Thyroid-like low-grade nasopharyngeal papillary adenocarcinoma with squamous differentiation in the posterior nasal septum
A rare case report
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Abstract
Rationale: Thyroid-like low-grade nasopharyngeal papillary adenocarcinoma (TL-LGNPPA) is a rare neoplasm characterized by papillary epithelial proliferation and abnormal expression of thyroid transcription factor-1. To our knowledge, squamous differentiation in TL-LGNPPA is an unusual phenomenon, and only 1 case has been reported to date. The tumor occurs mainly on the roof of the nasopharynx. Herein, we report a case of TL-LGNPPA with squamous differentiation located on the posterior nasal septum.

Patient concerns: A 45-year-old Chinese woman presented to our hospital with a complaint of nasal obstruction for 10 years and the symptom has been getting worse for recent 3 years.

Diagnoses: Microscopic examination of the tumor tissue revealed findings typical of TL-LGNPPA, and in addition to the typical components, squamous differentiation could be found in some areas within the fibrovascular cores of the papillary components. Immunohistochemically, the squamous cells were positive for P40 and P63 but negative for thyroid transcription factor-1.

Interventions: The patient underwent complete resection of the tumor.

Outcomes: Follow up results showed that the patient had no recurrence 41 months after removal of the tumor.

Lessons: Our report anticipates that, although TL-LGNPPA with squamous differentiation in the posterior nasal septum is rare, this report will expand the existing knowledge associated with TL-LGNPPA.

Abbreviations: PTC = papillary thyroid carcinoma, TTF1 = thyroid transcription factor-1, TL-LGNPPA = thyroid-like low-grade nasopharyngeal papillary adenocarcinoma.

Keywords: nasopharynx, posterior nasal septum, squamous differentiation, TL-LGNPPA

1. Introduction
Epithelial tumors of the nasopharynx are classified as either nasopharyngeal carcinoma, nasopharyngeal papillary adenocarcinoma, or salivary gland-type carcinoma according to the 2017 World Health Organization Classification of Head and Neck Tumors. Thyroid-like low-grade nasopharyngeal papillary adenocarcinoma (TL-LGNPPA) is an extremely rare type of nasopharyngeal papillary adenocarcinoma. Nine cases were first described and characterized by Wenig et al. who named LGNPPA in 1988. Carrizzo and Luna described 2 pediatric cases, indicating that these tumors were similar to papillary thyroid carcinoma (PTC) for histological morphology; they documented the expression of thyroid transcription factor-1 (TTF-1) in tumor cell nuclei, and renamed them TL-LGNPPA. Because of the rarity of the neoplasm, TL-LGNPPA is still considered to be at the stage of case collection. Only 1 case of TL-LGNPPA exhibiting a squamous differentiation component has been reported previously in the English literature. This tumor is typically located in the nasopharynx arising from the roof or lateral wall of the nasopharynx, and in a few cases, it has been noted to be attached to the posterior nasal septum. Herein, we present another case of TL-LGNPPA with squamous differentiation located on the posterior nasal septum. Interestingly, the TL-LGNPPA exhibited a different histopathological finding in addition to different morphological and immunohistochemical features compared to previous cases.

2. Case report
A 45-year-old woman visited our hospital with a 10-year history of nasal obstruction that worsened over the past 3 years. On head
and neck computed tomography, a horizontal view and a coronal view image exhibited a protruding mass on the posterior edge of the nasal septum (Fig. 1A and B). No swollen lymph nodes were observed. The mass was clinically diagnosed as nasopharyngeal carcinoma. Finally, the patient underwent complete resection of the tumor. Several days after surgery, the patient’s nasal obstruction was relieved, and she was discharged from hospital. Three months later, computed tomography examination and pathological biopsy revealed no residual neoplastic tissue. The patient remained free of recurrence or metastasis after 41 months of follow-up.

Macroscopically, the tumor was described as an ash red polypoid with dimensions of 4 × 3.5 × 2 cm. Histopathological examination confirmed that the tumor was partially covered with normal respiratory epithelium and growth under the mucosa (Fig. 2A). The tumor was mainly composed of a papillary growth pattern; moreover, the fibrovascular cores of the papillary showed hyalinization (Fig. 2B). Each papilla was covered by cuboidal or columnar epithelial cells with tiny bland, round to oval nuclei. In some areas, the nuclear appearance was very similar to that of PTC, including irregular thickened nuclear membranes, nuclear grooves, intranuclear cytoplasmic pseudo-inclusions, and ground-glass appearance (Fig. 2C). Necrosis was not observed, whereas scattered psammoma bodies were observed (Fig. 2D). In addition to the above common pathological features, the case showed an additional characteristic called squamous differentiation. Squamous cells were observed within some of the fibrovascular cores of the papillary configuration, packed closely, grown in a solid pattern with a lobular configuration, and prominent peripheral palisading; atypia and keratinization were not found (Fig. 2E and F).

On immunohistochemistry, all tumor cells were positive for pan cytokeratin and epithelial membrane antigen, but negative for thyroglobulin. Additionally, TTF-1 exhibited strong nuclear immunoreactivity for glandular tumor cells but was negative for squamous cells (Fig. 3A). In addition, these squamous cells were positive for p63 and p40 (Fig. 3B). The Ki-67 index reached 3% in the area of the greatest concentration, and the squamous cells also showed low proliferating activity (Fig. 3C). We performed polymerase chain reaction to amplify exons 11 and 15 of 1v-Raf murine sarcoma viral oncogene homolog B1 (BRAF), and the results were negative.

3. Discussion

Most TL-LGNPPAs have a pedunculated and polypoid shape, and occur in any part of the nasopharynx, commonly observed on the top, lateral, and posterior walls of the nasopharynx as well as on the posterior edge of the nasal septum.[5] The mean age of patients with TL-LGNPPA is 32.3 years (range: 9–68 years), and the incidence does not differ according to sex; the tumor size ranges from 0.3 to 4.0 cm.[6] Because of its location, patients usually present with nasal obstruction and/or nasal discomfort, epistaxis, and headache. In this case, the neoplasm was located on the posterior edge of the nasal septum, which is a rare location. The other clinical manifestations were not distinct.

The presence of squamous differentiation in TL-LGNPPA is unusual. Only 1 case report describing TL-LGNPPA exhibiting squamous differentiation has been published to date.[4] Histologically, the tumor showed the typical appearance of TL-LGNPPA. In addition, the presence of squamous differentiation is a major feature. The case described by Oide et al[4] showed that scattered squamous cell foci were intermixed with glandular tumor cells. However, our case showed additional characteristics: the squamous cells were isolated within the fibrovascular cores and grew in a solid pattern with a lobular configuration and prominent peripheral palisading, but not intermixed with glandular tumor cells. Keratinization was not observed; however, the squamous cells showed no obvious nuclear atypia or proliferating activity in either of these 2 cases.

The immunohistochemical features of this case were consistent with those of previously reported cases of TL-LGNPPA: the glandular tumor cells exhibited diffuse strong immunoreactivity for TTF-1, but no immunoreactivity for thyroglobulin. The difference is that the squamous cells exhibited both p40 and TTF-1 immunoreactivity in Oide’s case, but in our case, the squamous cells were positive for p63 and p40, and negative for TTF-1. BRAF mutations were negative in both cases, which is consistent with results reported previously in the literature.[5,7]

TL-LGNPPA with squamous differentiation is morphologically similar to the squamous metaplasia observed in PTC and needs to be differentiated from squamous cell carcinoma. Cytological atypia of the squamous epithelium is the key to differentiating carcinomas. In addition, squamous morules should be distin-
guished from squamous differentiation. Squamous morules are characterized by rounded aggregates or syncytial sheets of cells with bland round, ovoid, or spindle-shaped nuclei with indistinct cell borders; p63 is focally or diffusely positive in most of the squamous differentiation elements, whereas morules are usually negative. In the present case, the squamous cells showed no obvious nuclear atypia and were diffusely positive for p63 and p40; therefore, we believe that it is squamous differentiation.

The prognosis of TL-LGNPPA as a low-grade malignancy is excellent. Thus far, no case of TL-LGNPPA has been associated with lymphovascular spread or metastasis with follow-up periods of up to 15 years. Local recurrence may occur because of inadequate surgical resection. After resection of the tumor, the patient in our case was followed up for 41 months, with no recurrence. Follow-up information for the other case of TL-LGNPPA with squamous differentiation was unavailable for comparison. Some studies have found that squamous differentiation is associated with a poor prognosis of PTC. Because the number of cases reported is small, we do not know what this means for TL-LGNPPA with squamous differentiation. This suggests that we should follow up our patient for a long period of time.

The etiopathogenetic mechanisms responsible for the development of TL-LGNPPA remain to be elucidated. The majority of nasopharyngeal malignant tumors are squamous cell carcinomas and are associated with Epstein Barr virus (EBV). Therefore, many studies have used in situ hybridization and polymerase chain reaction to detect EBV in tumor cells of TL-LGNPPA, but no evidence of EBV infection has been found. Kakkar et al reported a case of TL-LGNPPA as a second malignancy five-and-a-half years after the patient was diagnosed and treated with radiotherapy for a diffuse astrocytoma in the frontal lobe. Therefore, they suggested that exposure to radiation could be a shared pathogenic factor. Considering the morphological similarity to PTC, some investigators have performed sequencing for mutations in BRAF and NRAS in cases of TL-LGNPPA, but

Figure 2. (A) The tumor growth under mucosa ×25. (B) The tumor was mainly composed of complex papillae with hyalinized fibrovascular cores ×40. (C) Nuclear overlapping, irregular thickened nuclear membranes, nuclear grooves, and ground-glass appearance were frequently seen ×400. (D) Scattered psammoma bodies appeared ×100. (E) Low-power view of the squamous cells gathered into clusters in the fibrovascular cores ×100. (F) The squamous cells showed no obvious atypia ×200.
the results were negative.\textsuperscript{15,16} Clearly the pathogenesis of TL-LGNPPA requires further research.

4. Conclusion

In summary, our case exhibited a rare histopathological finding in addition to morphological and immunohistochemical features typical of TL-LGNPPA with squamous differentiation. Although the posterior nasal septum has been involved in previous cases of TL-LGNPPA, this is the first case of TL-LGNPPA with squamous differentiation in this rare site. Hence, the report of this case extends the understanding of this tumor, in order to aid both clinicians and pathologists in making appropriate final diagnoses and increases awareness of this possibility.

Author contributions

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