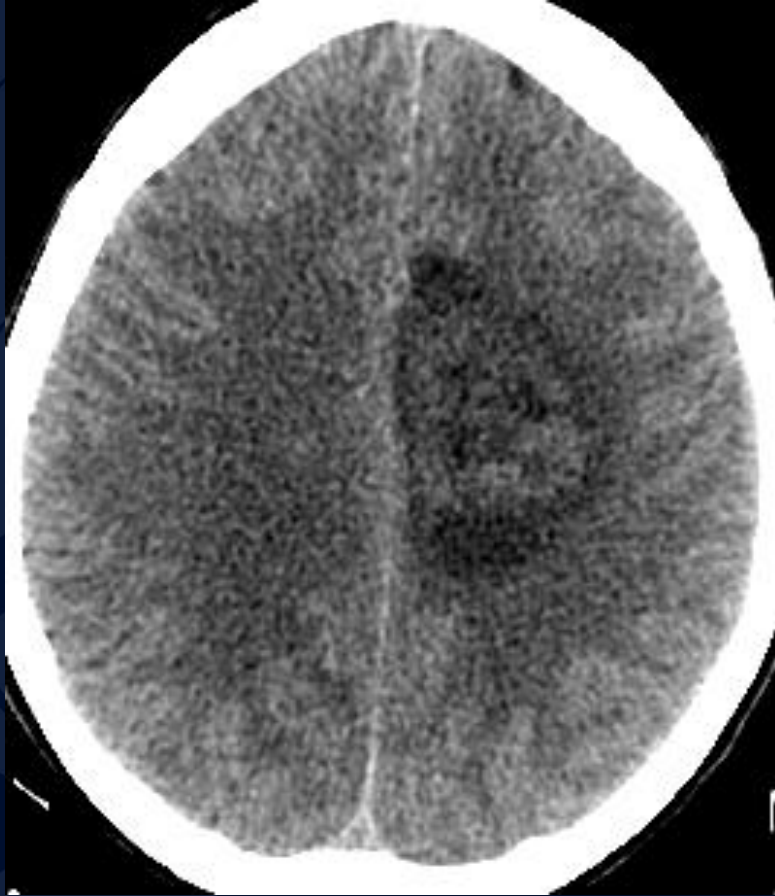


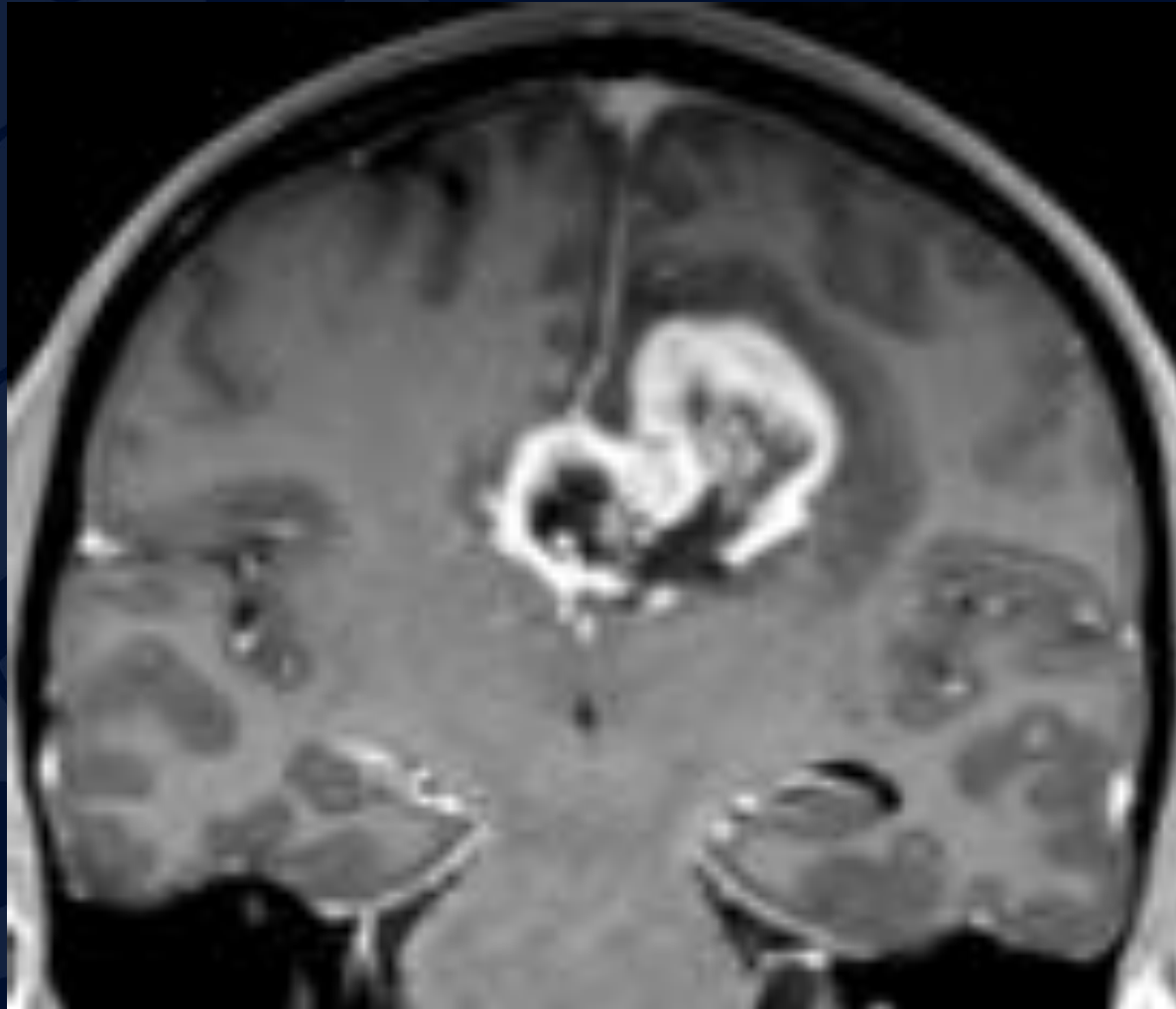
42 y/o female with 2 weeks of
ataxia, 2 days of nausea/vomiting

Daniel Chen, MD
Leo Wolansky, MD

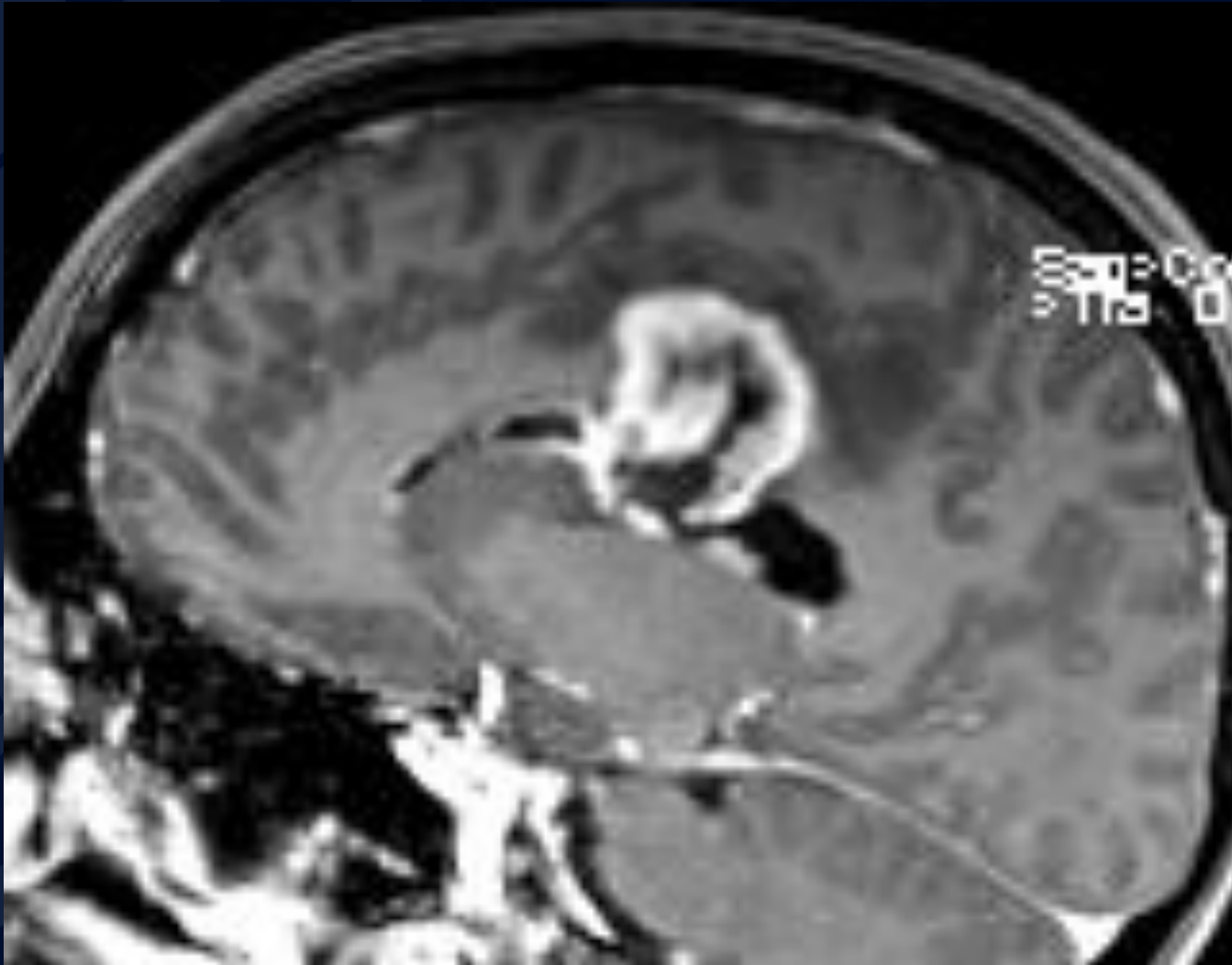
CT w/o contrast



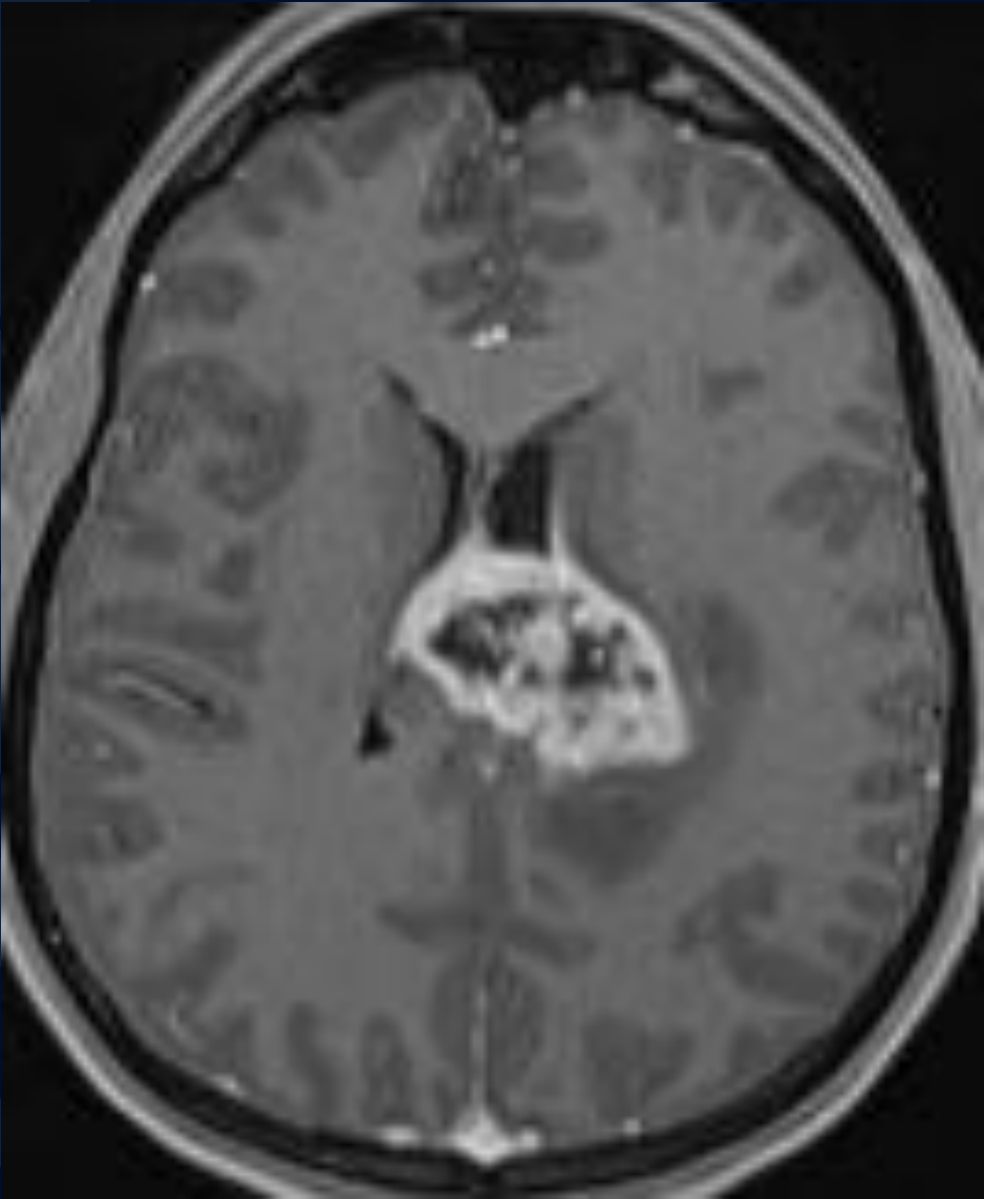
T1-Gd Coronal



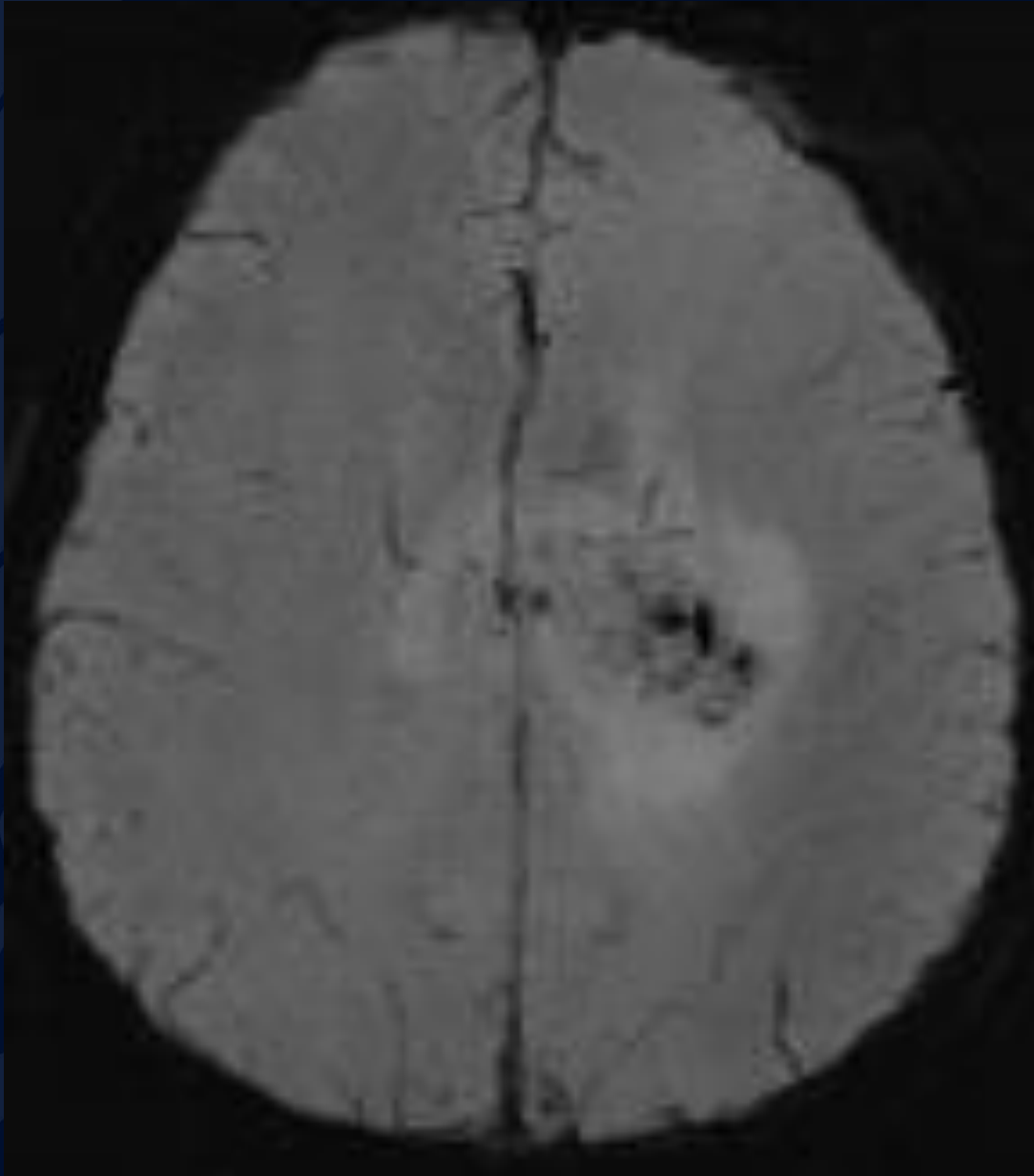
T1-Gd Sagittal



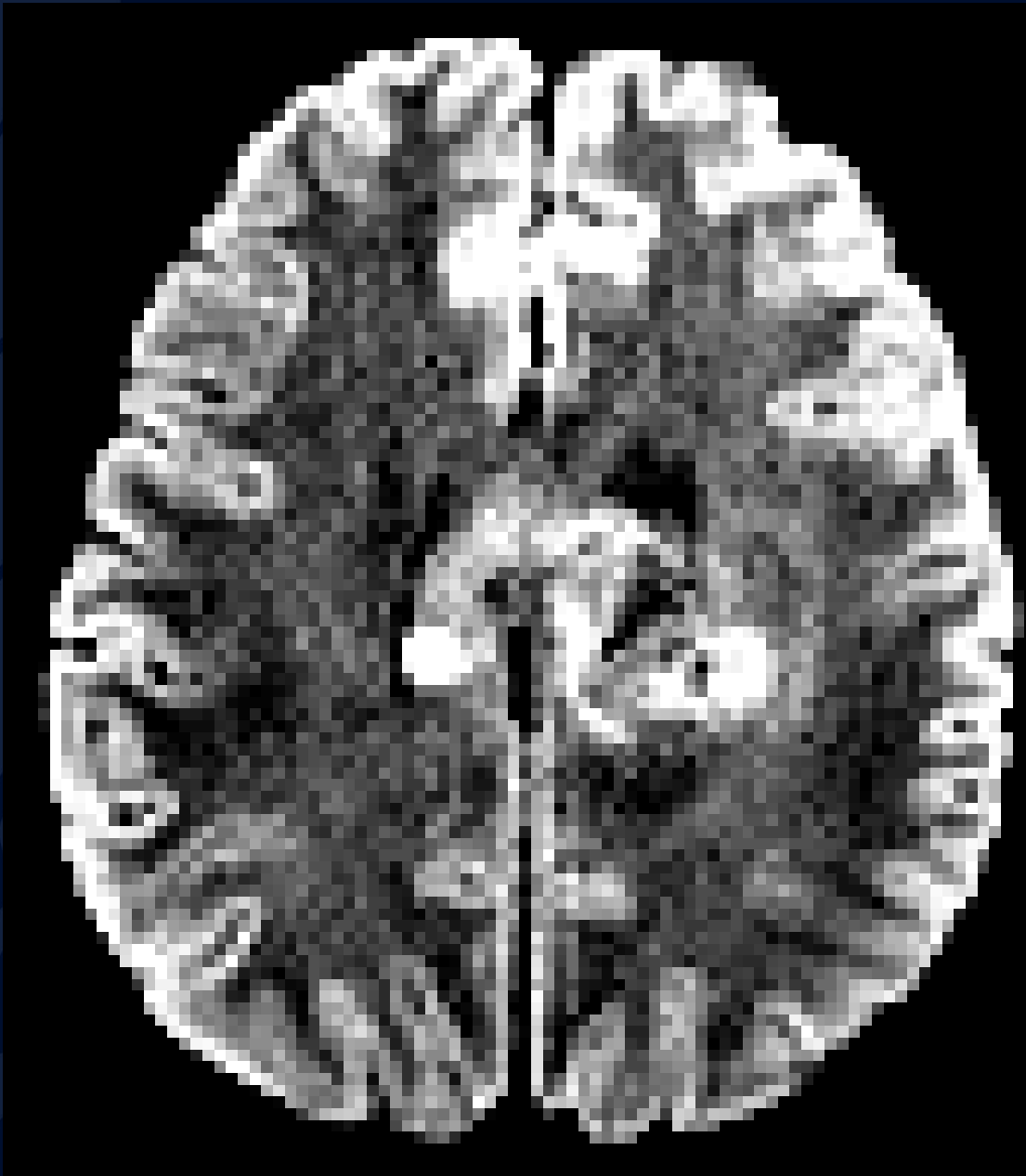
T1-Gd Axial



SWI Axial

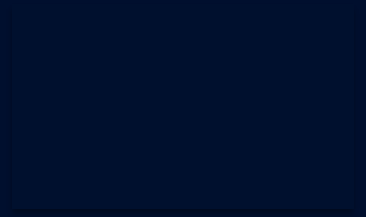


DWI Axial





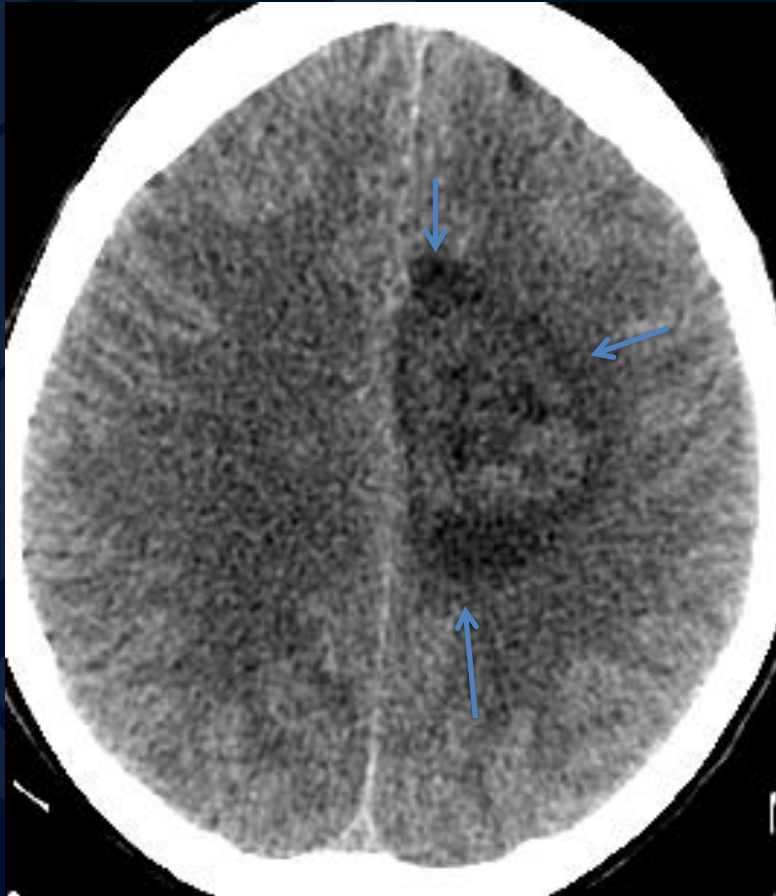
?



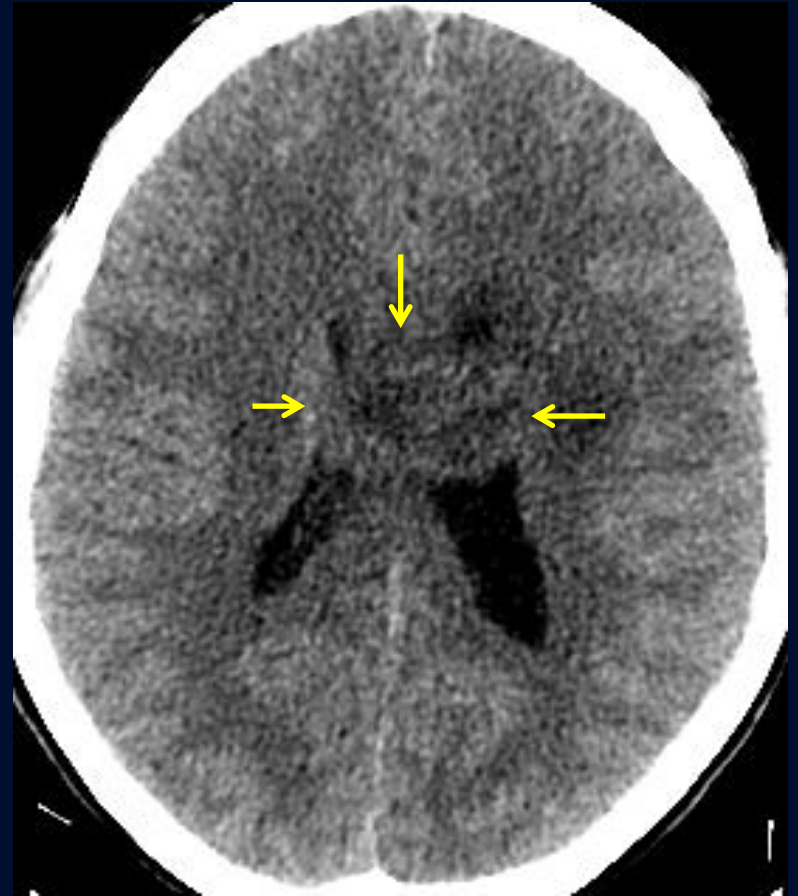
The background of the slide features a dark blue, stylized illustration of oak leaves on the left side, extending towards the center. The leaves are layered and have a textured, vein-like appearance. The rest of the background is a solid, dark blue color.

Glioblastoma (multiforme)

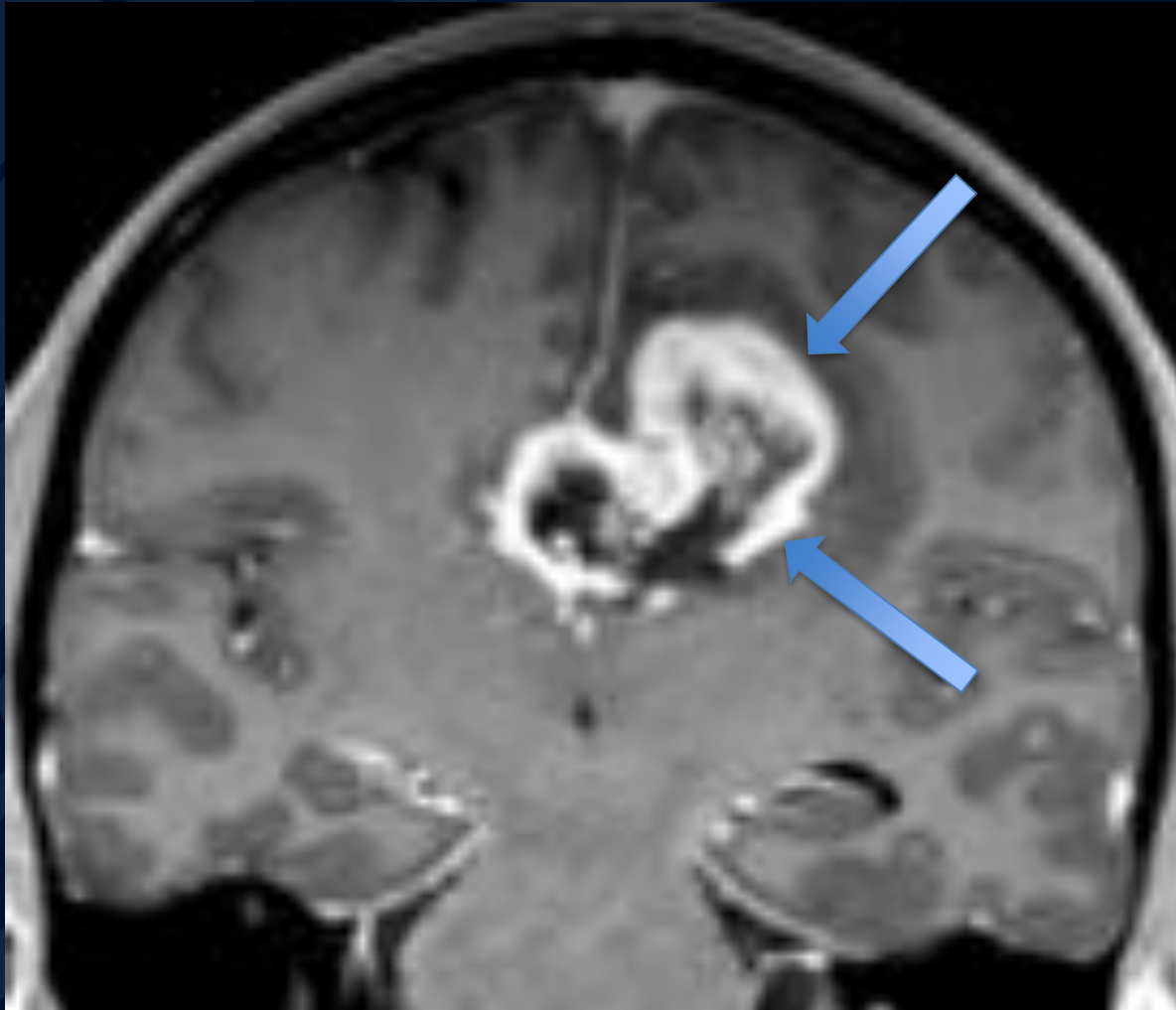
Corpus callosum involvement (yellow arrows)



Surrounding vasogenic
edema (blue arrows)

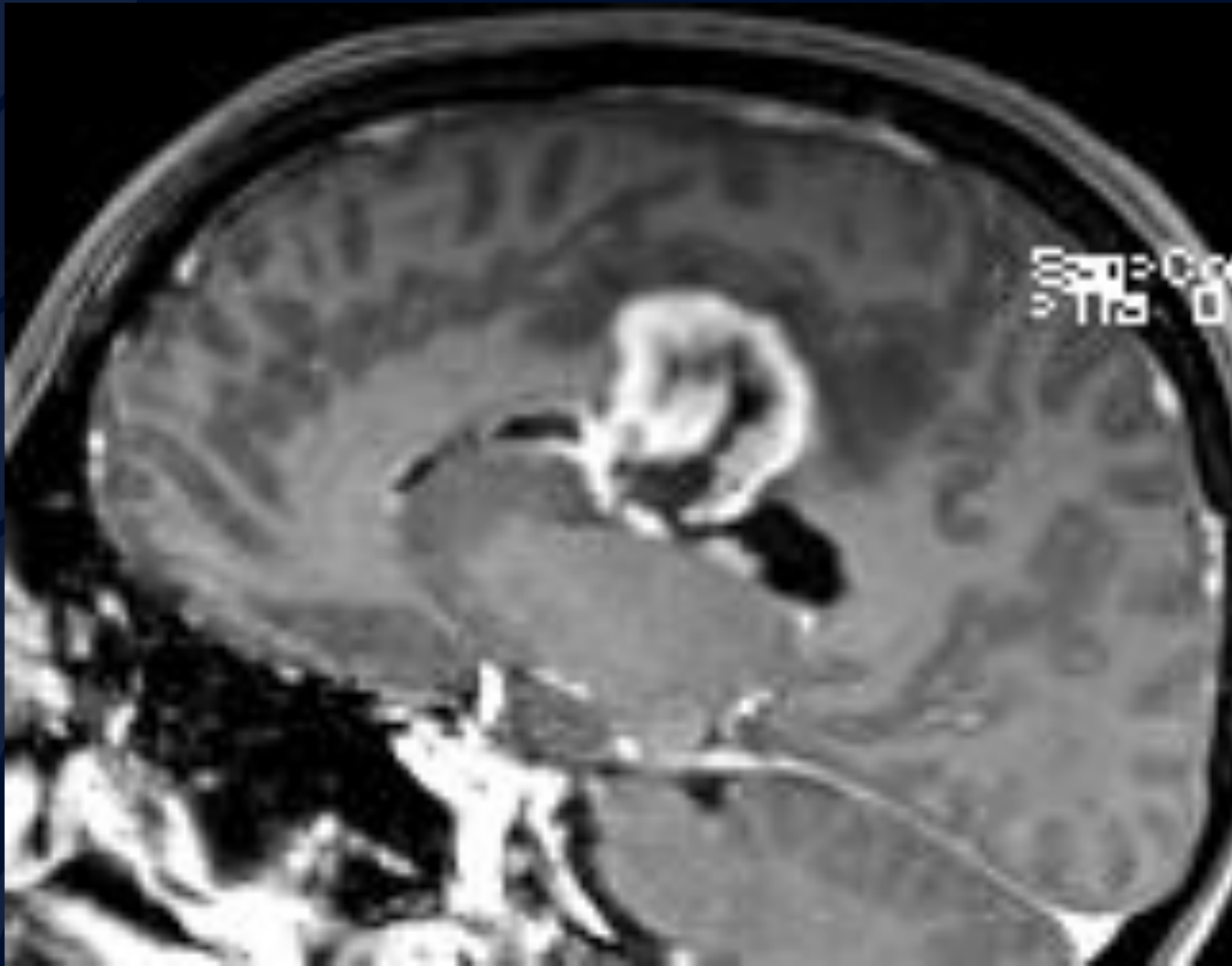


Thick, irregular rim-enhancement

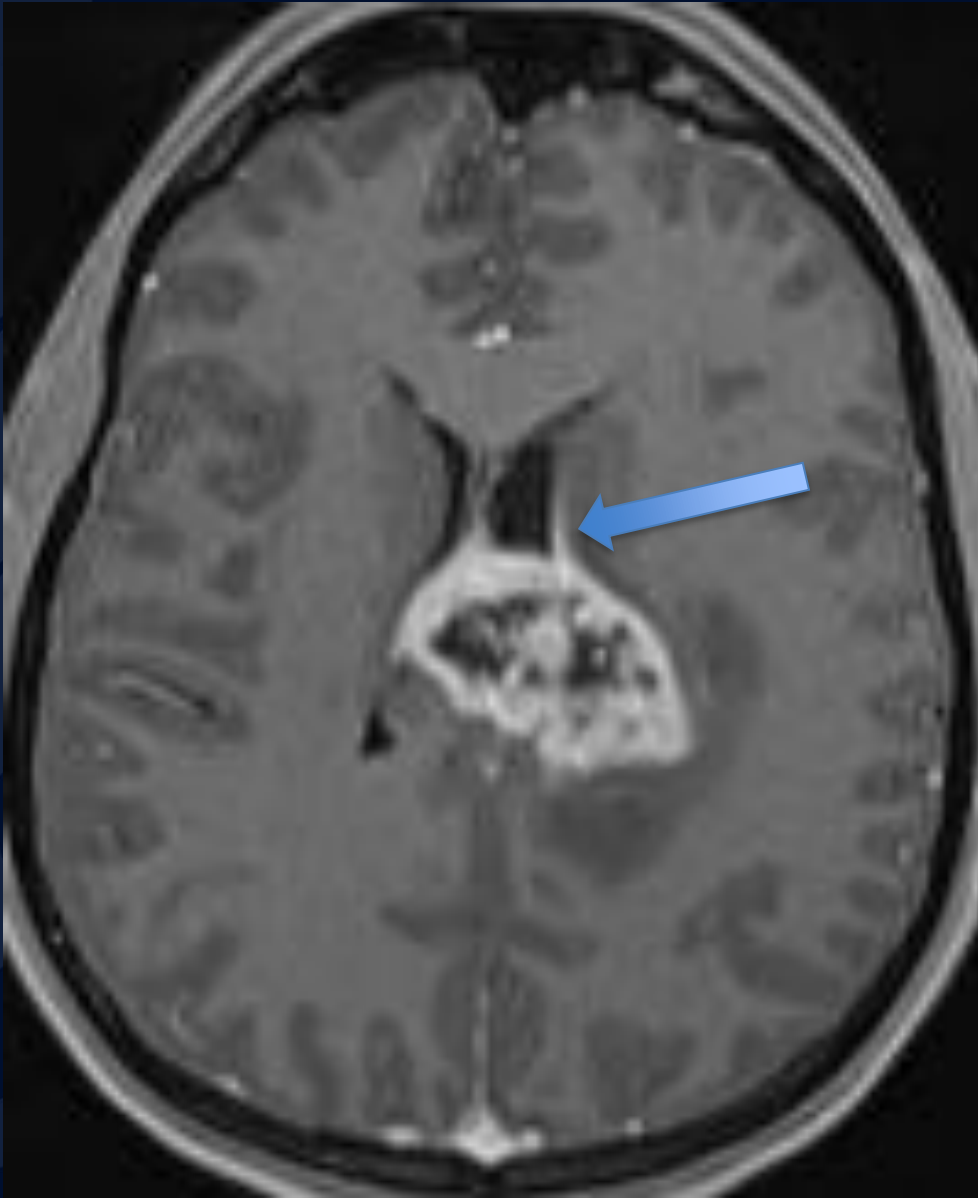


Varying thickness of enhancing rim is characteristic of GBM (arrows)

Coronal view is excellent for demonstrating corpus callosum involvement.



Central areas
of diminished
enhancement
are
frequently
necrotic, a
hallmark of
Grade IV
Glioma



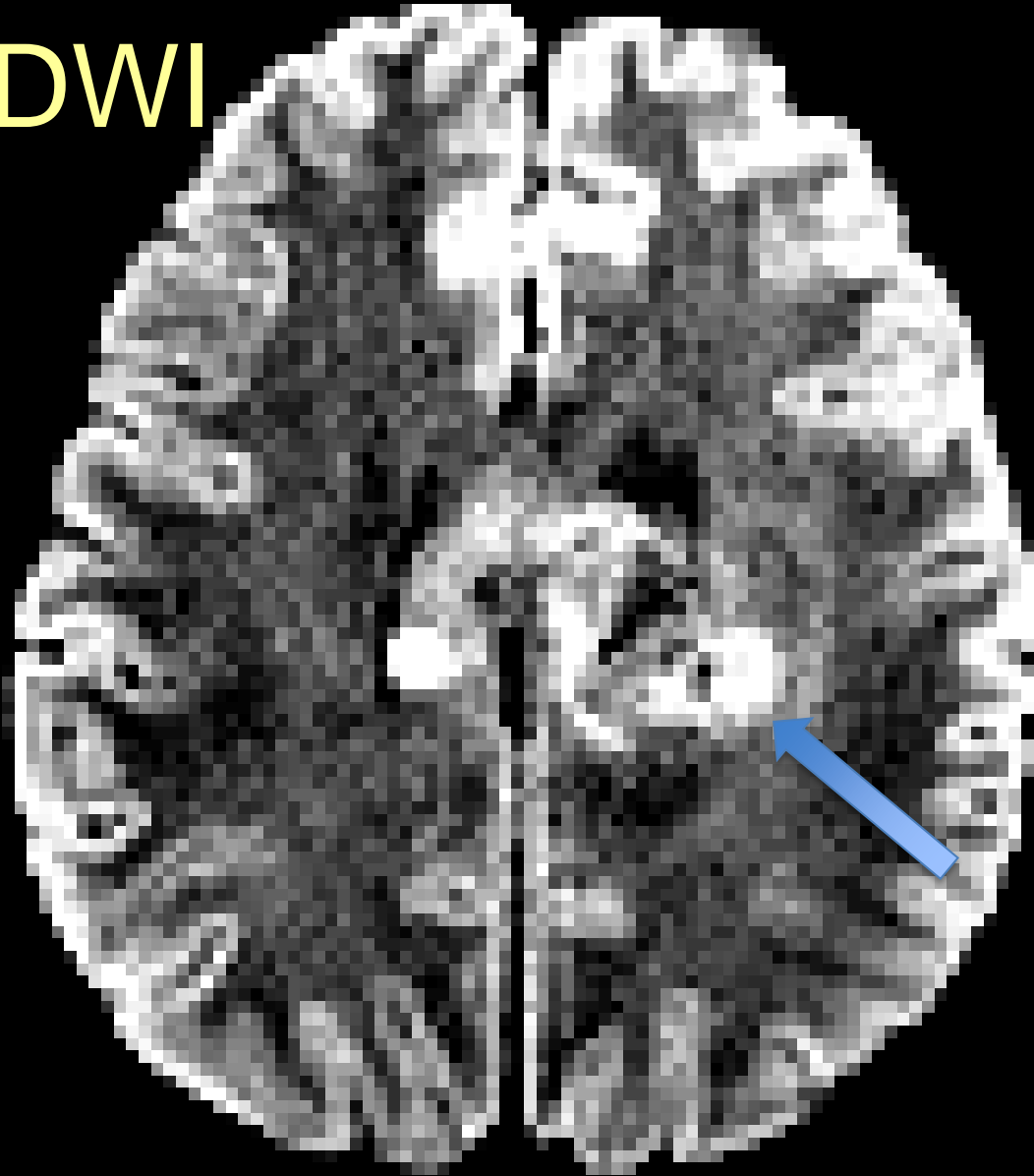
Enhancement
extending
along
ependyma

SWI



Presence of magnetically susceptible blood products favors high-grade glioma (arrow).

DWI



Moderate
diffusion
restriction
(hyperintense
area indicated
by arrow) is
characteristic

Glioblastoma (multiforme) (GBM)

- WHO 2016 classification names this Glioblastoma (without the “multiforme”)
- WHO 2016 emphasizes genetic mutations IDH-wild type (poor prognosis) vs. IDH-mutant type (better prognosis).
- Glioblastomas are most commonly in supratentorial white matter.
- Infrequently seen in brainstem or cerebellum

Epidemiology

- 3-4/100,000/yr incidence
- Highest incidence of any primary brain tumor
- Peak incidence: 45-75 years

Radiologic/Pathologic Correlation

- Tumor cells are most abundant in the thick, irregularly enhancing areas (e.g. rim)
- Necrosis is most abundant in nonenhancing center
- Surrounding FLAIR hyperintense area typically has both vasogenic edema as well as tumor cells (which contribute to high frequency of recurrence).
- Can have macroscopic vessels, hemorrhage, due to neovascularity

Treatment

- Maximum safe resection
- 6 week course of 60 Gy
- Concomitant & adjuvant Temozolomide (Temodar)
- 14-1/2 month median survival
- Bevacizumab (Avastin) for recurrence
- Ongoing clinical trials

References

- Salzman K. “Glioblastoma.” statdx.com.
- Stupp R, et al. Radiotherapy plus concomitant and adjuvant temozolomide for glioblastoma. NEJM 2005 Mar 10:352(10):987-96
- Chen D, Wolansky L. Glioblastoma. Radiology Online. (2021).