Basal cell adenoma misdiagnosed as an adenoid cystic carcinoma in the parotid gland

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Basal cell adenoma (BCA) of the parotid gland is a rare benign tumor. In the parotid gland, BCA is occasionally difficult distinguish from adenoid cystic carcinoma in terms of clinical and pathological perspectives. An adenoid cystic carcinoma of the parotid gland grows slowly but spreads persistently to the surrounding tissues, particularly along the perineural spaces. In the present case, BCA of the parotid gland was misdiagnosed as an adenoid cystic carcinoma. We discuss the reason for such a misdiagnosis, and present a method for making a correct diagnosis.

Key words: Adenoma, Parotid gland

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I. Introduction

Basal cell adenoma (BCA) of the salivary glands is a rare benign neoplasm having a monomorphous histological appearance dominated by basaloid cells¹. The reported data state the incidence of BCA in all salivary neoplasms to be 1-3%^{2,3}. It appears most frequently in the parotid gland in adults^{1,4}. Clinically, it is generally a slow-growing, asymptomatic, freely movable mass^{1,4}.

As one of the most common and best recognized malignant salivary tumors⁵, adenoid cystic carcinoma (ACC) was first reported in 1853 by Robin, Lorain, and Laboulbene⁶. ACC was originally called cylindroma because of its histopathological morphology. ACC was recorded to be located in the major and minor salivary glands; usually small with an incomplete capsule, it has a propensity toward perineural spread⁵. It has high, almost inevitable predisposition to recur in a person with old age, occurring

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even when radical excision has been performed⁷.

Clinically and histopathologically, there are similar features between BCA and ACC. The treatment plan is completely changed by the result of diagnosis, especially when malignant or benign; hence the need for different diagnoses. In this paper, we report BCA misdiagnosed as ACC in the parotid gland, review literature, and discuss the different diagnoses of similar cases.

II. Case Report

In December 2010, a 50-year-old man was referred to our hospital for evaluation of a palpable mass in the left parotid region. He complained of severe burning and pulling sensation in the left pre-auricular area. The painful sensation began about 2 years ago, continuing intermittently but not worsening. The physical examination revealed a mass measuring about 5×4 cm, which was hard, tender, and movable. (Fig. 1) There was no symptom on facial nerve function and cervical lymphadenopathy. The magnetic resonance image showed a well-defined, non-homogeneously enhanced mass on the deep portion of the left parotid gland. (Fig. 2) The T1W1 image revealed moderate to low signal intensity (Fig. 2. A), but the T2W1 image showed higher signal intensity portion in the middle and medial site of the lesion suspected to be a necrotic lesion. (Fig. 2. B) There was

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no evidence of infiltrating margin.

Partial parotidectomy was performed. We approached the mass by pre-auricular incision.(Fig. 3. A) The superficial parotid layer of the parotid gland was removed, and the mass was excised.(Fig. 3. B) The facial nerve trunk was conserved. (Fig. 3. C)

Macroscopically, an encapsulated whitish lesion measuring 4×4×2 cm was observed. The microscopic examination showed infiltrating epithelial strands with multiple cystic changes and solid pattern. The tumor cells rarely showed keratinization and consisted almost exclusively of intermediate cell type. The tumor cells were small and cuboidal, exhibiting deeply basophilic nuclei and little cytoplasm; mitotic activity was rarely seen. There was no perineural invasion (Fig. 4), however. Immunohistochemical examination was done as well. The immunostain of



Fig. 1. Left preauricular area swelling and painless movable mass. Chan-Woo Kim et al: Basal cell adenoma misdiagnosed as an adenoid cystic carcinoma in the parotid gland. J Korean Assoc Oral Maxillofac Surg 2012

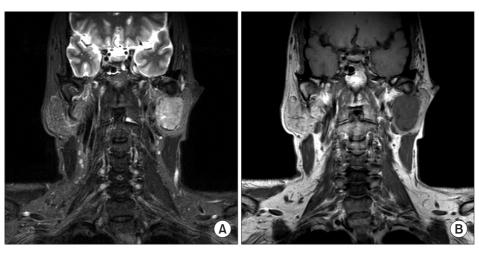


Fig. 2. A. In T1W1 magnetic resonance imaging, there is a well-defined and non-homogeneously enhanced mass on the deep portion of left parotid gland. B. There is higher signal intensity portion in middle and medial site of lesion.

Chan-Woo Kim et al: Basal cell adenoma misdiagnosed as an adenoid cystic carcinoma in the parotid gland. J Korean Assoc Oral Maxillofac Surg 2012



Fig. 3. A. Pre-auricular incision was done. B. Superficial parotid gland was removed and the mass was excised. C. Facial nerve trunk was conserved.

Chan-Woo Kim et al: Basal cell adenoma misdiagnosed as an adenoid cystic carcinoma in the parotid gland. J Korean Assoc Oral Maxillofac Surg 2012

proliferating cell nuclear antigen was frequently positive in tumor cells, but that of p53 was hardly visible. The immunostain of cytokeratin-7 was frequently positive in glandular structure, but the same cannot be said for cytokeratin-14. The immune stain of snail was frequently positive in tumor cells, but that of beta-catenin was hardly visible. Pan-K and S-100 were positively stained, but wnt-1

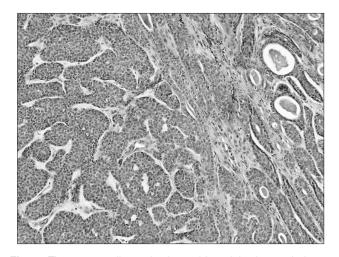


Fig. 4. The tumor cells rarely showed keratinization and almost consisted of intermediate cell type. There were small and cuboidal, exhibiting deeply basophilic nuclei and little cytoplasm and mitotic activity was rarely seen (H&E staining, x100). Chan-Woo Kim et al: Basal cell adenoma misdiagnosed as an adenoid cystic carcinoma

in the parotid gland. J Korean Assoc Oral Maxillofac Surg 2012

was negative. (Fig. 5) Initially, the pathologist considered the lesion to be ACC.

Nonetheless, we had doubts on the pathological diagnosis because it was not in accord with the patient's clinical symptoms and radiological feature. Thus, we requested a reexamination on the section to another clinical pathologist. The final diagnosis was BCA, not ACC. The cells of the islands were palisaded and cuboidal in shape, with the trabecular subtype demonstrating narrow, cordlike epithelial strands. There was no evidence of malignancy.

There was slight facial weakness in one third of the lower left part of the face. The symptom persisted for 6 months but eased gradually. The patient was monitored for 1 year, and there was no evidence of the tumor or signs and symptoms recurring.

Discussion

BCA in the salivary gland is a rare benign neoplasm, consisting of isomorphic basaloid tumor cells^{4,8}. Constituting only 1% of all salivary neoplasms^{2,9}, it is controversial for its gender predominance⁸. The tumor can grow at any age but is most common among old-age adults⁸. BCA occurs in epithelial cells, usually in the terminal duct⁸. Histologically, it has many variants such as solid, tubular, trabecular, and

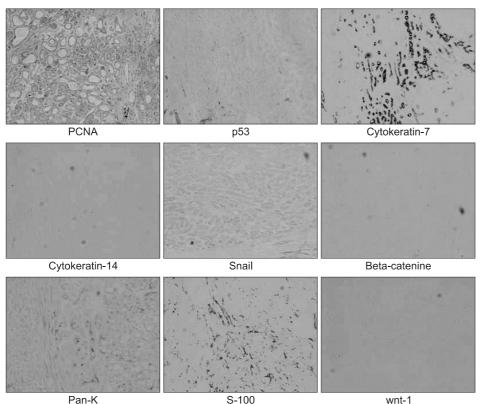


Fig. 5. PCNA was frequently positive in tumor cells, but the immunostain of p53 was rarely stained. Cytokeratin-7 was frequently positive in glandular structure, but the cytokeratin-14 was rarely stained. Snail was frequently positive in tumor cells, but the beta-catenine was rarely stained. Pan-K and S-100 was positively stained, but wnt-1 was negative (x100). (PCNA: proliferating cell nuclear antigen, Pan-K: pancytokeratin)

Chan-Woo Kim et al: Basal cell adenoma misdiagnosed as an adenoid cystic carcinoma in the parotid gland. J Korean Assoc Oral Maxillofac Surg 2012 membranous¹⁰. The most common variant is the solid type, but each tumor has combination-type variants¹⁰. BCA consists of 2 types of cells¹⁰: one is a small cell with insufficient cytoplasm and round-shaped nucleus¹⁰, and the other type has large eosinophilic cytoplasm and ovoid-shaped nuclei¹⁰.

There are some benign and malignant neoplasms that must be differentially diagnosed with BCA. Pleomorphic adenoma is clinically similar to BCA. It appears as a slowly growing, freely movable mass¹, typically appearing as a painless, firm mass¹⁰. Pleomorphic adenoma of the parotid gland mostly occurs in the superficial lobe, manifesting swelling on the preauricular area¹⁰. Accounting for 53-77% of parotid tumors¹⁰, pleomorphic adenoma originated with a mixture of ductal and myoepithelial elements¹⁰. In contrast, the basic tumor pattern is very variable, but the individual cells are rarely pleomorphic¹⁰. When we evaluated the patient clinically, we easily supposed that the lesion to be pleomorphic adenoma because it is the most common benign tumor in the parotid gland.

ACC accounts for 10% of all tumors in salivary glands^{5,11}. The parotid gland area is the most common site in the head and neck⁵. Histopathologically, ACC can be classified into three morphological patterns including cribriform, tubular, and solid⁵. The most important and unique feature of ACC is the tendency of perineural invasion, even in earlystage tumors^{5,10}. In most cases, the cytological typing of ACC is distinguished by the detection of large globules of extracellular matrix surrounding the basaloid tumor cells⁵. ACC shows the most histological similarities to BCA, since both have the same developmental origin¹⁰. Note, however, that their characters are very much different in terms of the integrity of the basal layer, number of mitoses, and growth speed. In the parotid gland, ACC is rare, accounting for only 2-3% of all tumors 10,11. There is fairly equal gender distribution, although some studies have shown slight female predilection¹⁰.

In our case, the lesion was growing slowly, and it was separated with adjacent normal tissue. The mass was firm,

painless, and freely movable. The patient did not complain about his lesion until 2 years ago. He had no facial palsy and lymphadenopathy. Therefore, clinically, it had a benign character, so we believed it to be a benign tumor. Nonetheless, the histopathological result was ACC, although we could not believe the result. We requested for the examination of the sample to another pathologist, expecting a different result. True enough, it was BCA; we accepted the result because of many clinical similar evidences.

It is difficult to diagnose basal cell adenoma in the parotid gland, since it is uncommon in the parotid gland and it seems to be just another tumor. The surgeon must consider the patient's clinical symptom, radiological symptom, and pathological symptom.

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