Are your MRI contrast agents cost-effective? Learn more about generic Gadolinium-Based Contrast Agents.





Intracranial Osteochondroma: MR and CT Appearance

David W. Beck and Gregg N. Dyste

AJNR Am J Neuroradiol 1989, 10 (5 suppl) S7-S8 http://www.ajnr.org/content/10/5_suppl/S7.citation

This information is current as of May 2, 2024.

Intracranial Osteochondroma: MR and CT Appearance

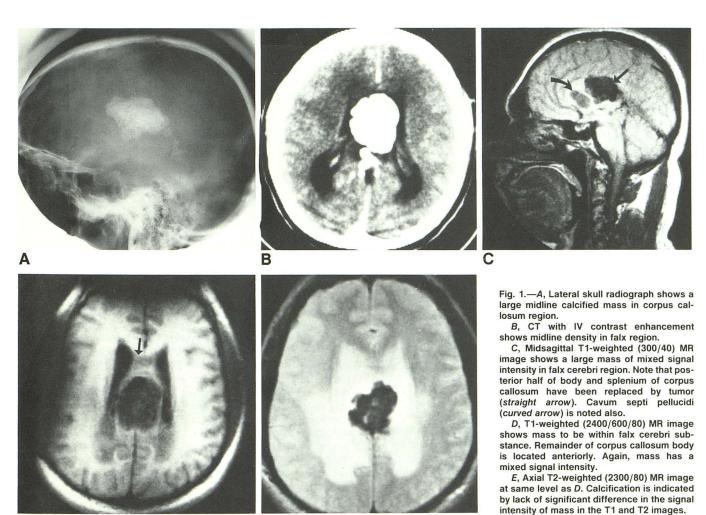
David W. Beck1 and Gregg N. Dyste

D

Osteochondromas are rare intracranial lesions that have been reported to arise from the skull base and dura. We report a case of a midline osteochondroma arising from the inferior portion of the falx and replacing the midportion of the corpus callosum, and we describe the MR and CT appearances.

Case Report

A 48-year-old right-handed man presented to the University of lowa for evaluation of unsteadiness of gait over the previous 6 months. He also reported a 1-year history of bifrontal headaches that were worse in the morning and improved during the day. He had experienced good health all his life, and his past history was unre-



E

markable. His general physical examination was normal. His neurologic examination revealed a wide-based, unsteady gait. He had no other neurologic findings. Neuropsychological testing revealed left-hand tactile anomia and a mild alexia and agraphia. In addition, he had a mild left ear extinction on dichotic listening.

Skull radiographs revealed a large calcified mass in the region of the corpus callosum (Fig. 1A). On CT, the mass did not enhance with contrast infusion. The mass was somewhat asymmetric with a larger portion of the tumor on the left side (Fig. 1B). MR imaging localized the mass to the region of the distal body and splenium of the corpus callosum (Fig. 1C). The MR appearance did not change between T1 and T2 images, consistent with the calcification within the tumor (Figs. 1D and 1E).

The patient underwent a left frontoparietal craniotomy with total excision of the tumor via an interhemispheric approach. At surgery, the mass was very hard and required a CO_2 laser for excision. The tumor appeared to rise from the inferior portion of the falx. No corpus callosum was seen in the area of tumor, and it appeared that the mass had completely replaced the posterior body of the corpus callosum. After surgery, the patient had a mild weakness in his right foot, which improved to normal over the next 6 weeks. His neuropsychologic testing 3 months after surgery revealed improvement of his left hand anomia. He had a mildly impaired left-handed agraphia, but this also was improved from preoperative testing. His gait at 3 months was normal.

Discussion

Osteochondroma is the most common benign bone tumor, accounting for 40% of the benign lesions in the Mayo Clinic series [1]. Intracranial osteochondromas are rare. In Cushing's series of 2023 intracranial tumors, only three were osteochondromas [2]. Most of these tumors arise from bones that are embryologically derived from cartilage [3, 4], which accounts for their predilection for the base of the skull [5, 6]. Only some 15% of intracranial cartilaginous tumors have a parafalcial dural attachment as seen in this case.

The radiologic appearance of this tumor is fairly typical but not diagnostic. Skull radiographs show evidence of bony changes caused by local tumor growth, increased intracranial pressure, and areas of tumor calcification [5, 7]. The skull radiographs in this case showed the midline location of the tumor well. The CT appearance of these tumors is variable. Matz et al. [8] reported a large frontoparietal osteochondroma demonstrating a high-density mass with foci of lower densities, producing a honeycomb appearance. The lesion showed

a modest degree of enhancement after contrast infusion. Ikeda [9] reported a middle fossa osteochondroma that demonstrated a high-density, irregularly shaped mass on CT. These findings are similar to those of the present case, in which the mass did not enhance after contrast infusion. The MR appearance of an intracranial osteochondroma has not been reported previously. The appearance of extracranial osteochondromas consists of a mixed signal intensity on both T1- and T2-weighted images [10], which is similar to the MR appearance in this case. The honeycomb appearance on MR conformed well to the findings at surgery of a mixture of soft tumor alternating with prominent areas of calcification. MR was clearly superior to CT in this case in respect to delineation of the local anatomy; this shows the importance of MR in preoperative surgical planning.

Surgical resection is the treatment of choice for cartilaginous falcial tumors [11, 12], and long-term survival is expected [11, 13].

REFERENCES

- Dahlin DC. Bone tumors. General aspects and data on 6,221 cases, 3rd ed. Springfield: Charles C Thomas, 1978:18
- Cushing H. Intracranial tumours. Notes upon a series of two thousand verified cases with surgical-mortality percentages pertaining thereto. Springfield: Charles C Thomas, 1932:8, 133
- Ettore G, Benedittis G, Bernasconi V. Tumors of the fifth cranial nerve. Acta Neurochir 1977;38:37–64
- Sarwar M, Swischuk LE, Schechter MM. Intracranial chondromas. AJR 1976;127:973–977
- Berkmen YM, Blatt ES. Cranial and intracranial cartilaginous tumours. Clin Radiol 1968:19:327–333
- Gabrielsen TO, Kingman AF Jr. Osteocartilaginous tumors of the base of the skull. Report of a unique case and review of the literature. AJR 1964:91:1016–1023
- Minagi H, Newton TH. Cartilaginous tumors of the base of the skull. AJR 1969:105:308–313
- Matz S, Isreali Y, Shalit MN. Computed tomography in intracranial supratentorial osteochondroma. J Comput Assist Tomogr 1981;1:109–115
- 9. Ikeda Y. Intracranial osteochondroma. No Shinkei Geka 1980;3:263-269
- Hudson TM, Hamlin DJ, Enneking WF, Pettersson H. Magnetic resonance imaging of bone and soft tissue tumors. Skeletal Radiol 1984;13:134–136
- Ozgen T, Necmettin P, Akalan N, Bertan V, Onol B. Intracranial solitary chondroma. Case report. J Neurosurg 1984;61:399–401
- Yang PJ, Seeger JR, Carmody RF, Fleischer AS. Chondroma of falx: CT findings. J Comput Assist Tomogr 1986;10:1075–1076
- Hardy RW Jr, Benjamin SP, Gardner WJ. Prolonged survival following excision of dural chondroma. Case report. J Neurosurg 1978;48:125–127