UNUSUAL EPITHELIAL-MYOEPITHELIAL CARCINOMA IN PALATE-CASE REPORT AND IMMUNOHISTOCHEMICAL STUDY

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Abstract
Epithelial-myoepithelial carcinoma (EMC) is a neoplasm of salivary gland extremely rare and low-grade malignant, exhibiting predilection for major salivary gland, especially parotid gland. This paper described a case of EMC localized in soft palate. The initial clinical diagnosis were mucoepidermoid and adenoid cystic carcinoma. A incisional biopsy was performed, and the histological diagnosis was adenocarcinoma not otherwise specified. The patient underwent excisional biopsy, where was realized immunohistochemical analysis, staining the duct-like structures, where the inner layer of epithelial cells positively for CK 14, whereas outer layer of myoepithelial cells stained positively for smooth muscle actin (SMA) and vimentin antibodies. Based on these findings, the final histological diagnosis was EMC. The patient was submitted 30 sessions of radiotherapy, and she is being follow-up about 2 years without evidence recurrences.

Key words
Epithelial-myoepithelial carcinoma, oral cavity, soft palate, immunohistochemistry.
with a central hyperchromatic nucleus and scant eosinophilic cytoplasm. The myoepithelial cells were polygonal with more abundant, clear, vacuolated and well-defined cytoplasm, and an eccentric irregular nucleus. The fibrous connective tissue band contained numerous blood vessels and necrosis was found locally (Fig. 2).

Immunohistochemical staining was carried out with the streptavidin-biotin method. Immunoreactivity for CK7 and CK14 was observed in inner layer of epithelial cells, as well as in mucous acinus and duct salivary. The clear cells of outer layer were found to stain positively for smooth muscle actin (SMA), which is consistent with a myoepithelial phenotype. In solid areas observed immunoreactivity for vimentin in muscle cells around blood vessels and small clear cells nests, confirming the myoepithelial characteristic (Fig. 3).

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Discussion
Epithelial-myoepithelial carcinoma, recognized by WHO in 1991, is a rare malignant neoplasm in the salivary glands, comprising approximately 1% of all salivary gland tumors (9). The case related was localized in the soft palate, a rare area of onset, where it showed histologic characteristics of a low-grade behavior of aggressiveness. Rarely, high-grade or dedifferentiated EMC has been described (10,11).

These tumors arise in the salivary glands, predominantly the major salivary gland and the parotid gland (6). However, they have rarely been observed in tissues other than the salivary glands, such as the breast, lacrimal gland, lung and nasal cavity (6,12,13).

It is more frequent in elderly women, especially in the seventh decade (1). Approximately 60%
of patients with EMC are female (2). The patient here was 72 years-old female, such as in the mean age reported in the literature. She is being follow-up about 2 years, without any evidence recurrences. In the series reported by Seethala et al. (3), the recurrence rate was 36.3% in the 11-year disease-free survival, but death was rare with 5-year (93.5%) and 10-year (81.8%) disease-specific survivals. However, Fonseca and Soares (14) reported that the 5-year survival rate was 6% and 40% of patients died of their disease. In these cases, the authors suggest that the classification of epithelial-myoepithelial carcinoma as low-grade should be reviewed, as the clinical behavior of the tumor contrasts with its histopathologically bland appearance.

EMC has been postulated to be derived from the intercalated ducts of salivary glands, because the tubular growth pattern of this tumor recapitulates this phenotype. More current concepts of salivary gland tumorigenesis avoid attributing tumors to a specific structure or cell type of origin, focusing instead on classification based on the morphologic and immunophenotypic differentiation (3). Studies have reported that pleomorphic adenoma cell culture are monoclonal, probably deriving from a common precursor of myoepithelial and ductal cells. In contrast, epithelial-myoepithelial carcinoma cell culture present two different cell populations, already committed to luminal and myoepithelial differentiation (15). Although the histological appearance of the EMC is usually characteristic, immunohistochemical studies may prove to be useful in separating of others similar tumors, as adenoid cystic and pleomorphic adenoma (8).

In the present case, the tumor was composed of two types of cells, one type of duct-forming cell and other type of cells surrounding these cells. However, the pattern of the cellular arrangement found in this tumor excludes the possibility of a adenoid cystic carcinoma. The tumour also exhibiting greater epithelial cellularity with less stroma compared to pleomorphic adenomas. Immunohistochemically, in this case was observed that the myoepithelial cells were positive for vimentin and smooth muscle actin (SMA) and the epithelial cells were positive for CK 7 and CK 14. Seethala et al. (3) also observed that vimentin and SMA were positive in 75% and 81.8% for myoepithelial cells, respectively. According to this author, among of the myoepithelial markers, p63 has the best performance, showing diffuse strong staining in large number of cases, with no staining of epithelial cells.

In some cases, adenoid cystic carcinoma (ACC) can also show a double-layered, duct-like structure, it can be mistaken for ECM. But, the inner ductal cell layer in ACC stain positive for the S-100 protein, whereas only the outer ductal cell layer in ECM is positive for the S-100 protein. The different immunostaining characteristics will be useful for distinguishing between EMC and ACC when necessary (2).

Furuse et al. (5) observed nuclear staining of myoepithelial cells for β-catenin protein in 10 cases of ECM, suggesting that it may be related aggressive biologic behavior of the tumor. According these authors, the universal expression of E-cadherin and β-catenin in salivary gland tumors may explain the slow growth and later metastasis seen in these tumors compared with other carcinomas, which confirms the role of both E-cadherin and β-catenin as tumor-suppressing and invasion-suppressing molecules. In the case of ECM, when myoepithelial cells participate in tumor pathogenesis, the expression of E-cadherin/β-catenin complex should be considered a useful prognostic marker.

Actually, there is no consensus regarding the optimal treatment of ECM. In our case, surgical excision was done with the margin all around the tumor. No recurrences or metastasis were found 24 months after the surgery. The adjunctive radiation therapy is sometimes used, and in this case, the patient was submitted 30 sessions of radiotherapy, of 100 grays each, and she is making prosthodontic rehabilitation.

References

Epithelial-Myoepithelial Carcinoma