Basal Cell Adenoma in the Parotid Gland: A Case Report of Rare Salivary Gland Tumor

Chonji Fukumoto*, Kazuya Hiroshima, Toshihide Watanabe

Department of Dental and Oral Surgery, Kimitsu Chuo Hospital, 1010 Sakurai, Kisarazu, Chiba 292-8535, Japan

Abstract

Salivary gland tumors are rare and constitute about 2 to 6.5% of all head and neck neoplasms. Basal cell adenoma (BCA) is a rare benign tumor in salivary gland, accounting for 1 to 3% of all salivary gland tumors. Histopathologically, BCA is composed of basaloid cells and can be classified into solid with most common type, trabecular, tubular, and membranous types. It is mainly found in the parotid gland, and it is commonly treated via a parotidectomy. In this study, we report a case of surgical excision trabecular-tubular mixed type BCA in the surface of the parotid gland, with preservation of the facial nerve and body of the parotid gland.

Keywords
Salivary Gland Tumor, Basal Cell Adenoma, Parotid Gland, Facial Nerve Preservation

1. Introduction

Salivary gland tumors are uncommon and constitute about 2 to 6.5% of all head and neck neoplasms. Basal cell adenoma (BCA), a rare benign tumor in salivary gland, accounts for 1 to 3% of all salivary gland tumors [1-3]. BCA was first reported by Kleinsasser and Klein [4] in 1967 and was later described as a separate salivary gland tumor in the 1991 WHO classification. BCA has been classified into four histological types: solid, trabecular, tubular, and membranous types [3,5]. Its most frequent location is the parotid gland, and a parotidectomy is commonly performed. Here, we report a case of surgical excision trabecular-tubular mixed type BCA in the surface of the parotid gland, with preservation of the facial nerve and body of the parotid gland.

2. Case Presentation

A 44-year-old woman presented to the Division of Dental and Oral Surgery, Kimitsu Chuo Hospital, with a history of painless swelling in her right cheek. The mass slowly increased in size and became asymptomatic. Facial examination revealed a well-circumscribed elastic, hard, and movable mass in the right malar region measuring approximately 25 mm in diameter (Figure 1). There was no palpable lymph node in the neck region, and facial nerve function was intact. She had no relevant medical history.

Figure 1. Facial preoperative view
On further diagnosis, the Gd-DTPA-enhanced magnetic resonance imaging (MRI) showed a well-defined round mass with moderate signal intensity on the T1-weighted image (Figure 2A) and high signal intensity on fat suppression T1-weighted image (Figure 2B) in the superficial part of right parotid. The clinical diagnosis was suspected to be a benign tumor arising in the superficial portion of the parotid gland.

An excisional biopsy was performed under general anesthesia for further confirmation. The tumor was surrounded by a thin capsule and was removed with preservation of the facial nerve and no excision of the body of the parotid gland (Figure 3). The enucleation was relatively easy due to the lack of adhesions between the tumor and the parotid gland. Macroscopically, the spherical surgical specimen was measured to be $13 \times 13 \times 12$ mm and encapsulated by a thin fibrous tissue (Figure 4).

**Figure 2.** Gd-DTPA enhanced MRI sequence (A) T1-weighted, (B) fat suppression T1-weighted

**Figure 3.** Intra-operative appearance of tumor mass

**Figure 4.** Surgical specimen
The histopathological examination of the section stained with hematoxylin & eosin showed the proliferation of basaloid cells with trabecular and tubular patterns and a peripheral layer of cells (Figure 5). No mitotic activity, necrosis, or nuclear pleomorphism was noticed in the tumor. Immunohistochemically, the inner cells of tubular structures were strongly positive for Cytokeratin (CK) AE1/AE3 (Figure 6A), and the nuclear of the outer cells were positive for p63 (Figure 6B). The MIB-1 (Ki-67) labeling index was extremely low, i.e., 2.5% (Figure 6C). On the basis of these features, the final diagnosis was confirmed as trabecular-tubular mixed type BCA.

3. Discussion

Previously, BCA was considered to be a type of “monomorphic adenoma” [4]. However, the name of this tumor has been changed to “BCA” in the WHO salivary gland tumor classification since 1991. The most frequent location of BCA is the parotid gland (75%), followed by the minor salivary gland from the upper lip with 6%, and submandibular gland with 5% [6-8]. Clinically, BCA frequently occurs in people aged more than 50 years as a unilateral tumor with round or oval swelling. This tumor tends to be an asymptomatic, slowly growing and movable mass with a maximum diameter of 3 cm [9,10].

Histopathologically, BCA is composed of basaloid cells, classified based on their morphologic pattern into four subtypes: solid with most common type, trabecular, tubular, and membranous [5]. The recurrence and malignant transformation are more common in the membranous type than in other subtypes [9,11,12]. In the present case, the tumor was histopathologically classified as tubular-trabecular mixed type BCA.

The differential diagnoses of BCA are pleomorphic
adenoma (PA), adenoid cystic carcinoma (ACC), and basaloid cell adenocarcinoma (BCAC). BCA has no rich stroma and the myxoid or chondroid elements as characteristics of PA [8,9,13]. Although ACC and BCAC have clinical and histological similarities with BCA, these malignant tumors exhibit invasive, unencapsulated growth into adjacent soft tissue, often with associated vascular or perineural invasion [7-9]. In addition to cytology differences, the high MIB-1 (Ki-67) labeling index helps distinguish ACC and BCAC from BCA [8].

The primary treatment of BCA is surgical excision by means of a parotidectomy. In the present case, we performed local excision because the tumor existed in the superficial portion of the parotid gland and was easily detached from the parotid gland. The patient’s condition was well without facial paralysis and free of disease at 3 years after the surgery.

4. Conclusions

In conclusion, we report a case of surgical excision trabecular-tubular mixed type BCA in the surface of the parotid gland, with preservation of the facial nerve and body of the parotid gland. Though regarded as rare, BCA should be kept in mind in the management of salivary gland tumors.

Conflict of Interest

The authors declare that there is no conflict of interests regarding the publication of this paper.

REFERENCES


