# 10 y/o with blurry vision, vomiting, and headache

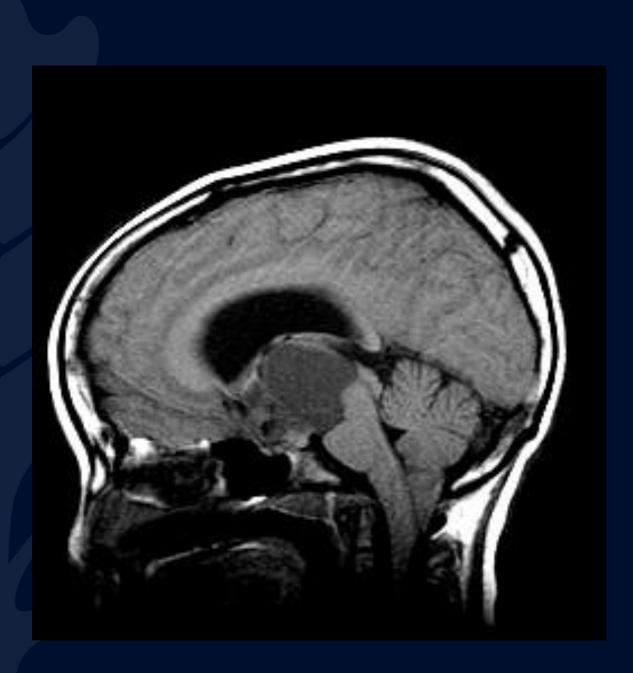
Jignesh Modi, MD







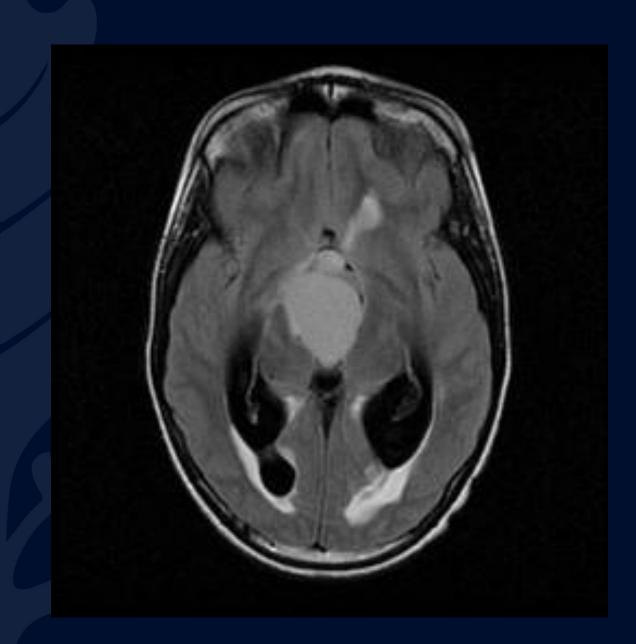
RADIOLOGY



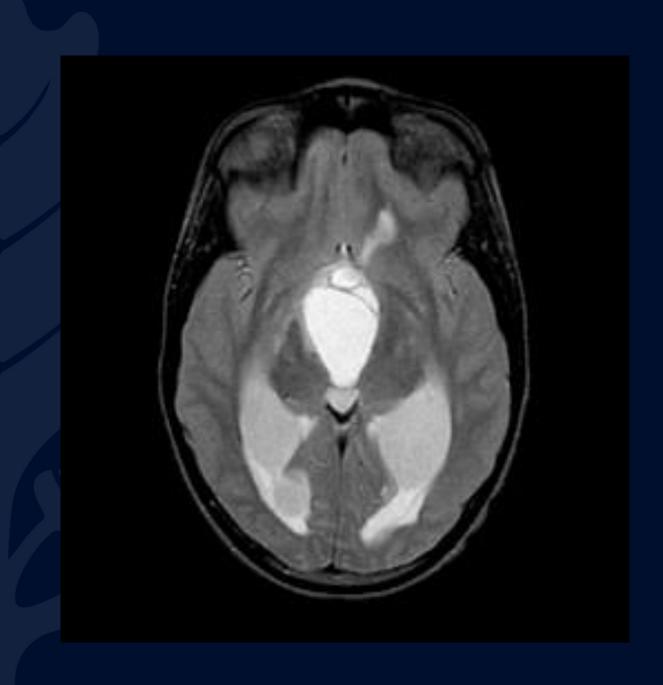












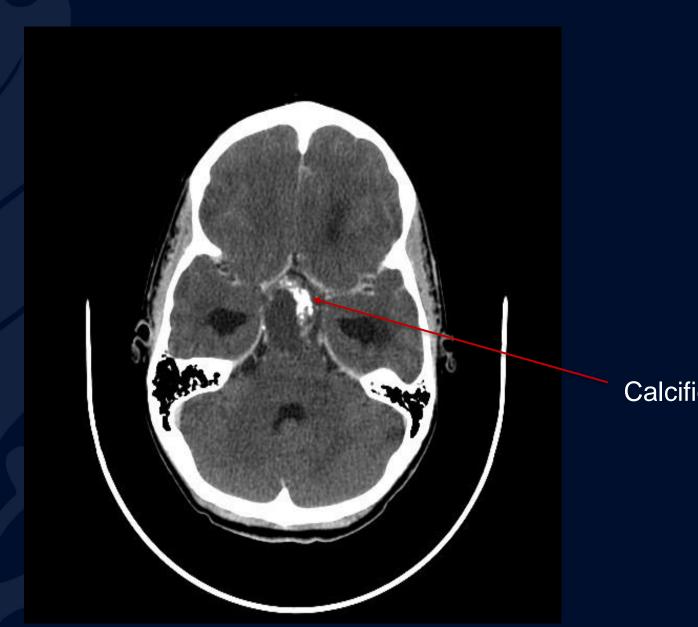






# Craniopharyngioma





Axial contrast enhanced CT

Calcifications





#### Sagittal T1 Pre-Contrast

#### Cystic Suprasellar Mass



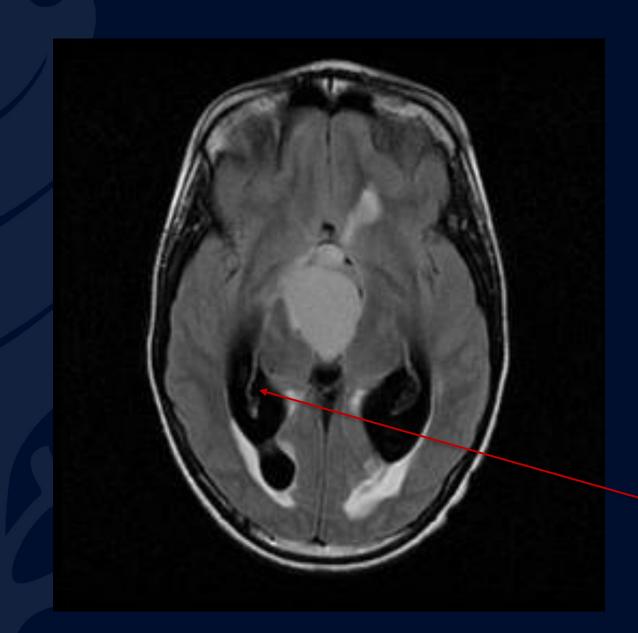
#### Sagittal T1 Pre-Contrast



#### Cystic Suprasellar Mass



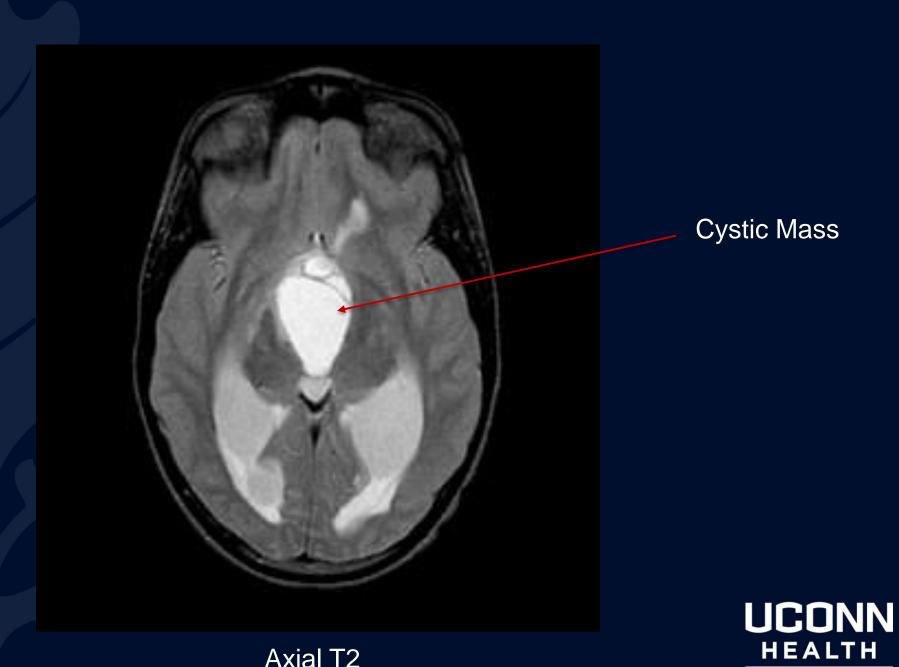
#### Sagittal T1 Post-Contrast



Hydrocephalus



Axial Flair



Axial T2

RADIOLOGY



#### Suprasellar region

#### Pituitary

Schematic of pituitary anatomy



# Craniopharyngioma

- Slow-growing, benign, dysontogenetic epithelial tumor derived from Rathke's cleft.
- Two histologic subtypes
  - Papillary: less prevalent, typically seen in adults.
  - Adamantinomatous: more prevalent in all age groups, typically seen in children. Classic presentation is a child with morning headache, visual disturbance, and short stature.
- Approximately 54% of all pediatric sellar and suprasellar lesions, peak age is 5-10 yrs, second small peak in 6th decade of life.



# Imaging Features

- Adamantinomatous
  - CT: multilobulated, multicystic, partially calcified mass.
  - CTA: vascular displacement and encasement.
  - MR: hyperintense cysts and heterogenous nodule.
- Papillary
  - CT: predominantly solid +/- cystic component, isodense, rarely calcify. Solid component and cyst wall avidly enhance.



## Treatment

- Cranipharyngiomas are generally benign and rarely undergo malignant transformation.
- Both subtypes are considered WHO grade I tumors.
- Complete resection can be curative, however, treatment often also incorporates irradiation.
- Surgical resection via craniotomy or endoscopically— via endonasal/transsphenoidal approach, when feasible, however endoscopic transsphenoidal approach can be more challenging given incomplete development of the sphenoid sinus.
- Surgical management, especially in children, remains controversial.
- Imaging is critical in evaluating this tumor with regards to tumor location and adjacent structure involvement
  – tumor involvement of and/or proximity to frontal, temporal lobes, ventricle, optic chiasm and nerves, hypothalamus, pituitary gland, circle of willis, and brain stem all impact treatment plan



### References

 Barkovich, A. James and Raybaud, Charles, "Pediatric Neuroimaging, 6th ed." (2019). *Faculty Bookshelf*. 121.
 Applied Radiology. 2021;50(5);58-60. cdn.agilitycms.com

