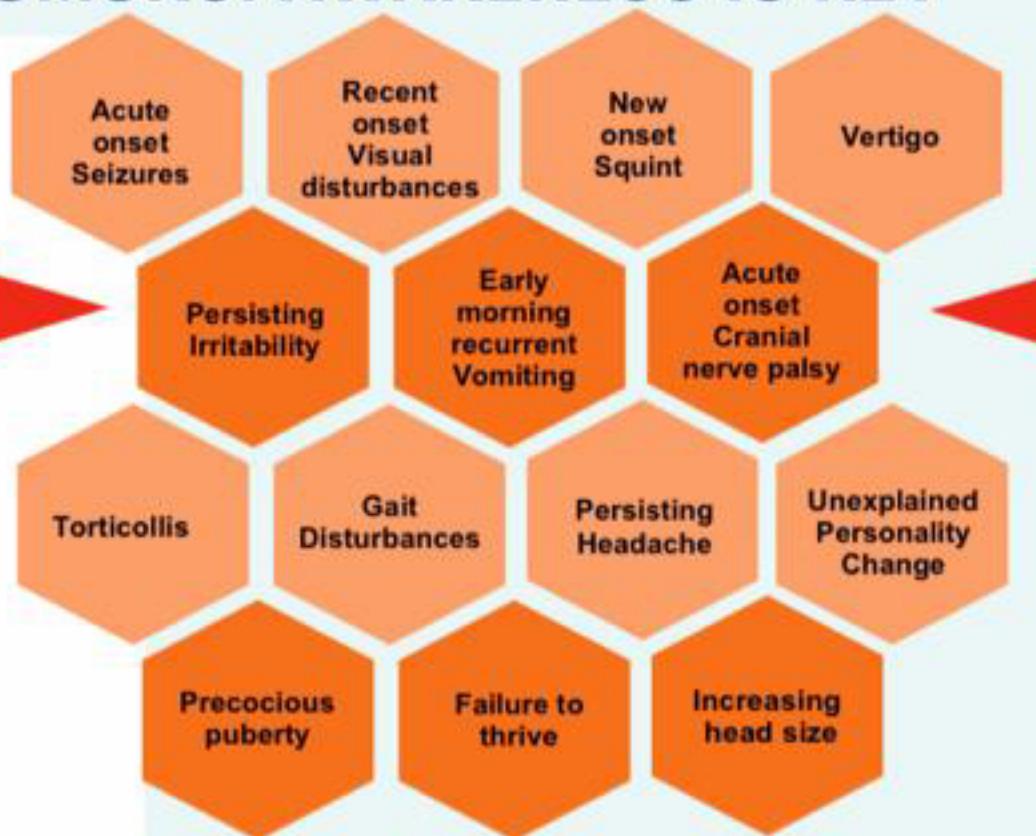


UNMASKING PEDIATRIC BRAIN TUMORS: AWARENESS IS KEY

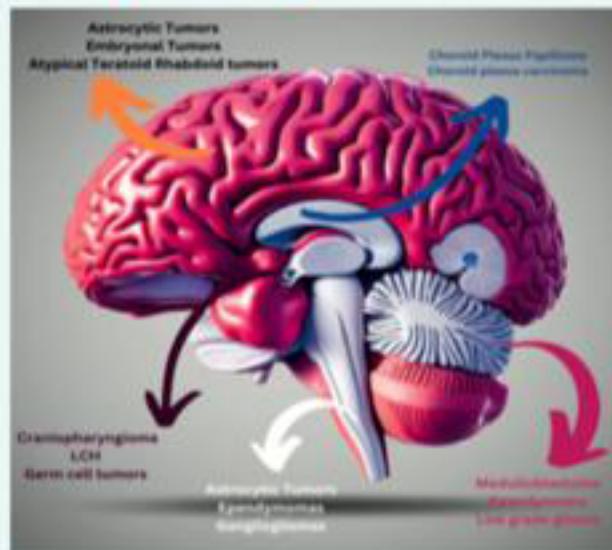
Fact File

- Brain tumors are the most common type of solid tumor in children
- Second to leukemia as a cause of childhood malignancies
- Estimated incidence in India - 12,900 cases/year
- Leading cause of childhood cancer-related deaths
- Prognosis depends on the age at presentation, histological type, and extent of resection

Red Flag Signs - Indication to do Neuro Imaging



Differential diagnosis depending on location



Infants & Toddlers

- Teratoma
- Medulloblastoma
- Embryonal tumors
- Rhabdoid tumor
- Low grade glioma
- Choroid plexus papilloma

Young children (<10 y)

- Pilocytic astrocytoma
- Medulloblastoma
- Ependymoma
- Germ cell tumors
- Craniopharyngiomas

Older children & teenagers

- Germ cell tumors
- Embryonal tumors
- Ependymomas
- High-grade glioma
- Pineoblastoma
- Meningioma

Differential diagnosis depending on age

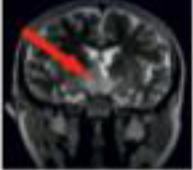
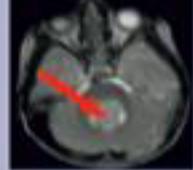
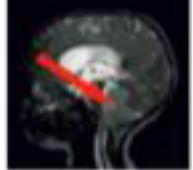
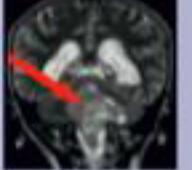
Initial investigations: an outline

Investigation	Purpose/Rationale
Ultrasound head	Accessible and useful in infants
CT Brain	Quickest and most readily available modality Can pick up haemorrhage, hydrocephalus and calcifications
MRI of Brain with Spine	Preferable modality of choice prior to definitive surgery or biopsy
Serum electrolytes	Associated diabetes insipidus or SIADH
Serum AFP & β-HCG	Diagnostic tumor markers for intracranial Germ cell tumors can possibly avoid biopsy/surgery if elevated

Genetic syndromes associated with brain tumors

Li-Fraumeni syndrome	Lynch syndrome	Rhabdoid tumor predisposition syndrome	DICER1 syndrome
Gorlin syndrome	Von Hippel-Lindau syndrome		Neurofibromatosis type 1 and type 2
			Tuberous sclerosis

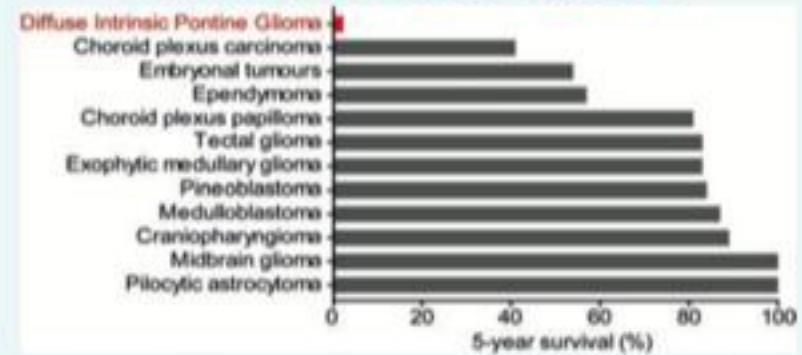
Common 4 tumors encountered

LOW GRADE GLIOMA	MEDULLOBLASTOMA	EPENDYMOMA	HIGH GRADE GLIOMA
<ul style="list-style-type: none"> ▪ Most common brain tumor in children ▪ Can be associated with NF-1 ▪ Biopsy can be avoided in optic pathway glioma 	<ul style="list-style-type: none"> ▪ Second most common brain tumor ▪ Arises from posterior fossa ▪ Gait disturbances and signs of raised ICP 	<ul style="list-style-type: none"> ▪ Can arise from either supratentorial, infratentorial or spinal cord ▪ Symptoms depends on site of tumor 	<ul style="list-style-type: none"> ▪ Can arise from any location ▪ Associated with mismatch repair deficiency if presents early in life ▪ Prognosis is generally guarded 
<ul style="list-style-type: none"> ▪ If resectable, complete resection if sufficient ▪ Chemotherapy in younger children ▪ Radiotherapy in older children- Avoid if possible in younger children 	<ul style="list-style-type: none"> ▪ Surgery followed by adjuvant radiotherapy and chemotherapy ▪ In those younger than 3 years, radiation sparing chemotherapy can be considered 	<ul style="list-style-type: none"> ▪ Surgery – Near total resection preferred ▪ Radiation in certain grades of ependymoma ▪ Role of chemotherapy is limited 	<ul style="list-style-type: none"> ▪ Surgery – Near total resection preferred ▪ Adjuvant therapy with focal radiation and temozolamide ▪ Immunotherapy in children with mismatch repair deficiency

Treatment principles

Surgery	Biopsy Hydrocephalus treatment – Shunt / Omaya Debulking or definitive surgery Least morbid surgery with maximal safe resection
Radiotherapy	Definitive treatment in >3 years
Chemotherapy	Least toxic regimen Targeted therapy in low grade glioma Concurrent chemoradiotherapy in high grade glioma

Survival in brain tumors



Survival for most brain tumors have improved over the past decade due to early diagnosis, better understanding of disease biology and improvement in treatment protocols

