Pleomorphic adenoma with extensive squamous metaplasia and keratin cyst formations in minor salivary gland: a case report

Maria Carolina Vaz Goulart1, Patricia Freitas-Faria2, Gláuter Rodrigues Goulart3, Adriano Macedo de Oliveira4, Roman Carlos-Bregni5, Cleverson Teixeira Soares6, Vanessa Soares Lara7

1- DDS, MSc, PhD student, Department of Stomatology (Oral Pathology), Bauru School of Dentistry, University of São Paulo, Bauru, SP, Brazil.
2- DDS, PhD student, Department of Stomatology (Oral Pathology), Bauru School of Dentistry, University of São Paulo, Bauru, SP, Brazil.
3- DDS, Specialist in Oral and Maxillofacial Surgery and Traumatology, Private Practice, Piumhi, MG, Brazil.
4- MD, Specialist in Pathological Anatomy, José do Rosário Vellano University, Unifenas, Alfenas, MG, Brazil.
5- DDS, Director, Division of Pathology, Centro Clínico de Cabeza y Cuello, Guatemala City, Guatemala.
6- MD, Pathologist, Lauro de Souza Lima Institute and Institute of Anatomopathology of Bauru (ANATOMED), Bauru, SP, Brazil.
7- DDS, MSc, PhD, Department of Stomatology (Oral Pathology), Bauru School of Dentistry, University of São Paulo, Bauru, SP, Brazil.

Corresponding address: Maria Carolina Vaz Goulart - Faculdade de Odontologia de Bauru - Departamento de Estomatologia - USP - Alameda Octávio Pinheiro Brizolla 9-75 - Vila Universitária - Bauru-SP, Brasil, CEP: 17012-901 - Fone: +55-14-32358251 - Fax: +55-14-32234679 - e-mail: mariacarolinaodont@ yahoo.com.br

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ABSTRACT

Pleomorphic adenoma (PA), the most common salivary gland tumor, accounts for 54 to 65% of all salivary gland neoplasias and 80% of the benign salivary gland tumors. It most frequently affects the parotid gland, followed by the submandibular and the minor salivary glands. Microscopically, mucous, sebaceous, oncocytic and squamous metaplasia, sometimes with the formation of keratin pearls, may be present, but the latter rarely results in the formation of extensive keratin-filled cysts lined by squamous epithelium. Extensive squamous metaplasia can be mistaken for malignancy, including mucoepidermoid carcinoma and squamous cell carcinoma. Here, we present an unusual case of PA with extensive squamous metaplasia and keratin cyst formations in a minor salivary gland, and discuss its microscopic features, including the immunohistochemical characteristics, and differential diagnosis of this uncommon presentation.

Key words: Pleomorphic adenoma. Diagnosis. Salivary gland neoplasms.

INTRODUCTION

Pleomorphic adenoma (PA) affecting minor salivary glands presents a female predilection29. It occurs over a wide age range, but the mean age is 43.6 years and the peak incidence is between the fourth and fifth decades of life30. PA is clinically characterized by a slow growing, sessile-based, firm, painless mass, which occasionally has an ulcerated surface22. The most common site for minor salivary gland tumors is the palate4.

Microscopically PA exhibits a great diversity of morphological aspects20, showing varying combinations of epithelial and myoepithelial cells in a mesenchymal or stromal background. The duct-like formations exhibit ductal luminal cells in the inner layer and abluminal cells (myoepithelial cells and myoepithelial-like cells derived from them) in the outer layer. The capsule varies in thickness and presence, and many tumors show finger-like processes projecting into the capsule13. Squamous metaplasia, with the formation of keratin pearls may be present13, but rarely results in extensive keratin-filled cysts lined by squamous epithelium5,7,28. This microscopic finding may represent a diagnostic dilemma for pathologists.

Wide local excision is recommended as the treatment of choice11. Recurrence, which is estimated to occur in 5-30% of cases, is associated with incomplete surgical excision due to the presence of finger-like tumor extensions outside the capsule or satellite tumors13,22. PA presents a risk of malignant transformation, usually giving rise to a carcinoma in situ.
ex pleomorphic adenoma and the frequency of this transformation increases as a result of the tumor persisting without being treated. A close follow-up is necessary postoperatively.

Reports of PA presenting multiple and large squamous epithelium-lined keratin cysts are scarce. Here, we present an unusual case of PA with extensive squamous metaplasia and keratin cyst formations in a minor salivary gland. We discuss the microscopic features, including the immunohistochemical characteristics, and differential diagnosis of this unusual presentation.

CASE REPORT

A 37-year-old white man presented at the clinic with a 10-year history of a painless mass in the upper vestibule. There was no history of local surgery, trauma, or infection. On intraoral physical examination, there was a soft tissue mass in the right upper vestibule, near lateral incisor and canine teeth. The nodule was movable and firm on palpation. No abnormality was detected on radiographic examination.

A provisional diagnosis of benign salivary gland lesion was made. At surgery, the solid mass was encapsulated, well circumscribed and not adhered to the surrounding tissues, and was easily and completely excised. The specimen was submitted for microscopic examination.

Pathological features

Histochemistry

Hematoxylin and eosin (H&E)-stained sections revealed a well circumscribed encapsulated lesion composed of multiple squamous epithelium-lined cysts. The epithelium of the cystic spaces was covered by layers of para- and orthokeratotic cells, usually without a granular layer. Focally, the outer layer of epithelium demonstrated bud-like protrusions. Scattered rare mitotic figures were present but limited to the outer layer of the epithelium. The cystic spaces contained keratotic lamellae, which, in some areas, exhibited small calcification foci or cholesterol crystals. Although the cystic structures were variable in size and shape, most of them were large and round. Rare areas of mucous cells were observed in the cystic epithelial lining, which were confirmed by Periodic acid-Schiff (PAS) with or without diastase digestion and Alcian blue stains.

In other areas, tumoral epithelial cells formed solid sheets, nests, cords and ductal structures, the latter presenting a lumen lined by a double layer of cells and frequently containing eosinophilic material. Rare areas of mucous cells were observed in the cystic epithelial lining, which were confirmed by Periodic acid-Schiff (PAS) with or without diastase digestion and Alcian blue stains.

Immunohistochemistry

The reactions were performed using the streptavidin-biotin-peroxidase method. The primary antibodies, their clones and dilutions are summarized in Figure 3. Negative controls were obtained by the omission of primary antibodies, which were substituted by 1% PBS (phosphate buffered saline)-BSA (bovine serum albumin).

The squamous epithelium covering the keratin cysts was homogeneously positive for high-molecular-weight cytokeratins (Figure 2C). Only
in focal areas of the cystic epithelial lining did cells present immunostaining for CK7, CK19 and EMA in the inner layers, and p63 in the basal layer.

In the ductal structures, the luminal cells were identified by intense and frequent positivity for CK7 (Figures 2F and 4A), CK19 (Figure 4B), EMA and S-100, as well as focal immunostaining considered moderate for CK8 (Figure 4C) and intense for both polyclonal and monoclonal antibodies to CEA. Some abluminal cells were frequently positive for myoepithelial cell markers such as high-molecular-weight CKs (Figure 4D), p63 (Figure 4E), S-100 and vimentin (Figure 4F), and occasionally for SMA, MSA and GFAP. The luminal and abluminal cells, i.e. glandular phenotype-cells, were negative for CK20.

Solid sheets, nests and cords presented immunopositivity that was either focal for CK7, CK19, vimentin, GFAP and S-100, or frequent for high-molecular-weight CKs. Cells in the basal layer of the epithelial cords were positive for p63.

The epithelium lining the cysts and the solid areas were surrounded by collagen IV-positive material. The proliferative index of the lesion, expressed as the percentage of Ki-67-positive neoplastic cells among the total number of cells counted in 10 randomly selected fields, was low (2.63%) and proliferation was mainly localized in the outer layers of the epithelial lining of the large keratin cysts. Expression of p53 was observed in rare neoplastic cells, throughout the lesion.

The mesenchymal areas presented S-100- and vimentin-positive cells and the blood vessels were collagen IV-, MSA-, SMA- and vimentin-positive. Most inflammatory cells were Ki-67 positive. The immunohistochemical features of the lesion are summarized in Figure 5.

Considering the morphological and

| Antibody Specificity | Clone     | Dilution |
|----------------------|-----------|----------|
| CK7*                 | OV-TL12/30| 1:200    |
| CK8*                 | 35BH11    | 1:150    |
| CK19**               | B170      | 1:100    |
| CK20*                | KS20.8    | 1:50     |
| Hmw CKs*             | 34BE12    | 1:200    |
| EMA*                 | E29       | 1:200    |
| CEA*                 | 11-7      | 1:200    |
| CEA*                 | Polyclonal| 1:400    |
| GFAP*                | 6F2       | 1:100    |
| S-100*               | COW S-100 | 1:1400   |
| SMA*                 | 1A4       | 1:200    |
| MSA*                 | HHF35     | 1:5000   |
| p63*                 | 4A4       | 1:150    |
| Vimentin*            | V9        | 1:400    |
| Collagen IV*         | CIV22     | 1:80     |
| p53*                 | DO-7      | 1:50     |
| Ki-67*               | KI-55     | 1:100    |

CK (cytokeratin), Hmw (high-molecular-weight), EMA (epithelial membrane antigen), CEA (carcinoembryonic antigen), GFAP (glial fibrillary acidic protein), SMA (smooth muscle actin), MSA (muscle specific actin)

Source: * Dako, Carpinteria, CA, USA
** Novocastra Laboratories Ltd, Newcastle, UK

Figure 3—List of antibodies used for immunohistochemical staining
Antibody Specificity

| Antibody Specificity | Cystic lining | Luminal cells | Abluminal Cells | Solid areas | Mesenchymal cells |
|----------------------|--------------|---------------|----------------|-------------|------------------|
| CK7                  | +++          | +++           | -              | +++         | -                |
| CK8                  | -            | ++            | -              | -           | -                |
| CK19                 | +++          | +++           | -              | +++         | -                |
| CK20                 | -            | -             | -              | -           | -                |
| Hmw CKs              | +++          | -             | ++            | +++         | -                |
| EMA                  | +++          | +++           | -              | -           | -                |
| CEA(mono)            | -            | +++           | -              | -           | -                |
| CEA (poly)           | -            | +++           | -              | -           | -                |
| GFAP                 | -            | -             | +++           | +++         | -                |
| S-100                | -            | +++           | ++            | +++         | +++              |
| SMA                  | -            | -             | +             | -           | -                |
| MSA                  | -            | -             | +             | -           | -                |
| p63                  | +++          | -             | ++            | +++         | -                |
| Vimentin             | -            | -             | ++            | +++         | +++              |

Immunostaining was graded as (−) negative; (+) weakly positive; (++) moderately positive; (+++) strongly positive.

Figure 4- Immunoprofile of duct-like structures. Luminal cells intensely positive for cytokeratin (CK) 7 (A) and CK19 (B) and moderately positive for CK8 (C). Abluminal cells moderately positive for high-molecular-weight (CKs) (D), p63 (E) and vimentin (F). Immunoperoxidase stain.

Figure 5- Immunohistochemical features of the neoplastic cells.

Figure 6- Immunohistochemical features of the lesions included in the differential diagnosis of PA with extensive squamous metaplasia.

PA (Pleomorphic adenoma), MEC (Mucoepidermoid carcinoma), SCC (Squamous cell carcinoma), KC (Keratocystoma), Vim (Vimentin)

(−) negative; (+) positive; (a) variable expression; (nf) data not found.
immunohistochemical findings a final diagnosis of PA with extensive squamous metaplasia was made. The postoperative course was uneventful and the patient has been reviewed regularly, with no evidence of recurrence 3 years after excision.

DISCUSSION

PA presenting extensive squamous metaplasia is uncommon and can signify a potential pitfall in the histopathological diagnosis.

Focal squamous metaplasia, probably related to ischemia, may be present in about 25% of the PA. The cases of PA presenting extensive keratin-filled cysts lined by squamous epithelium are referred to in the literature as “cystic PA with extensive adnexa-like differentiation”, since the histologic features mimic cutaneous appendages.

In our case, many glandular cells were transformed into squamous cells through a process of squamous metaplasia, resulting in multiple squamous epithelium-lined cysts containing keratotic lamellae and some solid squamous cell islands presenting keratin pearls. The epithelium lining the keratin cyst formations was homogeneously positive for a set of high-molecular-weight CKs, commonly expressed in squamous epithelium, while some cells presented immunostaining for p63, EMA and low-molecular-weight CKs (CK7 and CK19), representing glandular cells which probably had not yet undergone squamous metaplasia.

Extensive squamous metaplasia in PA, especially in the absence of chondromyxoid stroma, can mistakenly lead to a diagnosis of benignity, such as choristoma or keratocystoma, and malignancy, including mucoepidermoid carcinoma (MEC) and squamous cell carcinoma, especially when sampling the tissue by fine-needle aspiration biopsy (FNAB) and incisional biopsy, due to limited and selective sampling. It is important to discuss the diagnostic pitfalls of this unusual presentation of a common benign entity, particularly the differential diagnosis with these malignant lesions.

Microscopically, MEC presents mucous, intermediate and squamous (epidermoid) cells and is usually multicystic. Unlike our case, the cystic spaces of MEC are usually lined by mucous cells and prominent keratinization is rare, with scarce epidermoid cells associated with keratin production including keratin pearl formation. Furthermore, the presence of a fibrous capsule in our case favors the diagnosis of a benign lesion. In addition to the above mentioned morphological findings, the immunopositivity for S-100, SMA and vimentin found in the reported lesion could help to exclude the diagnosis of MEC.

Regarding the differential diagnosis with squamous cell carcinoma, the absence of cytological atypia, metastasis, necrosis, invasion, as well as minimal cellular proliferative activity and the presence of a fibrous capsule in the case presented here weigh against the diagnosis of malignancy. Moreover, the neoplastic cells of CEC, differently from PA, are immunonegative for CK7, S100, SMA and vimentin.

A choristoma resembling a trichoadenoma (a type of hair follicle tumor), described in the parotid gland by Seifert, Donath and Jautzke (1999), also presents multiple cystic spaces limited by a multilayered squamous epithelium and filled with keratotic lamellae. It also exhibits solid squamous cell islands, keratinized masses outside the cysts with multinucleated giant cells and focal calcification. Nagao, et al. (2002) described two cases histologically identical to the choristoma previously reported, both of the parotid gland, but the authors proposed a new designation for the lesions: keratocystoma, since they believe that the choristoma described by Seifert, Donath and Jautzke (1999) is a peculiar variety of salivary gland tumor. Although our case presents similar morphological aspects to those described for choristoma or keratocystoma, these lesions are mainly composed of squamous cells and lack the neoplastic structures with glandular phenotype, including duct-like formations and myoepithelial cells, which were demonstrated in our case by means of morphology and immunohistochemistry. Moreover, the cystic spaces of the keratocystoma are lined by a stratified squamous epithelium without a granular cell layer, unlike our case, which showed some areas of granular layers. Similar to our findings, the squamous cells lining the cystic formations were negative for CK8, focally immunoreactive for CK19, and surrounded by collagen-IV material. Based on the morphological aspects, we are inclined to believe that the PA reported here and the keratocystoma may constitute related lesions, representing different stages in the evolution of a specific type of salivary gland tumor, although the latter previously demonstrated immunonegativity for S100 and SMA. Immunohistochemical features of the lesions included in the differential diagnosis of PA with extensive squamous metaplasia are included in Figure 6.

The immunohistochemical analysis of the current case helped to define the nature of the tumor cells, adding support to the diagnosis of PA. It identified the luminal cells of the duct-like structures by immunopositivity for low-molecular-weight CKs, including CK7, CK8 and CK19, as well as CEA and EMA, whereas the abluminal cells were identified by heterogeneous positivity for myoepithelial cell markers such as p63, high-molecular-weight CKs, S-100, vimentin, SMA, MSA and GFAP.
immunoprofile of the luminal and abluminal cells of the duct-like formations observed is similar to those described in other articles in the literature\(^6\)\(^,\)\(^14\). In addition to the abluminal cells, the luminal cells were also positive for S-100, which is in agreement with the findings of other authors\(^4\).

The solid sheets, nests and cords presented immunoreactivity either for CK7 and CK19, suggesting a luminal cell phenotype, or for vimentin, GFAP and p63, in this case demonstrating a myoepithelial cell phenotype. Although not accompanied by myxochondromatous elements, the glandular phenotypic profile associated with the morphological features reinforces the diagnosis of PA. Degrees of stromal changes, such as chondroid, osseous, myxoid or mucoid changes vary widely in PA\(^6\),\(^13\), however the absence of these changes, as in our case, is an unusual finding.

Ki-67 expression was sporadic (2.63\%), which is consistent with the literature data describing PA as having a low proliferative rate\(^2\),\(^27\). Interestingly, the epithelial lining of the large keratin filled cysts presented a higher proliferative index than the other areas. It may signify that the squamous metaplasia resulting in large keratin filled cysts in PA may be clinically significant, probably related to an important growth potential, although our data are insufficient to support this hypothesis. Further studies are required to clearly establish this relationship. Our case, as also indicated in previous reports\(^1\)-\(^2\), presented scarce p53-positive cells, which are associated with changes in the apoptotic mechanism.

CONCLUSION

In conclusion, the localization, gender and microscopic features of the presented case are unusual. PA with extensive squamous metaplasia and keratin cyst formations can pose a significant diagnostic challenge. It is important to be aware of this possibility to distinguish it from malignant lesions and to avoid unnecessarily aggressive therapy. The morphological features, as discussed above, must be carefully evaluated in order to exclude other lesions and to define the diagnosis of PA. Additionally, the immunohistochemical analysis, in addition to establishing the nature of the tumor cells, may also be helpful for the differential diagnosis, reinforcing the morphological diagnosis, especially when PA has an unusual presentation.

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