

British Journal of Medicine & Medical Research

21(5): 1-6, 2017; Article no.BJMMR.32449 ISSN: 2231-0614, NLM ID: 101570965

Cementoblastoma in the Maxilla: An Unusual Case Report

Delise Pellizzaro^{1*}, Guilherme Monteiro Tosoni¹, Marcelo Gonçalves¹, Valfrido Antônio Pereira Filho¹, Oscar Fernando Muñoz Chávez¹ and Andrea Gonçalves¹

¹Faculdade de Odontologia de Araraquara, Universidade Estadual Paulista, FOAr- UNESP, Araraquara - SP, Brazil.

Authors' contributions

This work was carried out in collaboration between all authors. Authors DP, AG and MG managed the radiographic and histopathological analysis and wrote this manuscript. Author GMT wrote this manuscript. Authors VAPF and OFMC did surgical treatment. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/BJMMR/2017/32449

<u>Editor(s):</u>

(1) Emad Tawfik Mahmoud Daif, Professor of Oral & Maxillofacial Surgery, Cairo University, Egypt.
 (2) James Anthony Giglio, Adjunct Clinical Professor of Oral and Maxillofacial Surgery, School of Dentistry, Virginia Commonwealth University, Virginia, USA.

(3) Salomone Di Saverio, Emergency Surgery Unit, Department of General and Transplant Surgery, S. Orsola Malpighi University Hospital, Bologna, Italy.

Reviewers:

(1) Deepak Pandiar, Banaras Hindu University, India.

(2) V. K.Vaishnavi Vedam, Asian Institute of Medicine, Science and Technology (AIMST) University, Malaysia.
(3) Kotya Naik Maloth, Kaloji Narayana Rao University of Health Sciences, India.
Complete Peer review History: http://www.sciencedomain.org/review-history/19009

Case Study

Received 26th February 2017 Accepted 6th May 2017 Published 11th May 2017

ABSTRACT

Cementoblastoma is characterized by the formation of cementum-like tissue attached to the root of a tooth. It is a benign neoplasm with odontogenic mesenchymal origin, and is considered the only true neoplasm of cemental origin. Cementoblastomas are usually located in the mandible, the majority of which affect the permanent first molar. There is no distinct gender preference and the mean age of occurrence is approximately 20 years of age. The lesion is usually identified as well defined, radiopaque, delimited by a cortical border, and with a well-defined radiolucent band just inside this cortical border. Root resorption and loss of the root outline are commonly associated with the cementoblastoma. The only treatment is enucleation of the lesion with removal of the associated tooth. Here, we present an unusual cementoblastoma case in a 52-year-old woman

^{*}Corresponding author: E-mail: delisepellizzaro@yahoo.com.br;

with different radiographic findings from those frequently reported in the literature. The patient has been followed for 2 years with no sign of recurrence.

Keywords: Cementoblastoma; panoramic radiography; cone-beam computed tomography; maxilla.

1. INTRODUCTION

The cementoblastoma is considered to be a rare benign neoplasm of odontogenic mesenchymal origin [1], characterized by the formation of cementum-like tissue contiguous to the root of a tooth [2,3,4]. This tumor manifests as a bulbous growth surrounding and attached to the apex of a tooth root [5].

The majority of cementoblastomas are located in the mandible, most commonly in the molar region [1,2,4,5,6,7]. There is no gender predisposition [2,6,7,8], and the age of occurrence ranges from 8 to 44 years [2], with the highest incidence in the second and third decades of life [2,4,6,7,8,9].

This paper describes an unusal case of a patient who was diagnosed with cementoblastoma based on radiographic, surgical, and histological findings.

2. PRESENTATION OF CASE

A dental practitioner referred a 52-year-old Caucasian female to the Division Dentomaxillofacial Radiology for panoramic imaging. This image revealed a 2 cm x 1.2 cm round, well-defined, radiopaque mass attached to the apex of the root of the left maxillary second molar, (Fig. 1A). The density of the cemental mass obscured the outline of the root apex, and no radiolucent halo was evident. Given these our rule-out diagnosis included: findings, cementoblastoma, periapical cemental dysplasia. and periapical idiopathic osteosclerosis. A periapical image was requested to allow more detailed evaluation of the region (Fig. 1B).

The patient reported no history of pain and her medical history showed no evidence of systemic diseases. Panoramic and periapical images taken 23 years prior were available in her dental records. The panoramic image (Fig. 1D) did not show any evidence of a radiopaque mass attached to the left maxilla, and the periapical image (Fig. 1C) showed widening of the periodontal ligament space and slight hypercementosis.

The patient was referred to an oral maxillofacial surgeon who requested a cone beam computed

tomography (CBCT) scan (Figs. 2A and B). This scan showed the presence of a well-defined, high-density mass of $1.53~\rm cm \times 1~\rm cm \times 1~\rm cm$ in the left maxilla. This lesion was attached to the roots of the second molar. The radiology report of the CBCT, which was carried out in a private radiological practice, suggested the following potential diagnoses: (1) periapical idiopathic osteosclerosis, (2) retained teeth, or (3) complex odontoma.

On clinical examination, there was no intraoral swelling in the buccal gingiva of the left maxillary second molar (Fig. 3), and tooth was vital. According to the clinical examination and radiological findings, a presumptive diagnosis of cementoblastoma was made. Considering the patient's cooperative behavior, a decision to perform excision of the lesion, tooth extraction and an autogenous ramus bone graft under local anesthesia was planned. The excised mass was histopathological sent for examination. Macroscopically the tooth was observed to be irregular, white, and with an adherent hard tissue mass (3 cm in width). Microscopy the tumor was composed of dentin with cementum containing basophilic incremental lines, multinucleated giant cells, cementoblasts, and cementocytes (Fig. 4). The histopathology diagnosis was cementoblastoma.

The post-operative course was uneventful and the patient's condition at the 6-month follow-up was excellent. Her dental practitioner restored the maxilla with a dental prosthesis. The patient has been followed clinically and radiologically for the past 2 years and no recurrence of the tumor. (Figs. 2C and D).

3. DISCUSSION

The typical clinical presentation of cementoblastoma includes: male or female under 30-years-old, can be painful or asymptomatic, swelling, located in the premolar/molar region, [1,4,6,10,11,12,13], more frequently in the mandible [1,4,6,7,10,12,11,12,13,14,15,16]. Previous studies have reported the occurrence of cementoblastoma in the right maxillary and mandibular canine [6,8], right upper posterior tooth region [1], maxillary central incisor [7],

mandibular first molar [14] and maxillary right first molar [4]. In the case described here, a tumor was identified in the left posterior maxilla of a 52year-old female. This is now the oldest reported age occurrence for cementoblastoma in the Western population.

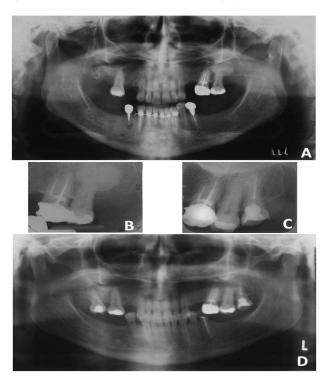


Fig. 1. (A) Panoramic image showing a round, well defined, radiopaque mass attached to the root of the left maxillary second molar. (B) Periapical image allowing more detailed evaluation. (C) Periapical image showing widening of the periodontal ligament space. (D) Panoramic image showing the absence of a radiopaque mass attached to the left maxilla

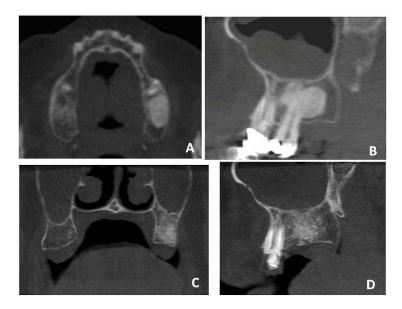


Fig. 2. (A) Axial and (B) sagittal slice images from cone beam CT. (C) Coronal and (D) sagittal images performed during follow-up showing no signs of recurrence

Pain, tenderness and swelling are present in most cases of cementoblastoma [2,8,11]; however, some cases can be asymptomatic [1,4,6,7,8]. In the present case, the tooth was vital and asymptomatic, however, unlike most cases described in the literature, no swelling was observed and the lesion was only detected from the radiographic examination.



Fig. 3. Intraoral appearance



Fig. 4. Histopathological image

Radiographically, most cementoblastomas show opacity surrounded by a radiolucent halo [1,2,7,10,11,13,14,15], and this thin, uniform, radiolucent border has been observed to surround lesions that are primarily radiopaque [14]. This present case is consistent with previous reports that showed a radiopaque mass [6,12]; however, it was not surrounded by a radiolucent halo. The root contour was lost due to fusion with the tumor [2,10], in addition to elimination of the periodontal ligament space [2]. Table 1 shows comparative clinical and radiographic features described for several authors [1,3,4,5,6,7,12,13,15,17,18].

Intraoral and panoramic radiography are highly effective methods for detecting abnormal

maxillo/mandibular lesions and examining the relationship between the lesions and teeth [2,19]. Although these images are an essential component of the diagnostic protocol, CBCT images were also helpful for surgical planning in the case described here.

The combination of clinical, radiographic and histopathological methods is recommended for the diagnosis of this type of lesion, even when pathognomonic signs are present [20]. This protocol was followed in the present case.

Cementoblastoma is occasionally difficult to distinguish from other lesions, such as periapical cemental dysplasia, periapical idiopathic osteosclerosis. hypercementosis, focal sclerosing osteitis, osteoid osteoma and osteoblastoma [3]. Differentiating between periapical cemental dysplasia and cementoblastoma may be difficult in some cases, requiring observation over an extended period. radiolucent band The around cementoblastoma is usually better defined [5]; however, a radiolucent halo was not present in this case, and the existence of periapical idiopathic osteosclerosis may not be correlated with the presence of teeth [5].

Hypercementosis may resemble a small benign cementoblastoma [5]. In the present case, the cementoblastoma caused elimination of the periodontal ligament space of the left maxillary second molar, and did not induce expansion of the jaws. The differential diagnosis of focal sclerosing osteitis may rely solely on clinical examination, including a test of tooth vitality [5]. Osteoid osteoma is a benign tumor that is rare in the jaw, and presents central radiolucency [5].

According to previous reports [2,14], the histopathological presentation of cemento-blastoma is identical to osteoblastoma, and the primary distinguishing feature is fusion of the tumor with the involved tooth [1].

Two methods have been reported for treatment of cementoblastoma. The more invasive consists of excision and extraction of the associated tooth [1,2,4,5,6,7,8,14]. The least invasive being apical resection and root canal treatment of the affected tooth [1,2,20]. In the case reported here, the most invasive method was chosen because it presents a lower rate of recurrence [1,2]. The reported recurrence rate ranges from 9-37% [4,6,8,20].

Table 1. Comparison the maxillary cementoblastomas clinical and radiographic features reported in literature

Author	Gender/Age	Clinical features	Radiographic features
Present case	Female/52 years	Asymptomatic;	Well-defined;
		No swelling;	No radiolucent halo;
		Mucosa appeared normal;	Radiopaque mass;
Dadhich and Nilesh,	Female/23 years	Asymptomatic;	Well-defined;
2015 [3]		Swelling;	Hyper dense mass;
White and Pharaoh,	General description:	Pain frequently;	Well-defined;
2014 [5]	male and female/ 12 to 65 years		Radiolucent halo;
Neelakandan et al.,	Female/11 years	Pain;	Well-circumscribed radio-
2012 [1]		Diffuse smooth surfaced	opacity;
		swelling;	Radiolucent halo;
Harada et al., 2011	Male/8 years	Asymptomatic;	Hard mass;
[4]		Slight redness;	
Costa et al., 2011	Female/11 years	Slightly painful;	Radiolucent halo;
[7]	·	Swelling;	Radiopaque mass;
		Mucosa was normal;	
Hirai et al., 2010 [6]	Female/15 years	Asymptomatic;	Well-defined;
		Mucosa was normal;	Radiopaque mass;
Infante-Cossio et al., 2008 [17]	Male/20 years	Swelling;	Radiopaque mass;
Ohki et al., 2004	Male/12 years	Asymptomatic;	Well-defined;
[12]	•	Swelling;	Radiopaque mass;
		Mucosa was normal;	
Garlick et al., 1990 [13]	Male/19 years	Pain;	Radiolucent halo;
	·	Swelling;	Radiopaque mass;
		Mucosa was normal;	
Puterman et al.,	Female/14 years	Large swelling;	Radiolucent halo;
1988 [18]	·	Normal mucoperiosteum;	Radiopaque mass;
Adkins et al., 1973	Male/24 years	Pain;	Radiopaque mass;
[15]	•	Swelling;	
		Mucosa was normal;	

4. CONCLUSION

In conclusion, the present case report describes an unusual cementoblastoma case in a 52-year-old woman, the radiographic findings of which differ from those frequently reported in the literature. The patient has been followed for 2 years with no sign of recurrence.

CONSENT

All authors declare that written informed consent was obtained from the patient for publication of this case report and accompanying images.

ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- Neelakandan RS, Deshpande A, Krithika C, Bhargava D. Maxillary ementoblastoma

 A rarity. Oral Maxillofac Surg.

 2011:16:119-21.
- Barnes L, Eveson JW, Reichart P, Sidransky D, editors. World Health Organization classification of tumours. Pathology and genetics of head and neck tumours. Lyon: IARC Press; 2005.
- 3. Dadhich AS, Nilesh K. Cementoblastoma of posterior maxilla involving the maxillary sinus. Ann Maxillofac Surg. 2015;5:127-9.
- Harada H, Omura K, Mogi S, Okada N. Cementoblastoma arising in the maxilla of an 8-year-old boy: A case report. Int J Dent; 2011.
- 5. White SC, Pharoah MJ. Oral radiology: Principles and interpretation. 7th ed. St Louis: Mosby; 2014.
- 6. Hirai E, Yamamoto K, Kounoe T, Kondo Y, Yonemasu H, Kurokawa H. Benign

- cementoblastoma of the anterior maxilla. Oral Maxillofac Surg. 2010;68:671-4.
- Costa FW, Pereira KM, Magalhães Dias M, da Costa Miguel MC, de Paula Miranda MA, Studart Soares EC. Maxillary cementoblastoma in a child. J Craniofac Surg. 2011;22:1910-3.
- Çalışkan A, Karöz TB, Sumer M, Açıkgöz A, Süllü Y. Benign cementoblastoma of the anterior mandible: An unusual case report. J Korean Assoc Oral Maxillofac Surg. 2016;42(4):231-5.
- Nuvvula S, Manepalli S, Mohapatra A, Mallineni SK. Cementoblastoma relating to right mandibular second primary molar. Case Rep Dent. 2016;23.
- Sankari LS, Ramakrishnan K. Benign cementoblastoma. J Oral Maxillofac Pathol. 2011;15:358-60.
- Rubino I, Cudia G, Ficarra G. Benign cementoblastoma. A case report. Minerva Stomatol. 1999;48:539-41.
- Ohki K, Kumamoto H, Nitta Y, Nagasaka H, Kawamura H, Ooya K. Benign cementoblastoma involving multiple maxillary teeth: Report of a case with a review of the literature. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 2004:97:53-8.
- Garlick AC, Newhouse RF, Boyd DB. Benign cementoblastoma: Report of a case. Mil Med. 1990;155:567-70.
- Brannon RB, Fowler CB, Carpenter WM, Corio RL. Cementoblastoma: An innocuous neoplasm? A clinicopathologic

- study of 44 cases and review of the literature with especial emphasis of recurrence. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 2002;93:311-20.
- Adkins KF, Monsour FN. Benign cementoblastoma in the maxilla. Aust Dent J. 1973;18:99-101.
- Mosqueda-Taylor A, Ledesma-Montes C, Caballero-Sandoval S, Portilla-Robertson J, Ruíz-Godoy Rivera LM, Meneses-García A. Odontogenic tumors in Mexico. A collaborative retrospective study of 349 cases. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 1997;84:672-5.
- Infante-Cossio P, Hernandez-Guisado JM, Acosta-Feria M, Carranza-Carranza A. Cememtoblastoma involving the maxillary sinus. Br J Oral Maxillofac Surg. 2008;46(3):234-6.
- 18. Puterman M, Fliss DM, Sidi J, Zirkin H. Giant cementoblastoma simulating a peridental infection. J Laryngol Otol. 1988;102(3):264-6.
- Kaneda T, Minami M, Kurabayashi T. Benign odontogenic tumors of the mandible and maxilla. Neuroimag Clin N Am. 2003;13:495-507.
- Costa BC, de Oliveira GJ, Chaves Md, da Costa RR, Gabrielli MF, Guerreiro-Tanomaru JM, et al. Surgical treatment of cementoblastoma associated with apicoectomy and endodontic therapy: Case report. World J Clin Cases. 2016;4(9):290-5.

© 2017 Pellizzaro et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here:
http://sciencedomain.org/review-history/19009