

**Squamous cell carcinoma ex pleomorphic adenoma; a case report**

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**Abstract**

A rare case of squamous cell carcinoma ex pleomorphic adenoma in maxillary sinus is presented. The patient is a 50-year-old woman presenting with a slow-progressive swelling in left side of her face that she has noticed 2 years earlier. The lesion was not painful and caused asymmetry and mild exophthalmos. Microscopic examination revealed that the tumor was composed of two components; partly of a pleomorphic adenoma and partly of a squamous cell carcinoma. Immunohistochemical examination for Ki-67 and SMA and mucicarmine staining were also done and confirmed the diagnosis of carcinoma ex pleomorphic adenoma. There is no evidence of recurrence 12 months after operation.

**Keywords:** Pleomorphic adenoma, Malignant transformation, Squamous cell carcinoma, Case report

**Introduction**

Among different salivary gland tumors, carcinoma ex pleomorphic adenoma (Ca ex PA) accounts for approximately 3.6% (1). It constitutes 6.2% of all pleomorphic adenoma (PA) and 11.7% of all malignant salivary gland neoplasms (1, 2). Furthermore, the risk of malignant transformation in PAs in the first five years after evolution of the lesion is 1.6% (3). Car ex PA is defined as an epithelial malignant transformation within a primary or previous PA, and is often considered as a diagnostic challenge for the pathologists and clinicians, since it is difficult to

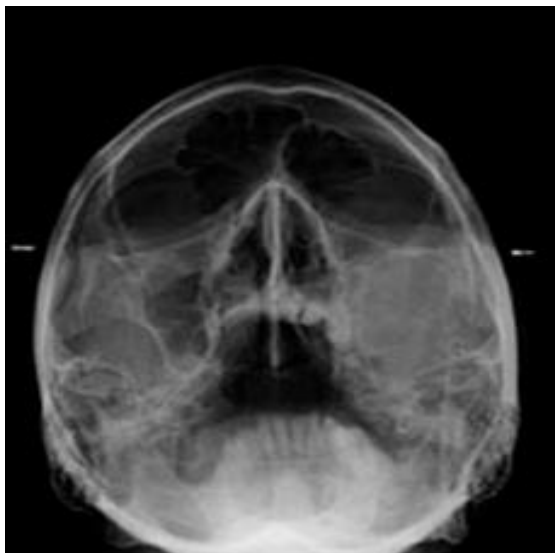
diagnose this entity pre-operatively due to unspecific clinical presentation [4, 5]. In fact, most of Ca ex PA tumors show clinical presentation similar to PA. It means that they can be asymptomatic and not invasive in gross examination. However, early diagnosis and accurate surgical intervention is important to increase the patients' survival rates because of aggressiveness and destructive behavior of this tumor (1-6). WHO states that the malignant component of Ca ex PA is mostly adenocarcinoma, not otherwise specified (NOS), and sometimes salivary

duct carcinoma (SDC), undifferentiated carcinoma, adenoid cystic carcinoma, or mucoepidermoid carcinoma (7).

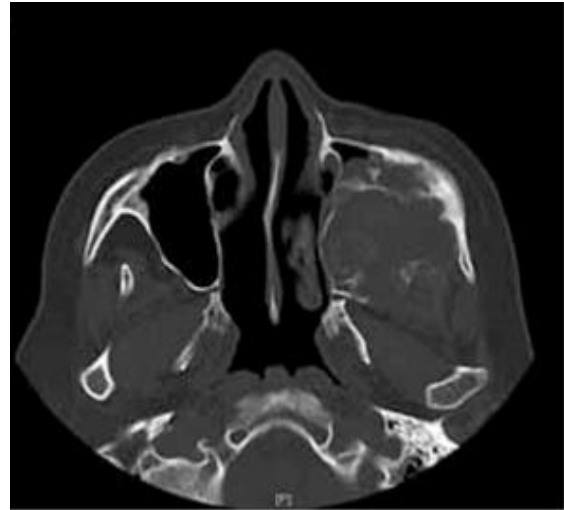
We discuss the presentation, histopathologic picture, and management of a rare case of squamous cell carcinoma (SCC) ex pleomorphic adenoma originating in the maxillary sinus.

### Case report

A 50-year-old woman presented to our clinic with a chief complaint of asymmetry due to a long-standing swelling on the left side of her face which had first appeared 2 years earlier. There were no other symptoms such as pain and facial paralysis. The swelling was bony hard and non-tender in palpation. There was also a mild exophthalmos without pain and visual disturbance. Clinical examination of cervical lymph nodes showed no abnormal findings. Waters radiograph revealed a space-occupying destructive mass filling the left antrum completely. Computed tomography scans depicted a large ill-defined mass in left maxillary sinus causing destruction of medial, infraorbital and floor of the sinus with extension to the left orbital region (Figures 1, 2 and 3).



**Figure 1.** Waters view. Complete opacification of left maxillary sinus.

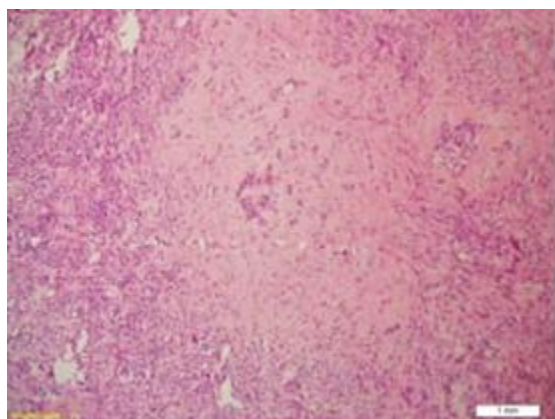


**Figure 2.** Computed tomography. Axial view; a destructive lesion in left antrum.

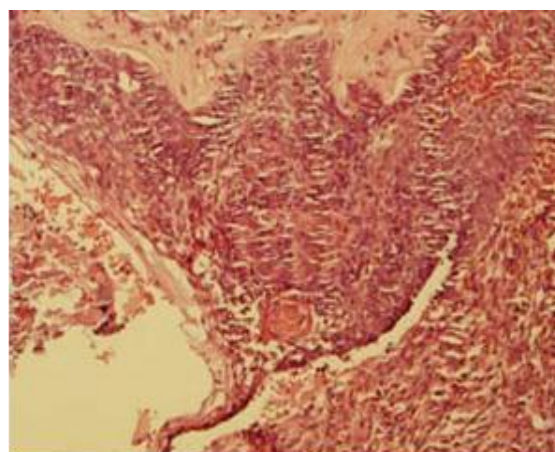


**Figure 3.** Computed tomography. Coronal view; distruction of palate, lateral nasal wall and maxillary sinus.

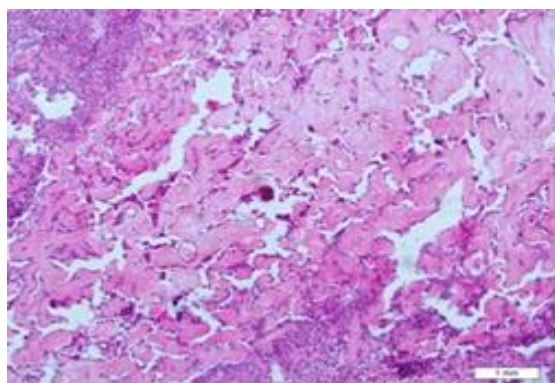
Incisional biopsy of the lesion was performed and histopathological examination revealed a case of pleomorphic adenoma in a hyalinizing fibrotic and chondromyxoid stroma with the malignant areas of carcinomatous component consisting dysplastic squamous cells which exhibited large nuclei with pleomorphic shapes, prominent nucleoli and few abnormal mitotic figures. Focal areas of calcification were present. Moreover, clear cells arranged in small nests or as individual cells intermixed with squamous cells were also detected (Figures 4, 5 and 6).



**Figure 4.** Histopathologic features of PA component of tumor (Hematoxylin-Eosin, original magnification  $\times 100$ ).



**Figure 5.** Squamous cell carcinoma cell component of the tumor (Hematoxylin-Eosin, original magnification  $\times 100$ ).



**Figure 6.** Extensive hyalinization and areas of calcification close to the malignant cells (Hematoxylin-Eosin, original magnification  $\times 100$ ).

Mucicarmin staining was done and all clear cells were negative, therefore, the

diagnosis of mucoepidermoid carcinoma was ruled out for the malignant component of the tumor. Immunohistochemical staining was positive for ki-67 in more than 60% of squamous cells whereas pleomorphic adenoma component was negative for this marker. Additional immunohistochemical staining was also conducted. The clear cells in malignant component were negative for SMA, but clear cells of PA showed reactivity for SMA which proved the myoepithelial origin of these cells. These findings were consistent with the diagnosis of squamous cell carcinoma ex pleomorphic adenoma for this case.

### Discussion

Based on World Health Organization classification in 2005, Ca ex PA is the most common malignant change in PA and accounts for 11.7% of salivary malignancies (3, 8). Malignant mixed tumors are divided into three categories: 1. carcinoma ex pleomorphic adenoma, 2. Carcinosarcoma and 3. metastasizing mixed tumor. (9)

Although PA mostly arise in the parotid gland or minor salivary glands, it is also the most frequent tumor of the seromucinous glands of other organs such as nasal cavity, sinuses and trachea (10). Any type of carcinoma can arise in a PA, but adenocarcinoma NOS and salivary duct carcinoma are the most common types. (4, 5, 11). The frequency of malignant components of Ca ex-PA is 43–90% in adenocarcinoma, not otherwise specified (NOS), 10–34% in SDC, and rare in mucoepidermoid carcinoma and SCC (9, 11). In sinonasal tract, Ca ex PA is considered as the least common malignancy (12).

Generally, there is a history of a painless and slow-growing mass which start to enlarge rapidly over a short period of time (13). Patients usually present with signs and symptoms indicating malignant transformation (e.g., facial nerve palsy, pain, skin perforation and trismus).

Approximately 23% to 40% of cases shows facial nerve weakness or palsy. (14) Cellular pleomorphism, cellular anaplasia, atypical mitosis, destruction of normal tissues, invasiveness and abnormal architectural patterns are defined by Gnepp and Wenig as the criteria of malignancy in a PA. (15). Gerugthy et al. suggested that microscopic evidence of an invasive growth pattern, neural or vascular invasion, focal areas of calcification and necrosis are related to a poor prognosis. (16) In agreement with Gnepp and Wenig and according to the criteria proposed by Gerugthy, this patient is definitely a case of carcinomatous transformation in the PA.

Several cases of Ca ex PA have been described in previous reports, (17-21) but SCC ex pleomorphic adenoma is appear to be rare as mentioned by Peel and Gnepp (22). Seifert et al. recognized SCC components in 4 out of 38 (10%) Ca-ex-PA patients (23). In contrast, Lewis et al. (11) found no cases with SCC components among 73 patients, nor did Matsubayashi et al. among 1010 cases. (9). In addition, in 37 cases of Car ex PA reported by Tortoledo et al. (24), no malignant components were diagnosed as SCC. A case of Ca ex PA with the presence of SCC and salivary duct carcinoma was reported by Nakamori et al. (25). Therefore, SCC as the malignant component of Car ex PA seems to be rare (25-28).

In differential diagnosis, it is important for the pathologists to distinguish SCC from squamous cell metaplasia, as well as SCC from mucoepidermoid carcinoma (29, 30). Ki-67 staining reported by Zhu et al. as a useful diagnostic marker to discriminate between malignant and benign tumors of salivary gland (31). In the current patient, the immunostaining of Ki-67 helped us to differential SCC from squamous cell metaplasia. Mucicarmin staining was also used to detect mucin cells in order to distinguish SCC from mucoepidermoid carcinoma.

In the present case, partial maxillectomy as well as left eye exenteration was done. As achieving a complete resection with wide margins in the head and neck region is not always possible, in order to reduce the probability of recurrence, the primary site received external irradiation postoperatively. The patient was then referred to a prosthodontist to fabricate maxillary and midface prosthesis. There was no evidence of recurrence 12 months after operation.

### Conclusion

This is a rare case of carcinoma ex pleomorphic adenoma of the antrum with an unusual malignant component of squamous cell carcinoma. The SCC component has originated from the epithelial part of PA.

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