**ALADE EMMANUEL ADEMOLA**

**18/MHS07/006**

**PHARMACOLOGY**

**PHS 212**

**Question**

**Discuss The somatosensory pathways**

The **somatosensory system** is a part of the [sensory nervous system](https://en.m.wikipedia.org/wiki/Sensory_nervous_system). The somatosensory system is a complex system of [sensory neurons](https://en.m.wikipedia.org/wiki/Sensory_neuron) and [neural pathways](https://en.m.wikipedia.org/wiki/Neural_pathway) that responds to changes at the surface or inside the body. The [axons](https://en.m.wikipedia.org/wiki/Axon) (as [afferent nerve fibers](https://en.m.wikipedia.org/wiki/Afferent_nerve_fiber)) of sensory neurons connect with, or respond to, various receptor cells. These sensory receptor cells are activated by different stimuli such as heat and [nociception](https://en.m.wikipedia.org/wiki/Nociception), giving a functional name to the responding sensory neuron, such as a [thermoreceptor](https://en.m.wikipedia.org/wiki/Thermoreceptor) which carries information about temperature changes. Other types include [mechanoreceptors](https://en.m.wikipedia.org/wiki/Mechanoreceptor), [chemoreceptors](https://en.m.wikipedia.org/wiki/Chemoreceptor), and [nociceptors](https://en.m.wikipedia.org/wiki/Nociceptor) which send signals along a [sensory nerve](https://en.m.wikipedia.org/wiki/Sensory_nerve) to the [spinal cord](https://en.m.wikipedia.org/wiki/Spinal_cord) where they may be processed by other sensory neurons and then relayed to the [brain](https://en.m.wikipedia.org/wiki/Brain) for further processing. Sensory receptors are found all over the body including the [skin](https://en.m.wikipedia.org/wiki/Skin), [epithelial tissues](https://en.m.wikipedia.org/wiki/Epithelium), [muscles](https://en.m.wikipedia.org/wiki/Skeletal_muscle), [bones](https://en.m.wikipedia.org/wiki/Bone) and [joints](https://en.m.wikipedia.org/wiki/Joint), internal [organs](https://en.m.wikipedia.org/wiki/Organ_%28anatomy%29), and the [cardiovascular system](https://en.m.wikipedia.org/wiki/Cardiovascular_system).

Somatic senses are sometimes referred to as **somesthetic senses**, with the understanding that somesthesis includes the **sense of touch**, [proprioception](https://en.m.wikipedia.org/wiki/Proprioception) (sense of position and movement), and (depending on usage) [haptic perception](https://en.m.wikipedia.org/wiki/Haptic_perception).

The mapping of the body surfaces in the brain is called [somatotopy](https://en.m.wikipedia.org/wiki/Somatotopy%22%20%5Co%20%22Somatotopy). In the cortex, it is also referred to as the [cortical homunculus](https://en.m.wikipedia.org/wiki/Cortical_homunculus). This brain-surface ("cortical") map is not immutable, however. Dramatic shifts can occur in response to stroke or injury.

### Sensory receptors

The four [mechanoreceptors](https://en.m.wikipedia.org/wiki/Mechanoreceptor) in the [skin](https://en.m.wikipedia.org/wiki/Human_skin) each respond to different stimuli for short or long periods.

[Merkel cell](https://en.m.wikipedia.org/wiki/Merkel_cell) nerve endings are found in the basal [epidermis](https://en.m.wikipedia.org/wiki/Epidermis) and [hair follicles](https://en.m.wikipedia.org/wiki/Hair_follicles); they react to low vibrations (5–15 [Hz](https://en.m.wikipedia.org/wiki/Hertz)) and deep static touch such as shapes and edges. Due to having a small receptive field (extremely detailed info), they are used in areas like fingertips the most; they are not covered (shelled) and thus respond to pressures over long periods.

[Tactile corpuscles](https://en.m.wikipedia.org/wiki/Tactile_corpuscle) react to moderate vibration (10–50 Hz) and light touch. They are located in the [dermal papillae](https://en.m.wikipedia.org/wiki/Dermal_papillae); due to their reactivity, they are primarily located in fingertips and lips. They respond in quick [action potentials](https://en.m.wikipedia.org/wiki/Action_potentials), unlike Merkel nerve endings. They are responsible for the ability to read Braille and feel gentle stimuli.

[Pacinian corpuscles](https://en.m.wikipedia.org/wiki/Pacinian_corpuscle) determine gross touch and distinguish rough and soft substances. They react in quick action potentials, especially to vibrations around 250 Hz (even up to centimeters away). They are the most sensitive to vibrations and have large receptor fields. Pacinian corpuscles react only to sudden stimuli so pressures like clothes that are always compressing their shape are quickly ignored. They have also been implicated in detecting the location of touch sensations on handheld tools.

[Bulbous corpuscles](https://en.m.wikipedia.org/wiki/Bulbous_corpuscle) react slowly and respond to sustained skin stretch. They are responsible for the feeling of object slippage and play a major role in the [kinesthetic](https://en.m.wikipedia.org/wiki/Proprioception) sense and control of finger position and movement. Merkel and bulbous cells - slow-response - are [myelinated](https://en.m.wikipedia.org/wiki/Myelinated); the rest - fast-response - are not. All of these receptors are activated upon pressures that squish their shape causing an action potential.

### Somatosensory cortex



 the sensory tract, showing the pathway (blue) up the spinal cord, through the somatosensory thalamus, to S1 (Brodmann areas 3, 1, and 2), S2, and BA7



: Detail showing path adjacent to the [insular cortex](https://en.m.wikipedia.org/wiki/Insular_cortex) (marked insula in this figure), adjacent to S1, S2, and BA7

The postcentral gyrus includes the [primary somatosensory cortex](https://en.m.wikipedia.org/wiki/Primary_somatosensory_cortex) ([Brodmann areas](https://en.m.wikipedia.org/wiki/Brodmann_area%22%20%5Co%20%22Brodmann%20area) 3, 2 and 1) collectively referred to as S1.

BA3 receives the densest projections from the [thalamus](https://en.m.wikipedia.org/wiki/Thalamus). BA3a is involved with the sense of relative position of neighboring body parts and amount of effort being used during movement. BA3b is responsible for distributing somatosensory information, it projects texture information to BA1 and shape and size information to BA2.

Region S2 ([secondary somatosensory cortex](https://en.m.wikipedia.org/wiki/Secondary_somatosensory_cortex)) divides into Area S2 and parietal ventral area. Area S2 is involved with specific touch perception and is thus integrally linked with the amygdala and hippocampus to encode and reinforce memories.

Parietal ventral area is the somatosensory relay to the premotor cortex and somatosensory memory hub, BA5.

BA5 is the topographically organized somato memory field and association area.

BA1 processes texture info while BA2 processes size + shape info.

Area S2 processes light touch, pain, visceral sensation, and tactile attention.

S1 processes the remaining info (crude touch, pain, temperature).

BA7 integrates visual and proprioceptive info to locate objects in space.

The [insular cortex](https://en.m.wikipedia.org/wiki/Insular_cortex) (insula) plays a role in the sense of bodily-ownership, bodily self-awareness, and perception. Insula also plays a role in conveying info about sensual touch, pain, temperature, itch, and local oxygen status. Insula is a highly connected relay and thus is involved in numerous functions.

##  General somatosensory pathway

All afferent touch/vibration info ascends the spinal cord via the posterior (dorsal) column-medial lemniscus pathway via gracilis (T7 and below) or cuneatus (T6 and above). Cuneatus sends signals to the cochlear nucleus indirectly via spinal grey matter, this info is used in determining if a perceived sound is just villi noise/irritation. All fibers cross (left becomes right) in the medulla.

A somatosensory pathway will typically have three neurons: first-order, second-order, and third-order.

1. The **first-order neuron** is a type of [pseudounipolar neuron](https://en.m.wikipedia.org/wiki/Pseudounipolar_neuron%22%20%5Co%20%22Pseudounipolar%20neuron) and always has its [cell body](https://en.m.wikipedia.org/wiki/Cell_body) in the [dorsal root ganglion](https://en.m.wikipedia.org/wiki/Dorsal_root_ganglion) of the [spinal nerve](https://en.m.wikipedia.org/wiki/Spinal_nerve) with a peripheral [axon](https://en.m.wikipedia.org/wiki/Axon) innervating touch [mechanoreceptors](https://en.m.wikipedia.org/wiki/Mechanoreceptor) and a central axon synapsing on the second-order neuron. If the somatosensory pathway is in parts of the head or neck not covered by the cervical nerves, the first-order neuron will be the [trigeminal nerve ganglia](https://en.m.wikipedia.org/wiki/Trigeminal_ganglion) or the ganglia of other sensory [cranial nerves](https://en.m.wikipedia.org/wiki/Cranial_nerves)).
2. The **second-order neuron** has its [cell body](https://en.m.wikipedia.org/wiki/Cell_body) either in the spinal cord or in the brainstem. This neuron's ascending [axons](https://en.m.wikipedia.org/wiki/Axons) will cross ([decussate](https://en.m.wikipedia.org/wiki/Decussate)) to the opposite side either in the [spinal cord](https://en.m.wikipedia.org/wiki/Spinal_cord) or in the [brainstem](https://en.m.wikipedia.org/wiki/Brainstem).
3. In the case of touch and certain types of pain, the **third-order neuron** has its [cell body](https://en.m.wikipedia.org/wiki/Cell_body) in the [ventral posterior nucleus](https://en.m.wikipedia.org/wiki/Ventral_posterior_nucleus) of the thalamus and ends in the [postcentral gyrus](https://en.m.wikipedia.org/wiki/Postcentral_gyrus) of the [parietal lobe](https://en.m.wikipedia.org/wiki/Parietal_lobe) in the [primary somatosensory cortex](https://en.m.wikipedia.org/wiki/Primary_somatosensory_cortex) (or S1).

Photoreceptors, similar to those found in the [retina](https://en.m.wikipedia.org/wiki/Retina) of the [eye](https://en.m.wikipedia.org/wiki/Eye), detect potentially damaging [ultraviolet radiation](https://en.m.wikipedia.org/wiki/Ultraviolet_radiation) ([ultraviolet A](https://en.m.wikipedia.org/wiki/Ultraviolet_A) specifically), inducing increased production of [melanin](https://en.m.wikipedia.org/wiki/Melanin) by [melanocytes](https://en.m.wikipedia.org/wiki/Melanocytes). Thus tanning potentially offers the skin rapid protection from DNA damage and sunburn caused by [ultraviolet radiation](https://en.m.wikipedia.org/wiki/Ultraviolet_radiation) (DNA damage caused by [ultraviolet B](https://en.m.wikipedia.org/wiki/Ultraviolet_B)). However, whether this offers protection is debatable, because the amount of melanin released by this process is modest in comparison to the amounts released in response to DNA damage caused by [ultraviolet B](https://en.m.wikipedia.org/wiki/Ultraviolet_B) radiation.

### Tactile feedback

The tactile feedback from [proprioception](https://en.m.wikipedia.org/wiki/Proprioception#Basis_of_proprioceptive_sense) is derived from the proprioceptors in the skin, muscles, and joints.

### Balance

The receptor for the sense of balance resides in the [vestibular system](https://en.m.wikipedia.org/wiki/Equilibrioception#Vestibular_system) in the ear (for the three-dimensional orientation of the head, and by inference, the rest of the body). Balance is also mediated by the kinesthetic reflex fed by [proprioception](https://en.m.wikipedia.org/wiki/Proprioception) (which senses the relative location of the rest of the body to the head). In addition, proprioception estimates the location of objects which are sensed by the [visual system](https://en.m.wikipedia.org/wiki/Visual_system) (which provides confirmation of the place of those objects relative to the body), as input to the mechanical reflexes of the body.

### Fine touch and crude touch

See also: [Two-point discrimination](https://en.m.wikipedia.org/wiki/Two-point_discrimination)



The [cortical homunculus](https://en.m.wikipedia.org/wiki/Cortical_homunculus), a map of somatosensory areas of the brain, was devised by [Wilder Penfield](https://en.m.wikipedia.org/wiki/Wilder_Penfield).

Fine touch (or discriminative touch) is a sensory modality that allows a subject to sense and localize touch. The form of touch where localization is not possible is known as crude touch. The [posterior column–medial lemniscus pathway](https://en.m.wikipedia.org/wiki/Posterior_column%E2%80%93medial_lemniscus_pathway) is the pathway responsible for the sending of fine touch information to the [cerebral cortex](https://en.m.wikipedia.org/wiki/Cerebral_cortex) of the brain.

Crude touch (or non-discriminative touch) is a sensory modality that allows the subject to sense that something has touched them, without being able to localize where they were touched (contrasting "fine touch"). Its fibres are carried in the [spinothalamic tract](https://en.m.wikipedia.org/wiki/Spinothalamic_tract%22%20%5Co%20%22Spinothalamic%20tract), unlike the fine touch, which is carried in the dorsal column.  As fine touch normally works in parallel to crude touch, a person will be able to localize touch until fibres carrying fine touch ([Posterior column–medial lemniscus pathway](https://en.m.wikipedia.org/wiki/Posterior_column%E2%80%93medial_lemniscus_pathway)) have been disrupted. Then the subject will feel the touch, but be unable to identify where they were touched.

### Neural processing of social touch

The somatosensory cortex encodes incoming sensory information from receptors all over the body. Affective touch is a type of sensory information that elicits an emotional reaction and is usually social in nature, such as a physical human touch. This type of information is actually coded differently than other sensory information. Intensity of affective touch is still encoded in the primary somatosensory cortex and is processed in a similar way to emotions invoked by sight and sound, as exemplified by the increase of adrenaline caused by the social touch of a loved one, as opposed to the physical inability to touch someone you don't love.

Meanwhile, the feeling of pleasantness associated with affective touch activates the anterior cingulate cortex more than the primary somatosensory cortex. [Functional magnetic resonance imaging](https://en.m.wikipedia.org/wiki/Functional_magnetic_resonance_imaging) (fMRI) data shows that increased blood-oxygen-level contrast (BOLD) signal in the anterior cingulate cortex as well as the prefrontal cortex is highly correlated with pleasantness scores of an affective touch. Inhibitory [transcranial magnetic stimulation](https://en.m.wikipedia.org/wiki/Transcranial_magnetic_stimulation) (TMS) of the primary somatosensory cortex inhibits the perception of affective touch intensity, but not affective touch pleasantness. Therefore, the S1 is not directly involved in processing socially affective touch pleasantness, but still plays a role in discriminating touch location and intensity.