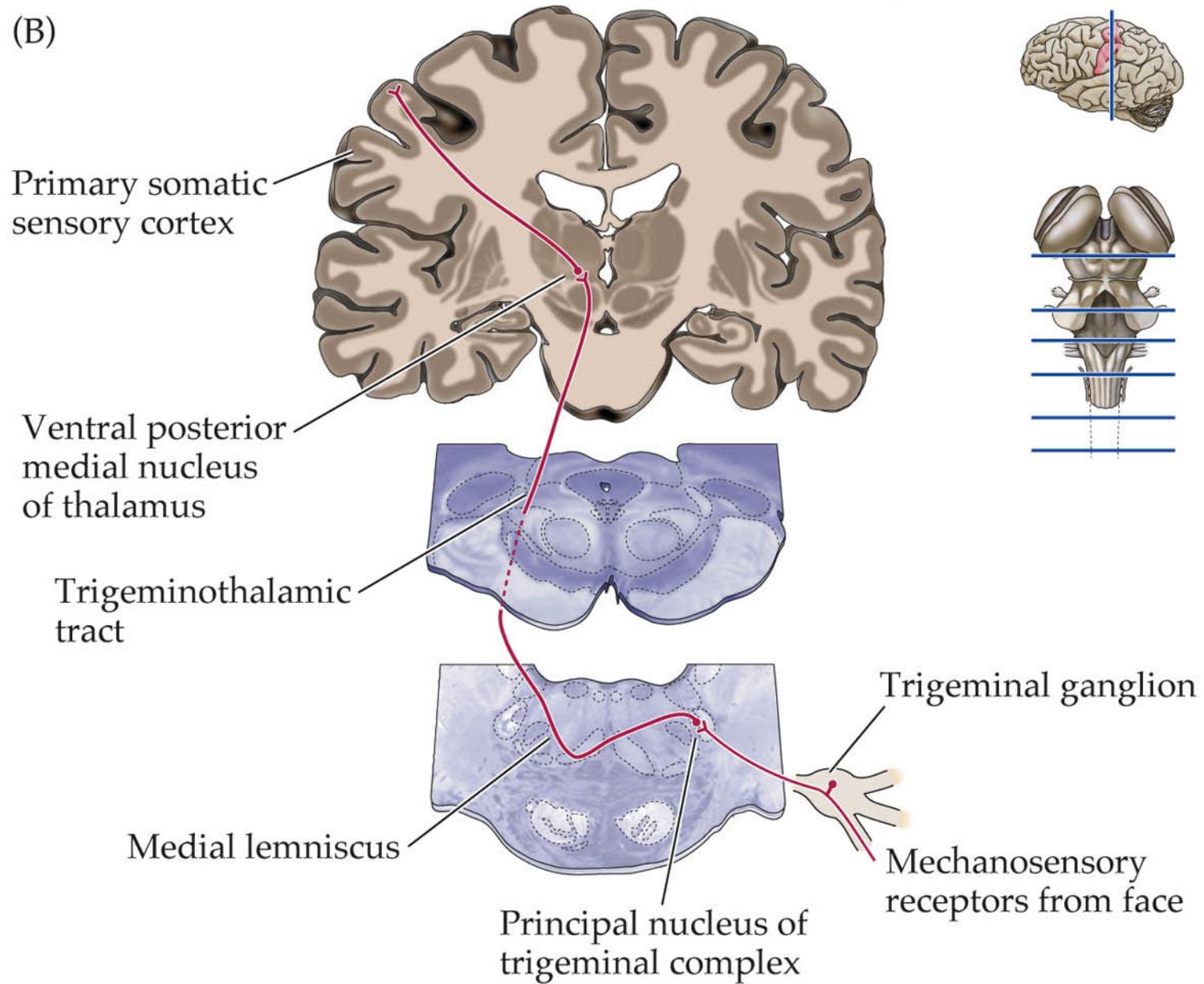
- processing - lateral inhibition to sharpen spatial localization.
- (This is the first mention of lateral inhibition, a fundamental mechanism of sensory processing.)
- If you tap your forearm, there are big waves but you feel localized touch.



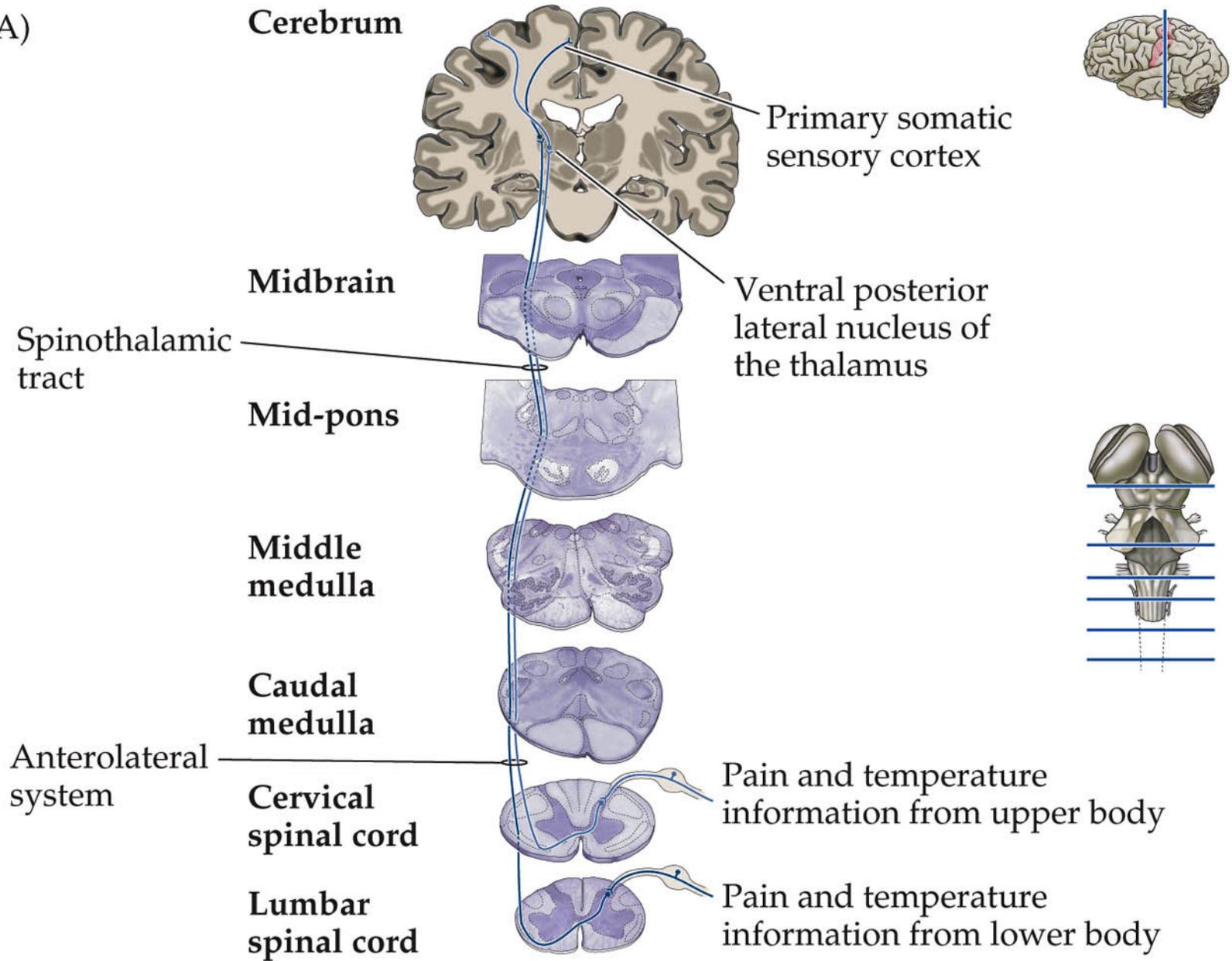
Dermatomes

- segmental organization of spinal cord -
- the dorsal root ganglion where input is translated into dermatomes -
- which place is innervated
- herpes zoster "shingles" reactivated virus - localized to one sensory ganglion

(B)



(A)

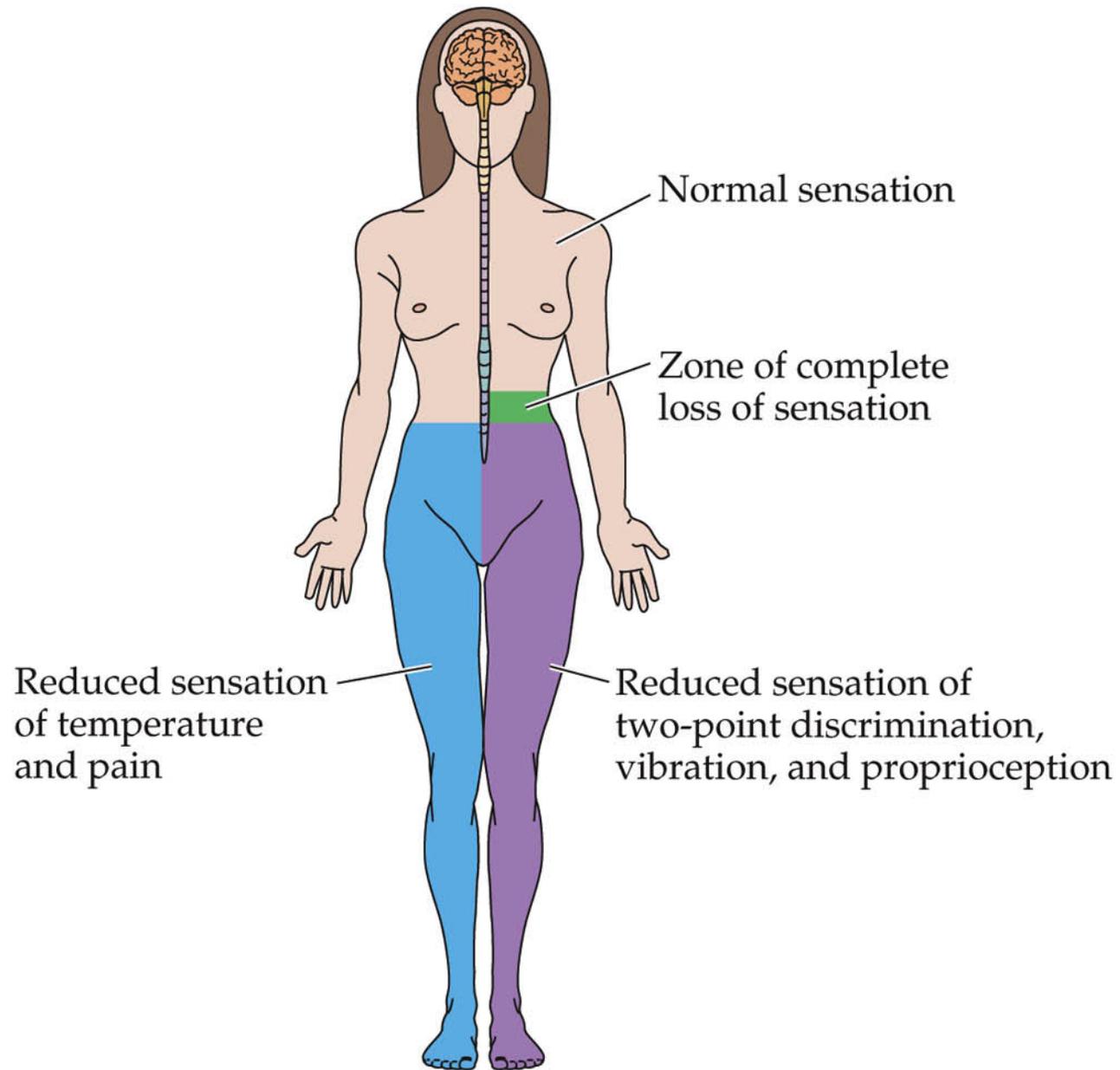


From outline

- spinothalamic snapse and decussation at entry point.
- There are separate tracts in spinal cord.
- lateral portion is for pain and temperature.
- ventral (anterior) part is for gross tactile sense.
- Hence the nomenclature "anterolateral."

More

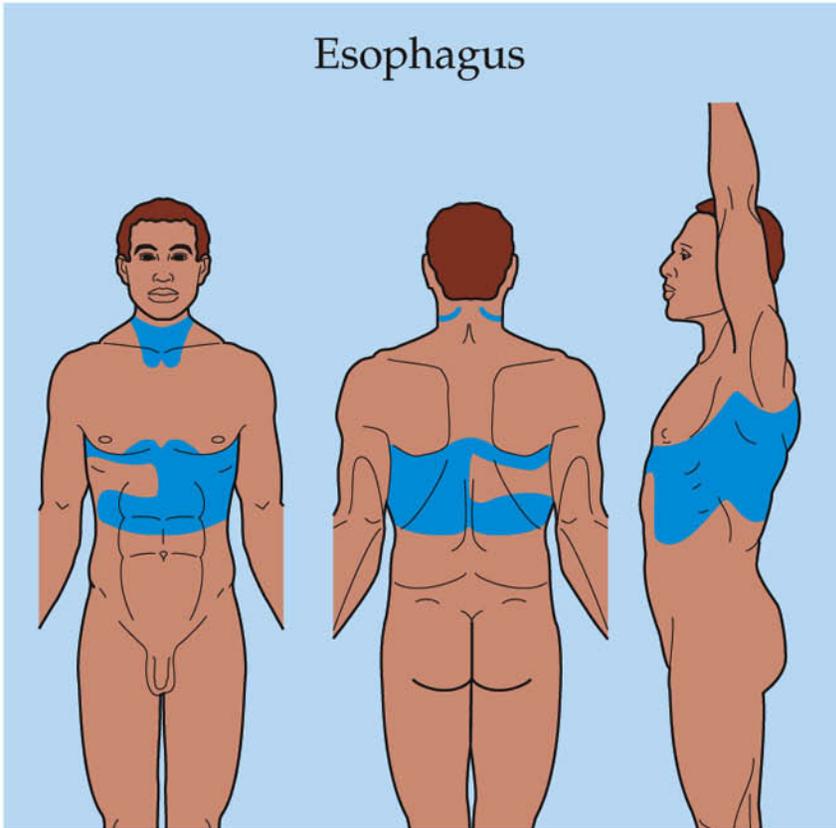
- Sharp pain inhibits worse pain
- (example: a hard touch to a door knob makes an electric shock less annoying)
- "neospinothalamic" (more recently evolved) A-delta
- "paleospinothalamic" (more ancient) C fibers
- injury to the former intractable pain,
- "psychosurgery" can be helpful.
- Dull pain diffuse less localized.



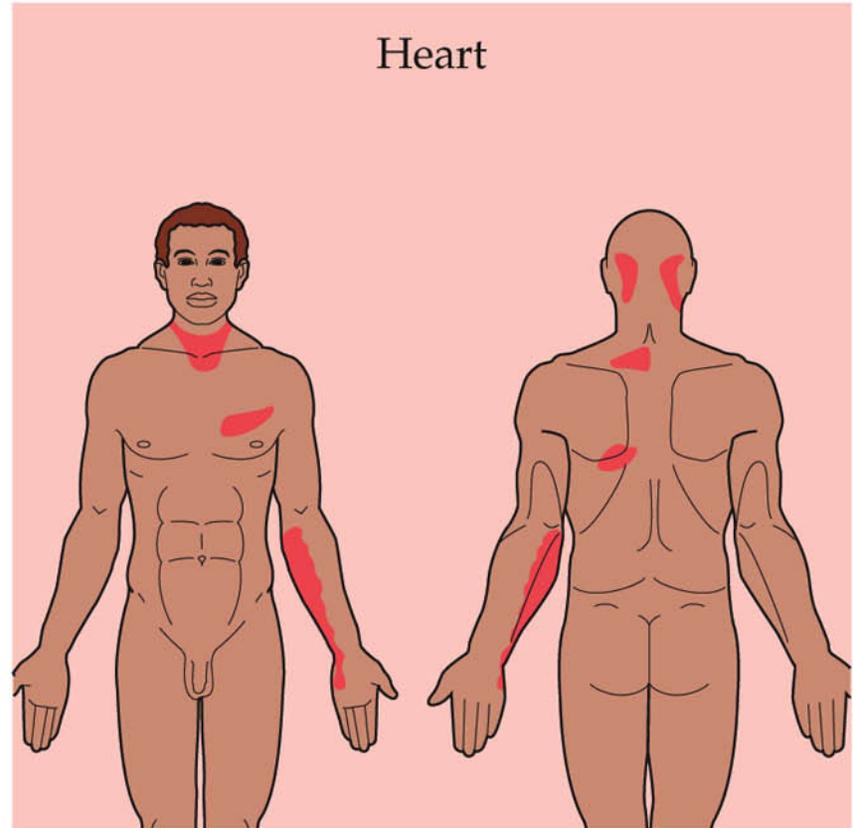
Loss from hemisection

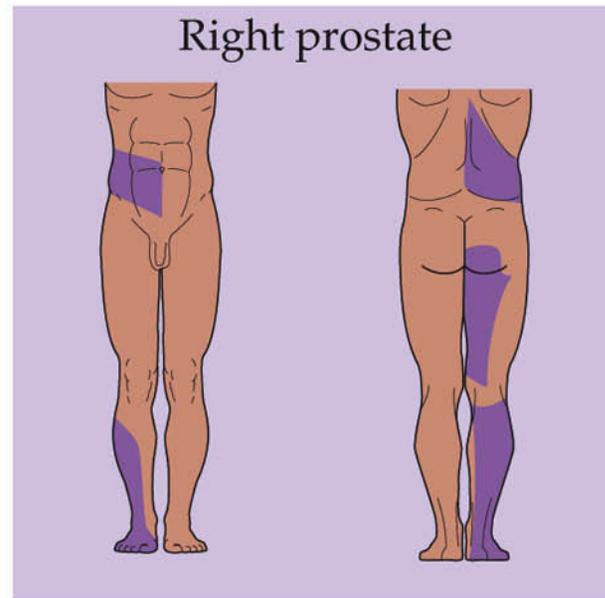
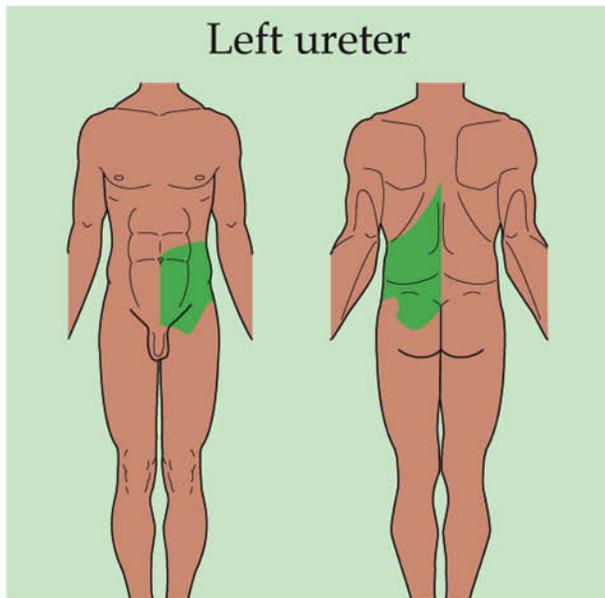
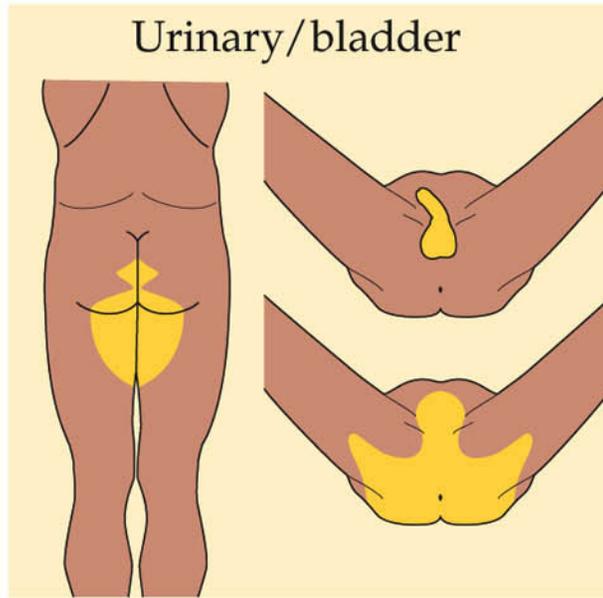
- A half spinal cord injury:
- contralateral loss of spinothalamic below injury
- ipsilateral loss of lemniscal.
- Brown-Sequard syndrome include motor (ipsilateral impairment)

Esophagus

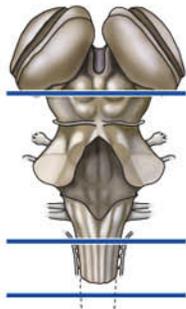
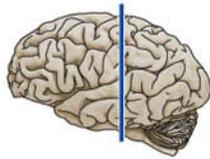


Heart





(A)



Gastrointestinal tract

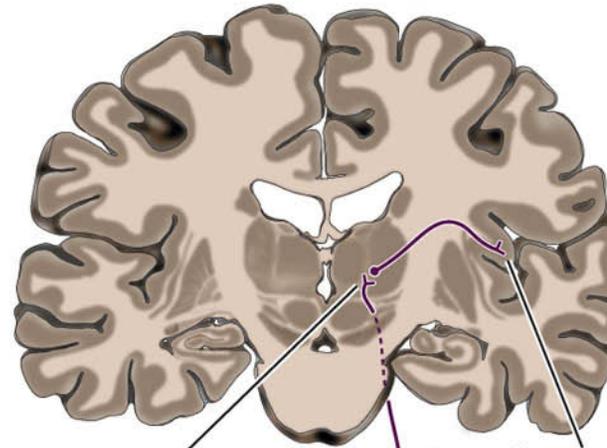
Ventral posterior nuclear complex of thalamus

Gracile nucleus

Cuneate nucleus

Dorsal root ganglion cells

Cerebrum



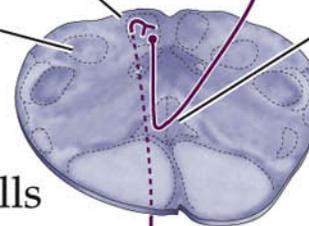
Insular cortex

Midbrain

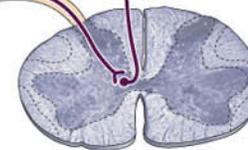


Medial lemniscus

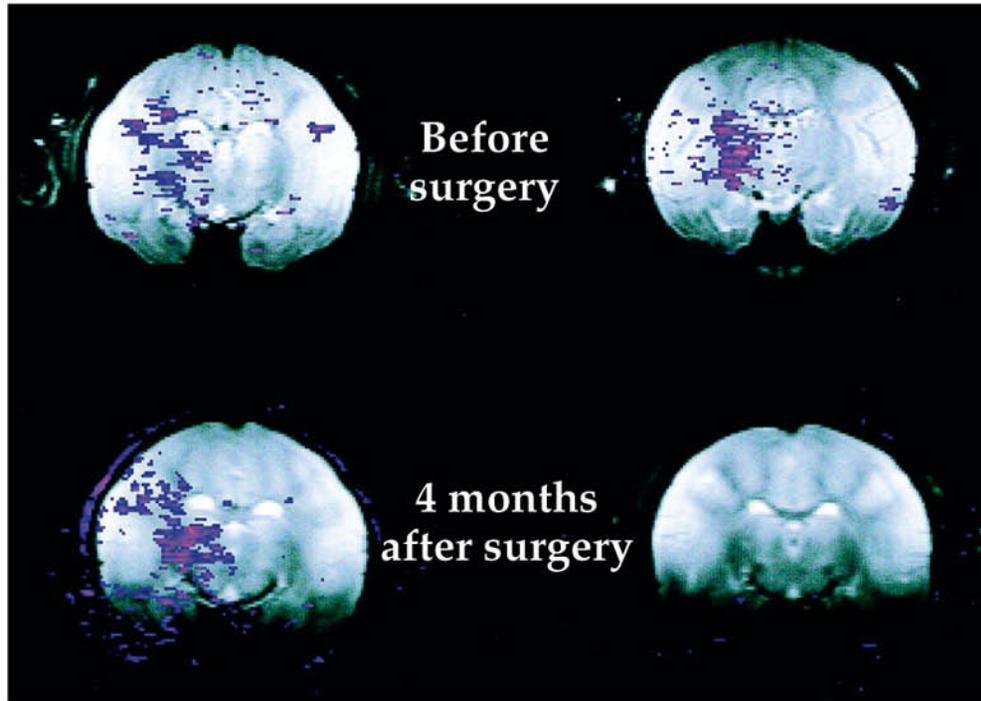
Medulla



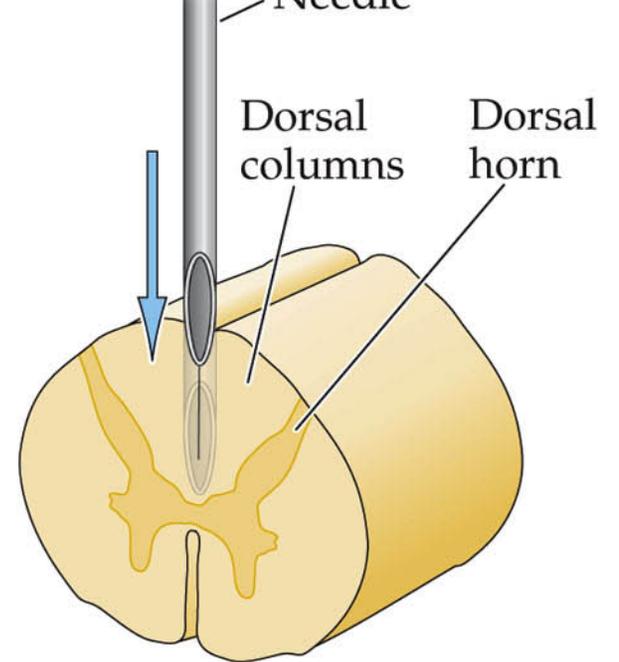
Spinal cord



(B) Sham lesion Dorsal column lesion



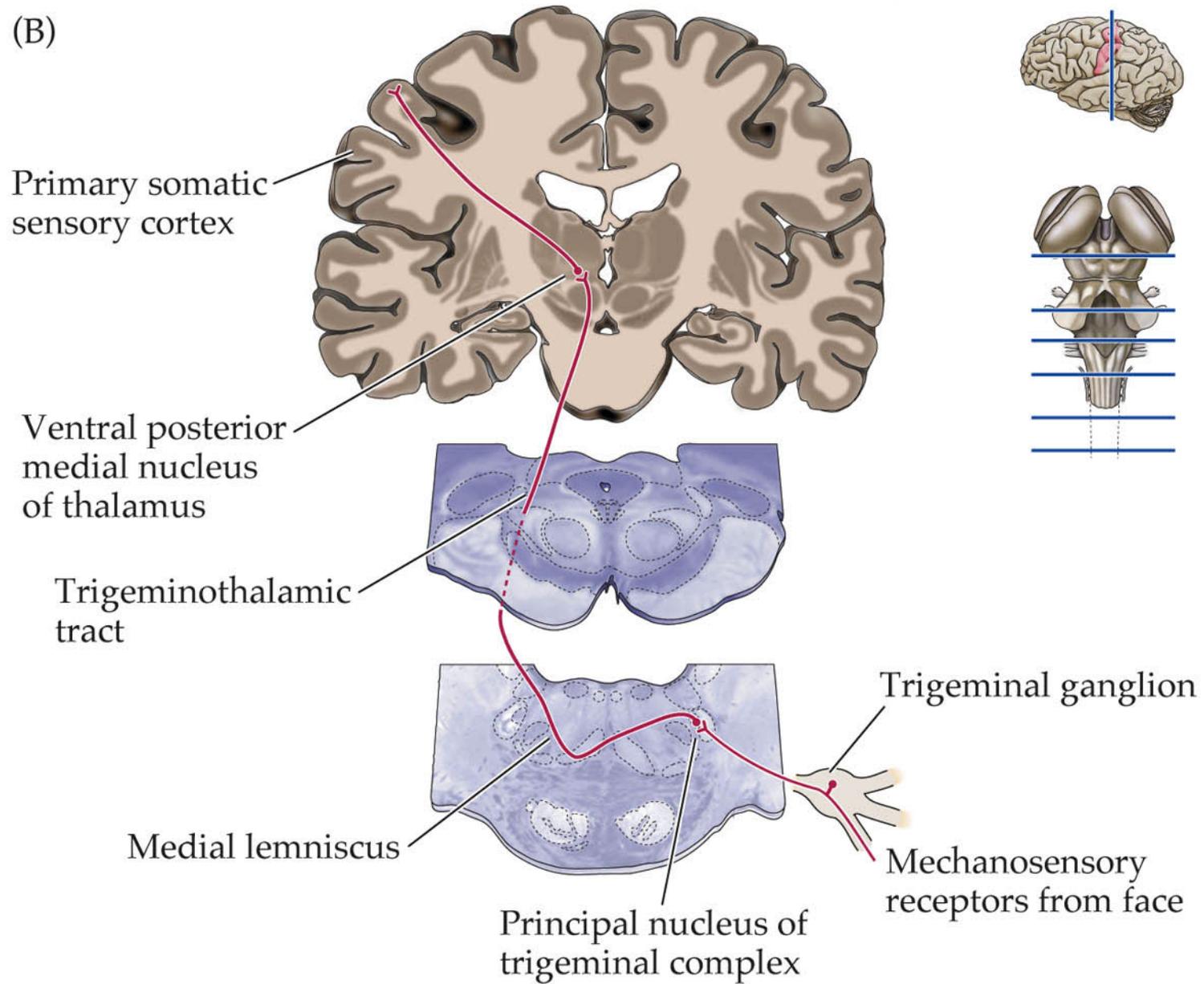
(C)

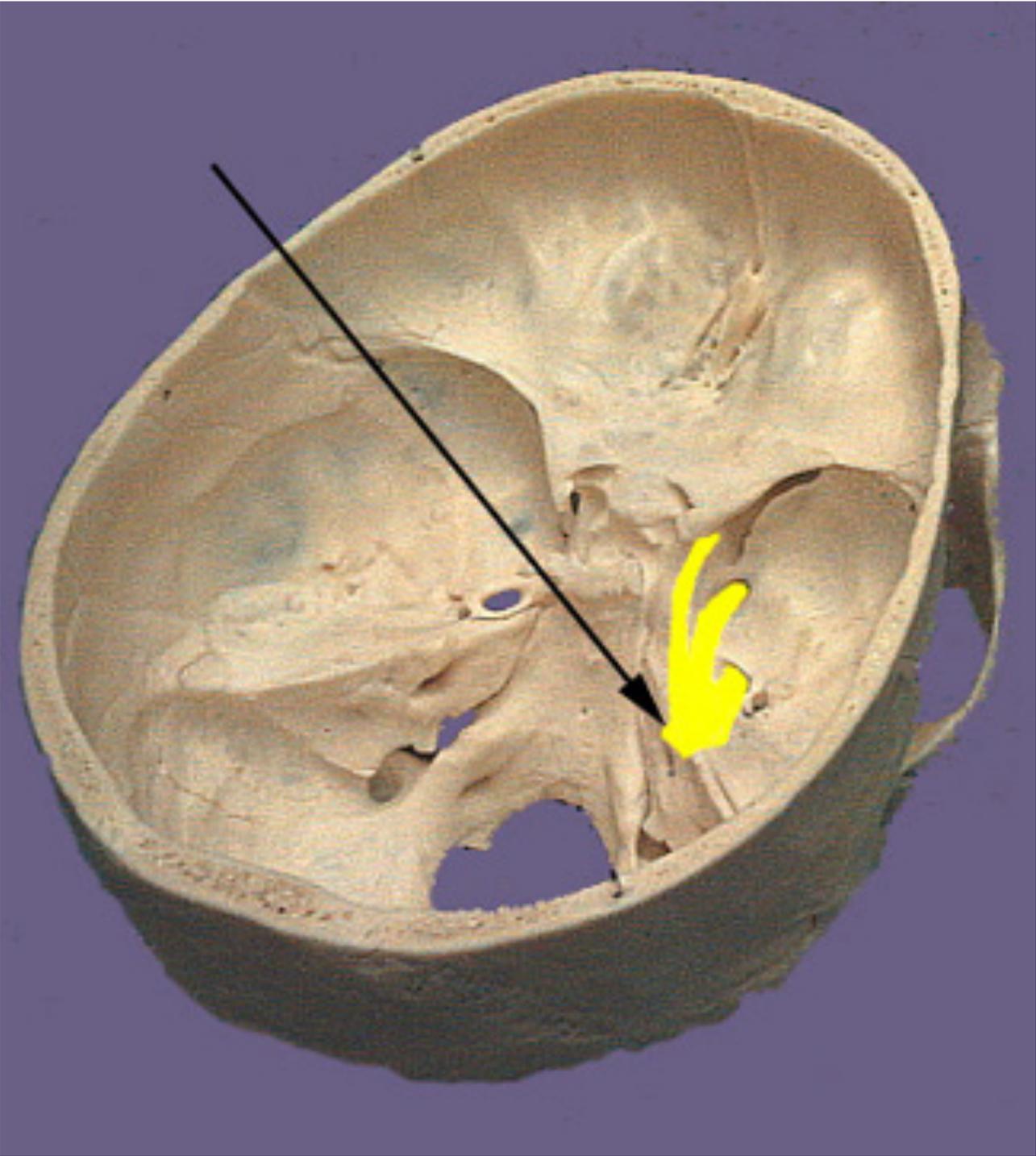


visceral pain

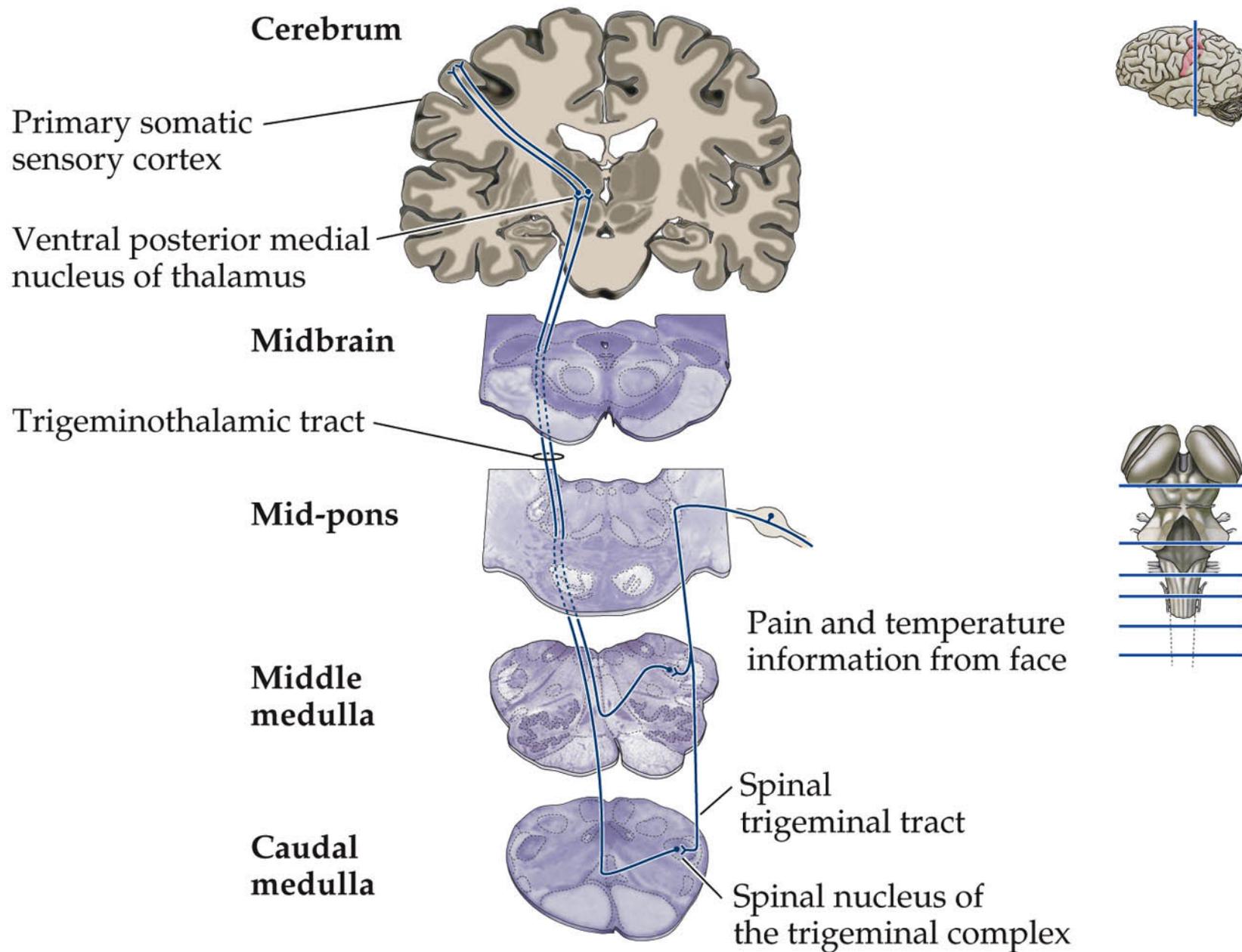
- Interestingly, visceral pain goes in dorsal columns.
- Very useful since midline myelotomy for palliative treatment in terminal and painful cancer.

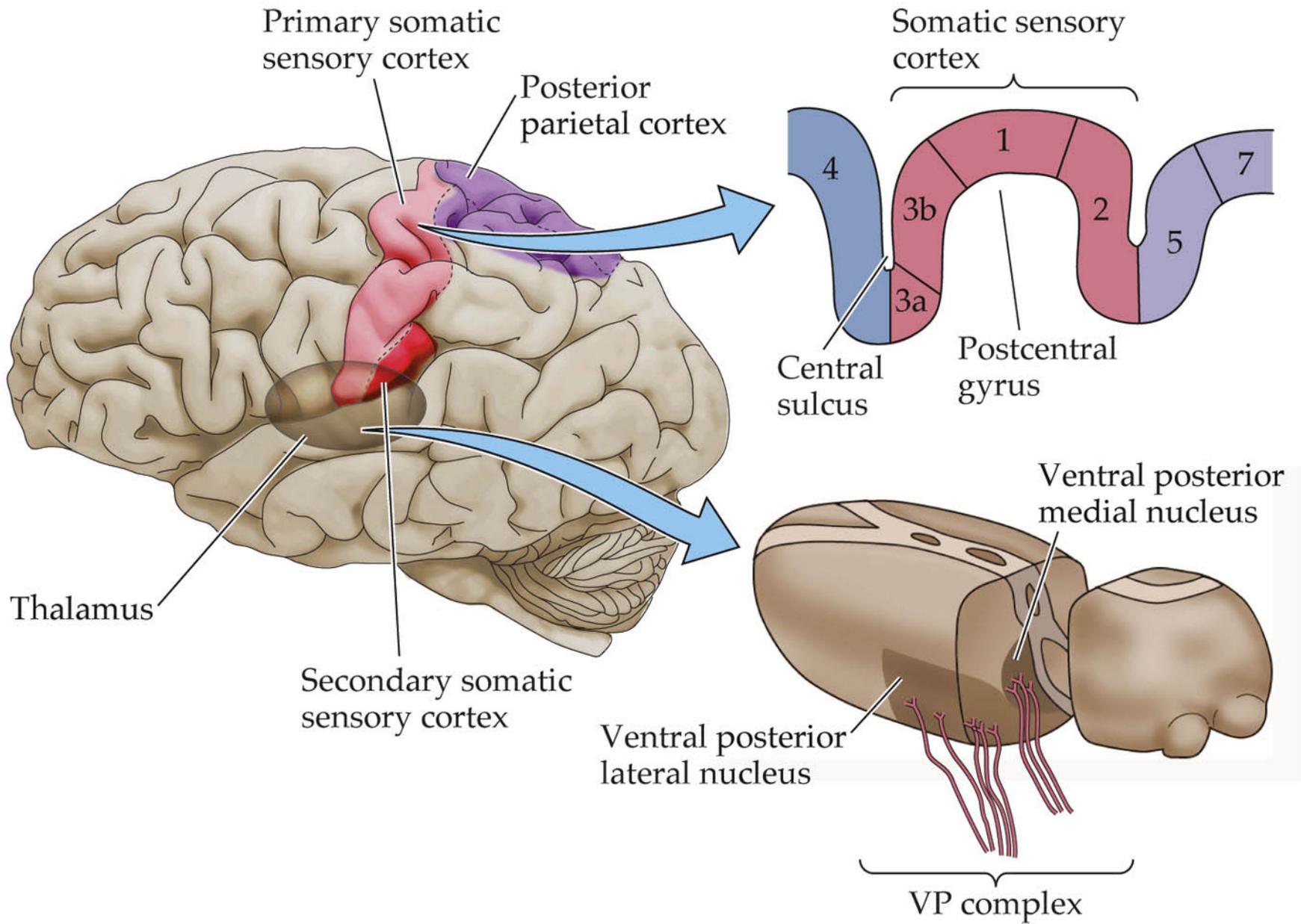
(B)





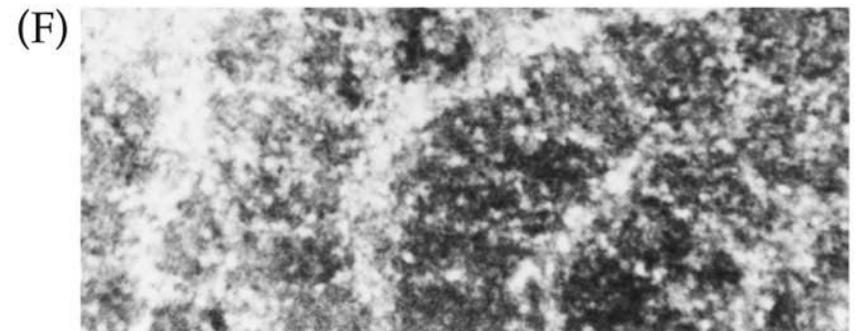
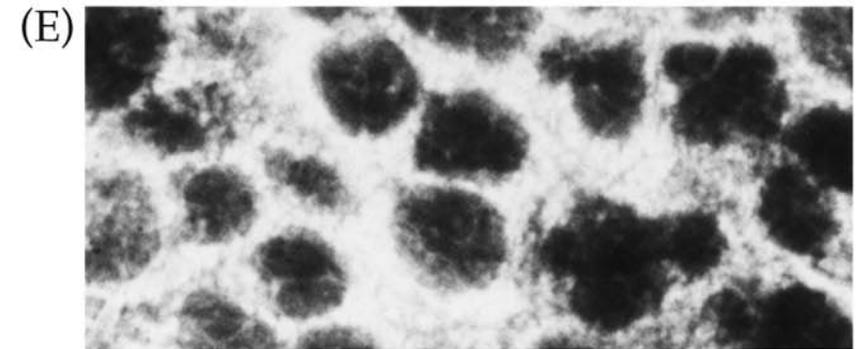
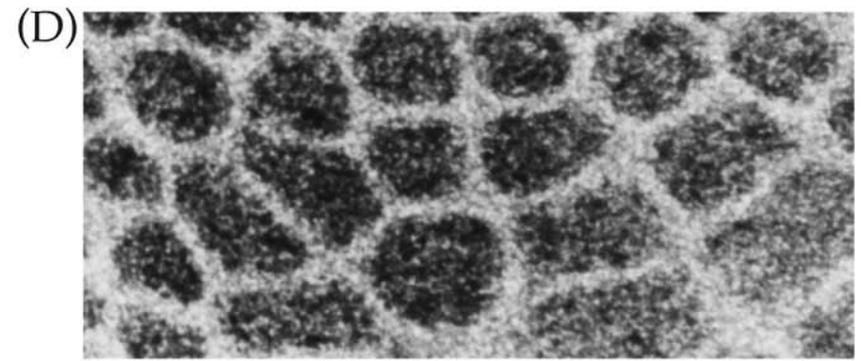
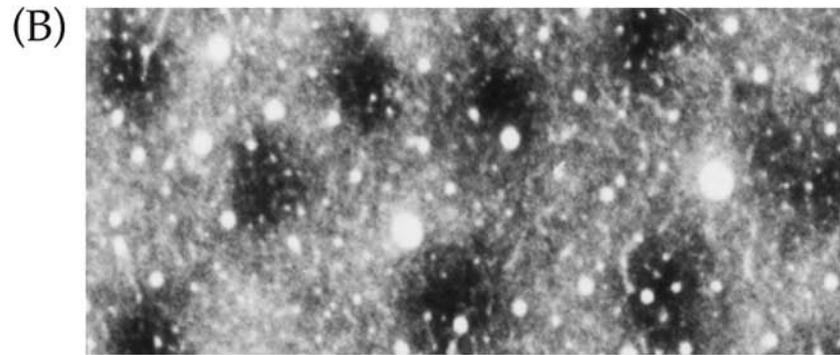
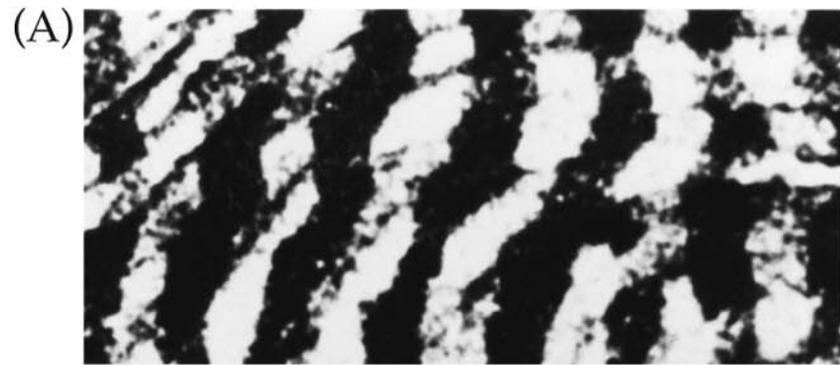
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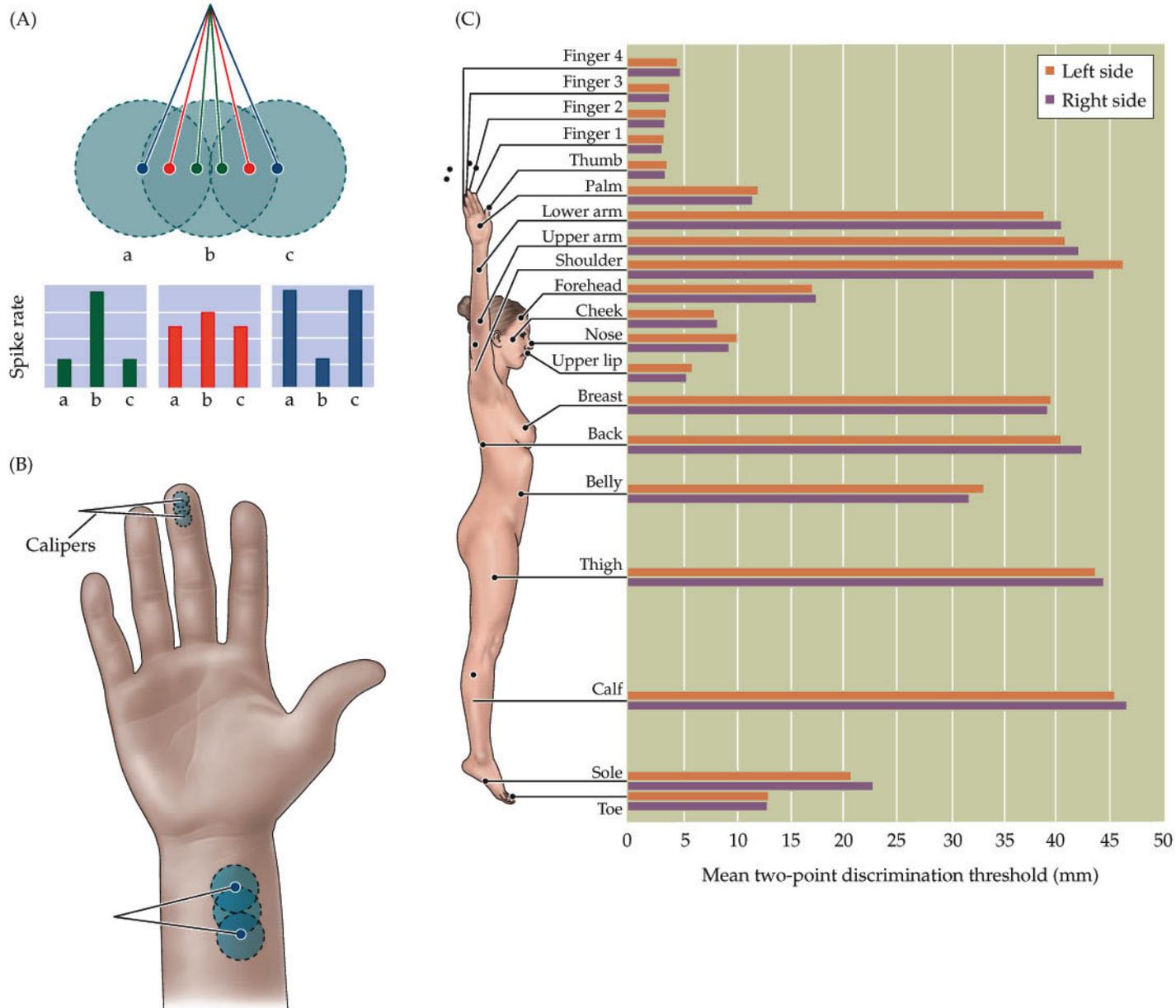




Thalamus and cortex

- VPL of thalamus to Postcentral gyrus-
- S1 = areas 1, 2, 3a & 3b
- arranged in columns
- - a vertical electrode penetration same submodality
- each S1 nerve responds to only one receptor type



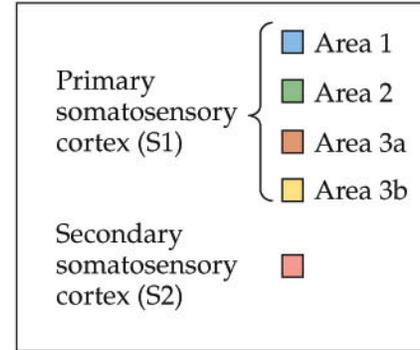
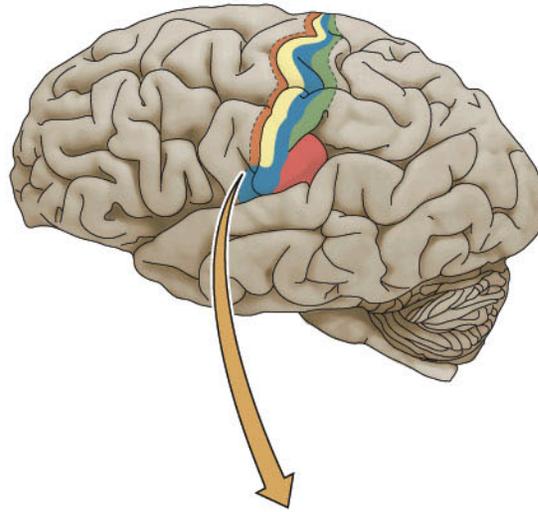


NEUROSCIENCE, Fourth Edition, Figure 9.3

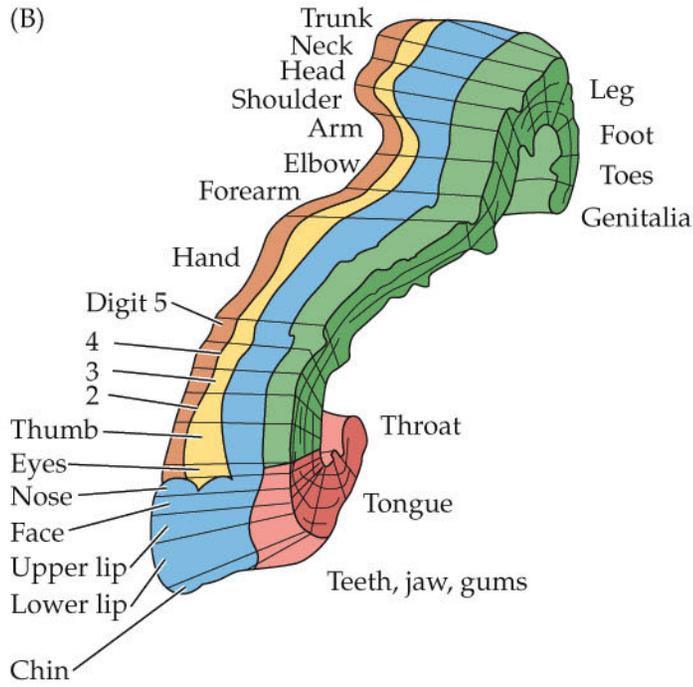
Sensory mapping

- In sensory map of cortex, all cells as electrode penetrates vertically are from one area (Mountcastle)
- (a) Ocular dominance columns for vision (Hubel and Wiesel) [Nobel](#) 1981
- (d) Woolsey - (box) "barrels" from vibrissae (whiskers)

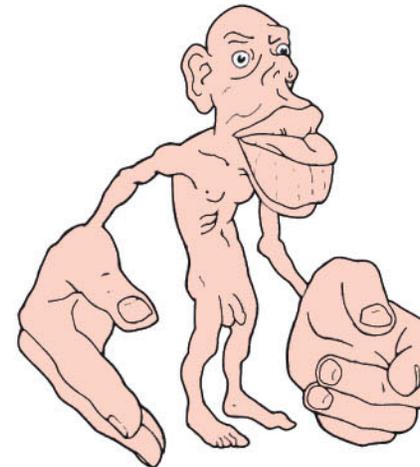
(A)



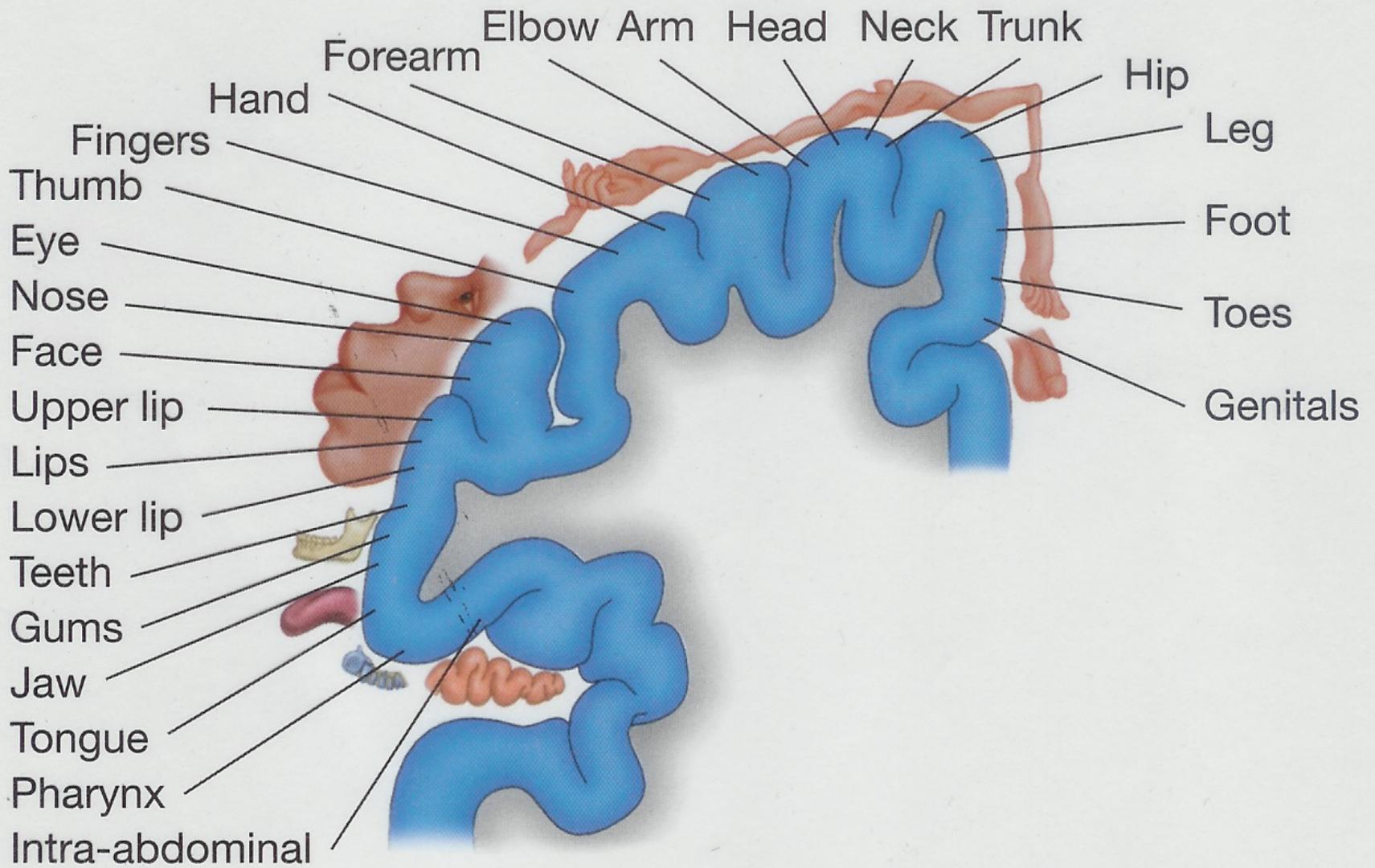
(B)



(C)

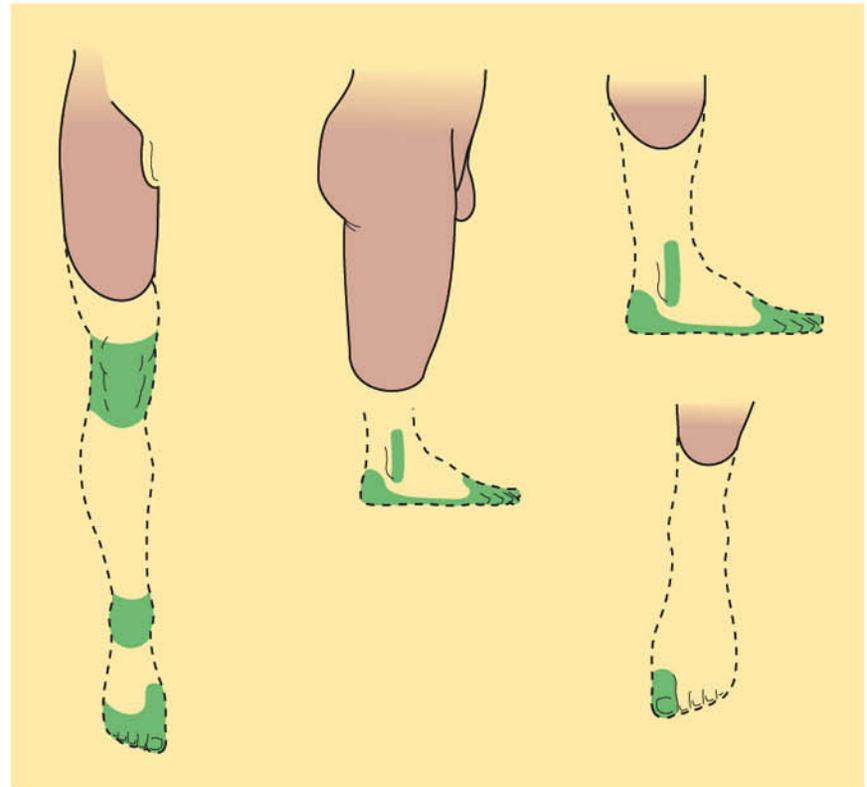
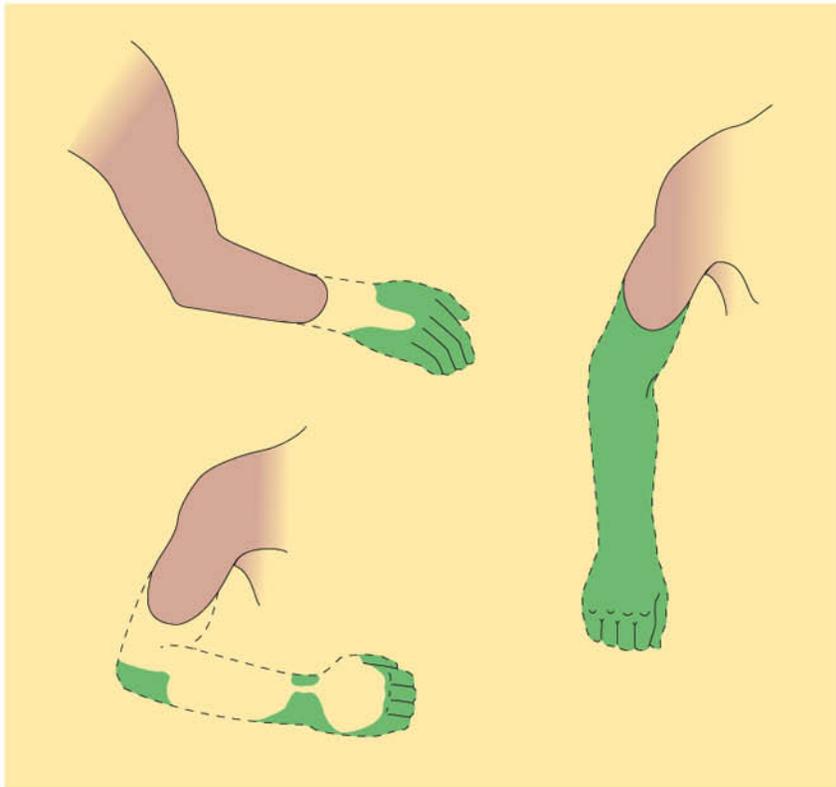


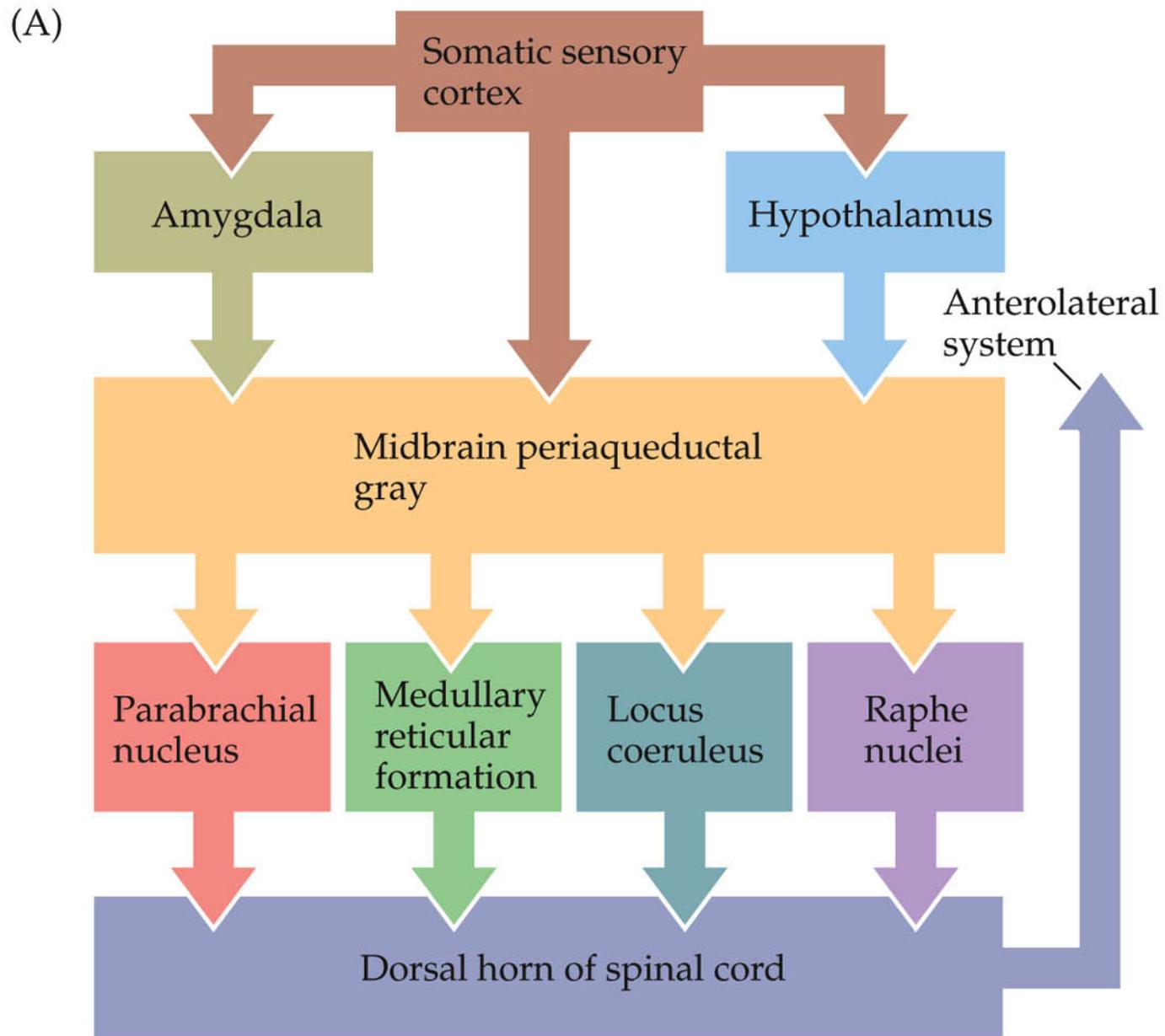
NEUROSCIENCE, Fourth Edition, Figure 9.11

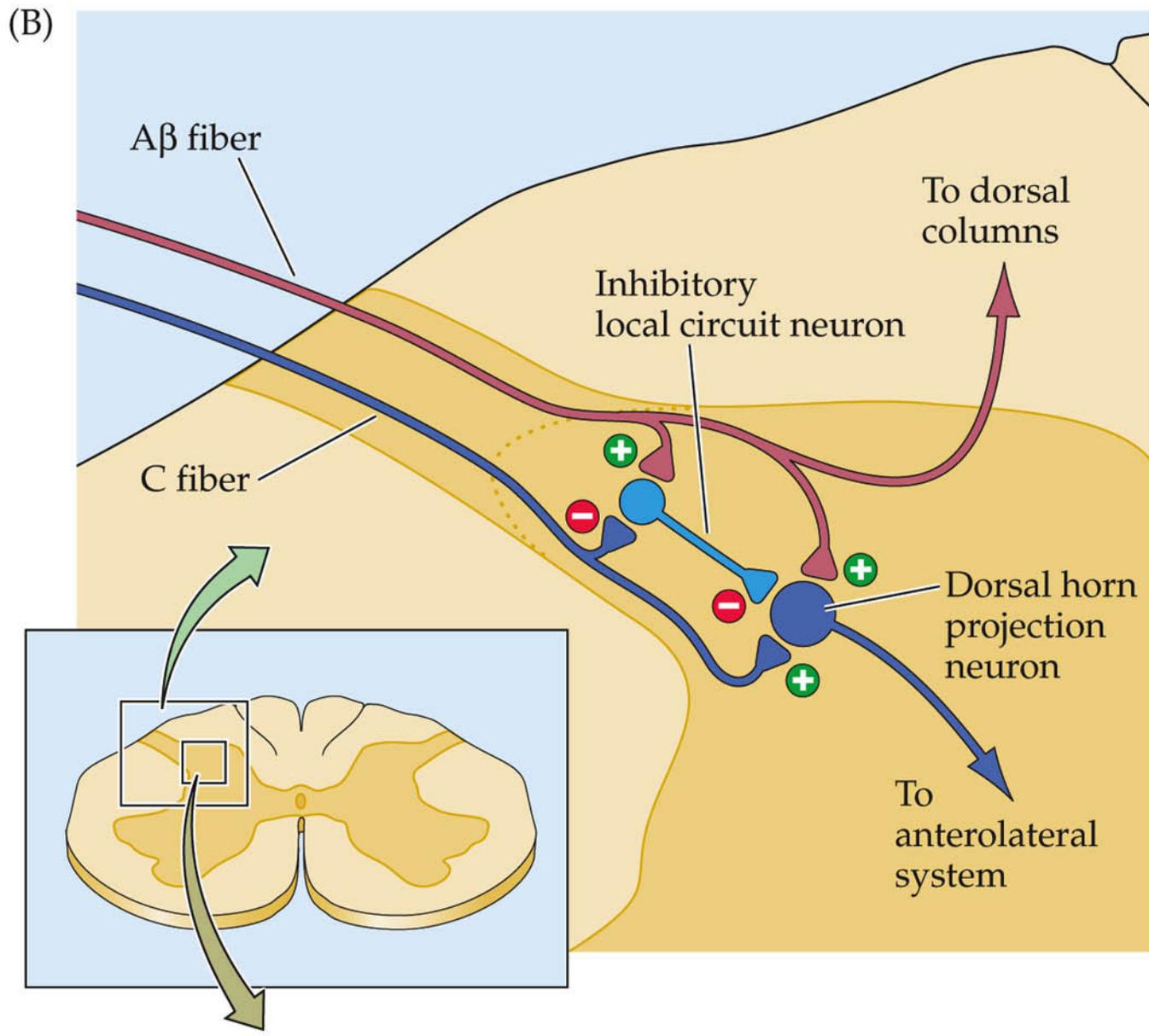


From outline

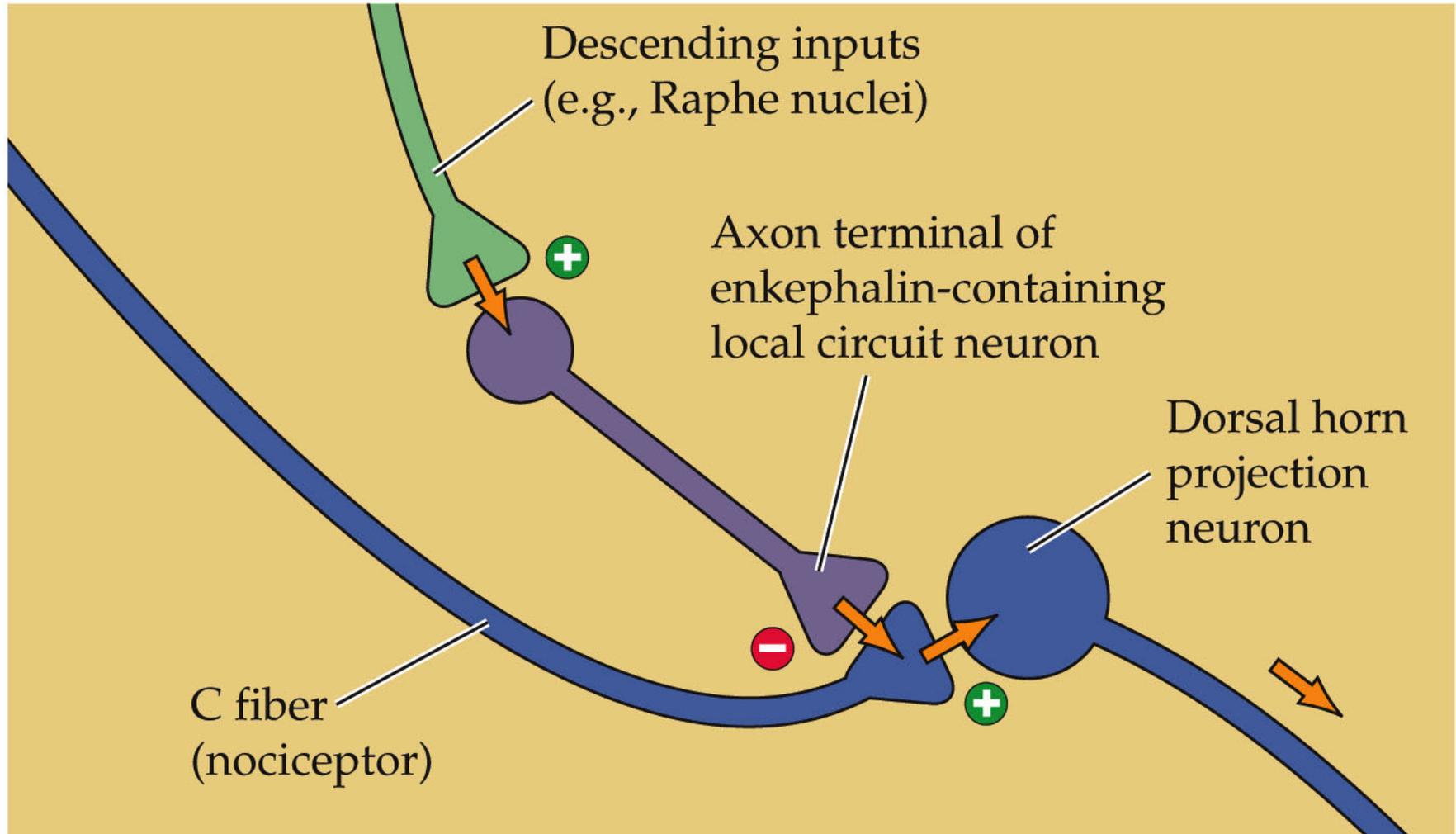
- two point threshold
- 2 mm fingertips, 30 arm, 70 back
- this relates to the cortical projection
- sensory magnifications
- Penfield - homunculus







(C)



From outline

- "microcircuits" in dorsal (posterior) horn
- all sensory input uses glutamate
- pain also uses substance P
- capsaicin causes release of substance P
- enkephalin from Substantia Gelatinosa interneuron - presynaptic
- (opiates are narcotic analgesics) stimulate - cause analgesia
- connect to Raphe
- Itch: only skin, mucous - opiates not suppress