

# Case 4

# Patient Profile

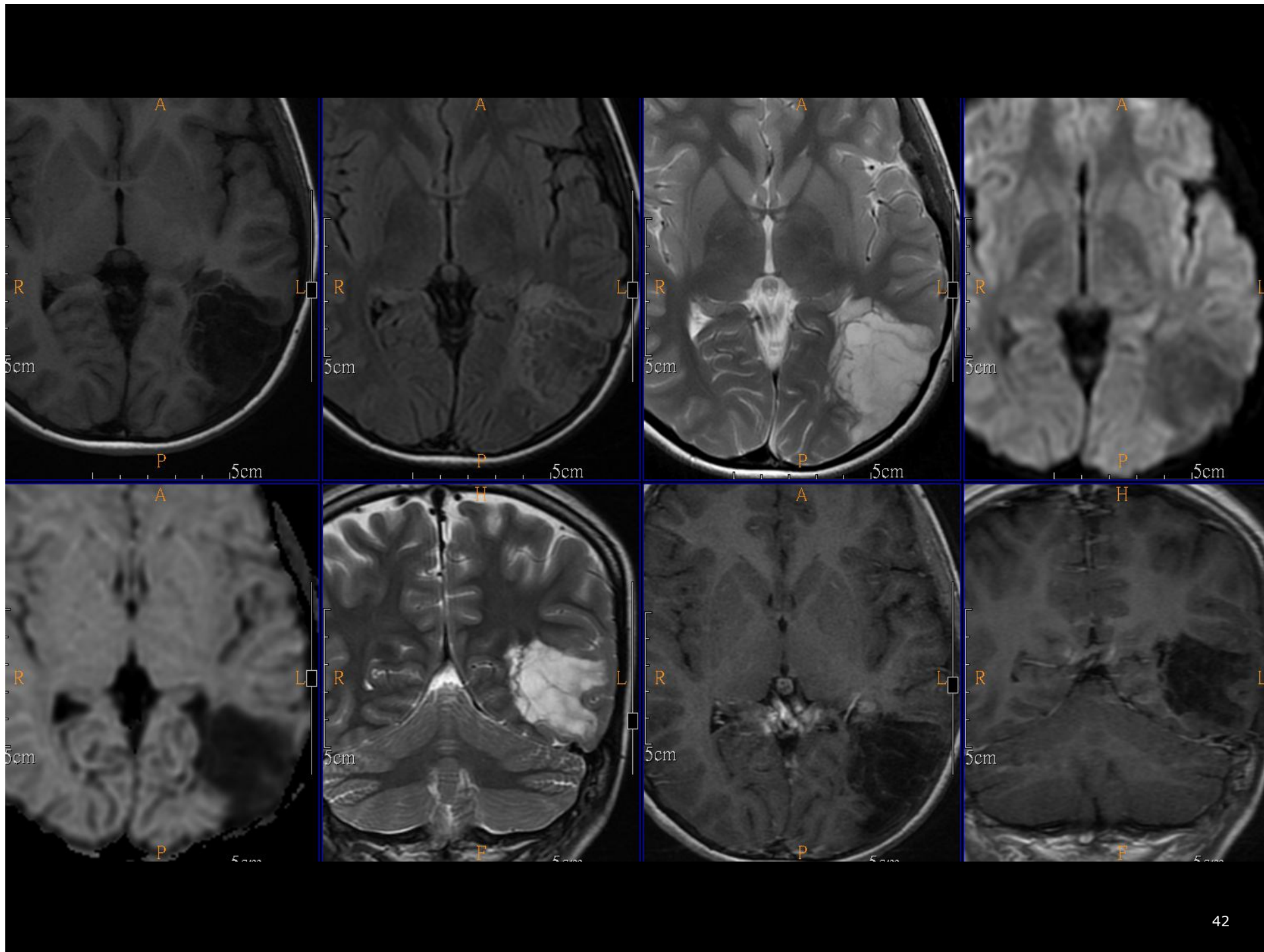
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- 11 years old girl
- Chief complaint
  - Seizure for years
- Past history
  - nil

# Image

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- 2018-03-27 Brain MRI

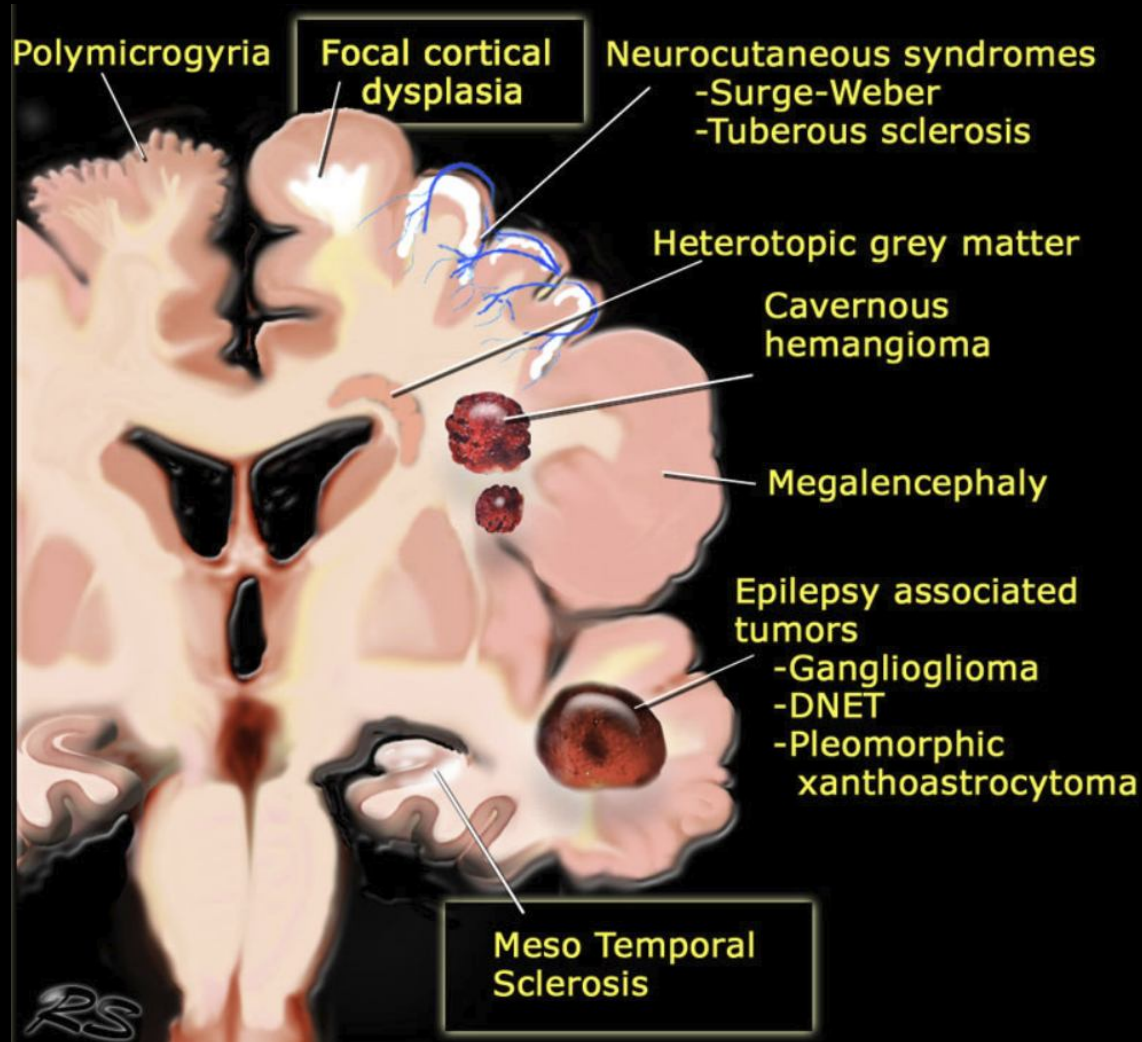


# Clinical Course

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- Diagnosis
  - Previous diagnosis of dysembryoplastic neuroepithelial tumor in other hospital

# Common Cause of Seizure



Radiology Assistant  
<http://www.radiologyassistant.nl>

# Dysembryoplastic Neuroepithelial Tumor

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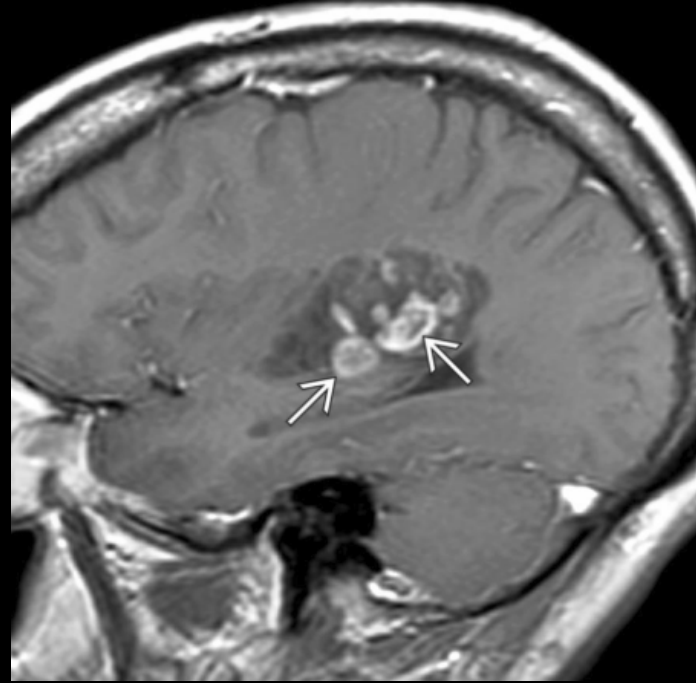
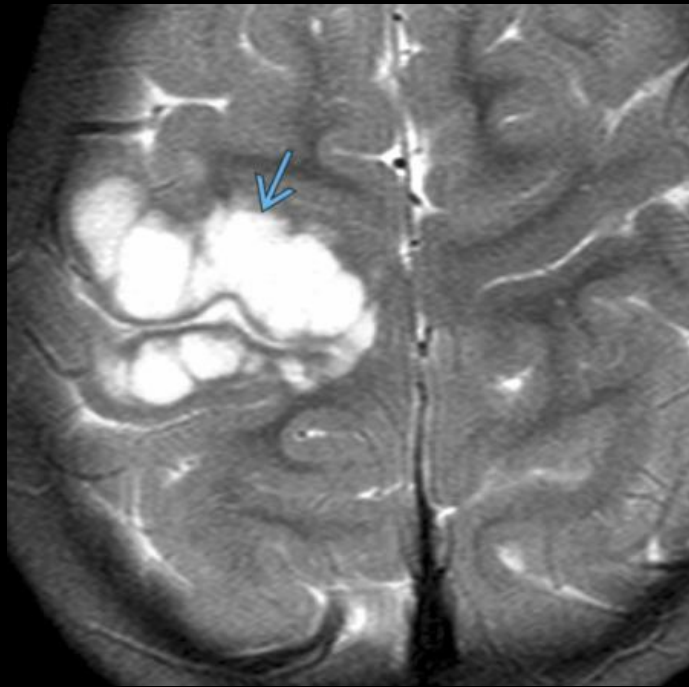
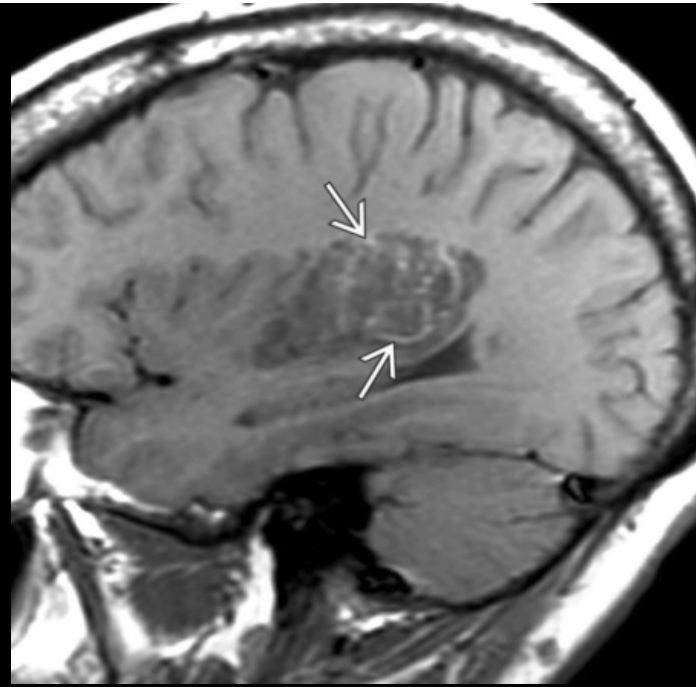
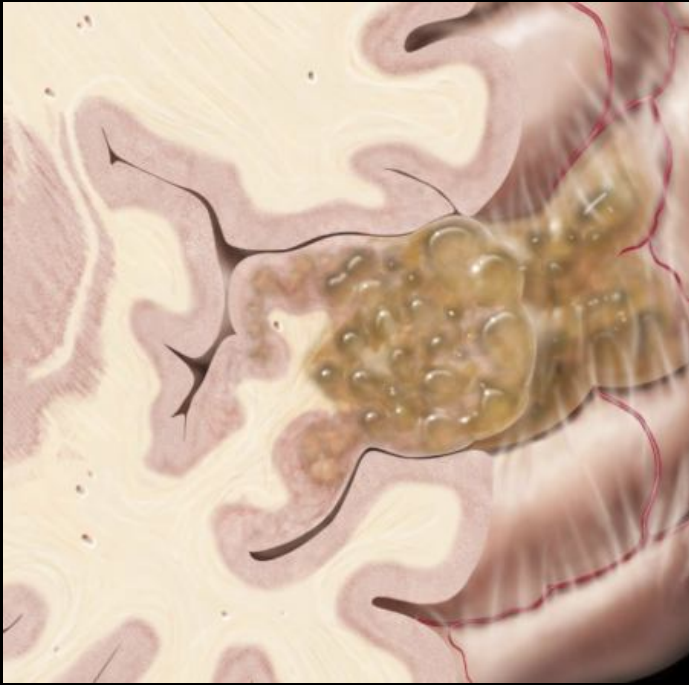
- Benign mixed glial-neuronal neoplasm
- Frequently associated with cortical dysplasia
- WHO grade I brain tumor
- Clinical symptoms
  - Longstanding drug-resistant **partial complex seizures** in child/young adult
  - Slow growth over years

# Image Findings

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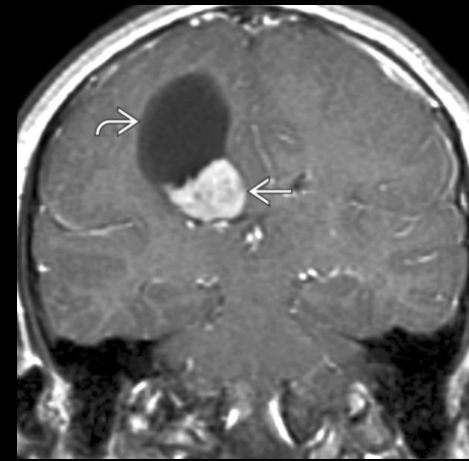
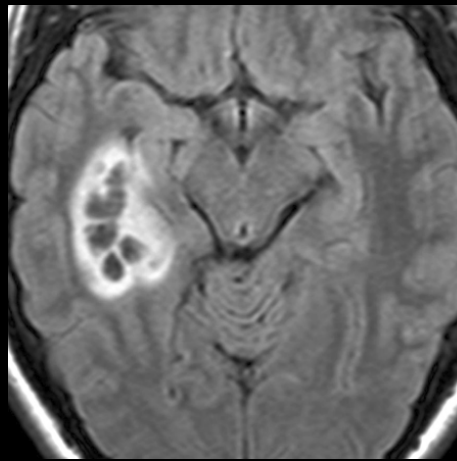
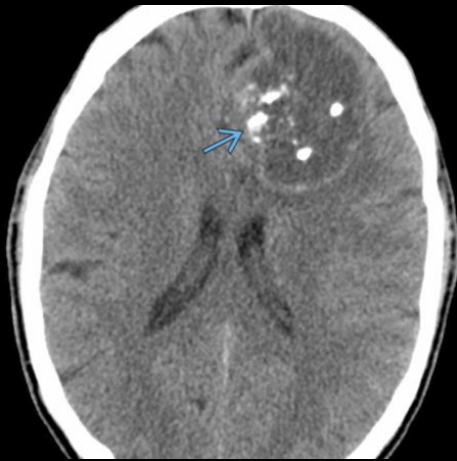
- May occur in any region of supratentorial cortex
  - Temporal lobe most common, followed by frontal lobe
  - Mass frequently "points" toward ventricle
- Sharply demarcated, wedge shaped
  - Cystic ("bubbly") intracortical mass
  - Minimal/no mass effect
  - No surrounding edema
  - Usually does not enhance





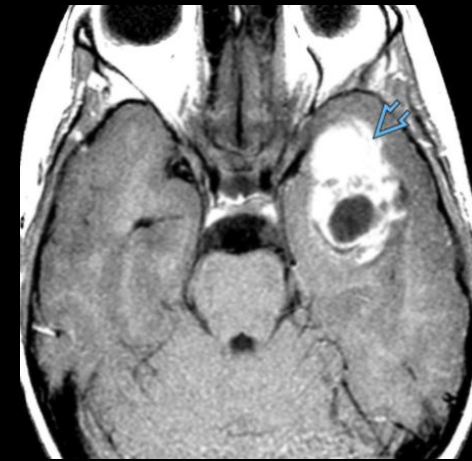
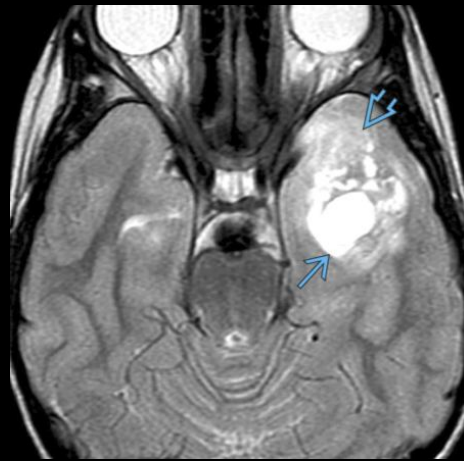
# Differential Diagnosis

- Ganglioglioma
  - Ca++ common
  - Frequently solid and cystic components
  - Solid components avidly enhance



# Differential Diagnosis

- Pleomorphic Xanthoastrocytoma (PXA)
  - Enhancing nodule abuts pia
  - May have pial enhancement
  - Look for dural "tail"



# Seizure Related of Brain Tumor

Seizure 44 (2017) 98–107



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journal homepage: [www.elsevier.com/locate/yseiz](http://www.elsevier.com/locate/yseiz)



Review

Seizures caused by brain tumors in children<sup>☆</sup>

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# Supratentorial Brain Tumor

**Table 1**  
Main characteristics of the most frequent pediatric brain tumors.

	Frequency	Location	Pathology	5-year survival
<b>Supratentorial brain tumors</b>				
Diffuse astrocytoma (grade II)	++	Cerebral hemispheres: frontal and temporal lobes (less frequently, brainstem and spinal cord)	Meningeal fibrils. No mitoses. No vascular proliferation	+++
Anaplastic astrocytoma (grade III)	+	Supratentorial	Increased cellularity, pleomorphism, mitoses	-
Glioblastoma multiforme (grade IV)	+	Supratentorial	Increased cellularity, pleomorphism, mitoses. Vascular proliferation. Necrosis	--
Ganglioglioma	+	Cerebral hemispheres: temporal > frontal	Mixture of neoplastic glial and neuronal elements	++++
Dysembryoplastic neuroepithelial tumor (DNET)	-	Cerebral hemispheres: temporal > frontal > parietal lobes. Centered in the cerebral cortex, although it might extend to the white matter	Multinodular microcystic lesions with floating neurons. Pleomorphism, glial-neural elements. May be surrounded by areas of dysplasia. Limited mitoses.	++++
Supratentorial primitive neuroectodermal tumor	+	Cerebral hemispheres	Undifferentiated or poorly differentiated neuroepithelial cells. Pathology similar to neuroblastoma.	+
Oligodendroglioma	-	Cerebral hemispheres (rarely medulla)	Honeycomb appearance: repetitive pattern of similar rounded cells with perinuclear halo	Variable depending on pathology and age. Higher risk of relapse than medulloblastoma. Variable depending on the other histologic components of the tumor

# Supratentorial Brain Tumor

**Table 1**  
Main characteristics of the most frequent pediatric brain tumors.

	Frequency	Location	Pathology	5-year survival
Posterior fossa tumors				
Pilocytic astrocytoma	++++	Cerebellum (less frequently cerebral hemispheres, optic pathway, hypothalamus, and brain stem)	Compact areas with Rosenthal fibers and spongy areas with microcysts	++++
Medulloblastoma	++++	Posterior fossa	Classic variant: densely packed cells with hyperchromatic nuclei and scant cytoplasm Desmoplastic/nodular variant: nodular, reticular-free zones of neuronal maturation surrounded by densely-packed mitotically active cells Extensive nodularity variant: expanded lobular architecture Large cell variant: monomorphic cells with large, round, vesicular nuclei with prominent nucleoli Anaplastic variant: cells with marked nuclear pleomorphism and nuclear moulding	+++  Variable depending on pathology and age
Ependymoma	+++	2/3 infratentorial, 1/3 supratentorial	Rosettes and pseudorosettes. Histologic features predict degree of malignancy poorly	++
Hypothalamic tumors				
Hypothalamic hamartoma	–	Hypothalamus	Abnormally distributed, but cytologically normal small neurons and glia	++++