

CASE REPORT

DOI: 10.5336/caserep.2025-110557

Tongue Schwannoma in an Adolescent Female

¹Yusuf AYDIN^a, ²Fatma ATALAY^a, ³Fatma AKYÜREK TAŞÇI^b, ⁴Nurtaç SARIKAŞ^c

^aKastamonu University Faculty of Medicine, Department of Ear Nose Throat Diseases, Kastamonu, Türkiye

^bKastamonu Training and Research Hospital, Clinic of Ear Nose Throat Diseases, Kastamonu, Türkiye

^cKastamonu University Faculty of Medicine, Department of Medical Pathology, Kastamonu, Türkiye

ABSTRACT Schwannoma is a benign tumor originating from the Schwann cells of peripheral nerves. Although schwannomas are usually located intracranially, extracranial occurrences are most commonly found in the head and neck region (25-45%). It is rarely observed in the oral cavity (1%). The most common location in the oral cavity is the tongue. In this article, we report a rare case of tongue schwannoma in a 16-year-old female patient who presented to our center with a 2-year history of tongue swelling. After the patient was evaluated with magnetic resonance imaging, the mass was excised and sent to pathology for histopathological examination. Pathology results revealed schwannoma.

Keywords: Tongue; neurilemmoma; schwann cells; schwannomatosis; neurofibromatosis 2

Schwannoma or neurilemmoma is a benign tumor originating from the Schwann cells of peripheral nerves. Although malignant transformation is possible, it is exceedingly rare.¹ It is a slow-growing, solitary, well-circumscribed and encapsulated tumor. The exact etiology of schwannoma remains unclear.^{1,2} It is usually sporadic and can rarely be detected in association with neurofibromatosis type 2 or schwannomatosis.³ The majority of extracranial schwannomas (25-45%) are detected in the head and neck region, and 1% are seen in the oral cavity. In the oral cavity, it is most frequently seen in the tongue.^{4,5} Clinical presentation varies depending on the tumor's location, but it is generally asymptomatic when present in the oral cavity.⁶

CASE REPORT

A 16-year-old female patient presented to Kastamonu Education and Research Hospital due to a painless

swelling in the tongue. The patient reported being aware of the swelling for approximately 2 years, with a noticeable increase in size over the past 6 months, accompanied by difficulty in speaking (Figure 1). She reported no history of trauma, systemic illnesses, or associated genetic disorders. There was no family history of a similar head and neck mass. Head and neck examination was unremarkable, with no additional masses identified. A painless, well-circumscribed, approximately 1.5 cm diameter mass with no color change was detected on the right lateral side of the ventral part of the tongue. The patient underwent contrast-enhanced magnetic resonance imaging (MRI), which revealed a well-circumscribed 15×12 mm mass demonstrating good contrast enhancement, isointense with muscle on T1-weighted images and hyperintense on T2-weighted images (Figure 2). The patient underwent intraoral mass excision under general anesthesia and the mass was completely excised

Correspondence: Yusuf AYDIN

Kastamonu University Faculty of Medicine, Department of Ear Nose Throat Diseases, Kastamonu, Türkiye

E-mail: yaydin87@gmail.com

Peer review under responsibility of Türkiye Klinikleri Journal of Case Reports.

Received: 19 Mar 2025

Accepted: 29 May 2025

Available online: 11 Jul 2025

2147-9291 / Copyright © 2025 by Türkiye Klinikleri. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).



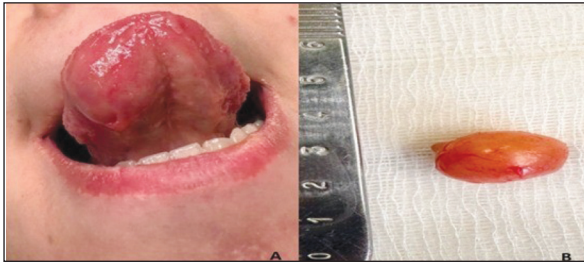


FIGURE 1: A: Well-circumscribed swelling on the ventral surface of the tongue; **B:** Excised mass

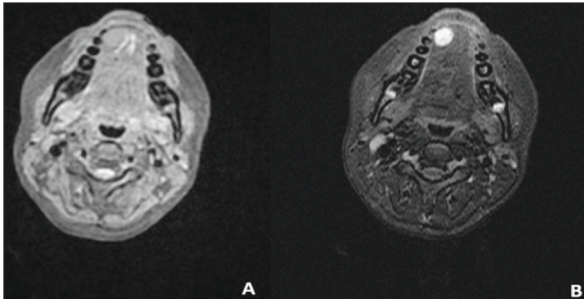


FIGURE 2: A: Aksial T1 image of lingual schwannoma; **B:** Aksial T2 image of lingual schwannoma

and sent to pathology for histopathological examination. No complications developed during or after the operation. The pathology report confirmed schwannoma. Focal degeneration and bleeding areas were noted in macroscopic examination. Immunohistochemical staining showed strong positivity with S-100 and faint positivity with CD 56. Smooth Muscle Actin, Desmin and CD 34 were negative. Written informed consent was obtained from the patient.

DISCUSSION

Schwannoma is a benign tumor originating from Schwann cells and can arise from any myelinated nerve fiber. Although rare, malignant transformation has been reported in approximately 8-10% of cases.^{1,7} Schwannomas most commonly occur intracranially, typically arising from the vestibulocochlear (VIIIth) nerve. Extracranial schwannomas are predominantly found in the head and neck region (25-45%), with only 1% occurring in the oral cavity.⁸ The tongue is the most common location in the oral cavity.⁸⁻¹⁰ It is detected equally in both sexes and is seen more frequently in the 2nd-4th decades.^{3,7,11}

Patients with oral cavity schwannoma may present with symptoms such as sore throat, dysphagia, dysphonia, infection or bleeding in that region, and sleep apnea, depending on the size of the mass, but the most common reason for presentation is a painless growing mass.^{6,7,11} In this case, the patient's reason for presentation was a painless growing mass in the tongue and dysphonia. Multiple schwannomas can be detected in patients with neurofibromatosis type 2 or schwannomatosis, and malignant transformation is more common (15%).^{3,10}

In differential diagnosis, soft tissue tumors such as hemangioma, lymphangioma, salivary gland tumors, fibroma, dermoid cyst, lipoma, squamous cell carcinoma, rhabdomyosarcoma and sarcoma should be considered. Regular borders, submucosal localization and slow growth are helpful in excluding malignant tumors. Computed Tomography (CT) and MRI can be preferred for diagnosis, MRI is more useful than CT due to its higher sensitivity to soft tissues. In MRI, isointense image is obtained with muscle in T1 sequence, while hyperintense image is obtained in T2.^{1,10,12} Invasion of surrounding tissues is not observed. Definitive diagnosis is made histopathologically. Histologic sections reveal an encapsulated neoplasm composed of cytologically bland spindle cells arranged in short fascicles. The tumor exhibits alternating patterns, with hypercellular areas showing nuclear palisading (Antoni A) interspersed with hypocellular regions (Antoni B). Immunohistochemical staining typically reveals strong S-100 and CD56 positivity, while focal degeneration and hemorrhagic areas may be observed macroscopically.^{3,13,14} These indicators were also found histopathologically in our case (Figure 3).

Treatment of tongue schwannomas is intraoral excision. Recurrence after total excision is very rare and the risk of complications is low.^{3,7,15} In this case, the patient was scheduled for monthly follow-up visits after the 1st-month check-up (Figure 4). During the 6-month follow-up period, no neurodeficits were noted, and the patient did not develop any speech or diction problems.

In conclusion, schwannoma of the oral cavity and tongue is extremely rare. It should be considered in patients presenting with a slowly growing, well-

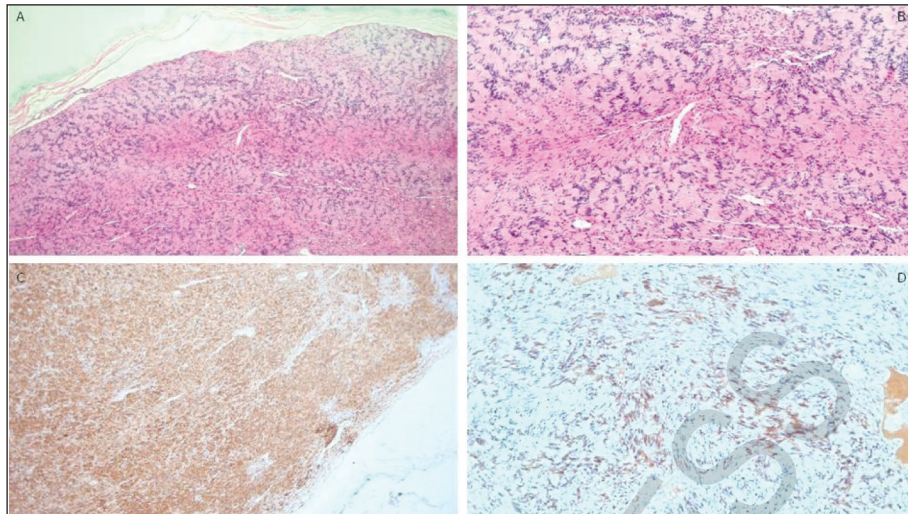


FIGURE 3: A: Hypercellular Antoni A areas (Hematoxylin-Eosin, original magnification x40); B: Hypocellular Antoni B areas (Hematoxylin-Eosin, original magnification x100); C: Strong and diffuse S 100 positivity (immunoperoxidase, original magnification x40); D: CD 56 positivity (immunoperoxidase, original magnification x100).



FIGURE 4: Postoperative 1st month image

circumscribed mass on the tongue. Preoperative imaging is essential to assess the lesion and its relationship with adjacent anatomical structures prior to biopsy. The definitive diagnosis is established through histopathological examination. No recurrence has been reported after intraoral total excision and the possibility of complications is minimal.

Source of Finance

During this study, no financial or spiritual support was received neither from any pharmaceutical company that has a direct connection with the research subject, nor from a company that provides or produces medical instruments and materials which may negatively affect the evaluation process of this study.

Conflict of Interest

No conflicts of interest between the authors and / or family members of the scientific and medical committee members or members of the potential conflicts of interest, counseling, expertise, working conditions, share holding and similar situations in any firm.

Authorship Contributions

Idea/Concept: Yusuf Aydın; **Design:** Yusuf Aydın; **Control/Supervision:** Fatma Atalay; **Data Collection and/or Processing:** Nurtaç Sarıkaş, Fatma Akyürek Taşçı; **Analysis and/or Interpretation:** Yusuf Aydın, Fatma Atalay; **Literature Review:** Fatma Akyürek Taşçı; **Writing the Article:** Yusuf Aydın; **Critical Review:** Fatma Atalay; **References and Fundings:** Fatma Akyürek Taşçı; **Materials:** Yusuf Aydın.

REFERENCES

1. Enoz M, Suoglu Y, Ilhan R. Lingual schwannoma. *J Cancer Res Ther.* 2006;2(2):76-8. PMID: 17998681.
2. Colreavy MP, Lacy PD, Hughes J, Bouchier-Hayes D, Brennan P, O'Dwyer AJ, et al. Head and neck schwannomas--a 10 year review. *J Laryngol Otol.* 2000;114(2):119-24. PMID: 10748827.
3. Cohen M, Wang MB. Schwannoma of the tongue: two case reports and review of the literature. *Eur Arch Otorhinolaryngol.* 2009;266(11):1823-9. PMID: 19130068; PMCID: PMC2758150.
4. Harada H, Omura K, Maeda A. A massive pleomorphic adenoma of the submandibular salivary gland accompanied by neurilemmomas of the neck misdiagnosed as a malignant tumor: report of case. *J Oral Maxillofac Surg.* 2001;59(8):931-5. PMID: 11474457.
5. Spandow O, Fagerlund M, Bergmark L, Boquist L. Clinical and histopathological features of a large parapharyngeal neurilemmoma located at the base of the tongue. *ORL J Otorhinolaryngol Relat Spec.* 1999;61(1):25-30. PMID: 9892866.
6. Sawhney R, Carron MA, Mathog RH. Tongue base schwannoma: report, review, and unique surgical approach. *Am J Otolaryngol.* 2008;29(2):119-22. PMID: 18314023.
7. Lira RB, Gonçalves Filho J, Carvalho GB, Pinto CA, Kowalski LP. Lingual schwannoma: case report and review of the literature. *Acta Otorhinolaryngol Ital.* 2013;33(2):137-40. PMID: 23853407; PMCID: PMC3665385.
8. Carinci F, Carls FP, Grasso DL, Pelucchi S, Pastore A. Schwannoma of the parapharyngeal space. *J Craniofac Surg.* 2000;11(4):367-70. PMID: 11314385.
9. Ağırbaş S, Bozan N. Tongue base schwannoma presenting with dyspnea. *J Craniofac Surg.* 2025. PMID: 40009465.
10. Chandra M, Singh P, Venkatchalam V. Tongue schwannoma: a case report with review of literature. *JK-Practitioner.* 2013;18(1-2):28-34. http://scholar.google.com/scholar_lookup?&title=Tongue%20Schwannoma%3A%20a%20case%20report%20with%20review%20of%20literature&journal=JK-Practitioner&volume=18&issue=1%E2%80%932&pages=28-34&publication_year=2013&author=Chandra%2CM&author=Singh%2CP&author=Venkatchalam%2CV
11. Nakasato T, Kamada Y, Ehara S, Miura Y. Multilobular neurilemmoma of the tongue in a child. *AJNR Am J Neuroradiol.* 2005;26(2):421-3. PMID: 15709149; PMCID: PMC7974093.
12. Ülkü ÇH, Demir H, Yeşildemir HS, Esen H. Lingual schwannom. *Kulak Burun Bogaz Ihtisas Dergisi.* 2014;24(2):97-9. doi: 10.5606/kbbihtisas.2014.79346
13. Hsu YC, Hwang CF, Hsu RF, Kuo FY, Chien CY. Schwannoma (neurilemmoma) of the tongue. *Acta Otolaryngol.* 2006;126(8):861-5. PMID: 16846930.
14. Ulusoy B, Bozdemir K, Ersoy Çallıoğlu E, Kutluhan A, Korkmaz MH. Dilde submukozal yerleşimli schwannom [Submucosal lingual schwannoma]. *Kulak Burun Bogaz Ihtis Derg.* 2015;25(5):315-8. Turkish. PMID: 26476523.
15. Alrohaimi FA, Alsadah SA, Althaqib GA. Lingual schwannoma in an adolescent boy: a case report. *Ann Med Surg (Lond).* 2021;65:102216. PMID: 33981419; PMCID: PMC8085895.