CASE REPORT

MYOPERICYTOMA OF THE TONGUE: A CASE REPORT

Sevtap Akbulut†, Derya Berk†, Mehmet G Demir†, Sibel Kayahan‡

Dr. Lutfi Kirdar Kartal Training and Research Hospital, Istanbul, Turkey: Department of Otolaryngology Head and Neck Surgery†, Department of Pathology‡

Summary: Myopericytoma is a rare benign tumour composed of pericyctic cells that show myoid differentiation and have a tendency for concentric perivascular growth. It belongs to a spectrum of perivascular myoid cell neoplasms. To date, only a small number of cases of myopericytoma involving the oral cavity have been reported. We describe a case of myopericytoma presenting as a slowly growing tongue nodule in a 61-year-old woman. A diagnosis of myopericytoma was established with the histopathological findings combined with immunohistochemical staining. Myopericytoma should be included in the differential diagnosis of well-circumscribed, slow-growing lesions of the oral cavity.

Key words: Myopericytoma; Tongue; Oral cavity; Histopathology

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Introduction

Myopericytoma is a benign mesenchymal tumor of pericytic cells demonstrating myoid differentiation (1). Myopericytoma is composed of oval to spindle-shaped myoid-appearing cells with a con to extend the myringotomy patency centric perivascular growth (2). It mostly arises within the skin and subcutaneous tissues of extremities (3). Myopericytoma of the oral cavity is extremely rare, only several cases have been reported in the literature (2, 4–7). To the best of our knowledge, there is only two reported cases of myopericytoma localized to the tongue (4, 5).

Case History

A 61-year old woman complained of a slow-growing, painless tongue nodule that had been present for more than a year. Clinical examination revealed 2-cm well-circumscribed firm nodule on the left mid-lateral tongue. The patient’s past medical history was unremarkable. She had been using dental prosthesis for the last 10 years. On MRI there were low signal intensities on T1 weighted images and high signal intensities on T2 weighted images. After the administration of contrast material, the lesion showed homogenous contrast enhancement and invasion of tongue muscles. An excisional biopsy of the nodule was performed for microscopic examination.

Histopathologic Findings

Microscopic examination of hematoxylin-eosin-stained sections revealed a circumscribed benign neoplasm characterized by the presence of cells arranged in concentric islands. The single-walled endothelium-lined vessels were surrounded by a concentric arrangement of myoid-appearing oval-to-spindle cells presenting bland nuclei with eosinophilic cytoplasm (Fig. 1).

Immunohistochemically, the perivascular cells were positive for smooth muscle actin, but negative for desmin and S-100 (Fig. 2). No reactivity for CD34 was evident. The Ki-67 labeling was 5%.

After correlating the immunohistochemical pattern with the histopathological findings a diagnosis of myopericytoma was rendered. No further treatment was necessary as the lesion was totally excised, and 18 months later, there was no evidence of recurrence.

Discussion

Myopericytes are employed as cells of uncertain position in the morphologic spectrum between pericytes and vascular smooth muscle cells, and share many features with myofibroblasts (1, 7–9). Myopericytoma belongs to a spectrum of disease entity called as perivascular myoid cell neoplasm. The spectrum includes myopericytoma, solitary myofibroma, infantile-type myofibromatosis in adults, glomangiopericytoma, angioleiomyoma and glomangiomyoma (1). Myopericytoma is known for tumors derived from, or differentiating along myopericytes (9). The defining histologic feature of myopericytoma is a distinctive, concentric perivascular proliferation of myoid tumor cells. These cells are characteristically reactive for smooth muscle actin and negative for CD34, cytokeratin and S100 protein. Vimentin and desmin are usually negative, or only focally positive, in the myopericytoma, which helps to distinguish it from other perivascular myoid neoplasms (1, 7, 8).

Myopericytoma presents as slow-growing, well-circumscribed nodule with less than 2 cm in size. It most frequently affects the skin and superficial soft tissues of the distal extremities in adults; but with increased recognition, lesions in
head and neck region and trunk have been described (3, 7).

There are very few oral lesions reported. The very first case of myopericytoma involving the oral cavity was described by Datta et al. as a nodule on the tongue (4). Afterwards Lau et al. reported two cases with myopericytoma occurred in patients with HIV infection, among which one presented with a tongue mass (5). Sapelli et al. reported a case of myopericytoma originated from the lip (6).

Ide et al. described a rare intravascular type of myopericytoma presenting as a mass in buccal mucosa (2). After a clinicopathologic re-evaluation of 35 perivascular myoid tumors, Ide et al. revised the pathologic diagnosis as myopericytoma in 2 more cases involving the lips which were originally diagnosed as angioleiomyoma. They suspected that more intra-oral myopericytoma could be found in the literature under the diagnosis of angioleiomyoma (7, 10). Because of morphologic similarity between myopericytoma and other perivascular myoid neoplasms, it was supposed that myopericytoma might be under-recognized by surgical pathologists.

Most cases of myopericytoma behave in a benign fashion, although a few malignant and/or recurring cases were described (6, 9). In the oral cavity only a case of myopericytoma of low grade malignancy was reported (11).

**Conclusion**

Myopericytoma is a rare benign tumor which shares morphological features with other perivascular myoid cell neoplasms. Histologic findings should be confirmed with appropriate immunohistochemical staining. Myopericytoma should be kept in mind in the differential diagnosis of nodular lesions in the oral cavity. The authors presented a rare case of myopericytoma in the tongue.

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**References**


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Sevtap Akbulut, M.D., Dr. Lutfi Kirdar Kartal Eğitim ve Arastırma Hastanesi KBB Bolumu, Denizer Cd., E-5 Karayolu Cevizli Mevki, 34890, Cevizli, Istanbul, Turkey; e-mail: sevtap.akbulut@gmail.com