Low Grade Central Mucoepidermoid Carcinoma

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ABSTRACT

Mucoepidermoid carcinoma (MEC) is the most common malignant salivary gland tumor, comprising about 15% of all salivary gland tumors and 30% of all salivary malignancies. Most of the MEC arise in the parotid gland. Rarely, it originates in the mandible and maxilla as an intraosseous variant, referred to as 'central mucoepidermoid carcinoma' or 'intraosseous mucoepidermoid carcinoma'. Central mucoepidermoid carcinomas (CMECs) are extremely rare, but well-known entity, comprising 2 to 3% of all MECs reported. Histopathologically, this malignant neoplasm is characterized by mucous, intermediate and epidermoid cells. In this report, we present a case of a male patient diagnosed as low grade CMEC.

Keywords: Central mucoepidermoid carcinoma, Intraosseous, Mandible, Low grade.

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INTRODUCTION

Mucoepidermoid carcinoma (MEC) is the most common malignant salivary gland tumor, comprising about 15% of all salivary gland tumors and 30% of all salivary malignancies. This entity was initially proposed by Masson and Berger in 1924. Stewart et al, in 1945, described its mucus secreting and epidermal cellular elements, thus, establishing it as a distinct pathologic entity. The involvement of parotid glands predominate, representing 45%. Palate is the most frequently involved minor salivary gland site. Mucoepidermoid carcinoma can also arise from ectopic salivary gland tissue in periparotid lymph nodes as well as in the larynx, lacrimal gland, nose, paranasal sinuses, lung and trachea. Rarely, it originates in the mandible and maxilla as an intraosseous variant, referred to as 'central mucoepidermoid carcinoma (CMEC)’ or ‘intraosseous mucoepidermoid carcinoma’. Central mucoepidermoid carcinomas are extremely rare, but a well-known entity, comprising 2 to 3% of all MECs reported. Lepp in 1939 first reported of a CMEC of the mandible in a 66-year-old woman, and Bhaskar (1963) analyzed the criteria for their central origin, histology and pathogenesis. Waldron and Mustoe (1989) suggested that CMEC should be included in primary intraosseous carcinoma of jaws as type 4.

Classification of primary intraosseous carcinomas (Waldron and Mustoe) is as follows:

- **Type 1:** Primary intraosseous odontogenic carcinoma ex odontogenic cyst
- **Type 2A:** Malignant ameloblastoma
- **Type 2B:** Ameloblastic carcinoma
- **Type 3:** Primary intraosseous odontogenic carcinoma developed de novo
  - a. Keratinizing type
  - b. Nonkeratinizing type
- **Type 4:** Central intraosseous mucoepidermoid carcinoma

Mucoepidermoid carcinoma is defined by WHO (2005) as a malignant glandular epithelial neoplasm characterized by mucous, intermediate and epidermoid cells, with columnar, clear cell and oncocytoid features. Previous literature indicates over 150 reported cases of CMEC. Here, we report a case of a 27-year-old male patient with a chief complaint of painful slow-growing swelling on left posterior aspect of lower jaw diagnosed as low grade CMEC.

CASE REPORT

Clinical and Radiographic Features

A 27-year-old male patient reported to the Department of Oral and Maxillofacial Surgery, Mahatma Gandhi Missions Dental College and Hospital, with a complaint of tender, slow-growing swelling on the left posterior region of lower jaw since 2 months. He gave history of dull, continuous pain related to fully erupted 38 which gradually increased over 2 months. He was advised extraction of 38 by a general dental physician. Patient noticed subsequent increase in pain postextraction, with no change in swelling. He gave no history of consumption of any deleterious substances. Extraoral clinical examination revealed a diffused, tender, nonfluctuant, noncompressible and nonreducible swelling measuring...
around 3 × 2 cm on left mandibular body extending posteriorly to the angle of the mandible (Fig. 1). Oral examination disclosed an incompletely healed, tender extraction socket of 38, with no other significant findings (Fig. 2). Left submandibular lymph nodes were palpable and tender, and no signs of paresthesia were evident. His medical and family history was not contributory.

Orthopantomograph revealed an ill-defined unilocular radiolucency in the left body of the mandible extending from the distal root of 37 to the angle of the mandible and till the lower border of the mandible inferiorly. Resorption of the distal root of 37 and the extraction socket of 38 were also evident (Fig. 3).

Clinical and radiological differential diagnosis included cyst or tumor of odontogenic origin. An incisional biopsy was performed, after obtaining consent.

**Histopathological Features**

Histopathologically, the hematoxylin and eosin-stained soft-tissue section showed small and large closely packed nests of epidermoid, intermediate and mucous cells in a fibrous connective tissue stroma (Figs 4A and B). The mucous cells of varied shapes and sizes had abundant pale foamy cytoplasm. Epidermoid cells were polygonal in shape with ovoid to elongated nuclei and abundant eosinophilic cytoplasm. Areas of microcyst formation (Fig. 5) and focal areas of clear cells were also evident. Normal mucous acini and lymphoid aggregates were seen in the vicinity of the lesional tissue. The histopathological features were suggestive of low grade variant of CMEC. Alcian blue staining performed indicated positive mucous cell population (Fig. 6).

**DISCUSSION**

Aberrant salivary gland tissue may be found in numerous regions of the body to include lymph nodes, ear, thyroglossal duct and pituitary gland. However, central salivary gland tumors are extremely rare. Central mucoepidermoid carcinoma tend to affect the mandible (premolar-molar region) thrice more than the maxilla, with majority of the cases occurring in 4th to 5th decades of life. Generally, main presented symptoms are swelling with bony expansion, in posterior mandible with or without paresthesia. Our case, though presenting with intraosseous swelling involving the posterior mandible, was in a male patient of age 27 years.

The diagnostic criteria for CMEC were laid down by Alexander in 1974, later modified by Browand and Waldron in 1975 which included the following:

- Intact cortical plate
- Radiological evidence of bone destruction
- Histologic confirmation
- Positive mucin staining
- Absence of primary lesion in salivary gland
- Exclusion of an odontogenic tumor
- Our case fulfilled the diagnostic criteria for CMEC that was laid down by them.
Brookstone and Huvos et al in 1992 proposed a staging system based on the condition of the overlying bone.\(^5\)

- **Stage I:** Lesions with intact cortical plates with no evidence of bone expansion.
- **Stage II:** Tumors with intact plates but intraosseous expansion.
- **Stage III:** Lesions associated with cortical perforation or nodal disease.

Our case was in staged II, due to intact cortical plates with evident bony expansion.

The origin of CMEC is still controversial and some theories have been introduced, including:\(^{11}\)

- Ectopic salivary glands tissue that was developmentally entrapped in the jaws.
- Neoplastic transformation of the sinus epithelium.
- Neoplastic transformation of the epithelial lining of an odontogenic cyst (especially dentigerous cysts).

Eversole et al found that approximately 50% of mandibular CMEC were associated with dental cysts or impacted teeth, whereas Brookstone and Huvos report a rate closer to 32%.\(^{11}\)

Mucoepidermoid carcinoma is thought to arise from pluripotent reserve cells of the excretory ducts of salivary gland that have the potential to differentiate into squamous, columnar and mucous cells.\(^2,4\) Microscopically, these tumors are characterized by the presence of two populations of cells—the mucus cells and the epidermoid cells, the proportion of which helps to define the grade of the tumor. The epidermoid cells are most easily detected by its appearance. Mucous cells may contain microscopic granules filled with acid mucopolysaccharide, which yield the appearance of foam cells or goblet cells.\(^{2,4,11}\) The present case revealed a highly differentiated neoplasia with a predominance of microcysts and presence of mucin-producing cells. Special staining for mucicarmine or alcian blue can readily identify the mucous cell population, which is considered diagnostic since mucous cells are only rarely encountered in other salivary tumors.\(^5\) Our case revealed alcian blue positive
mucous cells population. Based on these findings, in the present case, a diagnosis of low grade central mucoepidermoid carcinoma was established.

Central mucoepidermoid carcinoma should be distinguished from various pathological entities, like residual cyst, chronic supplicative osteomyelitis, dentigerous cyst, odontogenic keratocyst, aneurysmal bone cyst, traumatic bone cyst, ameloblastoma, central giant cell granuloma and malignancies. Though there exist several systems to diagnose and stage this neoplasm, histopathological diagnosis remains the universally accepted golden standard.

Metastases are reported in 9% of CMEC, mainly to the regional lymph nodes. Nodal metastases have been described from 1 to 24 years after the time of initial management. The widely practiced main treatment for CMEC is wide local excision with sectional or radical neck dissection in case of cervical involvement. Cervical nodal involvement was ruled out in our case. Documented recurrence rate for radical excision is 13%. The present case underwent a posterior enblock resection with no neck dissection. The histopathological findings of the excisional specimen confirmed the diagnosis of low grade CMEC. Prognosis is difficult to evaluate due to the inadequacy of the staging and the disparity of treatment reported in the literature, whereas a well-differentiated, low-grade tumor without perineural invasion and with tumor-free margin indicates a good prognosis. The CMEC that are graded in the intermediate to high grade range imply a worse prognosis than low-grade tumors. The presented case has been disease free for past 14 months.

REFERENCES