Soft Palate Schwannoma: A Rare Case of an Intraoral Mass

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Abstract

Introduction: Schwannomas are benign encapsulated neoplasms that arise from proliferation of Schwann cells and can originate from any cranial, peripheral, or autonomic nerve. While most schwannomas present in the head and neck region, only 1% of these occur intra orally and these typically occur on the tongue or floor of mouth.

Case: We report here a 55-year-old woman seen at our institution who presented with a mass of the soft palate. Biopsy found it to be a schwannoma of the soft palate, which was surgically excised.

Conclusion: Because of the predilection of schwannomas to occur in the head and neck region, otolaryngologists must maintain a high level of suspicion for unique and rare presentations.

Keywords: Schwannomas; Neoplasms; Glossotonsillar sulcus

Case Presentation

A 55-year-old African American female with a history of obstructive sleep apnea presented for evaluation of a soft palate mass. During a dental procedure, a dome-shaped mass on the soft palate was noted, and a resultant CT scan of the neck with contrast showed a 1.6 x 2 cm heterogeneous soft tissue mass localized to the lateral oropharyngeal wall and glossotonsillar sulcus concerning for intraoral malignancy. The patient had not noticed the mass previously. She denied a history of smoking, excessive alcohol consumption, or radiation exposure. She reported no recent weight loss, no change in her voice, no difficulty breathing, and no otalgia. The only significant finding on exam was the submucosal, rubbery mass without mucosal changes or ulceration, which did not cross the midline (Figure 1). It was non-tender, non-cystic, and non-fluctuant. Flexible laryngoscopy was within normal limits.

A biopsy was taken and several small pieces of rubbery, yellow tissue were obtained. No fluid was expressed. Immunohistochemical studies revealed that the tumor cells were positive for S100 and negative for AE1/AE3 cytokeratin (Figure 2A and B). Permanent sectioning demonstrated classic well-formed nuclear palisades surrounding the fibrillary processes (Figure 2C) as well as large, irregular vessels with hyalinization and rare thrombi (Figure 2D). These findings were most consistent with a schwannoma.

Surgical excision was performed under general anesthesia and exposure was obtained using a McIvor mouth gag. The lesion dissected easily away from the posterior hard palate and the palatal musculature. The specimen was 3.5 x 2.2 x 1.9 cm in size. The defect was closed primarily with a combination of absorbable suture and a small piece of AlloDerm® (Acelity, San Antonio, TX) graft.
At follow-up eleven days later, the surgical site was healing well and the AlloDerm® was no longer identifiable within the wound bed.

**Results and Discussion**

This case represents an unusual presentation of a schwannoma, identified on biopsy before definitive excision. The initial biopsy revealed a spindle cell neoplasm composed of long, thin cells with tapered nuclei, coarse chromatin, and ill-defined cytoplasm proliferating in a biphasic pattern. Focal areas suggestive of the presence of Verocay bodies were confirmed on the surgical specimen. Classically, Schwannomas have an Antoni A palisading hypercellular component with Verocay bodies and an Antoni B hypocellular and myxoid component. Cytologically, the cells are elongated with hyperchromatic, wavy and tapered nuclei [2,5]. Immunohistochemical analysis (S-100, Leu-7) is often used to confirm and classify the diagnosis of nerve sheath tumors [5].

While this schwannoma was identified prior to surgery, diagnosis is not always possible until the tumor has been excised. Diagnosis and surgical planning can be aided by CT or MRI imaging to evaluate the extent of infiltration into surrounding tissues. The combination of imaging, biopsy, and immunohistochemical staining allows for appropriate diagnosis and surgical management. The definitive treatment of schwannoma is excision, and the causal nerve is often unlikely to be identified [5].

The differential diagnosis for a soft palate neoplasm is varied and includes minor salivary gland tumor, pyogenic granuloma squamous cell carcinoma, or lipoma [1]. Histologically, however, other neurogenic tumors such as neurofibroma, neuroma, myoblastoma of granular cells, neurogenic sarcoma, malignant schwannoma, neuroepithelioma and melanoma must also be excluded [5]. Schwannomas very rarely undergo malignant transformation [5].

**Conclusion**

While our original differential diagnosis included minor salivary gland tumors, torus palatinus, and squamous cell carcinoma, the histological findings allowed us to refine our diagnosis to a soft palate schwannoma. Schwannomas may present from any cranial, peripheral, or autonomic nerve. However, as nearly half of benign nerve sheath tumors occur in the head and neck, otolaryngologists must be particularly aware of the many unique presentations of schwannoma and maintain a high level of clinical suspicion. Early detection and biopsy can allow for appropriate surgical management. Schwannomas, therefore, should represent an important differential diagnosis for neoplasms in the head and neck.

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**References**