Unusual CT and MR Appearance of an Epidermoid Tumor of the Cerebellopontine Angle

Makoto Ochi, Kuniaki Hayashi, Tomayoshi Hayashi, Minoru Morikawa, Ayumi Ogino, Rashid Hashmi, Mitsuto Iwanaga, Akio Yasunaga, and Shobu Shibata

Summary: We present a case of an unusual epidermoid tumor of the cerebellopontine angle that appeared hyperdense on CT scans and hyperintense on T1- and T2-weighted MR images. We believe that these imaging characteristics were caused by a high protein concentration within the contents of the cyst.

Intracranial epidermoid tumors generally appear as well-defined lobulated hypodense masses on CT scans (1, 2). On MR studies, they typically display hypointensity on T1-weighted images and hyperintensity on T2-weighted images (2). Occasionally, they may appear hyperdense on noncontrast CT scans (3–14). Although many cases of hyperdense epidermoid tumors have been reported, few that appeared hyperintense on T1-weighted MR images have been documented (13, 14). We report a case of a dense epidermoid that displayed hyperintensity on T1-weighted MR images.

Case Report

A 49-year-old man was admitted to the hospital because of a left-sided facial paresthesia. Neurologic examination showed only hypesthesia on the left side of the face. A noncontrast CT scan disclosed a hyperdense mass of the left cerebellopontine angle (Fig 1A). Attenuation values of the lesion were 48 to 50 HU. On spin-echo MR images, obtained on a 1.5-T unit, the lesion was hyperintense on T1- and T2-weighted sequences (Fig 1B and C). A frequency-selective fat-suppression technique did not suppress the signal of the lesion (Fig 1D). Contrast-enhanced T1-weighted images showed no appreciable enhancement (Fig 1E).

At surgery, a shiny multilobulated cystic mass was seen filling the left cerebellopontine angle. The trigeminal nerve was surrounded by the tumor. The cyst comprised a greenish semiliquid material containing debris. At biochemical analysis, the total protein concentration was 29.1 g per 100 mL, and the triglyceride level was 32 mg per 100 mL. Further chemical analysis was not available.

Microscopically, the wall of the tumor was composed of a keratinizing squamous epithelium and a thin layer of fibrous connective tissue. The cyst was filled with keratinous debris arranged in laminated layers. There was no evidence of prior hemorrhage. Focal rupture of the cystic wall with foreign body reaction and fibrosis was seen. The tumor was diagnosed as an epidermoid cyst.

Discussion

Intracranial epidermoid tumors are congenital cystic lesions that arise from epithelial inclusions at the time of neural tube closure or during formation of the secondary cerebral vesicles (4, 15). They are most common in the cerebellopontine angle and suprasellar and parasellar regions (15). The cysts are consistently circumscribed, with a smooth or irregularly nodular capsular surface, and are usually filled with soft, white material, which is the result of progressive desquamation and breakdown of keratin from the epithelial lining (16). At CT, therefore, they generally appear as well-defined lobulated hypodense masses (1). On the other hand, the contents are sometimes described as thick, viscid, and dark brown or gray (16).

Many cases of intracranial dense epidermoid tumors have been reported in the English-language literature (3–14). According to one report (8), the hypodense epidermoids have a pearly appearance with white, waxy material rich in cholesterol crystals, while the hyperdense epidermoids are grossly cystic and contain fluid of various colors. The physiochemical basis of hyperdensity of the tumor is not clear; proposed causes include proteinaceous contents of the cyst (6, 8), prior bleeding into the cyst (5), abundance of polymorphonuclear leukocytes (12), saponification of the debris to calcium soaps (4), and deposition of ferrocalcium complex or iron-containing pigment (4).

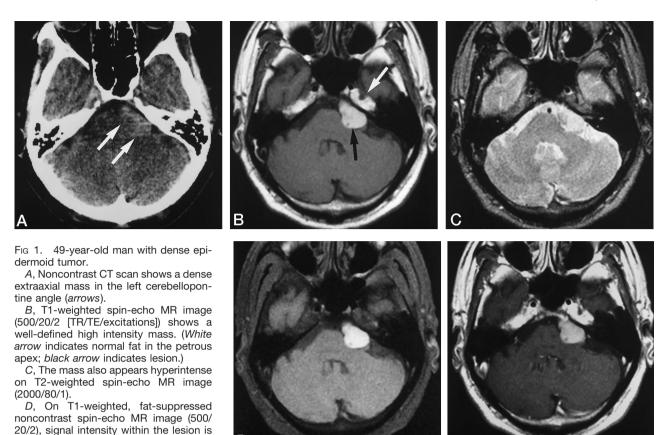
The linear relationship between CT density and protein concentration of fluid has been documented (17). Based on the results of biochemical analysis of the cystic fluid in our case, we believe that the high protein level of the cyst's contents was the primary cause of the hyperdensity seen on CT scans. Focal rupture of the cystic wall associated with foreign body reaction was seen histologically. Nagashima et al (8) suggested that high protein concentration might be

Received March 24, 1997; accepted after revision August 20.

From the Departments of Radiology (M.O., K.H., M.M., A.O., R.H.), Pathology (T.H.), and Neurosurgery (M.I., A.Y., S.S.), Nagasaki University School of Medicine, Nagasaki, Japan.

Address reprint requests to Makoto Ochi, MD, Department of Radiology, Nagasaki University School of Medicine, 1–7–1 Sakamoto, Nagasaki 852-8501, Japan.

1114 OCHI AJNR: 19, June 1998



due to a proliferative and exudative defense reaction to recurrent minor leaks of the lipidic material through the capsule. A similar explanation may be applicable in the present case.

not suppressed.

tration of contrast material.

E, Contrast-enhanced T1-weighted spin-echo MR image (500/20/2) shows no appreciable enhancement after adminis-

At MR imaging, epidermoid tumors are generally hypointense on T1-weighted sequences and hyperintense on T2-weighted sequences (2). MR signals depend on the relative composition of cholesterol and keratin of the cystic contents: generally, the cholesterol in an epidermoid is in a solid state and appears hypointense on T1-weighted images (1). Horowitz et al (18) found that epidermoid tumors with short T1 values had a high lipid content and were nonenhancing masses with negative Hounsfield numbers on CT scans whereas those with long T1 values had a much lower lipid content and appeared as a water-density mass on CT scans. To our knowledge, the MR imaging appearance of dense epidermoids has been infrequently reported. Gualdi et al (13) described a parasellar dense epidermoid that was inhomogeneously hypointense on T1-weighted MR images and had no signal on T2-weighted images. Another epidermoid of the cerebellopontine angle was reported to be hyperdense on CT scans, hyperintense on T1weighted MR images, and hypointense on T2weighted images (14). In that case, signal intensity on the T2-weighted images was different from the findings in our case. Suggested causes of the hyperintensity on T1-weighted images include high protein concentration, mild calcification, and paramagnetic effects (19-22). According to Ahmadi et al (19), a protein level of 9.0 g or greater per 100 mL can increase the signal intensity of the cystic fluid on T1-weighted MR images. As the protein concentration of lysozyme solution increases, there is a gradual increase and then decrease in signal intensity on T1weighted images. On T2-weighted images, the signal intensity decreases with increasing protein concentration (23). A relatively high protein concentration (20 to 25 g per 100 mL) of sinonasal secretions has high signal intensity on both T1- and T2-weighted images (23). The primary causes of the variable signal intensity of sinonasal secretions on T2-weighted images are the protein contents, the amount of free water, and the viscosity (23). Depending primarily on the percentage of protein and free water it contains, the proteinaceous fluid of epidermoids can have variable signal intensity on T2-weighted images. In this case, MR studies showed a hyperintense mass on T1- and T2-weighted images, and protein concentration of the cystic contents was 29.1 g per 100 mL. Therefore, we assumed that a relatively high protein concentration was the major factor for the hyperintensity on T1- and T2-weighted sequences.

AJNR: 19, June 1998 **EPIDERMOID** 1115

Conclusion

In our case of an epidermoid tumor that was hyperdense on CT scans and hyperintense on both T1and T2-weighted MR images, a relatively high protein concentration of the cystic contents probably was responsible for the unusual MR appearance.

References

- 1. Gao PY, Osborn AG, Smirniotopoulos JG, Harris CP. Epidermoid tumor of the cerebellopontine angle. AJNR Am J Neuroradiol 1992; 13:863-872
- Tampieri D, Melanson D, Ethier R. MR imaging of epidermoid cysts. AJNR Am J Neuroradiol 1989;10:351-356
- 3. Naidich TP, Lin JP, Leeds NE et al. Computed tomography in the diagnosis of extra-axial posterior fossa masses. Radiology 1976;120:
- 4. Braun IF, Naidich TP, Leeds NE, et al. Dense intracranial epidermoid tumors. Radiology 1977;122:717-719
- 5. Dunn RC, Archer CA, Rapport RL II, et al. Unusual CT-dense posterior fossa epidermoid cyst. J Neurosurg 1981;55:654-656
- 6. Hasegawa H, Bitoh S, Nakata M, et al. Intracranial epidermoid mimicking meningioma. Surg Neurol 1981;15:372-374
- 7. Handa J, Okamoto K, Nakasu Y, Nakasu S, Nakano Y. Computed tomography of intracranial epidermoid tumours with special reference to atypical features. Acta Neurochir (Wien) 1981;58:221–228
- 8. Nagashima C, Takahama M, Sakaguchi A. Dense cerebellopontine epidermoid cyst. Surg Neurol 1982;17:172-177
- Rahman N. Atypical epidermoid cyst of the brain. Neurosurgery 1988;22:122-124

- 10. Lunardi P, Rizzo AA, Guidetti G. Unusual CT-dense transtentorial epidermoid cyst. Neurochirurgia 1988;31:219-221
- Ozgen T, Oge K, Erbengi A, Bertan V. Cranial dermoid and epidermoid cysts. Neurochirurgia 1990;33:16-19
- 12. Tekkök IH, Cataltepe O, Sağlam S. Dense epidermoid cyst of the cerebellopontine angle. Neuroradiology 1991;33:255-257
- Gualdi GF, Di Biasi C, Pingi A, Vignati A, Maira G. Unusual MR and CT appearance of an epidermoid tumor. AJNR Am J Neuroradiol 1991;12:771-772
- 14. Lo WWM, Solti-Bohman LG. Tumors of the temporal bone and the cerebellopontine angle. In: Som PM, Curtin HD, eds. Head and Neck Imaging. St Louis: Mosby-Year Book; 1996:1449–1534 Vion-Dury J, Vincentelli F, Jiddane M, et al. MR imaging of
- epidermoid cysts. Neuroradiology 1987;29:333-338
- Russell DS, Rubinstein LT. Pathology of the Nervous System. 5th ed. London: Edward Arnold: 1989:690-710
- 17. New PFJ, Aronow S. Attenuation measurements of whole blood and blood fractions in computed tomography. Radiology 1976;121:
- 18. Horowitz BL, Chaari MV, James R, Bryan RN. MR of intracranial epidermoid tumors: correlation of in vivo imaging with in vitro 13C spectroscopy. AJNR Am J Neuroradiol 1990;11:299-302
- Ahmadi J, Destian S, Apuzzo ALJ, Segall HD, Zee CS. Cystic fluid in craniopharyngiomas: MR imaging and quantitative analysis. Radiology 1992;182:783-785
- 20. Henkelman RM, Watts JF, Kurcharczyk W. High signal intensity in MR images of calcified brain tissue. Radiology 1991;179:199-206
- 21. Bradley WG. MR appearance of hemorrhage in the brain. Radiology 1993;189:15-26
- 22. Woodruff WW, Djang WT, McLendon RE, Heinz ER, Voorhees DR. Intracerebral malignant melanoma: high-field-strength MR imaging. Radiology 1987;165:209-213
- Som PM, Dillon WP, Fullerton GD, Zimmerman RA, Rajagopalan B, Marom Z. Chronically obstructed sinonasal secretions: observations on T1 and T2 shortening. Radiology 1989;172:515-520