Auricular Angioleiomyoma: A Case Report

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ABSTRACT

Angioleiomyoma (vascular leiomyoma or angiomyoma) is a rare, benign smooth muscle tumor that originates in the tunica media of blood vessels. These tumors may be found anywhere in the body. They usually occur in the lower extremity. Auricular angioleiomyoma is very rare, and only a few cases have been reported. We describe here a 38-year-old male patient with angioleiomyoma on the left auricular helix.

Key Words: Angioleiomyoma, Vascular, Leiomyoma, Ear

INTRODUCTION

Angioleiomyomas account for 5% of all benign neoplasms of soft tissues. They occur mostly in the lower extremity, some of them have been described in the head and neck region. There are only a few cases of auricular angioleiomyoma reported in the literature. The tumor was located over the helix, pinna, and lobule in these reports (1-4). Hachisuga et al. reported a large series of angioleiomyoma including 562 patients, where only 14 cases (2.8%) were in the auricle (5,6). We describe here a 38-year-old man who presented with an unusual lesion on the left auricular helix.

CASE REPORT

A healthy 38-year-old man presented to the ear-nose-throat surgery clinic at our hospital with a small, intermittent painful, slowly growing nodule located on the helix of his left ear. He had first noted the lesion 13 years ago. No other similar lesion was found on his neck, face and extremities. His medical and family histories were unremarkable. Clinical diagnosis was a vascular malformation. The lesion was completely excised under local anesthesia. Grossly, the specimen was an ovoid, firm, white-gray mass, measuring 1.0x1.0x0.7 cm. The cut surface was solid and white-gray. Microscopically, the tumor was composed of a well circumscribed proliferation of smooth muscle bundles that surrounded numerous thick-walled blood vessels with partially patent lumina (Figure 1). The smooth muscle cells showed no cellular atypia or mitoses. In immunohistochemical studies, the perivascular proliferating smooth muscle cells and the muscular wall

Figure 1: The tumor mass is composed of well-differentiated smooth muscle cells and numerous thick-walled vessels with some dilated vascular channels (H&E; x40).

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Grossly, the tumors are circumscribed, glistening, white-gray nodules. Microscopically, the tumors have a characteristic appearance that varies little from case to case. The usual appearance is a well-demarcated nodule of smooth muscle tissue punctuated with thick-walled vessels with partially patent lumens. Typically, the inner layers of smooth muscle of vessels are arranged in an orderly circumferential fashion and the outer layers spin or swirl away from the vessel, merging with the less well-ordered peripheral muscle fibers. The vessels in these tumors are difficult to classify as veins or arteries (7). Various stains have been used to identify vascular leiomyomas, including; desmin, vimentin, actin, myosin, and Masson's trichrome.

DISCUSSION

Benign tumors of smooth muscle cells can be categorized in three clinicopathologic varieties: 1- Cutaneous leiomyomas (pilar leiomyoma, genital leiomyoma); 2- Angiomyoma (vascular leiomyoma or angioleiomyoma); 3- Leiomyoma of deep soft tissue (somatic leiomyoma, gynecologic leiomyoma) (7).

Angioleiomyomas, a subtype of leiomyomas, are rare, benign, smooth muscle tumors that arise from tunical media layer of small arteries and veins (5). Leiomyoma was published by Aufrecht in 1868 and it was the first report of a case arising from blood vessels’ wall. Angioleiomyoma is classified into three histological types. 1-Capillary or solid: the most common type, which is closely compacted smooth muscle and many small, slit-like vascular channels. 2-Venous: thick, easily identifiable muscular walls distinguish this type. 3- Cavernous: the vascular channels are dilated with less smooth muscle. This is the least common of the three types (5,8).

It may occur anywhere in the skin or subcutaneous tissue. The tumors are usually small (2 to 15 mm), solitary, round, firm, skin-colored, and well-encapsulated (1).

Hachisuga reviewed 562 cases. Among those cases, 205 cases were male and 357 were female, with a ratio of 1:1.7. The peak incidence was between the fourth and sixth decades (5). Most common anatomical sites are lower extremity (67%), upper extremity, head and trunk. Solid angioleiomyoma occurs most frequently in the lower extremity of females. The venous type is the most frequent type found in the head and neck region usually in males, as occurred in our case. Pain and tenderness are the most characteristic subjective complaints in patients with angioleiomyoma (67%) (5,6). Exposure to wind, cold, local pressure and other imperceptible stimuli cause attacks of paroxysmal pain. Painful tumors are the most frequent type of solid tumors. Tumors occurring in the head and neck regions are usually not accompanied by pain (5).

The clinical presentation of angioleiomyoma is nonspecific. It is not easily differentiated from other benign subcutaneous tumors. The diagnosis of this lesion is therefore based on excisional biopsy (3).
In our patient, the tumor intensely stained with smooth muscle actin and Masson's trichrome.

Recurrence after excision is rare (5). Malignant changes have been reported in recurrent tumors (9). There is one case report that an angioleiomyoma can occur in association with a leiomyosarcoma (10).

In this article, we describe a case of angioleiomyoma arising from the auricle. This lesion should be considered in the differential diagnosis of nodular masses of the auricle. The pathological differential diagnosis must include subcutaneous leiomyoma, hemangioma, angiofibroma, fibromyoma, leiomyoblastoma, angiomyolipoma, vascular leiomyosarcoma and other tumors of perivascular cells (glomus tumors and myopericytoma) (11,12).

REFERENCES