Case Report

Central cementifying fibroma of maxilla

Mahnaz Sheikhi¹, Farzaneh Mosavat¹, Faranak Jalalian¹, Roghayeh Rashidipoor¹
¹Torabinejad Dental Research Center and Department of Oral and Maxillofacial Radiology, School of Dentistry, Isfahan University of Medical Sciences, Isfahan, Iran

ABSTRACT

Central cementifying fibroma is a bony tumor, which is believed to be derived from the cells of the periodontal ligament. Central cemento-ossifying fibroma behaves like a benign bone neoplasm. This bone tumor consists of highly cellular, fibrous tissue that contains varying amounts of calcified tissue resembling bone, cementum, or both. Central cemento-ossifying fibromas of the mandible are common; however, they are rare in the maxilla region. This tumor is most frequent between 35 and 40 years of ages. In this report we have described a 37-year-old male with cemento-ossifying fibroma of the maxilla region with the mass that had been appeared 2-3 months prior to his first referral. Radiologic imaging such as intra-oral, panoramic, and Cone Beam CT had been performed. Histological analysis was done and finally diagnosis of central cementifying fibroma was made. The postoperative follow up at 12 months revealed no recurrence.

Key Words: Central cementifying fibroma, fibro-osseous lesions, ossifying fibroma, radiology

INTRODUCTION

In 1971, World Health Organization (WHO) categorized four types of cementum-containing lesions: Fibrous dysplasia, ossifying fibroma, cementifying fibroma, and cemento-ossifying fibroma. According to the latest WHO categorization, benign fibro-osseous lesions in the oral and maxillofacial regions were divided into two categories, osteogenic neoplasm and non-neoplastic bone lesions; cementifying ossifying fibroma belonged to the former category. However, the term “cementifying ossifying fibroma” was altered to ossifying fibroma (OF) in the new WHO classification in 2005.[¹]

OF is a bony tumor with possible odontogenic origin. It is believed to originate from the cells of the periodontal ligament.[²] This is a layer of fibrous connective tissue surrounding the roots of teeth. It contains multipotential cells that are capable of forming cementum, lamellar bone, and fibrous tissue.[³,⁴]

The presence of cementum or bone classifies the lesion as cementifying fibroma or OF, respectively, whereas lesions with mixture of both cementum and bone are called cemento-ossifying fibroma.[⁵]

This tumor generally occurs in young and middle-aged adults, most frequent between 35 and 40 years of ages.[²,⁶] There is a marked predilection for the female sex, the female: male ratio is 2:1.[⁶]

Central cemento-ossifying fibromas of the mandible are common, but they are unusual in the maxilla.[⁷]

CASE REPORT

A 37-year-old man was referred to the department of oral and of maxillofacial radiology of dentistry school of Isfahan medical science university. The patient had a chief compliant of pain and swelling in the right maxilla and asymmetry in it [Figure 1]. The patient mentioned that the mass had first appeared 3 months ago and became slightly larger since then. Furthermore, the patient mentioned that...
he experienced pain in the right side of his face with referral to the right eye and also episodes of blur vision.

The intraoral examination demonstrated an enlargement of the buccal maxillary right region, extending to the area of the canine and first premolar. In addition, the patient had a slight swelling at the palatal region [Figure 2]. The swelling was bony hard, non-tender with no fluctuation on manual palpation. The covering mucosa and gingiva had normal color and smooth in texture. The first premolar appeared to have been displaced by the lesion. The first premolar and canine at the right side were positive to thermal testing; percussion and palpation tests were within the normal limits. The needle biopsy did not revealed any sign.

Radiographic evaluation showed the presence of a lesion extending throughout the maxillary right toberousity and right canine. The margins of the lesion appeared to be well-defined. Bone pattern is radiolucent with a few wispy trabeculae [Figures 3a and b]. The tumor caused displacement of first right premolar with disappearing of its lamina dura. The mass caused expansion in all direction, but the cortical plate of the bone remained intact. The lesion occupied maxillary sinus and expanded its walls outward. Meanwhile a bony partition existed between the walls of the remaining sinus and the mass.

The CBCT showed a multi locular mass with wispy trabeculaes (similar to the pattern seen on Central Giant Cell Granuloma), occupying and expanding the right maxillary sinus [Figures 4a-c].

The expansion of Central Ossifying Fibroma into the antrum of the maxilla was convex, which was different from the fibrous dysplasia that its borders are unrecognizable from the antrum.

A comprehensive explanation of treatment was explained to the patient and an informed consent was taken. The tumor was removed under local anesthesia and the gross measures of the specimen were approximately $45 \times 30 \times 15$ mm [Figure 5]. Histological analysis showed a cellular fibroblastic tissue characterized by bundles of collagen densely packed and proliferating fibroblasts. There was presence of spherules of acellular mineralization that resembled dental cementum (cementoid type) [Figures 6 a and b]. Diagnosis of central cementifying fibroma was made. The postoperative follow up was favorable, and 1 year later the patient reported no discomfort in the area. A control panoramic X-ray study confirmed good bone regeneration.

**DISCUSSION**

COF predominantly affects the craniofacial bone and rarely involves the long bones, among the craniofacial
Central cementifying fibroma of maxilla

Sheikhi, et al.: Central cementifying fibroma of maxilla

Dental Research Journal / January 2013 / Vol 10 / Issue 1

124

bones; The mandible is the most commonly involved site.\[8,9\]

COF in the maxilla most often appears in the canine fossa and zygomatic arch area. The clinical features of COF can vary from indolent to aggressive behavior. The disease usually is asymptomatic at the time of discovery.\[10\]

But growth of COF can produces a noticeable swelling and mild deformity; displacement of teeth may be an early clinical feature.\[6\] Central cemento-ossifying fibromas are typically solitary and well-defined lesions. In the early stages, it appears as a radiolucent lesion with no evidence of internal radiopacities. As the tumor matures, increasing calcification is obvious and the radiolucent area becomes flecked with opacities until ultimately the lesion appears as an extremely radiopaque mass.\[6\]

The differential diagnosis includes: Chondrosarcoma or osteosarcoma, fibrous dysplasia, odontogenic cysts, squamous cell carcinomas, calcifying odontogenic cysts (Gorlin cysts), and calcifying epithelial odontogenic tumors (Pindborg tumors). The well-defined border of the central cemento-ossifying fibroma helps differentiate it from the aggressive sarcomas and carcinomas. Fibrous dysplasia has a characteristic “ground glass” appearance not seen in the central cemento-ossifying fibroma. The radiologic differentiation of central COF from Gorlin cysts and Pindborg tumors is difficult; the final diagnosis is based on histologic appearance.\[11\]

The recommended treatment for this tumor is surgical excision. COF usually “shell out” easily in surgical procedure, but maxillary central COF are more difficult to remove completely than mandibular. This may be attributable to the difference in bone character between the mandible and maxilla and to the available space for expansion in the maxillary sinus.\[2\]

REFERENCES

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