

Schwannoma In Tongue

Üzeyir Gök*, İrfan Kaygusuz*, Şinasi Yalçın*, Erol Keleş*, Turgut Karlıdağ*, Nihat Susaman*

* Fırat University Medical Faculty, Department of Otorhinolaryngology, Elazığ, TURKEY

Schwannoma originates from neuroectoderm, derived from Schwann cells of myelin coats, and it is hard, with smooth borders, encapsulated, slowly growing and generally a yellowish coloured benign tumour. Etiology is unknown. It is most frequently observed in the head and neck. A majority of intracranial cases originate from the 8th nerve. It is very rarely found in oral cavity, especially in the tongue. Treatment is to remove the mass surgically. Since 6 cases were reported in English literature, clinical presentation and treatment of a Schwannoma case in the tongue of an 18 year old female patient was presented.

Key words: Schwannomma, Tongue

Dilde Schwannoma

Schwannoma, sinir kılıflarının Schwan hücrelerinden kaynaklanan, nöroektodermden ori jin alan, sert, düzgün sınırlı, kapsüllü, yavaş büyüyen ve genellikle sarımtırak renkte iyi huylu bir tümördür. Etiyolojisi bilinmemektedir. En sık baş ve boyun bölgesinde görülür. İntrakranial olguların büyük bir bölümü 8. kran ial sinirden kaynaklanmaktadır. Oral kavitede ve özellikle dilde oldukça nadir görülür. Tedavisi kitlenin cerrahi olarak çıkarılmasıdır. İngilizce literatürde dilde yerleşimli sadece 6 vaka bildirildiğinden, bu makalede 18 yaşında bir bayan hastada dilde schwannoma olgusunun kliniği ve tedavisi literatür bilgisiyle sunuldu.

Anahtar kelimeler: Schwannoma, Dil.

Schwannomma (neurilemmoma) is a benign tumour originating from neuroectoderm and derived from Schwann cell coat of motor and sensory nerves.¹ First, it was identified by Virchow in 1908.² Of all the Schwannoma cases, 25-40% take place in the head and the neck.³ More than 90% of intracranial Schwannomas originate from the 8th nerve.⁴ Areas, in which the head and the neck Schwannommas apart from acoustic neur inommas are localised, the neck (44%), the parotid gland (7%), the cheeks (7%), the tongue (6%), the haired skin (5%) and the nose (3%). ⁵ It is very rarely seen in the oral cavity. ⁶ Although they are found in all age groups, they are mo re often observed in the second and the third decade of life and do not have any discrimination between the sexes. ⁷

Since Schwannoma in the tongue is rare, a local Schwannoma in the tongue of a femal e patient was presented in this report.

CASE

In May, 1998, an 18 years-old- female applied to out-patient clinic of the ear nose throat department, with a mass on the left side of her tongue appeared approximately two months ago. The patient had a history of a mass, which was very small at onset and gradually became larger. Pain at the time of swallowing was described by the patient. At physical examination, a mass with slight pain in palpation, endurated, hard, with smoot h surface, in dimensions of 2X3 cm near the root of tongue on the left side (Figure 1). There was no abnormality in laboratory tests and conventional radiographs. The patient underwent an operation under general anae sthesia on May 28, 1998 and the mass was removed together with its capsule. In histopathological investigat ion of the mass, Verocay bodies consisted of nucleoli with polizad sequence in Antoni A zones (Figure 2). Therefore, it was diagnosed as Scwa nnoma Postoperatively, there was no functional deficit in the tongue.

DISCUSSION

Schwannoma, originating from neuroectoderm and derived from Schwann cells of myelincoats, is hard, with smooth borders, capsulated, slowly growing and generally a yellowish coloured benign tumour. ¹ It can appear alone or is associated with Von Recklinghausen disease. ⁸ In general, it takes place in the head and neck, and its presence in the mouth is very rare. ¹ They are rarely seen in the tongue and the base of mouth. ⁶ Apart from these, they could also occupy palatine, cheek mucosa, lips and gingiva, subsequently. ⁷

Etiology is still unknown, and the disease is generally asymptomatic.1 In general, it starts as a capsulated nodule and grows slowly. If it invades submucosal areas, it leads to pain and discomfort. 7, 9 Although encountered, lipid accumulation Schwannoma, cystic degeneration and hemorrhage into the cyst may be observed. Malignant transformation is mentioned in 8-10% of the cases.8 Schwannoma has several histological features; two different cellular zones, such as Antoni type A and type B, are seen in histological examination. Antoni A zone has parallel formed thin reticulin fibres, fusiform shaped cells and curled nucleu s. In general, the zone includes a variety number of different cells without an apparent borders amongst their cytosols. Amongst the sheats, there are acellular eosinophilic bodies called as Verocay bodies formed by thin cytoplasmic fibres. 1,10 Antoni B zone is more regular area with less differentiated cells taken place amongst the Antoni A zones. Many findings indicate that these zones are only degenerated Antoni A zones. 9

Lopez and his colleagues ¹¹ reported a serial cases of nine, including three in oral vestibules, two in the tongue, two in palatine, one in oral base and one in lower lip. Nakayama et al ⁶ reported one case, Rocca et al ¹² one and Kroll et al ⁷ one case reported Schwannoma in the tongue. Additionally, Okura and his colleagues ⁴ reported 42 cases whose hypoglossal nerves were invaded.

It is necessary to discriminate between Schwannoma and neurofibroma. However, some authors indicate that both cases are derived from Schwann cells and peripheral connective cells. On the other hand, some other authors suggest that Schwannomas are derived from Schwan cells, and neurofibromas are derived from perineural cells. ¹ Electron microscopy may help to discriminate Scwannomas from neurofibromas. ⁵



Figure 1. Schwannoma in the tongue appearance.

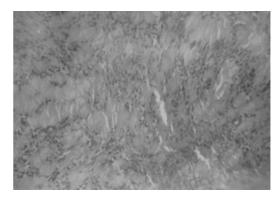


Figure 2. Histopathological investigation of the mass, Verocay bodies consisted of nucleoli with polizad sequence in Antoni A zones HEx200.

Schwannoma is diagnosed by histopathology in biopsy materials. ^{1, 12} Magnetic resonance imaging and computerised tomography are helpful in diagnosis. ^{3, 4, 13} These tumours are radioresistant. Surgical resection is only the treatment method to be chosen in all masses with marked sizes and giving symptoms. ¹⁴ Surgical removal of Schwannoma in the tongue is generally easy. Prognosis after surgical resection is good and remission is fairly rare. ¹⁵

Schwannoma In Tongue

In the presented case, complete surgical excision of Schwannoma in the tongue is sufficient for the treatment. Remission may be possible, unless the mass is not removed completely with its capsule. Schwannoma should be considered for differential diagnosis of hard masses with smooth surface, which is growing slowly in the tongue.

REFERENCES

- Chiapasco M, Ronchi P, Scola G. Neurilemmoma (schwannoma) of the oral cavity. A report of 2 clinical cases. Minerva Stomatol 1993; 42: 173 -78.

 Mosharrafa TM, Kuppersmith RB, Porter JP, Donovan DT. Pathologic quiz case
 1. Arch Otolaryngol Head Neck Surg 1997; 123: 654 -57.

 Younis RT, Gross CW, Lazar RH. Schwannomas of the paranasal sinuses. Arch Otolaryngol Head Neck Surg 1991; 117: 677 -80.

 Okura A, Shigemori M, Abe T, Yamashita M, Kojima K, Noguchi S. Hemiatrophy of the toract days the produced schwannome house. 1-
- 2-
- of the tongue due to hypoglossal schwannoma shown by MRI. Neu roradiology 1994; 36: 239-40.

 Adler ET, Cable BB, Casler JD, Greenspan RB. Burnun dış yüzünde kütle.
- Nature Er, Cabe DB, Casser JD, Giechard RD. Damini us yuzunde kule. Otolaringoloji Arşivi 1997; 15: 43-44. Nakayama H, Gobara R, Shimamoto F, Kajihara H. Ancient schwannoma of the oral floor and ventricular portion of the tongue: a case report and review of the literature. Jpn J Clin Oncol 1996; 26: 185-88.

- Krolls SO, McGinnis JP, Quon D. Multinodular versus plexiform neurilemoma of the hard plate. Report of a case. Oral Surg Oral Med Oral Pathol 1994; 77: 154 -57. Hibbert J. Laryngology and head and neck surgery. In: Kerra AG, ed. Scott -Brown's Otolaryngology. Sixty edition. Vol 5, Oxford, 1997; 16: 8, 22: 6. Donnelly MJ. Al Seder MH, Blayney AW. Benign nasal schwannoma. J Laryngol
- Otol 1992; 106; 1011-15.
- 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.. St. Louis: Mosby -Year Book, 1993: 1237-47.
- Dook, 1993: 1237-47. Lopez JI, Ballestin C. Intraoral schwannoma. A clinicopathologic and immunohistochemical study of nine cases. Arch Anat Cytol pathol 1993; 41: 18
- Rocca A, Leopoldi N, Bonneau HP, Ferra M. Lingual schwannoma. Apropos of 2 clinical cases. Ann Chir Plast Esthet 1993; 38: 167-71.
- cimicai cases. Ann Chir Piast Estrict 1993; 3:107-11.

 Sato M, Kanai N, Fukushima Y, Matsumoto S, Tatsumi C, Kitamura K, Ozaki M, Hayakawa T. Hypoglossal neurinoma extending intra and extracranially: case report. Surg Neurol 1996; 45: 172-75.

 Biliciler N, Gülle A, Cevanşir B, Başerer N, Güldike n Y, Oğuz A. İki olgu nedeniyle parafarengeal bölgedeki nörojenik tümörlerden schwannoma. Türk ORL
- Arsivi 1989: 27: 82-84.
- Arşıvı 1909; 27: 02-04. Yığıtbaşı OG, Ünlü Y, Kontaş O, Cüreoğlu S. Servikal sempatik zincir schwannoması. Türk ORL Arşivi 1993; 31: 57 -58

Correspondence Address:

Dr. Üzeyir GÖK

Fırat Üniversitesi Fırat Tıp Merkezi KBB Kliniği

23200 ELAZIĞ, TURKEY

Phone: 90. 424. 233 35 55 Fax: 90. 424. 238 80 96

E-mail: firdickbb@firat.edu.tr