ABSTRACT
Background: Idiopathic bone cavities (IBCs) are osseous pseudocysts with unclear etiology. Their clinical course and response to treatment are poorly understood.

Purpose: The aim of this study is to present a case of a patient with IBC with long-term follow-up.

Patients and Methods: A retrospective case of 16-years old male patient with IBC of the jaw is implemented. Medical records of the patient were reviewed for data on presentation and imaging. The patient underwent surgical exploration to confirm the diagnosis. Follow-up radiographs were evaluated for resolution or persistence after the procedure.

Results: The lesion was asymptomatic and discovered as incidental findings. Common radiographic features included scalloping without root resorption or displacement and cortical thinning in lingual aspect expansion. Intraoperative findings showed an empty bone cavity often filled with blood-tinged serous fluid or blood. The histopathology of scrapings from the bony wall showed benign mixed fibrous tissue and no epithelial lining. Radiographic follow-up was 3 years after surgery. After exploration and curettage, the patient had a complete bone fill.

Conclusion: IBC is a rare condition, and several differential diagnoses should be considered. A single procedure to confirm the diagnosis and curettage is sufficient for management.

Keywords: bone cyst, pseudocyst, mandible,

INTRODUCTION
Traumatic bone cysts may be characterized by the presence of an asymptomatic cavity in bone with no epithelial lining. Traumatic bone cysts were described in 1929; they are commonly found in the metaphysis of long bones but are rare in the jaws. [1, 2, 3]

Traumatic bone cysts may be classified as unicameral, simple, solitary, hemorrhagic, or idiopathic.[1, 4, 5, 6, 7, 8] They are usually asymptomatic and appear on routine radiographies. Because of a lack of unique clinical and radiographic features, it is important to establish the differential diagnosis between traumatic bone cysts and other bone lesions of the jaws, especially translucent lesions. [1]
Fig. 2. CT scan of the lesion

Fig. 3. CT scan of the lesion
Fig. 4. Postoperative OPG 8 month after surgery

Fig. 5. Postoperative OPG one and a half year after surgery

DISCUSSION

Traumatic bone cysts are rare lesions of the jaws. They are classified by the World Health Organization as part of a group of bone lesions that include the ossifying fibroma, fibrous dysplasia of bone, central giant cell lesions, aneurysmatic bone cysts, and cherubism. [1] Although traumatic bone cysts were described at the beginning of the 20th century, the pathogenesis remains unclear and speculative. [9] The most accepted version at present is the traumatic-hemorrhagic theory, which suggests that lesions develop if intramedullary clots due to trauma do not undergo lysis or resolution. [10] This theory explains why traumatic bone cysts occur more often in young individuals (an age at which trauma occurs more often) and also explain the presence of blood within the cavity at the time of surgical exploration. Reports of trauma at the site of lesions and the presence of blood in the cavities, however, are uncommon, as seen in our sample. This opens the possibility that microtrauma of teeth and the alveolar ridge are involved in the pathogenesis of traumatic bone cysts [11].

According to several authors, most cases of traumatic bone cysts present in young patients, although they may be detected at any age [12, 13, 14]; our case was similar – the patient was 16 years of old.

Most cases of maxillofacial traumatic bone cysts described in the literature are asymptomatic and do not cause expansion of the cortical area – these cysts are diagnosed as accidental findings in routine radiographs as in our case. [11, 13]

Traumatic bone cysts generally show up as unilocular radiolucent areas in the posterior portion of the mandible; its margins are scalloped among dental roots [10, 11, 12] Therefore, traumatic bone cysts should be part of the differential diagnosis of maxillary radiolucent lesions – together with dentigerous cysts, keratocystic odontogenic tumours, ameloblastomas, odontogenic myxomas, aneurysmatic bone cysts, focal osteoporotic bone marrow defect, intraosseous vascular malformations, central giant cell lesions, among others. [1]

A few authors have reported the occurrence of multiple traumatic bone cysts, and their association with fibrous/bony lesions – such as the florid cemento-osseous dysplasia – especially in older patients [15, 16, 17] Providers should be prepared for long-term follow-up in patients treated for multiple IBCs because of the high probability of recurrence.

The histology of traumatic bone cysts reveals only a connective tissue membrane lining the pathologic cavity, characteristic of pseudocysts. Cholesterol crystals, hemorrhagic foci, and osteoclasts may be found [1, 11, 18]. A final diagnosis of a traumatic bone cyst is almost invariably made at the time of surgery; the material available for histology is usually sparse because of the difficulty in removing the thin connective tissue membrane. Surgeons usually encounter an empty cavity, although there may be blood, serum, or both. [1, 19]

The treatment of choice for traumatic bone cysts is surgery for curettage of the bone walls, which generally results in short-term healing [1, 11]. Recurrences are rare and usually occur within three months of surgery. Cases of multiple cysts or those associated with florid cemento-osseous dysplasia have high recurrence rates – respectively, about 71% and 75% [1, 20].

CONCLUSION

The prognosis is usually excellent, although in some cases, IBCs are difficult to treat and require annual follow-up until radiographic healing has been completely verified.
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