# **Medical Neuroscience** | Tutorial Notes

## **Chemical Senses—Gustation**

## MAP TO NEUROSCIENCE CORE CONCEPTS<sup>1</sup>

- NCC1. The brain is the body's most complex organ.
- NCC3. Genetically determined circuits are the foundation of the nervous system.

### LEARNING OBJECTIVES

After study of the assigned learning materials, the student will:

- 1. Characterize the peripheral and central organization of the gustatory system.
- 2. Discuss sensory transduction in gustatory receptor cells.
- 3. Describe information coding in the gustatory system.

#### **TUTORIAL OUTLINE**

- I. Organization of the gustatory system
  - A. anatomical overview (see **Figure 15.17**<sup>2</sup>)
    - chemical constituents of foods interact with taste receptors cells located on epithelial specializations, called papillae, which contain the taste buds (see Figure 15.18)
    - 2. taste buds are distributed on the tongue, soft palate, epiglottis, pharynx and upper esophagus
    - taste receptor cells make synapses on the peripheral axons of cranial nerves VII, IX and X
      - a. anterior tongue (soft palate)  $\rightarrow$  facial nerve (CN VII)
      - b. posterior tongue  $\rightarrow$  glossopharyngeal nerve (CN IX)
      - c. epiglottis (esophogus)  $\rightarrow$  vagus nerve (CN X)
    - 4. these central axons project into the **solitary tract** and synapse in the *rostral* division of the **nucleus of the solitary tract** 
      - a. the posterior division of this nucleus is a visceral sensory relay (more on that in Unit 4 of this course)

<sup>&</sup>lt;sup>1</sup> Visit **BrainFacts.org** for *Neuroscience Core Concepts* (©2012 Society for Neuroscience ) that offer fundamental principles about the brain and nervous system, the most complex living structure known in the universe.

<sup>&</sup>lt;sup>2</sup> Figure references to Purves et al., *Neuroscience*, 5<sup>th</sup> Ed., Sinauer Assoc., Inc., 2012. [click here]

- b. integration of gustatory and visceral sensory inputs across the rostralcaudal axis of this nucleus facilitates appropriate visceral motor activities in response to harmful tastants, such as gagging or vomiting
- 5. the nucleus of the solitary tract projects to a distinct part of the **ventral posterior complex** in the thalamus
- this part of the ventral posterior complex projects to several cortical areas, including areas in the rostral **insula** and the frontal operculum (in the depth of the rostral lateral fissure)
- 7. these cortical regions are interconnected with the posterior **orbital prefrontal cortex**, where olfactory, gustatory and somatic sensory information is combined to produce *flavor* sensations from which the hedonic values of foods are represented
- B. sensory transduction (see **Figure 15.20**)
  - taste receptor cells have a variety of transduction mechanisms (see Figure 15.21)
    - a. some tastants interact with receptors on ion channels in the apical tip of the cell; the opening of Na<sup>+</sup> channels or the closing K<sup>+</sup> channels leads to the generation of a receptor potential
    - b. other tastants interact with G-protein coupled receptors that use second messengers to elevate intracellular Ca<sup>++</sup> concentrations and depolarize the membrane
  - 2. depolarization and elevated intracellular Ca<sup>++</sup> leads to the exocytosis of a chemical neurotransmitter (serotonin), which binds to receptors on the sensory axon
- C. gustatory coding & perception
  - 1. tastant identity is encoded by the activation of different receptors that are most sensitive to distinct chemical substances (see **Figure 15.19**)
    - a. there are receptors that are especially sensitive to:
      - i. sucrose (sweet)
      - ii. NaCl (salt)
      - i. HCl (acid)
      - ii. quinine (bitter)
      - iii. glutamate (Japanese, "umami" = "delicious")
    - b. detection thresholds are higher for substances that are important in our diet (e.g., salts and carbohydrates); in the *millimolar* range
    - c. detection thresholds are lower for potential harmful compounds (bitter-tasting alkaloids); in the *nanomolar* range
  - 2. the concentration of a tastant is generally correlated with the number of receptor cells activated and the intensity of afferent activation, as well as the perceived intensity of the taste

- 3. in central gustatory centers, as in olfactory centers, there are no obvious 'maps' or systematic representations of tastants or the sensory periphery
- 4. the nature of the gustatory code remains obscure, but it appears to operate according to a "labeled-line" code
  - i. different tastants activate distinct classes of receptor neurons that make synaptic contact onto peripheral processes of afferent neurons that are "labeled" by receptor specificity
  - ii. the responses of subcortical central neurons are tuned for tastant identity, reflecting receptor specificity
  - iii. nonetheless, at higher centers of processing (especially in the orbital cortex), the hedonic value of food becomes the most salient feature of representation, rather than the molecular identity of any particular tastant

### STUDY QUESTION

Which of the following statements concerning the encoding of gustatory signals is most accurate?

- A. Most gustatory receptor neurons express the full complement of genes that encode the five basic classes of gustatory receptors.
- B. Subcortical gustatory processing appears to implement a "labeled-line code" with the neural responses of sensory cells reflecting the molecular properties of the gustatory receptors that drive the response.
- C. There is a "combinatorial code" operating in the orbital cortex, since individual cortical neurons respond selectively to just one tastant.
- D. At higher stages of gustatory processing in the brain, the most salient property of food is the relative concentration of sour, bitter, salty, sweet, and "glutamate-like" tastants.