

Neurosurgical Oncology

- I. Anatomy and surgical approaches
 - A. Fundamental neuroanatomy
 - B. Cortical and functional anatomy: eloquent cortex, motor mapping, language/speech
 - C. Surgical approaches
- II. Neuro-imaging modalities and differential diagnosis
 - A. Tractography: white matter tracts, diffuse tensor imaging (DTI)
 - B. Metabolic imaging: positron emission tomography (PET), MR spectroscopy, perfusion
 - C. Functional Imaging: functional MRI, language mapping, motor mapping
 - D. Differential diagnosis
 1. Tumor imaging characteristics
 2. Radiation necrosis
 3. Non-neoplastic mass lesions: multiple sclerosis, tuberculoma, fungoma, neurocysticercosis
 - E. Intra-operative imaging: modalities, indications, extent of resection
- III. Clinical presentation and non-surgical management
 - A. Clinical presentation: signs and symptoms, syndromes (Gerstmann, SMA, Foster-Kennedy)
 - B. Medical management: steroids and steroid withdrawal, anti-epileptics
 - C. Radiation oncology
 1. Whole brain radiation and intensity-modulated radiation (gliomas, small cell, metastasis)
 2. Stereotactic radiosurgery and stereotactic radiation therapy (pituitary, meningioma, schwannoma, metastasis)
- IV. Gliomas and primary CNS lymphoma
 - A. High-grade gliomas: genetics, histopathology, imaging, surgical management, extent of resection, adjuvant therapy (Stupp protocol), prognosis
 - B. Low-grade gliomas: genetics, histopathology, imaging, surgical management, adjuvant therapy, prognosis
 - C. Other: juvenile pilocytic astrocytoma, pleomorphic xanthoastrocytoma, ganglioglioma, gliosarcoma, Lhermitte-duclos (dysplastic gangliocytoma of the cerebellum) - imaging, management, and prognosis
 - D. Primary CNS lymphoma: clinical presentation, imaging, management, prognosis
 - E. Large B cell lymphoma: clinical presentation, imaging, management, prognosis
- V. Skull base tumors
 - A. Meningioma: genetics, histopathology, classification, clinical presentation, imaging, surgical management/approaches, adjuvant therapies (radiation), prognosis
 - B. Vestibular schwannoma: genetics, histopathology, clinical presentation, imaging, surgical management/approach, adjuvant therapy, intra-operative monitoring (BAERs)
 - C. Other: epidermoid cyst, trigeminal schwannoma, glomus jugulare tumor, dermoid cysts, lipoma, paraganglioma, hemangiopericytoma

- VI. Pituitary tumors
 - A. Clinical syndromes: prolactinoma, Cushing's disease, acromegaly, MEN syndrome
 - B. Pituitary adenoma: histopathology, imaging, clinical presentation (visual field defect, hypopituitarism, apoplexy), medical and surgical management
 - C. Complications: DI, SIADH, panhypopituitarism, CSF leak, epistaxis
- VII. Pineal region tumors
 - A. Pineal parenchymal tumors: pineocytoma, pineoblastoma
 - 1. Imaging, presentation (Parinaud, nystagmus hydrocephalus), surgical management, adjuvant therapy
 - B. Germ cell tumors: germinoma, embryonal carcinoma, choriocarcinoma, yolk sac endodermal sinus tumor, teratoma
 - 1. Genetics, histopathology, tumor markers, surgical management, prognosis
- VIII. Intraventricular tumors
 - A. Central neurocytoma: histopathology, imaging, management
 - B. Ependymoma and subependymoma: histopathology, imaging, management
 - C. Colloid cyst: histopathology, imaging, management
 - D. Choroid plexus papilloma, choroid plexus carcinoma: histopathology, imaging, management
- IX. Pediatric tumors
 - A. Supratentorial: craniopharyngioma, pinealoma, DNET
 - B. Infratentorial: pilocytic astrocytoma, medulloblastoma, ependymoma, DIPG, atypical teratoid rhabdoid tumor
 - C. Other: neuroblastoma
- X. Brain metastases
 - A. Common primary tumors: lung, breast, melanoma, renal cell, adenocarcinoma incidence rates
 - B. Surgical indications: solitary vs. oligo metastasis
 - C. Medical Management: immune modulation therapies
 - D. Leptomeningeal disease: imaging, management, prognosis
- XI. Neurocutaneous syndromes
 - A. Neurofibromatosis: NF-2, genetics, presentation, imaging, management, prognosis
 - B. Von-Hippel Lindau: genetics, presentation, imaging, management, prognosis
 - C. Tuberous sclerosis: genetics, presentation, imaging, management, prognosis
 - D. Sturge-Weber: genetics, presentation, imaging, management, prognosis
 - E. Cowden disease: genetics, presentation, imaging, management, prognosis
- XII. Spinal cord tumors
 - A. Chordoma
 - B. Vertebral hemangioma
 - C. Osteoid osteoma
 - D. Other: schwannoma, meningioma, lipoma, filum terminale tumor, multiple myeloma
- XIII. Complication management
 - A. Intra-operative complications: mass effect, edema, hydrocephalus

B. Postoperative complications: infection, hydrocephalus, venous thrombosis