

Lingual Schwannoma: A Case Report

M Rahbar¹, K Mardanpour^{2*}

¹Department of Pathology, ²Department of Surgery, Kermanshah University of Medical Sciences, Kermanshah, Iran

Abstract

We describe an 18-year old male patient with a 5 months previous history of swelling at the posterior mobile part of the tongue. The patient had difficulty in swallowing and speaking at the referral time. Examination of the oral cavity showed a swelling of 2x2 cm on the posterior part of the tongue, more towards the right side. Macroscopically, the entire lesion was removed with its capsule. Histopathological examination of the surgical specimen showed a schwannoma.

Keywords: Schwannoma; Tongue; Intraoral mass; Encapsulated

Introduction

Schwannomas originate from Schwann cells of the nerve sheath which covers the myelinated nerve fibres.¹⁻⁴ It was first identified by Virchow in 1908.⁵ Approximately 25 to 40% of all schwannomas are seen in the head and neck region.⁵⁻⁷ Intraoral schwannoma accounts for 1% of all head and neck region tumors⁷⁻⁹ and are commonly seen at the base region of the tongue.^{7,10,11} We report a patient with a schwannoma of the posterior part of the tongue (rare location), that was excised intraorally. Immunohistochemistry showed positive staining for S-100 protein, vimentin and glial fibrillary acid protein. Based on these findings, a histopathological diagnosis of benign schwannoma of the tongue was made. The patient was followed up for one year and there was no evidence of recurrence.

Case Report

An 18-year old male patient is presented with a 5 months previous history of a small mass in the posterior mobile part of the tongue while the mass was small at the onset and grown later. The patient had

difficulty in swallowing and speaking at the referral time. Examination of the oral cavity showed a swelling of 2x2 cm on the posterior part of the tongue, more towards the right side. The swelling was non-tender, smooth and firm to elastic in consistency at the time of admission (Figure 1).

No regional lymphadenitis was detected. The patient's medical history was unremarkable. There was no need for radiological investigations, because the mass was easily visible and palpable on physical examination. The lesion was excised with a small border of clinically uninvolved surrounding tissue. Macroscopically, the entire lesion was removed with its capsule (Figure 2).

The postoperative course was uneventful. The mobility of the tongue was good. Histopathological examination of the surgical specimen showed a schwannoma (Figure 3 and 4).



Fig. 1: Swelling in the posterior part of the tongue

*Correspondence: Kaykhosrow Mardanpour, MD, Department of Surgery, Kermanshah University of Medical Sciences, Kermanshah, Iran. Tel: +98-831-7259480, Fax: +98-831-7259481, e-mail: kmardanpour@yahoo.com
Received: December 28, 2008 Accepted: May 17, 2009

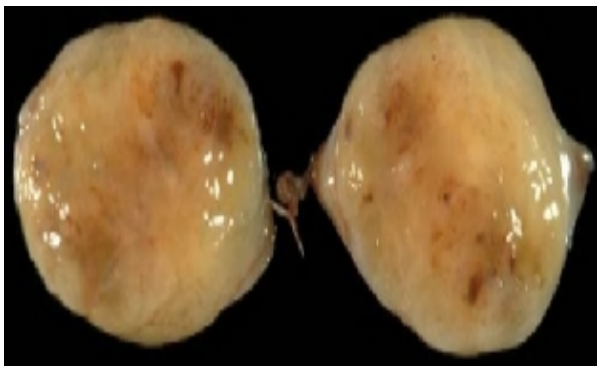


Fig. 2: The cut surface of a schwannoma with a "fish flesh" soft tan appearance.

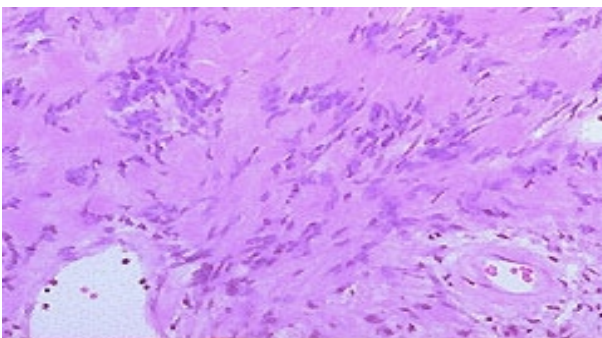


Fig. 3: The classic microscopic appearances of a schwannoma. The "Antoni A" pattern with palisading nuclei surrounding the pink areas (Verocay bodies).

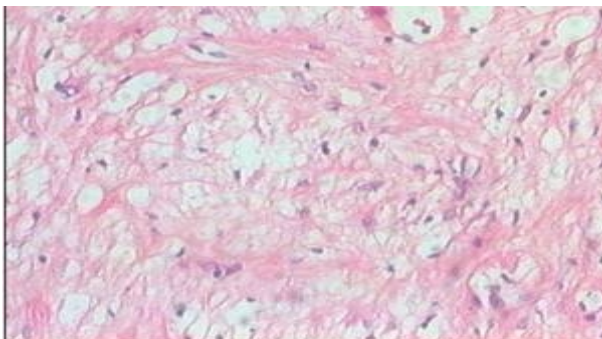


Fig. 4: The classic microscopic appearances of a schwannoma. The "Antoni B" pattern with a looser stroma, fewer cells, and myxoid change.

Immunohistochemistry showed positive staining for S-100 protein, vimentin and glial fibrillary acid protein. Based on these findings, a histopathological diagnosis of benign schwannoma of the tongue was made. The patient was followed up for one year and there was no evidence of recurrence.

Discussion

Schwannomas or neurilemmomas are benign, slow growing, usually solitary and encapsulated tumors, originating from Schwann cells of the nerve sheath.^{1,2} After a thorough review of the literature, we found only 6 previously reported cases of tongue base schwannomas.^{3,4} Approximately 25 to 40% of all schwannomas are seen in the head and neck region.^{5,6} Intraoral schwannoma accounts for 1% of all the head and neck region tumors⁷⁻⁹ and commonly seen at the base region of the tongue.^{7,10,11} Identification of the originating nerve may be difficult. In more than 50% of the intraoral lesions, it is not possible to differentiate between tumors of the lingual, hypoglossal and glossopharyngeal nerves.¹² It can be seen alone, or is associated with Von Recklinghausen disease.¹³ Etiology is still unknown and the disease is generally asymptomatic.¹⁴ In general, it starts as a capsulated nodule and grows slowly. If it invades the submucosal areas, it will lead to pain and discomfort.^{15,16} Malignant transformation is mentioned in 8-10% of the cases.¹³ The tumor develops in patients of all ages, without an obvious preference for either sex.⁷ The presenting feature of a tongue schwannoma is usually a tumor mass. Other symptoms include dyspnea or dysphagia and depend on the location and size of the tumor.¹⁰ Diagnosis is confirmed by histopathological studies. The tumor tissue consists of the so-called Antoni A and B type cells. Type-A tissue shows densely packed, elongated spindle cells, in the form of parallelly formed thin reticulin fibers, fusiform shaped cells and curled nuclei while type-B tissue has a more myxoid consistency. Amongst the sheets, there are acellular eosinophilic bodies called Verocay bodies, formed by thin cytoplasmic fibres. In addition, hemorrhage from adjacent tissue, necrosis, hyalinization and cystic degeneration may also occur in the tumor tissue.^{14,17,18} All histopathologic changes have been confirmed by immunohistochemistry staining such as S-100 protein, vimentin and glial fibrillary acid protein.¹⁹ Magnetic resonance imaging (MRI) is superior to other imaging modalities for the examination of the base of the tongue.²⁰ Malignant lesions such as squamous cell carcinomas and sarcomas and benign lesions as granular cell tumors, salivary gland tumors, schwannomas of the oral cavity, leiomyomas, rhabdomyomas, lymphangiomas, hemangiomas, dermoid cysts, lipomas, inflammatory

lesions and lingual thyroid are the differential diagnoses of this entity.²¹ Surgical excision or enucleation with preserve nerve function is the treatment of choice for this rare tumor. Excision is usually easy to perform and the prognosis is excellent as malignant transformation is rare.^{10,22}

Acknowledgement

We wish to thank for cooperation of Kermanshah University of Medical Sciences.

Conflict of interest: None declared.

References

- 1 Carinci F, Carls FP, Grasso DL, Pelucchi S, Pastore A. Schwannoma of the parapharyngeal space. *J Craniofac Surg* 2000;**11**:367-70. [11314385] [doi:10.1097/00001665-200011040-00016]
- 2 Cunningham LL Jr, Warner MR. Schwannoma of the vagus nerve first diagnosed as a parotid tumor. *J Oral Maxillofac Surg* 2003;**61**:141-4. [12524624] [doi:10.1053/joms.2003.50046]
- 3 Sawhney R, Carron MA, Mathog RH. Tongue base schwannoma: report, review, and unique surgical approach. *Am J Otolaryngol* 2008;**29**:119-22. [18314023] [doi:10.1016/j.amjoto.2006.08.003]
- 4 Enoz M, Suoglu Y, Ilhan R. Lingual schwannoma. *J Cancer Res Ther* 2006;**2**:76-8. [17998681]
- 5 Mosharafa TM, Kuppersmith RB, Porter JP, Donovan DT. Pathologic quiz case 1. Malignant peripheral nerve sheath tumor of the ethmoidal sinus. *Arch Otolaryngol Head Neck Surg* 1997;**123**:654,656-7. [9193232]
- 6 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**:931-5. [11474457] [doi:10.1053/joms.2001.25035]
- 7 Pfeifle R, Baur DA, Paulino A, Helman J. Schwannoma of the tongue: report of 2 cases. *J Oral Maxillofac Surg* 2001;**59**:802-4. [11429745] [doi:10.1053/joms.2001.24298]
- 8 Budde R, Brehmer D, Cantemir S, Laubert A. Schwannoma of the tongue. *Laryngorhinootologie* 2001;**80**:36-8. [11272245] [doi:10.1055/s-2001-11026]
- 9 Lacosta J, Zabaleta M, Sanchez Del Hoyo A. Extracranial schwannomas. Report of seven cases. *Acta Otorinolaringol Esp* 1999;**50**:587-9. [10619891]
- 10 de Bree R, Westerveld GJ, Smeele LE. Submandibular approach for excision of a large schwannoma in the base of the tongue. *Eur Arch Otorhinolaryngol* 2000;**257**:283-6. [10923945] [doi:10.1007/s004050050241]
- 11 Spandow O, Fagerlund M, Bergmark L, Boquist L. Clinical and histopathological features of a large parapharyngeal neurilemmoma located at the base of the tongue. *J Otorhinolaryngol Relat Spec* 1999;**61**:25-30. [9892866]
- 12 Dreher A, Gutmann R, Grevers G. Extracranial schwannoma of the ENT region. Review of the literature with a case report of benign schwannoma of the base of the tongue. *HNO* 1997;**45**:468-71. [9324502]
- 13 Hibbert J. Laryngology, head and neck surgery. Kerr AG, ed. Scott - Brown's Otolaryngology. 6th ed. Oxford 1997; pp. 8-22.
- 14 Chiapasco M, Ronchi P, Scola G. Neurilemmoma (schwannoma) of the oral cavity. A report of 2 clinical cases. *Minerva Stomatol* 1993;**42**:173-8. [8413099]
- 15 Krolls SO, McGinnis JP Jr, Quon D. Multinodular versus plexiform neurilemmoma of the hard palate. Report of a case. *Oral Surg Oral Med Oral Pathol* 1994;**77**:154-7. [8139833] [doi:10.1016/0030-4220(94)90278-X]
- 16 Donnelly MJ, al-Sader MH, Blayney AW. Benign nasal schwannoma. *J Laryngol Otol* 1992;**106**:1011-5. [1479267] [doi:10.1017/S0022215100121644]
- 17 Batsakis JG. Tumors of the head and neck. Clinical and pathological considerations. 2nd ed. Williams and Wilkins: Baltimore 1979; pp. 313-33.
- 18 Van der Wall I, Snow GB. Benign tumors and tumorlike lesions of oral cavity and oropharynx. In : Cummings CW, Fredrickson JM, Harker LA, Krause CJ, Schuller DE, eds. Otolaryngology Head and Neck Surgery. 2nd ed. Mosby -Year Book: St. Louis 1993; pp.1237-47.
- 19 Lopez JI, Ballestin C. Intraoral schwannoma. A clinicopathologic and immunohistochemical study of nine cases. *Arch Anat Cytol Pathol* 1993;**41**:18-23. [8517759]
- 20 Flickinger FW, Lozano RL, Yuh WT, Sachs MA. Neurilemmoma of the tongue: MR findings. *J Comput Assist Tomogr* 1989;**13**:886-8. [2778148]
- 21 Nelson W, Chuprevich T, Galbraith DA. Enlarging tongue mass. *J Oral Maxillofac Surg* 1998;**56**:224-7. [9461149] [doi:10.1016/S0278-2391(98)90873-4]
- 22 Gallo WJ, Moss M, Shapiro DN, Gaul JV. Neurilemmoma: Review of the literature and report of five cases. *J Oral Surg* 1977;**35**:235-6. [264530]