Research Paper

Notch as a Possible Cell Differentiation Factor in Pleomorphic Adenomas

Keisuke Takamine1, Yukiko Ueda1, Keisuke Nakano1, 2, Takanaga Ochiai1, Yoshihiko Sugita3, Katsutoshi Kubo3, Hatsuhiko Maeda3, Hiromasa Hasegawa3 and Toshiyuki Kawakami1

1. Hard Tissue Pathology Unit, Matsumoto Dental University Graduate School of Oral Medicine, Shiojiri, Japan
2. Department of Oral Pathology, Okayama University Graduate School of Medicine, Dentistry and Pharmaceutical Sciences, Okayama, Japan
3. Department of Oral Pathology, School of Dentistry, Aichi Gakuin University, Nagoya, Japan

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Abstract
The expression of Notch in 30 cases of pleomorphic adenoma was examined by immunohistochemistry. Comparing the results of our study with previous literatures, from the partial CK7 expression and substantial Notch expression in ductal epithelial cells as well as the Notch expression in solid tumor nests, it can be inferred that Notch is involved in cell differentiation. CK13 expression was observed in cells undergoing squamous metaplasia and Notch expression was seen in the nucleus of basal and squamous cells. The intense Notch expression in basal cells and weak expression in squamous cells suggests that Notch is involved in the differentiation from basal to squamous cell. Moreover, the loss of nuclear expression on the inner layer would signify that differentiation is about to end or has been terminated. Notch was expressed in the cytoplasm of cartilage cells and in the cell membrane of mucous cells but not in the nucleus indicating that differentiation has been concluded. Notch involvement is suspected in cell differentiation in areas showing ductal structures and squamous metaplasia. In summary, Notch is involved in cell differentiation of ductal cells in PA. Nuclear expression was shown in tumor cells in solid nests and surrounding structures. Moreover, Notch is expressed by basal cells undergoing squamous metaplasia suggesting the participation of Notch in cell differentiation in PA.

Key words: Notch1, pleomorphic adenoma, cell differentiation, immunohistochemistry

Introduction
Pleomorphic adenoma (PA) is the most common salivary gland tumor classified as benign epithelial tumors (1). Various cell types can be seen in the tumor indicating a high occurrence of cell differentiation. Okuda et al. performed immunohistochemistry on PA to determine the role of Wnt in cell differentiation (2). The results suggested that Wnt is involved in cell differentiation in PA. In the present study, we considered that Notch might also be involved in cell proliferation and differentiation in the same manner that Okuda et al. did on Wnt (3).

In this study, we focused on Notch expression in PA since it has been hypothesized to be strongly associated with neoplastic cell differentiation. Moreover, studies on the expression of Notch have been tremendously increasing.

Materials and Methods
Tissue samples from Department of Oral Pathology, School of Dentistry, Aichi Gakuin University, diagnosed histologically as PA were further re-evaluated for the purpose of this research. A total of 30 cases of PA based on WHO classification were used in this experiment. The specimens used in this study were the same as those used by Okuda et al. (2), described in Table 1.

Specimens were fixed in neutral buffered formalin solution, subjected into series of alcohol for dehydra-
stained with hematoxylin and eosin and examined under a light microscope.

**Table 1. Cases examined**

| Age | Sex | Location |
|-----|-----|----------|
|     | Male| 13       |
| 51.5|     | Palate   |
|     | Female| 17   |
|     |     | Parotid gland |
|     |     | Mandibular gland |
|     |     | Upper lip |
|     |     | Buccal mucosa |
|     |     | Other |

Immunohistochemistry was performed in the following manner. Briefly, deparaffinized sections were pre-treated for antigen retrieval in citrate buffer (Mitsubishi Chemical Medience, Tokyo, Japan) with a pH of 6.0 and placed in autoclave at 120 °C for 15 min. Then after, sections were covered with serum-free protein block (Dako, Japan Co., Ltd., Tokyo, Japan), incubated at room temperature for 30 min. Notch1 rabbit polyclonal antibody (Anti-Notch-1 intracellular domain antibody, ab83232, Abcam, Cambridge, UK) with a pH of 6.0 and placed in autoclave at 120 °C for 15 min. Then after, sections were covered with serum-free protein block (Dako). Antigenic sites were revealed using DAB.

Immunofluorescent staining was also performed using Notch1 rabbit polyclonal antibody, CK7 mouse monoclonal antibody (1:100; Abcam) and CK13 mouse monoclonal antibody (AE8; 1:100; Abcam). Double staining was done by combining Notch1 and CK7 as well as Notch1 and CK13. After deparaffinization, Notch1-CK7 and Notch1-CK13 were pre-treated in citric acid buffer (Mitsubishi Chemical Medience, Tokyo, Japan) with a pH of 6.0 and placed in microwave for 1 min. Then after, sections were covered with serum-free protein block (Dako). Antigenic sites were revealed using DAB.

Immunofluorescent staining was also performed using Notch1 rabbit polyclonal antibody, CK7 mouse monoclonal antibody (1:100; Abcam) and CK13 mouse monoclonal antibody (AE8; 1:100; Abcam). Double staining was done by combining Notch1 and CK7 as well as Notch1 and CK13. After deparaffinization, Notch1-CK7 and Notch1-CK13 were pre-treated in citric acid buffer (Mitsubishi Chemical Medience, Tokyo, Japan) with a pH of 6.0 and placed in microwave for 1 min. Then after, sections were covered with serum-free protein block (Dako), incubated at room temperature for 30 min. The primary antibodies Notch1 and CK7 (1:100; Can Get Signal, Toyobo Co., Osaka, Japan) were allowed to react for 4 °C for 16 hours. For the secondary antibody, Donkey anti-rabbit IgG H&L (1:200; Alexa Fluor 594; Abcam) and Donkey anti-mouse IgG H&L (1:200; Alexa Fluor 488; Abcam) were carried out after reaction with Can Get Signal (Toyobo) at 1:200 at room temperature for 60 min. DAPI was allowed to react for 3 min for nuclear clarity. The present study was approved by the ethics committee of Aichi Gakuin University, School of Dentistry under the title “Diagnosis and Clinico-pathological Study on the Elucidation of Salivary Gland Tumors’ (No. 284, December 5, 2011).

**Results**

**Histopathological examination**

Generally, the tumor consists of various tissue types with small and/or large duct-like/cystic spaces in the tissues. The tumor is round in shape and covered with relatively thin fibrous connective tissue. The substantial portion of the tumor consists of glandular structures of tumor nests surrounded by fibrous tissue. The tumor is clearly separated from the normal salivary gland tissues. The tumor stroma consists of duct-like structures predominantly ductal and myoepithelial cells forming cystic cavities.

The mesenchymal part consists of myoepithelial cells, spindle shaped cells and myxoma-like tissue formed by round or oval cells dissociated from the tumor growth. In some areas, cartilage-like tissues can be characteristically observed (Fig. 1-a). In myxomatous areas, the myoepithelial cells have sparsely formed cord-like structures and the presence of mucous was confirmed. Cartilage-like tissues can be seen in a wide range possibly due to the presence of cartilage-like matrix and cartilage-like cells. Neoplastic myoepithelial cells are dissociated from the mucous or cartilage-like substrate by their own secretions. The tumor parenchyma might have been formed by myxomatous cells, which migrated to seep into the structure of mesenchymal and cartilage-like cells. Thus, the epithelial component is dissociated and mixed into the mesenchymal-like tissue, giving rise to a ‘mixed appearance’. Moreover, tumor cells undergo remarkable squamous metaplasia (Fig. 1-b).

**Immunohistochemical findings**

Notch1 expression was observed in the cytoplasm of ductal epithelial cells as well as in some of the nuclei. Notch expression was also observed in the nucleus of tumor cells surrounding solid tumor nests (Fig. 1-c).

Notch was expressed in the cytoplasm of plasmacytoid myoepithelial cells but not in the nucleus. In areas undergoing squamous metaplasia, Notch was strongly expressed in the cytoplasm and nucleus of basal cells. However, the expression became weak towards the outer squamous cells (Fig. 1-d).

The cartilage-like cells representing the characteristic feature of PA showed strong cytoplasmic expression but not nuclear reaction (Fig. 1-e). Likewise, the myoepithelial cells in myxomatous area showed faint expression on the cell membrane but no cytoplasmic or nuclear reaction were detected (Fig. 1-f).

Ductal epithelial cells expressed CK7 (Fig. 2-a). Notch was also detected in the nucleus of some ductal epithelial cells (Fig. 2-b). Double immunofluorescent staining revealed that Notch and CK7 were co-expressed by tumor cells in ductal structures. Notch expression was observed in most of the nucleus of tumor cells in solid nests as well as those in ductal structures (Fig. 2-c, d).
The expression of CK13 in tumor cells undergoing squamous metaplasia was also observed. CK13 showed positive reaction on basal and prickle cell layers (Fig. 3-a). On the other hand, Notch was observed in the nucleus from basal cell to squamous cells (Fig. 3-b). Double immunofluorescent staining of Notch and CK13 revealed that most basal cells expressed Notch although expression decreased towards the squamous cells. Furthermore, nuclear expression was observed in basal cells and the expression became weak on the prickle cell layer (Fig. 3-c, d).

Figure 1. Histopathological features of typical PA (a: Case 9: 28 year-old female, the right palatal region; scale bar=200µm), squamous metaplasia with keratin pearl (b: Case 30: 40 year-old male, left palatal region; scale bar=100µm), immunohistochemical positive reactions in ductal areas (c: Case 2: 40 year-old female, right palatal region; scale bar=100µm), in squamous metaplasia area (d: Case 18: 69 year-old female, right buccal mucosa; scale bar=100µm), immunohistochemical features of cartilage-like area (e: Case 18: 69 year-old female, right buccal mucosa; scale bar=20µm) and myxomatous area (f: Case 18: 69 year-old female, right buccal mucosa; scale bar=50µm).

Figure 2. Immunofluorescent staining images of ductal structure forming area (a: CK7; b: Notch1; c: merged image of CK7 and Notch1; and d: merged image of CK7 and Notch1 and DAPI; Case 28: 39 year-old male, submandibular region; scale bar=100µm).
**Discussion**

PA is the most common benign epithelial tumor of salivary glands based on WHO classification (1). The tumor shows a wide variety of tumor development brought about by cell proliferation and differentiation as mentioned in previous studies (4-6).

Okuda et al. studied the expression of Wnt in 30 cases of PA. The results showed that tumor cells in ductal structure and in solid tumor nests expressed Wnt. It was inferred that β-catenin pathway was involved in cell differentiation indicated by Wnt expressions in cells undergoing squamous metaplasia (2). For this reason, we believed that Notch would have a role in tumor cell differentiation considering its association with other tumors (7, 8).

Notch appears as a typical signal that controls the growth of tissues responsible for the fate of cells (9). Notch is a single-pass transmembrane receptor with a domain outside and inside the cell membrane. Notch intracellular domain (NICD) is cleaved and binds to a ligand such as Jagged. Cleaved NICD is moved from the cell membrane into the cytoplasm, binds with Suppressor of Hair-less (Su(H)) to form a complex and acts on downstream target gene in the nucleus thus activating the expression (10). In addition to the role of transmitting instructions for various morphogenesis and tissue differentiation during development, Notch plays an important role in cell-cell signaling and is involved in stem cell maintenance, differentiation and neuronal function in adult.

When there is a disturbance in the activity of Notch signaling pathway, it becomes oncogenic detected in several cancers such as esophageal cancer, breast cancer and lymphoblastic acute leukemia (T-ALL) (11, 12). In addition, Notch has been the focus of researches on the metastasis of malignant tumors such as adenoid cystic carcinoma and malignant ameloblastoma (13, 14), believed to be a huge factor in the progress of malignant tumors.

Studies in Notch have been done in odontogenic tumors such as ameloblastoma (15-17). Notch is expressed in solid nests and in squamous metaplasia in ameloblastoma, implicating its contribution in cell differentiation and morphogenesis. Moreover, in calcifying cystic odontogenic tumor, calcifying epithelioma (pilomatrixoma), odontoma and cranio-opharyngioma, Notch expression in ghost or shadow cells was observed (18-20). Notch overexpression in ghost cells inhibited the development of neighboring cells. This means that Notch inhibits the production of a ligand in receiving cell signal between Notch ligand and Notch receptor to adjacent cells through a negative feedback mechanism. In contrast, during binary cell fate determination, Numb acts an inhibitor of Notch signaling cycle causing asymmetric distribution of the Notch pathway. This led to overexpression suggesting that Notch-Jagged1 is the primary mechanism in determining the fate of ghost cell (21).

Squamous metaplasia depicted as having a variety of cell differentiation is commonly observed in odontogenic tumors such as ameloblastoma. Squamous metaplasia can also be observed in PA. Muraki et al suggested a close relationship between Notch and ameloblastoma shown by the localization of Notch in peripheral tumor nests (15). From this aspect, we suspected that Notch might also be involved in cell differentiation in PA.
Comparing the results of our study with previous reports, from the partial CK7 expression and substantial Notch expression in ductal epithelial cells as well as the Notch expression in solid tumor nests, it can be inferred that Notch is involved in cell differentiation (2, 22). CK13 expression was observed in cells undergoing squamous metaplasia and Notch expression was seen in the nucleus of basal and squamous cells (23). The intense Notch expression in basal cells and weak expression in squamous cells suggests that Notch is involved in the differentiation from basal to squamous cell. Moreover, the loss of nuclear expression on the surface layer would signify that differentiation has been terminated. Notch is involved in cell differentiation in PA. The results coincided with our present study implicating the function of Notch in the same study (2). Nuclear expression was seen in the nucleus of basal and squamous undergoing squamous metaplasia and Notch expression was also confirmed in the basal cuboidal cells forming ductal structure expressed Wnt 5a. However, the expression was not confirmed in the basal cells surrounding those cells undergoing squamous metaplasia. The results coincided with our present study implicating the function of Notch in the same sites causing cell differentiation (2).

Notch involvement is suspected in cell differentiation in areas showing ductal structures and squamous metaplasia. In the study by Okuda et al, small cuboidal cells forming ductal structure expressed Wnt and the expression was also confirmed in the basal cells surrounding those cells undergoing squamous metaplasia. The results coincided with our present study implicating the function of Notch in the same sites causing cell differentiation (2).

In summary, Notch is involved in cell differentiation of ductal cells in PA. Nuclear expression was shown in tumor cells in solid nests and surrounding structures. Moreover, Notch is expressed by basal cells undergoing squamous metaplasia suggesting the participation of Notch in cell differentiation in PA.

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Competing Interests

The authors have declared that no competing interest exists.

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