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Summary: Aneurysmal bone cysts of the skull are rare, and orbital involvement of these cysts is even less frequent. We present CT, MR imaging, and histopathologic findings of an aneurysmal bone cyst of the orbit in a 13-year-old female adolescent. The tumor mainly involved the frontal bone. MR imaging findings of the aneurysmal bone cyst of the skull were highly suggestive of the diagnosis.

Aneurysmal bone cysts are uncommon benign, expansile, and lytic lesions that develop in childhood or early adulthood. An aneurysmal bone cyst usually occurs as a primary lesion. In one third of the cases reported in the literature, a preexisting condition or history of trauma is present. Pain and swelling are the main symptoms and usually last for as long as 6 months. Long bones are involved in more than half of the cases. Involvement of the spine and flat bones also has been reported (1, 2). Skull involvement is rare, occurring in less than 1% of all aneurysmal bone cysts (3). The orbit is much less frequently involved; a review of the literature revealed only 20 cases (3–7). Among them, we could find only two presentations with MR imaging findings, and these were not published in the radiologic literature (3, 4). We present CT and MR imaging findings in a case of an aneurysmal bone cyst of the orbit that was confirmed by histopathologic analysis.

Case Report

A 13-year-old female adolescent had rapidly growing, painful swelling in her right frontal region and diplopia over a 2-month period. She had no history of trauma or surgery. Physical examination revealed a firm, fixed mass situated just superior and posterior to the right orbital rim. Exophthalmos was present on the right. The lateral gaze of the right eye was restricted, but visual acuity was normal. On coronal and axial CT scans, a multiloculated mass with fluid-fluid levels was found. Bone destruction was present in the superior, lateral, and posterior walls of the right orbit (Fig 1). T1- and T2-weighted imaging revealed variable signal intensities of fluid-fluid levels, consistent with blood degradation products. The walls of the cystic spaces and surrounding rim showed contrast enhancement. The mass was compressing the optic nerve and ocular muscles inferomedially (Fig 2). A cystic mass, approximately 4.5 × 3 × 3 cm in size and containing brown-yellowish fluid, was surgically removed. Histopathologic analysis revealed multiple blood-filled cystic spaces. The borders of these spaces were lined by osteoclast type multinucleated giant cells, bone, and fibrous tissue. Endothelial cells were not present. These findings were consistent with those of an aneurysmal bone cyst (Fig 3).

Discussion

Aneurysmal bone cysts were first described in 1942 as “peculiar blood-containing cysts of large size” (8). They are composed of blood-filled, anastomosing cavernous spaces, separated by cystlike walls. The precise nature and histogenesis of the aneurysmal bone cyst remain unclear (2). However, the aneurysmal bone cyst is considered to be the result of a specific pathophysiologic change, which is probably caused by trauma or an anomalous vascular process. In one third of the cases, a preexisting lesion is present. Giant cell tumor, chondroblastoma, chondromyxoid fibroma, nonossifying fibroma, osteoblastoma, fibrosarcoma, fibrous histiocytoma, osteosarcoma, and fibrous dysplasia have been reported as antecedent lesions (1, 2).

Radiologic findings reflect the pathophysiologic properties of an aneurysmal bone cyst. Initially, well-defined osteolysis and periosteal elevation are present. The lesion grows rapidly and causes progressive destruction. Radiographs typically show an eccentric lesion with an expanded, remodeled bony contour or ballooned bony contour of the bone. The word aneurysmal refers to this appearance. The expanded contour is the result of bone production by the periosteum. Lesions frequently have a delicate trabeculated appearance (1). At histologic examination, these lesions are large, septated sinusoids filled with blood and lined by endothelium and multinucleated giant cells. Small cysts called diverticula project from larger cysts (9, 10). Hemorrhage of variable age within the cysts are present, and the degradation of blood products causes fluid levels. On MR images, all these structures are surrounded by a well-defined low-signal-intensity rim (11). Later, a soap bubble appearance is seen as the lesion stabilizes. Finally, calcification and ossification occur, and the lesion turns into a dense mass.

CT is especially useful in the evaluation of the lesions located in the regions that cannot be adequately assessed by radiography. On CT scans, a well-defined expansive mass causing cortical interruption is seen. Fluid levels are observed in 35% of the cases, and dependent layers show increased attenuation (1).

MR imaging features of an aneurysmal bone cyst have been described in several studies (9, 10, 12, 13). The hypointense rim surrounding the lesion is an...
important finding and suggests a benign process. This rim is composed of fibrous tissue (1, 10, 13). A fluid-fluid level is another finding and is more readily seen on MR images than on CT scans (1). Cavities that contain fluid-fluid level are conspicuous on T1-weighted images. Because of the degradation of blood products, the signal intensity of these levels vary (9, 10). Fluid-fluid levels strongly suggest the presence of an aneurysmal bone cyst (9), but this finding has been reported to occur in association with various conditions and is not pathognomonic. Chondroblastoma, osteosarcoma, giant cell tumor, fibrous dysplasia, osteoblastoma, and tumoral calcinosis are examples of pathologic abnormalities with which fluid-fluid levels are found (14, 15). Although it is not specific, its presence indicates an aneurysmal bone cyst if other suggestive features exist (11). Another important MR imaging feature of an aneurysmal bone cyst is the presence of small cysts projected from larger cysts; these are called diverticula (9, 10). These
structures contribute to the typical bubbly appearance. MR imaging features of calvarial lesions are not noticeably different from those of lesions occurring elsewhere in the body (3, 11).

Age, clinical course, and imaging findings in our patient suggested the diagnosis of aneurysmal bone cyst. Diverticula were seen on MR images. Although the hypointense rim was not obvious on T1-weighted images, fibrous tissue enhancement was observed. These MR imaging findings were considered typical for an aneurysmal bone cyst.

In conclusion, aneurysmal bone cyst should be in the differential diagnosis of rapidly growing calvarial masses in young patients. Although not pathognomonic, the presence of cysts and diverticula, fibrous tissue enhancement, and fluid-fluid levels with variable signal intensity on MR images strongly suggests the presence of an aneurysmal bone cyst. The multiplanar imaging capability of MR imaging is especially important in the evaluation of skull lesions for which the neighboring structures should be evaluated before surgical intervention.

References