ABSTRACT

Osteoblastoma is a rare benign bone tumor that usually arises in the vertebral column and long bones of young adults. Craniofacial involvement is extremely rare. To date, osteoblastoma of the frontal sinuses has not been reported in the English literature. We report an osteoblastoma of both frontal sinuses in a 23-year-old male who presented with headache and blurry vision in the left eye. Computed tomography (CT) demonstrated an expansile lesion involving both frontal sinuses with sclerotic and fibrous components, eroding into the roof of the left orbit. On magnetic resonance imaging (MRI) the dense portion of the lesion showed signal void on all sequences, while the fibrous matrix was isointense to grey matter on T1-weighted and T2-weighted images and showed avid enhancement following intravenous contrast administration. Surgical resection was performed and histology was consistent with osteoblastoma.

CASE REPORT

A 23-year-old white male patient presented to the emergency department of our institution following the onset of blurred vision in the left eye, worsening over the past 48 hours. He also described bilateral frontal headache over the past month associated with frontal fullness. The patient had a personal history of hypertension requiring medical treatment. On physical examination he had swelling and tenderness over the left facial area. There was minimal exophthalmos of the left eye but the movement of the extraocular muscles was intact. Visual field examination showed field cut in left lower quadrant. The physical exam was otherwise unremarkable. Results of routine laboratory tests were within normal limits.
Osteoblastoma was recognized as a separate pathologic entity, independently by Jaffe (3) and Lichtenstein (4) in 1956. It is an uncommon primary osseous tumor involving predominantly the vertebral column and the long bones (1). Unlike fibrous dysplasias, which may have a similar radiological appearance, craniofacial osteoblastoma is extremely rare with only twenty four cases of nasal cavity and paranasal sinus osteoblastomas reported (2,5-27) (Table 1). These included eight ethmoid sinus, seven maxillary sinus, five nasal cavity, and four frontal sinus lesions; none of the frontal sinus lesions was reported in the English literature (5,6,8,26). The affected population ages ranged between 3 and 69 years with only two patients older than 30 years (age is not available from one case (5)). This is similar to the reported age distribution of osteoblastoma elsewhere in the body (28). There was no sex predilection, however, (gender is not available from two cases (5,11)), in contradistinction to the reported male predominance of two to one in the vertebra and long bones (28). The reported presenting symptoms were headache, facial swelling, as well as nasal obstruction and epistaxis when the nasal cavity was involved, and signs and symptoms related to orbital involvement such as exophthalmos, epiphoria, and orbital pain. Our patient had in addition to headache, acutely blurred vision in the left eye, a previously unreported presentation. This may be due to the mass effect on the orbital structures, although the extraocular muscle movement was reported to be within normal on physical exam.

On imaging, osteoblastomas tend to be expansile and tend to remodel adjacent bone. The reported lesions of the nasal cavity and paranasal sinuses remained fairly well circumscribed without causing significant bone destruction like elsewhere in the skeleton (27). However due to its anatomic proximity to vital structures, paranasal osteoblastomas, when large, tend to be symptomatic. Lesions tend to have mixed components with a dense sclerotic portion and a fibrous portion (27). The main differential diagnosis includes fibrous dysplasia, which, due to its relatively higher incidence, is considered first radiologically when a similar such lesion is encountered. The entities may be difficult to differentiate on CT but the mixed osseous and fibrous appearance of an osteoblastoma tends to be more nodular and coarsely organized than that of most fibrous dysplasias (29). On MR imaging the sclerotic component of the lesion will demonstrate signal void and no enhancement (2). The fibrous component, on the other hand, will demonstrate low to intermediate signal intensity on both T1W and T2W images and intense enhancement following the administration of intravenous contrast (2). The enhancement pattern also helps differentiate osteoblastoma from fibrous dysplasia which generally enhances along the margin of the lesion (29). The major role of MRI is to carefully evaluate the involvement of the adjacent vital structures mainly the intra-orbital structures as part of pre-operative assessment.

Histologically, osteoblastoma resembles an osteoid osteoma, from which it is separated purely on the basis of size. Arbitrarily, a lesion that is smaller than 1.5 cm is considered an osteoid osteoma, and a lesion that is larger than 1.5 cm is considered an osteoblastoma. Both lesions are well circumscribed and associated with sclerosis.

TEACHING POINT

Osteoblastoma is a rare benign bone tumor arising predominantly in the vertebral column or long tubular bones, but may rarely be seen in craniofacial bones. We reported the first case in the English literature, arising in both frontal sinuses in a 23-year-old male patient. This entity may have similar radiologic appearance to fibrous dysplasia and histologic appearance to osteoid osteoma and should be included in the differential of a mixed sclerotic and fibrous lesion of the paranasal sinuses.
REFERENCES


**Figure 1:** 23-year-old male patient with frontal sinus osteoblastoma who presented with headache and blurry vision in left eye. Axial non enhanced CT image through the frontal sinuses shows a relatively well defined expansile lesion in both frontal sinuses with sclerotic and fibrous components. There is scalloping of the anterior frontal sinus cortex by the sclerotic component (open arrow) and erosion of the posterior cortex by the fibrous component (arrow). The remaining part of the frontal sinus is occupied by a soft tissue density (asterisk) that was found to represent entrapped secretions during surgery (Technique: KVP = 120; mA = 37; Slice Thickness = 5.00 mm).

**Figure 2:** 23-year-old male patient with frontal sinus osteoblastoma who presented with headache and blurry vision in left eye. Axial non enhanced CT image through the frontoethmoid recess and orbital roof shows extension of the fibrous component of the lesion into the left frontoethmoid recess (asterisk) and the sclerotic lesion into the left orbital roof (arrow). (Technique: KVP = 120; mA = 37; Slice Thickness = 5.00 mm).
Figure 3: 23-year-old male patient with frontal sinus osteoblastoma who presented with headache and blurry vision in left eye. Axial MRI images through the frontal sinuses, at same level of the displayed CT image in figure 1. a: Axial noncontrast spin echo T1W image [1.5 Tesla (TR/TE = 437 msec./14 msec.) Slice Thickness = 5.00 mm]. b: Axial noncontrast fast spin echo T2W image [1.5 Tesla (TR/TE = 4000 msec./112 msec.) Slice Thickness = 5.00 mm]. c: Axial post contrast spin echo T1W image [1.5 Tesla (TR/TE = 500 msec./14 msec.) Slice Thickness = 5.00 mm, 15 cc Gadobenate Dimeglumine, Multihance]. The sclerotic component on CT scan shows low signal intensity (open arrow) on T1W and T2W images and no enhancement. The fibrous component (dashed arrow) on the other hand is of intermediate signal intensity on both unenhanced sequences and shows avid enhancement following intravenous contrast administration. The soft tissue component seen on CT (Fig.1) consistent with secretions, has intermediate T1W and bright T2W signal intensity and shows faint peripheral enhancement (asterisks).

Figure 4 (left): Coronal T1W spin echo post contrast [1.5 Tesla (TR/TE = 500 msec./14 msec.) Slice Thickness = 6.00 mm, 15 cc, Gadobenate Dimeglumine, Multihance] clearly shows the enhancing fibrous component extending into the left frontoethmoid recess (asterisk) while the sclerotic non enhancing component extends through the left orbital roof (arrow) and causes mass effect on the superior rectus muscle (arrow head).
Figure 5: 23-year-old male patient with frontal sinus osteoblastoma who presented with headache and blurry vision in left eye. a: Axial non enhanced CT images through the orbits, performed on presentation shows that the posterior wall of the left globe (arrow) is at the level of the interzygomatic line (white line), while the normal right counterpart is slightly more posterior. Findings are those of proptosis of the left globe. (Technique: KVp = 120; mA = 37; Slice Thickness = 5.00 mm). b: Axial non enhanced CT image through the orbits (same level as in "a") done 2 months post operatively shows resolution of proptosis. The posterior wall of the left globe (arrow) is now posterior to the interzygomatic line and is now symmetric to the right counterpart. (Technique: KVp = 120; mA = 350; Slice Thickness = 5.00 mm)

Figure 6: 23-year-old male patient with frontal sinus osteoblastoma who presented with headache and blurry vision in left eye. Hematoxylin and Eosin stained section of the surgically resected frontal sinus tumor showing sclerotic lamellar bony trabeculae rimmed with osteoblasts (arrows) and intertrabecular loose fibrovascular stroma. (Original magnification x 400).
Osteoblastoma of the Frontal Sinuses Presenting with Headache and Blurred Vision: Case Report and Review of the Literature

<table>
<thead>
<tr>
<th>Osteoblastoma Epicenter</th>
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Table 1: Summary of the reported cases of paranasal sinus and nasal cavity osteoblastomas
Cases listed by date of publication starting with the oldest
M: Male. F: Female. ?: Not Available

**ABBREVIATIONS**

CC: Craniocaudal
CT: Computed tomography
MRI: Magnetic resonance imaging
FLAIR: Fluid-Attenuated Inversion Recovery
Cm: Centimeters
T1W: T1-weighted
T2W: T2-weighted
TR: Repetition Time
TE: Echo time
msec.: Millisecond

**KEYWORDS**

Frontal sinus, exophthalmos, osteoblastoma

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