INTRACRANIAL GLIOMAS-RADIOLOGY, DIAGNOSIS AND MANAGEMENT

Introduction

- WHO classification (2000) of CNS tumors:
- 1. Neuroepithelial tissue
- 2. Cranial an spinal nerves
- 3. Meninges
- 4. Hematopoietic system
- 5. Germ cell
- 6. Cysts and tumor like condition
- 7. Tumors of the sellar region
- 8. Local extension of regional tumors
- 9. Metastatic tumors

Neuroepithelial tumors

- Sub classification:
- 1. Astrocytic tumors:
 - A. Diffusely infiltrating:
 - 1. Astrocytoma- fibrillary

protoplasmic

gemistocytic

mixed

- 2. Anaplastic astrocytoma
- 3. Glioblastoma multiforme: Grade 1V

Giant cell

Gliosarcoma

- B. Circumscribed lesions:
 - 1. Pilocytic astrocytoma
 - 2. Plemorphic Xanthoastrocytoma
 - 3. Subependymal Giant cell astrocytoma

- 2. Oligodendrogliomas
- 3. Ependymal tumors: ependymomas

anaplastic

myxopapillary

subependymomas

- 4. Mixed gliomas
- 5. Choroid plexus tumors: papilloma

carcinoma

Tumors of uncertain origin: astroblastoma

polar spongioblastoma

gliomatosis cerebri

- 7. Neurons: a. ganglioglioma
 - b. dysplastic gangliocytoma
 - c. desmoplastic infantile ganglioglioma
 - d. dysplastic neuroepithelial tumors
 - e. gangliocytoma
 - f. central neurocytomas
 - g. olfactory neuroblastomas

- 8. Pineal tumors: pinealocytomas pinealoblastomas mixed pineal tumors
- 9. Embryonal tumors: medulloepithelioma

neuroblastoma

retinoblastoma

primitive neuroectodemal tumors(PNET):

- a. medulloblastoma
 - 1. desmoplastic
 - 2. medullocytoma
 - 3. melanotic
- b. cerebral and spinal PNET's

Low Grade Gliomas

Classified as:

- WHO grade II astrocytomas
- 2. Oligodendrogliomas
- 3. Mixed oligodendrocytic/ astrocytic
- 4. Gangliogliomas
- 5. Juvenile pilocytic astrocytomas
- 6. Pleomorphic astrocytomas
- 7. Dysembryoplastic neuroepithelial tumors

Spatial definition

- Type I:
 Solid tumor without infiltration:
 pilocytic astrocytoma,
 ganglioglioma
- Type II:
 Solid tumors with infitration:
 grade II astrocytoma
- Type III:
 Infilterative without solid component:
 oligodendroglioma

Grading of gliomas

WHO

Based on cellularity, pleomorphism, vascular, proliferation, necrosis

I Special tumors

Il Astrocytomas (low grade)

III Anaplastic Astrocytomas

IV Glioblastoma

- Kernohan
- St Anne/Mayo :atypia,mitosis,endothelial proliferation,necrosis

Grading by CT/MRI

- I: CT; Low density, MRI; Abnormal signal no mass effect, no enhancement
- II: CT; low density, MRI; abnormal signal mass effect, no enhancement
- III: Complex enhancements
- IV :Necrosis (ring enhancement)

Pathological features

Diffuse astrocytomas:

Gross findings: Adults hemispheric, children brainstem. Alter color, texture of tissue.

Microscopy: Hyper chromatic nuclei

GFAP containing cell process

Astrocytic cells without

macrophages/ microglia

Cells distributed in clumps(fibrillary/

protoplasmic, gemistocytic)

Anaplastic astrocytomas

- Margins better defined
- Cells populous/pleomorphic
- Perineural satellitoses
- Absence of necrosis/ vascular proliferation

Glioblastomas

- Occur denovo
- Dedifferentiation
- Macroscopic picture: grey fleshiness/ cystic / necrotic areas wide spread
- Microscopic picture: Astrocytes with long fine process with vascular proliferation/ necrosis
 large cells bizarre- giant cell GBM gliosarcoma
- Adequate pathological specimen required

Pilocytic astrocytoma

- Macroscopic picture:
 Optic nerve- fusiform dilatation
 Hypothalamic- soft lobular
 Cerebellar- well circumscribed
 Cerebral
- Microscopic picture:
 Elongated cells/ rosenthal fibres

Pleomorphic xanthoastrocytomas

- Young adults
- Superficial cortical areas
- Temporal lobe
- Cellular pleomorphism
- Lipidisation of cells
- Low mitotic rate
- Necrosis absent

Subependymal giant cell astrocytoma

- Tuberous sclerosis
- Abundant eosinophilic cytoplasm
- Neurofilament protein

Oligodendroglioma

- Cerebral hemispheres: frontal lobe
- Both grey and white matter
- Monotonous round cells
- Perineural satellitoses
- Calcification
- Perinuclear haloes
 - GFAP negative
 - High/ low grade

Ependymoma

- Frontal horns to filum
- IV ventricle/ filum
- Discrete
- Perivascular pseudo rosette
- Myxopapillary variant- caudal cell mass
- Gangliogliomas: both neural and glial cells

Clinical features

- Due to raised ICP: headache nausea/ vomiting nuchal rigidity papilloedema VI nerve palsy
- Progressive nature

Diagnosis

- Imaging- pre operative postoperative intra operative
- CT
- MRI
- SPECT
- Angiography

Imaging appearance

- Homogenous changes: reduced attenuation on CT hypo on T1W hyper on T2W
- Mixed/ inhomogenous changes
- Mildly/ homogenously increased density on CT/ T1W
- Hypo/ iso on T2W

Grading features

- Homogenity
- Margins
- Cysts/ necrosis
- Calcification
- Edema

Low Grade Astrocytomas

- CT: low density homogenous mass that is poorly marginated
- MRI: hypo on T1W/ hyper on T2W mass effect is less non enhancement
- D/D: cerebral infarct area of demyelination

Anaplastic astrocytomas

- Mass effect is greater
- Moderate contrast enhancement
- Absence of areas of necrosis and hemorrhage
- D/D solitary metastasis/ acute demylination

Glioblastoma multiforme

- Irregular homogenous areas of abnormality that are poorly marginated
- Hyper density on CT
- Hyper on T1W/ hypo on T2W
- Contrast enhancement
- Non enhancing areas
- Areas of hemorrhage
- D/D: Metastases/ abscess/ malignant meningioma

Oligodendroglioma

- Centrum semiovale of cerebral hemispheres/ found in adults
- CT. X-Rays prominent calcification
- MRI similar to astrocytoma
- D/D: central neurocytoma/ intraventricular meningiomas

Ependymoma

- Location
- Mimic low/ intermediate grade gliomas
- Homgenous
- Moderate enhancement
- Gangliogliomas:

temporal lobe/ posterior fossa calcification/ cysts

Other modalities

- Angiography: tumor/ giant aneurysm superficial/ intra axial/ extra axial masses vascularity
- FDG/ PET: grading/ treatment response/ recurrence/ radionecrosis
- MRS: lactate/ choline/ NAA/ creatine

- Diffusion perfusion weighted images
- SPECT, thallium201/ technetium99/ iodine
 123
- Functional imaging- MRI/ PET/ MEG

PreoperativeTreatment

- Anticonvulsants
- Peritumoral edema:

steroids/head end elevation/ mannitol/ diuretics/ hyper ventilation

Definitive treatment

- Surgery
- Radiotherapy
- Chemotherapy
- Others

Surgery

- Timing: early/ late
- Type of surgery: experience of surgeon attempt at gross total removal image guided surgery

 Indication: establishing diagnosis pilocytic astrocytoma mass effect hydrocephalus seizures to delay adjuvant therapy small tumors

 Role of surgery limited in: disseminated tumors

multifocal tumors

location in eloquent areas

Radiotherapy

- Controversial area
- No conclusive benefit
- Risk of radiation induced complications

- Recommendations:
 - in cases of gross total removal/ incomplete removal in pilocytic astrocytomas RT to be withheld
 - incomplete removal of low grade glioma 45 GY to tumor bed and 2 cm margin
 - malignant degeneration to be treated RT
 - Chiasmal/ hypothalamic tumors

Chemotherapy

- Not indicated except in tumor progression
- PCV regimen
- Pilocytic astrocytoma: cerebellum/ optic nerve/ hypothalamic glioma

Thank you