MR Spectroscopy of a Bright Cerebellar Lesion in a Patient with Langerhans Cell Histiocytosis

A. Munoz, R. Simon and G. Carmen

AJNR Am J Neuroradiol 2008, 29 (4) e14
doi: https://doi.org/10.3174/ajnr.A0911
http://www.ajnr.org/content/29/4/e14
MR Spectroscopy of a Bright Cerebellar Lesion in a Patient with Langerhans Cell Histiocytosis

We have read with interest the paper by Prosch et al.1 published in the American Journal of Neuroradiology concerning the long-term MR imaging course of 9 patients diagnosed with Langerhans cell histiocytosis (LCH). We were particularly interested in the evolution of MR abnormalities of cerebellum, pons, and basal ganglia structures. The authors observed no correlation between the progression of such abnormalities and clinical neurologic deterioration, and stated that the signal intensity persistence and progression were compatible with a neurodegenerative process, as reported by previous histopathologic studies. Furthermore, symmetric hyperintensities on T1-weighted images observed in these patients were first reported in other diseases, such as chronic liver failure. Such abnormalities could be found in up to 10% of patients with LCH. We have had the opportunity to study a patient similar to those presented in their report, a 3-year-old boy who was diagnosed with LCH because of diabetes insipidus and was treated only with antidiuretic hormone replacement and followed, both clinically and radiologically, over a course of 7 years. In the first MR imaging study, abnormalities encountered in our patient were those typically found in the pituitary-hypothalamic area, in patients with LCH and positive granuloma involvement, consisting of a lack of a hyperintense neurohypophyseal spot on noncontrast T1-weighted images and thickening and enhancement of the pituitary stalk and adjacent hypothalamic area. During this period of time, the MR enhancing pituitary lesions resolved with almost complete resolution. However, new progressive different lesions appeared, mainly in the basal ganglia and cerebellum, similar to those described in the report by Prosch et al.1 In the last study, we also performed monovoxel (TE, 35 ms) and multivoxel (TE, 144 ms) 1H-MR spectroscopy on the cerebellar lesions (point-resolved spectroscopy sequence acquisition; Fig 1). The spectrum showed a marked decrease of the N-acetylaspartate peak and an increase of the choline, as the most important findings. These findings would correlate with the histopathologic results found in other patients referred with similar lesions2 and would fit with the histopathologic findings of loss of neurons, demyelination, gliosis, and inflammation. Such a histopathologic-spectroscopic correlation has not been reported previously.

References


A. Munoz
Pediatric Neuroradiology
R. Simon
Pediatric Neurology
G. Carmen
Pediatric Endocrinology
Hospital Universitario “12 Doce de Octubre”
Universidad Complutense
Madrid, Spain

DOI 10.3174/ajnr.A0911

Fig 1. A, Axial noncontrast T1-weighted (500/12, TR/TE) MR image through the cerebellum shows hyperintense abnormal signal intensity lesions on both dentate nuclei of cerebellum. B–D, Monovoxel (B) 1H-MRS spectrum (TE, 35 ms) and multivoxel (C; TE, 144 ms) taken from the left dentate nucleus (D) shows a prominent Cho peak and decrease of the NAA peak.