Low-Grade Diffuse Astrocytoma

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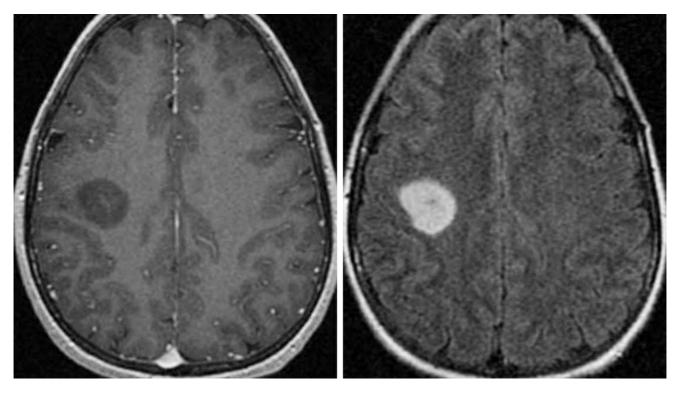


Figure 1: T1-weighted postcontrast (left) and axial FLAIR (right) images demonstrate a fairly circumscribed infiltrative lesion involving the cortex and white matter. This low-grade tumor is associated with no appreciable enhancement.

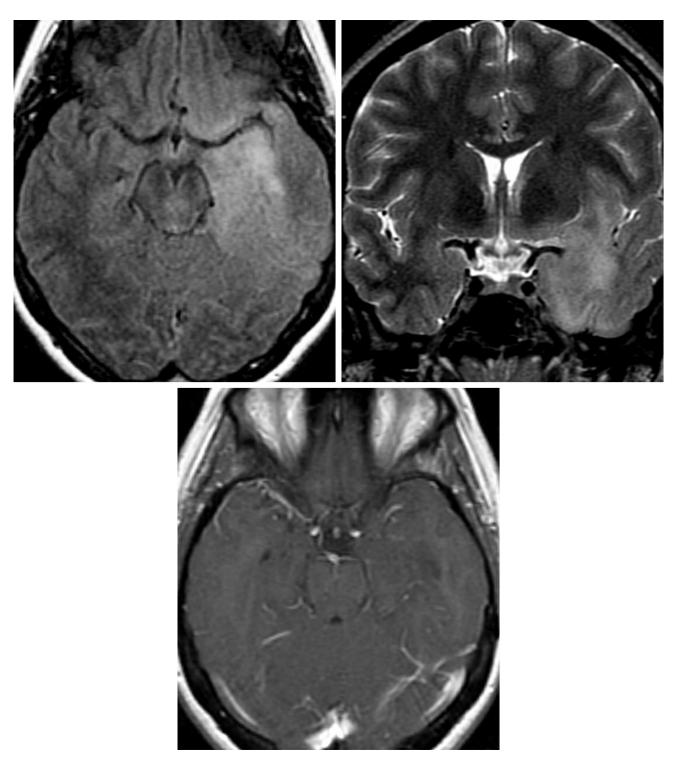


Figure 2: Axial FLAIR (top left) and coronal STIR (top right) images demonstrate a poorly defined infiltrative, hyperintense lesion involving the left temporal lobe, left insula, and inferior left frontal lobe. (Bottom) T1WI after contrast administration shows no contrast enhancement of this low-grade astrocytoma.

BASIC DESCRIPTION

Primary tumor arising from well-differentiated astrocytes

PATHOLOGY

- WHO grade II
- Well differentiated, infiltrating, slow growing
- Malignant degeneration into <u>anaplastic astrocytoma</u> is common

CLINICAL FEATURES

- Commonly presents with seizures
- Average patient age, 34 years (20–45 years)
- Median survival, 6–10 years
 - Survival greater in younger patients, gross-total resection,
 IDH1-, ARTX-, and MGMT-positive tumors
 - Pontine tumors are associated with decreased survival
- Sometimes associated with Li-Fraumeni syndrome and Ollier disease

IMAGING

- General
 - Infiltrating, focal, or diffuse white matter mass that distorts normal architecture
 - Variable size; frontal lobe masses can be large at presentation
 - Tumor commonly extends beyond region of signal abnormality
 - Expansion of involved cortex
 - Two-thirds are supratentorial; frontal lobe involvement is most common
 - One-third are infratentorial; brainstem is most common, cerebellum is uncommonly involved
 - Majority do not enhance
 - Greater degree of enhancement suggests malignant degeneration
 - ±Cysts, calcification (20%)
- CT

- Hypodense to isodense, poorly defined, homogenous mass
- ±Calcification
- Little to no enhancement on contrast-enhanced CT imaging
- MRI
 - T1WI: homogenously hypointense
 - T2WI: homogenously hyperintense
 - FLAIR: homogenously hyperintense
 - DWI: no restricted diffusion
 - T1WI+C: little to no enhancement; greater degree of enhancement suggests higher WHO grade
 - MR perfusion: low relative cerebral blood volume (rCBV)
 relative to anaplastic astrocytoma (AA) and <u>glioblastoma</u>
 <u>multiforme</u> (GBM); typically, the rCBV ratio to normal white
 matter is <1.8
 - MRS: mildly elevated choline, mildly depressed N-acetyl aspartate (NAA) peaks and usually no appreciable lactate peak

IMAGING RECOMMENDATIONS

• MRI with contrast; consider MR perfusion for equivocal cases

For more information, please see the corresponding chapter in <u>Radiopaedia</u>.

Contributor: Rachel Seltman, MD

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REFERENCES

Arevalo-Perez J, Peck KK, Young RJ. Dynamic contrast-enhanced perfusion MRI and diffusion-weighted imaging in grading of

- gliomas. *J Neuroimaging* 2015;25:792–798. doi.org/10.1111/jon.12239.
- Law M, Oh S, Babb JS, et al. Low-grade gliomas: dynamic susceptibility-weighted contrast-enhanced perfusion MR imaging—prediction of patient clinical response. *Radiology* 2006;238:658–667. doi.org/10.1148/radiol.2382042180.
- Louis DN, Ohgaki H, Wiestler OD, et al. The 2007 WHO classification of tumours of the central nervous system. *Acta Neuropathol* 2007;114:547. doi.org/10.1007/s00401-007-0243-4.
- Kleihues P, Cavenee WK. Pathology and genetics of tumours of the nervous system: diffuse astrocytoma. IARC Press, Lyon, France; 2000:22–26.
- Ogura R, Tsukamoto Y, Natsumeda M, et al. Immunohistochemical profiles of IDH1, MGMT and P53: practical significance for prognostication of patients with diffuse gliomas. *Neuropathology* 2015;35:324–335. doi.org/10.1111/neup.12196.
- Osborn AG, Salzman KL, Jhaveri MD. *Diagnostic Imaging* (3rd ed). Elsevier, Philadelphia, PA; 2016.
- Wessels PH, Weber WE, Raven G, et al. Supratentorial grade II astrocytoma: biological features and clinical course. *Lancet Neurol* 2003;2:395–403. doi.org/10.1016/s1474-4422(03)00434-4.