



- |  |  |
|--|--|
| 1. Optic sulcus                        | 24. Collateral sulcus                                      |
| 2. Olfactory bulb                      | 25. Pons   |
| 3. Olfactory tract                     | 26. Trigeminal nerve                                       |
| 4. Oculomotor nerve                    | 27. Abducens nerve   |
| 5. Anterior perforated substance       | 28. Facial nerve   |
| 6. Optic nerve                         | 29. Vestibulocochlear nerve                                |
| 7. Optic chiasma                       | 30. Flocculus of cerebellum                                |
| 8. Optic tract                         | 31. Choroid plexus from lateral recess of fourth ventricle |
| 9. Infundibulum (pituitary stalk)      | 32. Roots of glossopharyngeal, vagus, and accessory nerves |
| 10. Tuber cinereum and median eminence | 33. Spinal part of accessory nerve                         |
| 11. Mammillary body                    | 34. Rootlets of hypoglossal nerve (superficial to marker)  |
| 12. Posterior perforated substance     | 35. Vertebral artery                                       |
| 13. Anterior cerebral artery           | 36. Medulla oblongata                                      |
| 14. Middle cerebral artery             | 37. Tonsil of cerebellum                                   |
| 15. Internal carotid artery            | 38. Posterior inferior cerebellar artery                   |
| 16. Posterior communicating artery     | 39. Pyramid of medulla oblongata                           |
| 17. Posterior cerebral artery          | 40. Olive of medulla oblongata                             |
| 18. Oculomotor nerve                   | 41. Anterior inferior cerebellar artery                    |
| 19. Superior cerebellar artery         | 42. Basilar artery   |
| 20. Trochlear nerve                    |  |
| 21. Crus of cerebral peduncle          |  |
| 22. Uncus                              |  |
| 23. Parahippocampal gyrus              |  |

FIGURE 24-4 Inferior view of the human brain.

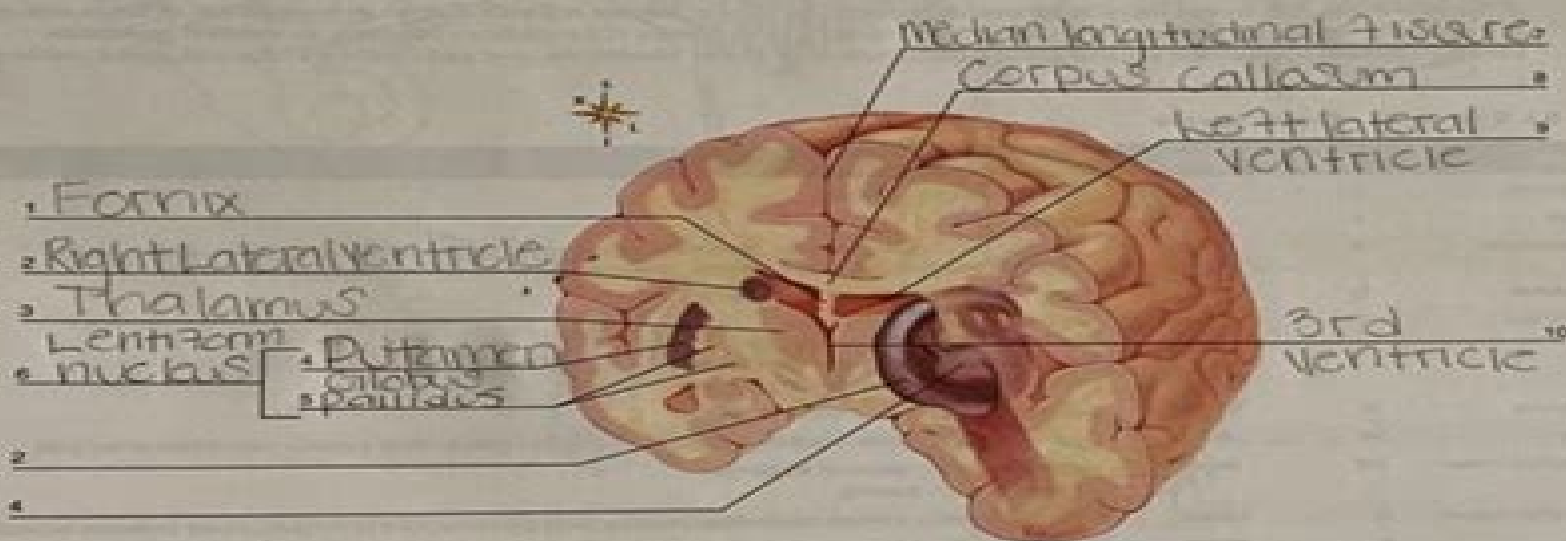


FIGURE 24-5 Label structures of the brain (frontal section) indicated on the lines provided and also on the blanks in the Lab Report at the end of this exercise.

# Laboratory Exercise 24 Surface Anatomy Answers

**Gerard J. Tortora**



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**Central Nervous System Tissue Engineering** Ashley E. Wilkinson, Aleesha M. McCormick, Nic D. Leipzig, 2022-06-01. Combating neural degeneration from injury or disease is extremely difficult in the brain and spinal cord, i.e. central nervous system (CNS). Unlike the peripheral nerves, CNS neurons are bombarded by physical and chemical restrictions that prevent proper healing and restoration of function. The CNS is vital to bodily function and loss of any part of it can severely and permanently alter a person's quality of life. Tissue engineering could offer much-needed solutions to regenerate or replace damaged CNS tissue. This review will discuss current CNS tissue engineering approaches integrating scaffolds, cells, and stimulation techniques. Hydrogels are commonly used CNS tissue engineering scaffolds to stimulate and enhance regeneration, but fiber meshes and other porous structures show specific utility depending on application. CNS-relevant cell sources have focused on implantation of exogenous cells or stimulation of endogenous populations. Somatic cells of the CNS are rarely utilized for tissue engineering; however, glial cells of the peripheral nervous system (PNS) may be used to myelinate and protect spinal cord damage. Pluripotent and multipotent stem cells offer alternative cell sources due to continuing advancements in identification and differentiation of these cells. Finally, physical, chemical, and electrical guidance cues are extremely important to neural cells serving important roles in development and adulthood. These guidance cues are being integrated into tissue engineering approaches. Of particular interest is the inclusion of cues to guide stem cells to differentiate into CNS cell types as well as to guide neuron targeting. This review should provide the reader with a broad understanding of CNS tissue engineering challenges and tactics with the goal of fostering the future development of biologically inspired designs.

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Introduction  
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