



IMAGES IN PAEDIATRICS

Key differential diagnosis in midline injuries

Diagnóstico diferencial clave en lesiones de línea media

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We present the case of a female adolescent aged 12 years admitted to hospital with fever and frontal headache of 5 days' duration associated with somnolence and vomiting. The salient findings of the physical examination were ptosis and drooping of the left corner of the mouth.

Papilledema was ruled out and the evaluation completed with head CT and MRI scans (Fig. 1). Based on the imaging findings, the principal diagnostic impression was diffuse midline glioma, although an inflammatory process could not be ruled out. This prompted extension of the evaluation with magnetic resonance spectroscopy (MRS), the findings of which were inconclusive for diagnosis of a tumor (Fig. 2). The surgical biopsy was deferred to perform a ¹¹C-methionine PET scan, the findings of which were not suggestive of high grade glioma. Suspicion of autoimmune encephalitis prompted testing for anti-MOG/AQP4 antibodies (which turned out negative) and initiation of intravenous steroid therapy.¹

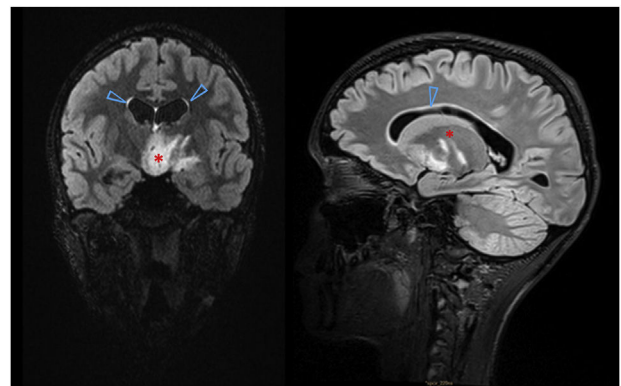


Figure 1 T2-weighted FLAIR images (coronal and sagittal planes, respectively) showing a poorly defined hypothalamic lesion with obstruction of the third ventricle (*) and moderate prominence of lateral ventricles with mild transependymal edema (Δ).

DOI of original article:

<https://doi.org/10.1016/j.anpedi.2024.503703>

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At 2 months of treatment, the neurologic symptoms had resolved and there was significant radiological improvement, with full resolution of the lesion at 5 months (Fig. 3), which confirmed the inflammatory etiology.

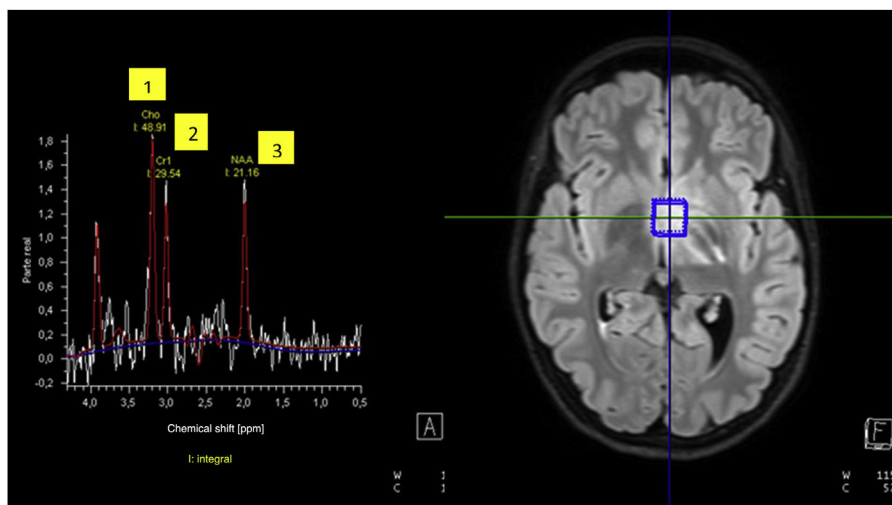


Figure 2 MR spectroscopy of the lesion showing a spectrum incompatible with a brain tumor: elevation of choline (1), creatine (2) and N-acetylaspartate (NAA) (3).

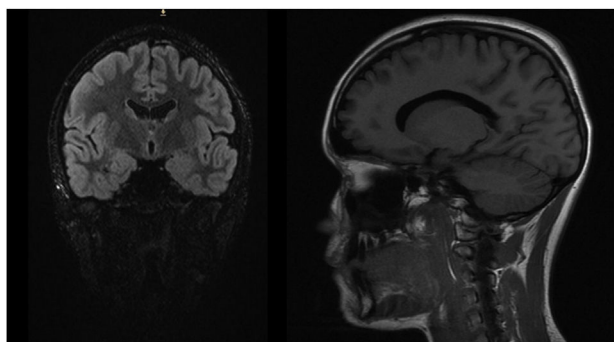


Figure 3 T2-weighted FLAIR images (coronal and sagittal planes, respectively) showing complete radiological resolution after steroid therapy.

Thus, we can conclude that not every intracranial mid-line lesion is a tumor. Tools such as MRS are useful for the differential diagnosis of these lesions.^{2,3}

References

1. Cellucci T, Van Mater H, Graus F, Muscal E, Gallentine W, Klein-Gitelman MS, et al. Clinical approach to the diagnosis of autoimmune encephalitis in the pediatric patient. *Neurol Neuroimmunol Neuroinflamm.* 2020;7:e663.
2. Tran D, Nguyen DH, Nguyen HK, Nguyen-Thanh VA, Dong-Van H, Nguyen MD. Diagnostic performance of MRI perfusion and spectroscopy for brainstem glioma grading. *Eur Rev Med Pharmacol Sci.* 2022;26:7938–48, <http://dx.doi.org/10.26355/eurrev.202211.30145>. PMID: 36394742.
3. Majós C, Pons-Escoda A, Naval P, Güell A, Lucas A, Vidal N, et al. Proton MR spectroscopy shows improved performance to segregate high-grade astrocytoma subgroups when defined with the new 2021 World Health Organization classification of central nervous system tumors. *Eur Radiol.* 2024;34:2174–82, <http://dx.doi.org/10.1007/s00330-023-10138-9>. Epub 2023 Sep 23. PMID: 37740778.