BRAIN TUMORS

- Most common : Secondary >> primary/ benign >>> primary / malignant

 Risk factors:
 Radiation, viruses (EBV, CMV), Trauma, immunosuppression, Racial, Genetic

 Hippel-Lindau disease (hemangioblastoma)/
 Li- Fraumeni syndrome (gliomas) / tuberous sclerosis (subependymal astrocytoma). Optic nerve gliomas in NF1 / multiple meningioma in NF2

Clinical Manifestations According to .. Site /size / grade/ hormones

- 1. Increased ICP >> From the mass itself or obstruction
- 2. NDs >> It depends on the site of the mass >> frontal lobe syndrome/ parietal lobe syndrome
- 3. CSF Obstruction... hydrocephalus
- 4. Seizures >> More from low grade & superficial tumors
- 5. Hormonal symptoms & Headache (dull & constant .. am)

Investigations :

- **1.** *X- ray* : limited help... widened sutures in children, calcifications, osteolytic lesions of the skull .
- 2. CT: It used to detect hemorrhage in them / if MRI is contraindicated
- 3. MRI : 🔶 T1 with C / T2/ FLAIR/MRA/ MRS -> to determine the chemical composition of the brain
- 4. PET: combined with 2 or 3 ... to show the result of Tx >> to differentiate between recurrent tumor and radionecrosis after radiotherapy.
- 5. Cerebral Angiography: to know which vessel supplies the tumor.

A. Metastasis

- From lung, breast, colon & kidney.
- After 45 yrs ... supratentorial (85%), cerebellum (15%)
- According to it's site ... headache, seizures,
- CT / MRI -> well enhance with C. -> with Vasogenic edema
- Tx : Steroids & anti-epileptic ... Remove or Debulk
- & take a **biopsy** to know it's primary origin +/- Chemotherapy/ Radiotherapy.

- with calcifications:
- 1. meningiomas
- 2. Craniopharyngioma
- 3. oligodendrogliomas

B. THE Gliomas 50%

- Arise from the neuroectoderm/ glial cells.
- Difficult in removal & Tx... 1 yr survival of pt
- Mostly in 50-70 yrs

1. Astrocytomas: MC.

- Arising from astrocytes.
- 1. Grade 1 : Pilocytic D Benign , Children Cyctic lesion in Cerebellum, in adult .. at brainstem

2. Grade 2 : Diffuse Fibrillary Denign , Children & Adult, slowly growing In Cerebellum , Optic nerve , Hypothalamus -> with no margins/ capsule

3. Grade 3 : <u>Anaplastic</u> Rapid growth , infiltrative. Highly vascular with areas of necrosis & high mitotic

4. Grade 4 : <u>Glioblastoma</u> Malignant, Rapid growth , solid & cystic With necrotic & hemorrhagic areas , new vs ... in cerebral hemispheres.

>> MRI, CT , Special dyes >> to localize it >> Remove or Debulk it >> reduce the symptoms, & radiotherapy dose

>> Steroids & Antiepilepic

2. Oligodendrogliomas

- ~ From oligodendrocytes, Younger
- Superficial areas -> Seizures
- •Well differentiated, Calcification -> on CT, MRI
- Surgery: totally *Restricted* + chemo/ radio

C. Medulloblastomas

-Childhood, in midline posterior fossa/ cerebellum

- > CSF obstruction, vomiting, headache, truncal Ataxia

- Malignant -> by CSF -> to spinal theca -> cauda equina
- MRI, CT >> enhance well, noncommunicating hydrocephalus
- TX: CSF diversion -> Complete Removal +Radiotherapy

D. Meningiomas

- Adult F> M ... from Arachnoids >> Extra axial
- Idiopathic & Due to ionizing radiation, NF2 associated
- Calcification / Hyperostosis of adjacent bone.

- Most common sites [olfactory groove, parasagittal area, convexity of the brain, supra sellar area, Intraventricular]

- MRI, CT >> enhance well with C. + Vasogenic edema - TX:
- Complete Removal -> Radiotherapy
- steroids +/- anti epileptics
- Endovascular embolization / highly vascular

E. Vestibular Shwanoma

- benign, posterior fossa tumor, from Schwann cells.
- from CN VIII, unilateral hearing loss, tinnitus, disequilibrium unsteady gait, dizziness ... late : CN V & VII involvement
- MRI/ CT with C. >> it well enhance

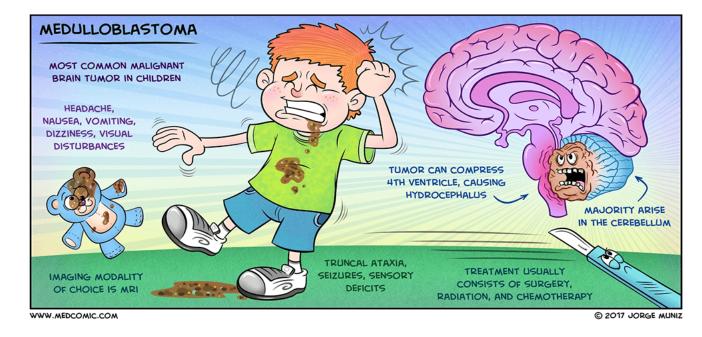
- CN testing >> Weber & Rinne test, decreased corneal Reflelx Facial twisting ~ ipsilateral

3. Ependymomas

- Around cavities, infratentorial -> Hydrocephalus.
- •Children , young adult.
- •CT, MRI >> in ventricular system, elk+/ calcification.
- Surgery : totally Restricted + chemo/ radio (seed tumor cells)

F. Pituitary Adenoma

- benign, from ant. lobe
- Macro vs. Micro ... Functional vs. Non-functional
- optic chiasm -> bitemporal hemianopia
- Cushing syndrome, acromegaly, hyperprolactinemia.
- hemorrhage or necrosis -> pituitary insufficiency
- MRI/ CT + Endocrine tests + Visual Field testing
- In micro- prolactinomas -> medications/dopamine agonist
- Surgery -> severe visual manifestations, failure of medical
- treatment -> transphenoidal or transethmoidal
- Post op endocrine replacement therapy.



С.

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E.

