Findings of Florid Cemento-Osseous Dysplasia: a Report of Three Cases

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ABSTRACT

Background: Florid cemento-osseous dysplasia (FCOD) is a benign jaw lesion originating from periodontal ligament tissues usually asymptomatic and diagnosed accidentally at routine dental radiographic examination. The purpose of this paper is to report three cases diagnosed as FCOD with their clinical, radiographic and histological findings.

Methods: Radiologic and clinical symptoms of three cases diagnosed as FCOD are presented. Serum alkaline phosphatase test and biopsy taken from two of the patients are discussed to eliminate the Paget’s disease.

Results: Three patients diagnosed as FCOD and called for routine follow-up. Because of no sign of infection or osteomyelitis, conservative treatment was applied.

Conclusions: Radiographic examination is significant for the diagnosis of florid cemento-osseous dysplasia, especially in the asymptomatic cases. The roles of the dentist are to ensure the follow-up of the diagnosed patients and to take the necessary measures for preventing from the infections.

Keywords: florid cemento-osseous dysplasia; panoramic radiography; bone diseases.

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INTRODUCTION

Florid cemento-osseous dysplasia (FCOD) is one of the subgroups of cemento-osseous dysplasia (COD) which was first described by Melrose [1], Abrams and Mills in 1976 [2]. FCOD was also reported as multiple cemento-ossifying fibroma, sclerosing osteomyelitis, sclerosing osteitis, multiple osteomas, periapical cementoblastoma, Paget’s disease of the mandible, multiple enostosis, gigantiform cementoma, florid osseous dysplasia [1,3]. The exact etiology of FCOD is still unknown [2]. Most authorities suggest that pathogenesis of these lesions come from periodontal ligament, because they are seen close to it and have similar histopathology [4,5]. Therefore, few authors believe that remains of cementum in bone after extraction might be a reason of FCOD [6,7].

FCOD affects mostly middle aged to older black women. Male:female ratio reported as 1:2,6 [4,5]. FCOD is usually asymptomatic and diagnosed accidentally at routine dental radiographic examination. Dull pain, drainage, exposure of the lesion in oral cavity, focal expansion and facial deformities are present when infection occurs [3,8]. Radiographic image of FCOD is lobular radiopacities that grow with the maturation of lesion surrounded with radiolucent area and located mostly in mandibular premolar-molar region [2,9]. This image is named “cotton wool” appearance and is also seen in Paget’s disease of bone [9]. The purpose of this paper is to report three cases diagnosed as cemento-osseous dysplasia with their clinical, radiographic and histological findings.

CASE DESCRIPTION AND RESULTS

Case 1

25 year-old Caucasian woman was referred to a dentist, because of periodontal complaints. Bilateral radiopaque lesions were revealed by dental radiographic examination and consequently, the patient, without any related clinical complaint, was directed to Istanbul University, Faculty of Dentistry. Panoramic radiograph showed ovoid radiopaque masses in wide radiolucent spaces in the periapical areas of all the molars in both quadrants of the mandible and in the maxillary right quadrant. The lesion, which located at the apex of #27 showed no internal calcification (Figure 1). There was no expansion of jaws. Electric pulp tests were applied to the involved teeth: teeth were vital except right mandibular first molar at which a big composite restoration and a poor root canal therapy was observed. It was recommended to patient that retreat her #46 tooth’s root canal therapy in order to prevent a possible infection. The result of the requested serum alkaline phosphatase (ALP) test was 88 IU/L (N: 25-94 IU/L). We diagnosed present pathology as FCOD lesion and decided to follow the patient without taking biopsy. For the patient, who did not have any clinical complaints, radiographic control was recommended twice a year. Patient’s follow-up radiograph, which was taken 6 months later, showed the same condition with the previous one (Figure 2).

Case 2

71 year-old Caucasian female patient applied to Istanbul University, Faculty of Dentistry, because of the pain in posterior region of the left mandible
when using her total dentures during mastication. She was systemically healthy and had no extraoral symptoms. Intraoral examination showed totally edentulous jaws and normal mucosa. Panoramic radiograph was taken and radiopaque masses were seen at four quadrants, no sign of infection or osteomyelitis (Figure 3). Initial diagnosis was FCOD, but there was no clinical explanation for the pain. Therefore, dental volumetric tomography was taken. In cross sectional images of the dental volumetric tomography a bone spine was detected and we considered that this spine was the reason of pain (Figure 4). Alveoloplasty was performed on left posterior mandible and during the surgical process, the biopsy was performed owing to lesion’s close adjacency to surgical area. The result of the histopathological examination was FCOD. No other treatment was thought. The patient was called for follow-up 6 months later.

**Case 3**

41 year-old systemically healthy Caucasian female patient was referred to Istanbul University, Faculty of Dentistry, Department of Oral and Maxillofacial Radiology for dental care. Panoramic radiograph obtained due to dental caries and prosthetic indications. Radiopaque ovoid masses were seen at left and right mandibular quadrants (Figure 5). To eliminate the diagnosis of Paget’s disease of bone, serum alkaline phosphatase level examination test was performed. Decayed roots of right second molar seen at radiography. Biopsy was performed at right left molar region. In histopathological examination, trabecular and compact bone tissue in collagen rich connective tissue stroma was seen. In some area spheroidal hard tissue islands like cementum was detected. Based on the result of the histopathological examination diagnosis the FCOD was established (Figure 6). #24 and #47 were extracted and patient was disposed for routine dental treatments. After 6 months patient was recalled for clinical and radiographic examination. Panoramic radiograph revealed no significant difference (Figure 7).

**DISCUSSION**

FCOD classified by World Health Organization in 2005 as bone related lesions and is a non-neoplastic fibro-osseous lesion [25]. FCOD is one of the subgroups of cemento-osseous dysplasia (COD) [12]. The other COD lesions are focal cemento-osseous dysplasia and periapical cemento-osseous dysplasia [1]. This classification based on clinical characteristic, location and radiographic features [24]. These clinical features include lesions’ localization and diffuseness.

![Figure 3. Panoramic radiograph of the patient shows radiopaque masses at four quadrants.](image)

![Figure 4. Cross-sectional image showing the FCOD lesion and the spine on the top of the crest.](image)

![Figure 5. Panoramic radiograph of the patient shows radiopaque masses at both mandibular quadrants.](image)
Periapical cemental dysplasia usually locates area of apices of anterior teeth and mostly doesn’t grow more than 1 cm, focal cemeto-osseous dysplasia usually appears at two or more mandibular anterior teeth or at the apices of molar region, and rarely grow more than 2 cm [5,10]. FCOD appears bilateral, often quite symmetrical location [3]. The density of the lesion has a wide range from radiolucent to almost complete radiopacity [9]. The term “florid” refers to its excessive and widespread location [5,11].

All types of COD lesions have similar histopathology. In dysplastic period normal bone is replaced with fibrous connective tissue [1]. This fibrous connective tissue contains woven bone, lamellar bone and cementum like particles [4]. As the lesions mature, calcification degree increases [1].

FCOD have three developmental stages and all stages have different radiographic images. First or osteolytic stage is seen as well-defined radiolucent area with loss of lamina dura and periodontal ligament. In the second or cementoblastic stage, small radiopacities appears in radiolucent area, because of deposition of cementum-like droplets in fibrous tissue. Last stage is described as definite radiopacity present in majority of the lesion [10].

Diagnosis is made by clinical and radiographic examination. If the radiological and clinical findings are enough to diagnose, biopsy should be avoided because of infection, sequestrum formation and osteomyelitis risks [2]. Generally, infection risk relies on poor vascularization of the lesion [1]. This condition requires surgical treatment of the lesion. Because of the poor healing, fracture or infection risks, extraction and excision of the lesion are not recommended [2]. For asymptomatic patients the treatment is not necessary and best choice is routine follow-ups and protection from infection [2,4]. Treatment options of the symptomatic patients are antibiotic therapy and sequestrectomy [4].

Another possible complication is difficult to control infections due to insufficient antibiotics concentration in the lesion area owing to reduced blood flow [1]. In our cases, there were no infections. Therefore we recommended routine follow-ups and to increase oral hygiene in order to prevent possible infections. Biopsies were taken in case two and three, since the operation areas were adjacent to the lesions.

Differential diagnosis of FCOD must be made with Paget’s disease of bone, chronic diffuse osteomyelitis, Gardner’s syndrome, familial gigantiform cementoma [5,2]. FCOD is not seen with skin tumours, dental anomalies

Figure 6. Trabecular and compact bone seen between loose connective tissue (hematoxylin and eosin stain, original magnification x40).

Figure 7. Follow-up radiograph taken 6 months later.
and skeletal changes like Gardner’s Syndrome [3]. In Paget’s disease of bone, no radiolucent capsule is present around the radiopaque masses, also it is usually polyostotic and causes raised alkaline phosphatase level [3,9]. Chronic diffuse osteomyelitis is not limited in tooth bearing areas and cause unilateral pain and swellings [3]. Familial gigantiform cementoma has familial basis and from this point it is differentiated from FCOD [5]. In our cases, serum alkaline phosphatese levels were in normal ranges and there were no signs of infection. Our cases had neither dental anomalies nor skin tumours and the patients had no familial history.

CONCLUSIONS

Radiographic examination is significant for the diagnosis of florid cemento-osseous dysplasia, especially in the asymptomatic cases. The roles of the dentist are to ensure the follow-up of the diagnosed patients and to take the necessary measures for preventing from the infections.

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There are no relevant conflicts of interest to disclose.
