Adenomatoid Odontogenic Tumor: Case Report and Literature Review

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Abstract
The Adenomatoid Odontogenic Tumor presents in low frequency and is originated from epithelial rests, specifically from the dental lamina, and shows similar characteristics to a wide variety of pathologies. It is a benign non-invasive lesion that can be managed by conservative treatment. Thus, it is important to recognize and differentiate it from other pathologies that required more aggressive surgical treatment. The objective of the present article is to report a case of an Adenomatoid Odontogenic Tumor along with its common features.

Keywords
Adenomatoid odontogenic tumor, Odontogenic tumors

Introduction
Odontogenic Tumors are a group of heterogenous uncommon lesions that affect maxillary and mandibular osseous tissue, due they originate from mesenchymal and/or epithelial components from the odontogenesis. OT are lesions that present a wide variety of clinical and histopathological features, developing from hamartomas to malignant lesions [1,2].

The AOT was first described more than 100 years ago by Harbitz as “cystic adamantoma”. Since then, it has received multiples denominations until 1969 when the World Health Organization accepted the term “Adenomatoid Odontogenic Tumor” proposed by Philipsen and Birn [3]. The WHO defines it as a tumor composed of odontogenic epithelium in a variety of histarchitectural patterns, embedded in a mature connective tissue stroma and characterized by slow but progressive growth [4].

The AOT is the fourth more common odontogenic tumor, appearing in 2-7% of cases [5,6]. It arises from odontogenic epithelium, which indicates it could originate from the enamel organ, reduced enamel epithelium, Mallasez rests or dental lamina rests, being this the most accepted theory [7]. Establishing this pathology as a hamartoma or a neoplasm is reason of great controversy.

Some authors name the AOT as the “two thirds tumor” because two thirds of the cases present in the anterior maxilla, in women, associated with an included tooth of which two thirds is the canine and two thirds of the intraosseous type present radiopacities within the lesion. Tooth resorption is rare; although, cortical expansion and adjacent tooth displacement are regularly seen [8-10].

Three types differentiate according to the location: Follicular and extrafollicular which are differentiated radiographically and peripheral which is observed only clinically [3]. The differential diagnosis with other pathologies such as ameloblastoma and keratocyst is important due to the aggressive treatment they required. Conservative treatment is indicated for AOT because of its low recurrence [6].

The aim of the present article is to report an Adenomatoid Odontogenic Tumor case, as well as describe its more common features allowing correct diagnosis and treatment.
Case Report

An asymptomatic 17-year-old female presented to the consult with facial asymmetry on the left side of the maxilla. On the extraoral exploration, an increase of volume of the left maxilla was observed, as well as no change of the skin coloration. Intraorally, it was crackling due to vestibular cortical expansion on left maxilla, extending from the left lateral incisor to the left molar region. Displacement of the left lateral incisor, absence of the left upper canine and persistence of the deciduous canine was also seen.

In the computed tomography was observed a 4 cm hypodense, unilocular, well-defined and non-invasive image located on the left maxilla. A hyperdense image, compatible with unerupted left canine was noted within the lesion (Figure 1), and displacement of the maxillary sinus.

The treatment, initially, consisted of an incisional biopsy which histopathologic study reported the definitive diagnosis of Adenomatoid Odontogenic Tumor. Posteriorly, under general anesthesia, enucleation and curettage was performed as definitive treatment (Figure 2 and Figure 3).

At 4-year follow-up, the patient showed no signs of clinical or imaging recurrence (Figure 4).

Discussion

The AOT is an uncommon tumor that originates from the odontogenic epithelium; specifically, from the rests of the dental lamina, which are found in bone, mucosa and the gubernaculum canal. There are three variants depending on the location: follicular, extrafollicular and peripheral [7,11].

The intraosseous variants (follicular and extrafollicular) correspond to 96% of the cases [12]. The follicular variant is associated to the gubernaculum canal, which
contains a fibrous band with rests of the dental lamina, connecting the peri coronal follicle of a permanent tooth with the alveolar mucosa. In occasions, the rests of the dental lamina from the deciduous teeth can migrate within de gubernaculum canal [3,7]. The extralocular variant is associated to rests of the dental lamina that remained in the bone after the development of the deciduous germ [7]. The peripheral variant is associated to rests of the dental lamina in the alveolar mucosa and is seeing clinically as a gingival overgrowth [11,13].

Radiographically, the intraosseous variants share some features as well-defined radiolucency with small radiopacities, adjacent tooth displacement, size usually no more than 3 cm; root resorption and cortical perforation are rare [12]. Despite their similarities, the follicular type is associated to an included tooth, being the canine the more frequent; while the extralocular type is not associated to an included tooth and can appear between, above, below or superimposed to the roots of permanent teeth [10]. In suspicion of presence of AOT, periapical radiography shows better the calcified deposits within the lesion [8,13].

Clinically, is an expansive and painless lesion that usually displace adjacent teeth. It is a slow growing lesion that shows facial asymmetry when growth is considerable [14]. In some case reports, facial asymmetry caused by growth of the tumor has been compared to an ameloblastoma [15-17].

Classify the AOT as a hamartoma [8,11,18] or as a non-aggressive, non-invasive neoplasm [5,13,19] has generated great controversy. By definition, a neoplasm has unlimited potential for growth; while, the hamartoma has limited potential for growth and is a developmental anomaly with tissue differentiation [13].

The differential diagnosis of the AOT depends on the variant found. The peripheral variant is differentiated only with gingival pathologies such as epulis and gingival fibroma. The diagnosis is purely histopathological [8,13]. As most of the cases of the AOT are associated to an unerupted tooth, the differential diagnosis most include the dentigerous cyst; nonetheless, when the radiolucency exceeds the cementum-enamel junction it has to be ruled out [10,19]. Because of the small radiopacities within the lesion, the epithelial odontogenic tumor and the epithelial odontogenic cyst should be considered [14]; as well as the ossifying fibroma and cemento-ossifying fibroma. Presenting as a unicocular radiolucency it could appear as an unicystic ameloblastoma. On the other hand, though rare, the AOT can present as a multilocular radiolucency, hence the keratocyst and ameloblastoma must be contemplated [10].

Conservative surgical management consisting of enucleation and curettage is the treatment indicated because of its low recurrence and favorable prognosis [9,11,14].

Conclusion

The reported case presented the usual features describe in the literature such as age (seventeen), gender (feminine), anatomic region (anterior maxilla), unerupted teeth (canine), slow growing, tooth displacement and no root resorption. Nevertheless, the size was bigger (4 cm). There was no recurrence after conservative treatment. Identifying the features of this tumor allows differentiating it from other entities that required a different surgical management.

Declarations

Consent for publication

Patient’s consent for publication was acquired.

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Competing interests

The authors declare they have no competing interests.

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