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Neuroepithelial (Colloid) Cyst of the Cerebellar Vermis Containing a Xanthogranuloma

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Summary: A case of a neuroepithelial cyst of the cerebellar vermis is presented. The cyst contained a viscous fluid and a xanthogranulomatous nodule. The fluid showed high signal on T1- and low signal on T2-weighted MR images. The xanthogranuloma showed mixed intensities with partial contrast enhancement. The correlation of the cyst contents and MR signals is discussed.

Index terms: Cerebellum, vermis; Brain, cysts; Cerebellum, magnetic resonance

Neuroepithelial (colloid) cysts represent 0.25% to 1% of intracranial tumors. Although most frequent in the third ventricle (1–9), they are found rarely in other parts throughout the neuraxis, including ventricular system, subarachnoid space, and brain parenchyma. Neuroradiologic findings vary depending on the location and cyst contents. We describe a case of a neuroepithelial cyst located in the cerebellar vermis, whose contents gave neuroradiologic findings simulating a cerebellar hemangioblastoma.

Case Report

The patient was a 44-year-old woman who presented with a dull pain in the occipital region, nausea, and unsteady gait for a month. Physical and neurologic examination revealed a port wine nevus in the right side of the upper...

Fig. 1. A, Sagittal T1-weighted MR image, 600/15 (TR/TE), shows hyperintense cyst in the cerebellar vermis. B, Axial T2-weighted image (3000/90) shows hypointense cyst content and nodule with mixed intensity. C, Axial T1-weighted image with Gd-DTPA enhancement shows a partial enhancement in the nodule in the cyst.
Cysts lined with a single layer of epithelial cells have been given a variety of names depending on their location and microscopic features, including "neuroepithelial cyst," "colloid cyst," "ependymal cyst," "paraphysial cyst," "gliopen-dymal cyst," "choroid plexus cyst," and merely "epithelial cyst." The nature and origin of these lesions are still debatable. Neuroepithelial cysts are found throughout the neuraxis, but most commonly appear adjacent to the ventricular system or in the subarachnoid space above the tentorium. Rarely, they are located in the posterior fossa in the quadrigeminal, preopticine, cerebellomedullary cistern (1-4), the pons (1, 5), the cerebellar hemisphere (6), and the fourth ventricle (7). To our knowledge, there have been only two reported cases of this lesion occurring in the cerebellar vermis (8, 9).

Formation of xanthogranuloma is rarely encountered in the "colloid" cyst of the third ventricle or lateral ventricle (10-15), although some cases have been reported to occur de novo from the choroid plexus in the absence of neuroepithelial cyst (16). Localization in the infratentorial fossa seems to be exceptional (17). As speculated by Shuangshoti et al, bleeding in cyst is probably responsible for the development of a xanthogranuloma containing cholesterol crystals, although some authors have suggested that such a reaction may result from desquamation of the cells lining the cyst or from the irritative action of the colloid material (10). The color of the cyst at the operation suggested old hemorrhagic episodes. The high density of the cyst content on CT scan is attributable to increased electron density material probably caused by high protein or metallic content (18), possibly derived from previous bleeding or desquamation of cells. The high signal on T1- and low signal on T2-weighted MR images may be caused by high cholesterol content, metallic (Mg, Ca, Cu, Fe) content (18, 19) or by the presence of mucinous materials (20). The signal
on MR image of the xanthogranuloma (ie, low- to iso-signal in T1- and mixed signal in T2-weighted images) contrasts to the signal of cyst fluid. This difference in relaxation times between the cholesterol-rich cyst fluid and the xanthogranuloma is attributed to states of cholesterol (ie, liquid versus solid) (21).

Although the present case showed a striking resemblance in radiologic images to a hemorrhagic cystic hemangioblastoma with a mural nodule (22), there were some features that were not typical of a hemangioblastoma. Our case lacked the intense contrast enhancement of a "mural nodule," a vascular stain on angiography, the presence of a nodule along a pial surface, or flow void on MR, which are common to a hemangioblastoma. Therefore, it might be possible to differentiate the rare lesion of this kind from that more common cystic lesion of the infratentorial fossa.

References