

Neuropathology

- I. General neuropathology and histology of CNS
 - A. Stains, tissue preparation
 - B. Cells, inclusion bodies, general
 - C. Cellular reactions to CNS injury
 - D. Edema
 - E. Tissue reaction to CNS injury and herniations
- II. Nervous system development, developmental pathology
 - A. Fetal development
 - B. Neural tube defects
 - C. Cleavage disorders
 - D. Migrational disorders
 - E. Cranium malformations
 - F. Chromosome disorders
 - G. Genetic disorders
 - H. Hydrocephalus
 - I. Perinatal brain injury
 - J. Disorders of hindbrain development
 - K. Lesions of developing grey matter
- III. Lesions of developing white matter infectious disease
 - A. Bacterial infections
 - B. Tuberculosis
 - C. Fungal and parasitic infections
 - D. Syphilis
 - E. Viral infections including HIV
 - F. Sarcoidosis
 - G. Opportunistic infections
 - H. Prion diseases
- IV. Neuro-oncology
 - A. Classification
 - B. Primary neoplasms (astrocytic, oligodendroglial, ependymal, choroid plexus, neuronal)
 - C. Molecular markers
 - D. Pineal tumors
 - E. Embryonal tumors
 - F. Schwannoma
 - G. Neurofibroma
 - H. Meningiomas
 - I. Non-meningothelial tumors
 - J. Lymphomas
 - K. Hematopoietic neoplasms
 - L. Metastases
 - M. Pituitary
 - 1. Normal pituitary
 - 2. Adenomas

3. Other sellar lesions

V. Encephalopathy due to intoxications, drugs, vitamin deficiencies, and metabolic diseases

- A. Cerebral hypoxia
- B. Electrolyte disturbances
 - 1. Hyponatremia
 - 2. Hypernatremia
- C. Vitamin deficiencies
- D. Toxic encephalopathies related to CNS toxins
- E. Secondary CNS involvement in systemic diseases
- F. Biochemical abnormalities
- G. Rare disorders
 - 1. Lysosomal storage disorders
 - 2. Sphingolipidoses
 - 3. Leukodystrophies
 - 4. Mucopolysaccharidoses
 - 5. Aminoacidopathies
 - 6. Peroxisomal disorders
 - 7. Mitochondrial disorders
 - 8. Glycogen storage diseases
 - 9. Enzyme deficiencies without storage problems
 - 10. Metabolic disorder
 - 11. Disorders of structural proteins

VI. Degenerative/ demyelinating diseases

- A. Multiple sclerosis
- B. Acute disseminated encephalomyelitis
- C. Neuromyelitis optica
- D. Dementia:
 - 1. Classification of dementia
 - 2. Alzheimer disease
 - 3. Dementia with Lewy bodies
 - 4. Vascular dementia
 - 5. Hippocampal sclerosis
 - 6. Movement disorders
 - 7. Cerebellar degenerations
 - 8. Motor neuron diseases
 - 9. Involvement of autonomic nervous system in degenerative disorders
 - 10. Hydrocephalus

VII. Cerebrovascular disease

- A. Aneurysmal subarachnoid and intraventricular hemorrhage
- B. Hypertensive cerebrovascular disease
- C. CAS
- D. Vascular malformations
 - 1. Aneurysms
 - 2. Arteriovenous malformation
 - 3. Cavernous malformation

- E. Small vessel disease
- F. CADASIL
- G. Vasculitis
- H. Microbleeds
- I. Stroke and lacunar infarcts
- J. Venous pathology

VIII. CNS Trauma – cranial/spinal

- A. Classification of TBI
- B. Scalp/skull lesions
- C. Concussion
- D. Contusions/lacerations
- E. Hemorrhage
 - 1. Epidural hematoma (EDH)
 - 2. Subdural hematomas (SDH)
 - 3. Traumatic SAH
 - 4. Intraventricular hemorrhage
- F. DAI
- G. Penetrating injuries, blast injuries
- H. Chronic traumatic encephalopathy
- I. Pediatric head injury
- J. Skull fractures
- K. Herniation syndromes
 - 1. Subfalcine
 - 2. Uncal
 - 3. Tonsillar
- L. Spinal shock
- M. Spinal cord syndromes
- N. Spine trauma – axial and subaxial fractures

IX. Spinal developmental lesions

- A. Lipoma
- B. Tethered cord
- C. Diastematomyelia
- D. Neuroenteric cysts
- E. Arachnoid cysts

X. Spinal tumors and spinal infections, infarcts, inflammatory diseases, degenerative

- A. Bone tumors
- B. Epidural lesions
- C. Intradural – extramedullary
- D. Intramedullary tumors
- E. Metastases
- F. Osteomyelitis
- G. Abscess
- H. Transverse myelitis
- I. ALS
- J. Polio

- K. Rheumatoid arthritis
- L. Spondylosis
- M. Stenosis
- N. OPLL
- O. Spinal infarcts
- P. Nerve root sleeve CSF cysts

XI. Peripheral nerve disorders, nm junction diseases/muscle diseases

- A. Skeletal muscle pathology
 - 1. Stains
 - 2. Fiber type
- B. Neurogenic atrophy
- C. Neuromuscular transmission defects
- D. Genetic diseases of skeletal muscle (muscular dystrophies)
- E. Congenital myopathies
- F. Metabolic myopathies
- G. Toxic myopathies
- H. Rhabdomyolysis
- I. Inflammatory myopathies
- J. Peripheral nerve histology
 - 1. Normal anatomy - axons, myelin, etc.
 - 2. Reaction of peripheral nerve to disease pathology
 - 3. Demyelination
 - 4. Axonal loss
- K. Inflammatory polyneuropathies
- L. Infectious/vasculitic paraneoplastic neuropathies
- M. Monoclonal gammopathies
- N. Nutritional/toxic neuropathies
- O. Hereditary motor-sensory neuropathies
- P. Porphyria