CASE REPORT

Schwannoma of the epiglottis: case report focusing on clinico-pathological aspects

Schwannoma dell’epiglottide: descrizione di un caso con considerazioni clinico-patologiche

V. SAITA, A. AZZOLINA, A. GALIA, F. FRAGGETTA
Phonosurgery Unit; Pathology Unit, “Cannizzaro” Hospital, Catania, Italy

Key words
Larynx • Benign tumours • Schwannoma • Epiglottis • Therapy • Differential diagnosis

Summary
Laryngeal schwannomas are uncommon lesions with only few cases reported. Herein we present a further case of a schwannoma of the epiglottis, occurring in a 62-year-old with a clinical history of a cutaneous malignant melanoma and laryngeal glottic keratosis. The schwannoma was incidentally discovered as a small polypoid lesion located on the laryngeal surface of the epiglottis and was removed endoscopically. The procedure was uneventful and the patient is well six months later. Authors focus on the diagnostic and therapeutic options for this unusual lesion and discuss the differential diagnosis of the spindle cell proliferation of the larynx.

Parole chiave
Laringe • Tumori benigni • Neurilemmoma • Epiglottide • Terapia • Diagnosi differenziale

Introduction
Neurogenic tumours of the larynx are rare with only a few cases reported in the literature. Benign (namely, schwannoma and neurofibroma) and malignant (namely, malignant peripheral nerve sheath tumour) neurogenic tumours of the larynx may occur as incidental lesions or in a setting of clinical syndromes, including neurofibromatosis. In particular, schwannomas account for 0.1% of all benign neoplasms of the larynx and are usually located in the aryepiglottic folds. Herein, we report an additional case of a small schwannoma located on the laryngeal surface of the epiglottis. Clinical and pathological features of this unusual lesion are discussed.

Case report
A 62-year-old male, with a clinical history of a malignant cutaneous melanoma, underwent a laryngoscopic exam on account of a long history of dysphonia. The patient had been treated endoscopically for bilateral laryngeal keratosis one year before. The examination revealed a polypoid lesion, measuring 0.5 cm, in the maximum diameter, located on the laryngeal surface of the epiglottis. The lesion was well circumscribed and was removed endoscopically and sent for histological examination. The procedure was uneventful and the patient was discharged a few hours later.
other antibodies tested. On the basis of morphological and immunohistochemical findings, a final diagnosis of laryngeal schwannoma was made.

Discussion

Mesenchymal tumours of the larynx are unusual with cartilaginous neoplasms being the most common. The neurogenic group is the most rare with the schwannoma representing 0.1% of all benign laryngeal neoplasms. It has been postulated that these neoplasms may arise from the internal branch of the superior laryngeal nerve and may have an insidious clinical course. Laryngeal schwannomas may approach a large size, causing upper airway obstruction, dysphonia and even vocal cord fixation, depending on their location. Recognition and treatment of laryngeal schwannomas is mandatory. In our case, the lesion was an incidental finding during scheduled laryngeal endoscopy, at clinical follow-up, on account of a laryngeal keratosis. Endoscopic recognition of laryngeal schwannoma is difficult or even impossible because of the rarity of the lesion and of the non-specificity of the endoscopic findings. Because of the small size of the lesion, endoscopic excision, in our case, had a diagnostic and therapeutic effect.

It has been suggested that imaging studies may be helpful in defining laryngeal schwannoma. Although magnetic resonance imaging is not diagnostic, CT scan images often exhibit heterogenic density, on contrast enhancement, with centrally distributed areas of low attenuation, surrounded by a peripheral enhancing ring thus suggesting a possible diagnosis of schwannoma. In these cases, an external surgical approach is indicated. This includes median thyrotomy, lateral pharyngotomy or lateral thyrotomy depending on the location and size of the neoplasm.

Faced with a spindle cell proliferation, the pathologist has to take into consideration benign and malignant conditions in the differential diagnosis, both epithelial, and mesenchymal. Neurofibroma is a non-encapsulated lesion consisting in a proliferation of schwann cells associated with strands of collagen; moreover, S-100 immunoreactivity is usually weaker than in neurilemmoma. From a clinical viewpoint, differential diagnosis is very important since evidence of a neurofibroma should lead the clinician to the exclusion of a neurofibromatosis.

Among the malignant lesions, spindle cell carcinoma and malignant melanoma have also to be excluded. Although sarcomatoid carcinomas usually show marked cellular atypia, necrosis and mitosis, they may also present as polypoid lesions showing minimal or no pleomorphism. This is supported by the fact that lesions referred to as benign neurogenic tumours were then considered to be spindle cell carcinomas, after histological reviewing. Meticulous clinical, morphological and immunohistochemical findings are of importance, in making the correct diagnosis.

Malignant melanomas, primary or metastatic, have to
be excluded, especially in those patients with a previous clinical history of melanoma. Laryngeal melanomas present as polypoid pigmented lesions, histologically composed of a proliferation of round to spindle cells showing marked atypia, mitoses and consistent immuno-staining for S-100 protein and HMB-45. In our case, the proliferation had no atypia and showed no immunostaining for HMB-45.

Since conservative surgery provides excellent results \(^1\), the possibility of a schwannoma should always be taken into account especially when dealing with spindle cell proliferation from a pre-operative biopsy sample. It should not be forgotten, however, that, in some cases, only electron microscopy observations may help to clarify the nature of spindle cell proliferation of the larynx \(^16\).

This case stresses, once again, the possibility of the occurrence of a common lesion (i.e., schwannoma) in an unusual site. Although it is not clear why a common lesion is extremely rare in peculiar sites, we believe that local or micro-ambiental factors may influence their development. Moreover, the possibility that a patient with a previous history of malignant cutaneous melanoma may harbour other neurogenic tumours, at a different site should also be taken into consideration \(^17\).

References


