Maxillary’s cystic lymphangioma: A rare location

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Abstract

Lymphangiomas are benign hamartomas of the lymphatic vessels. They are congenital malformations that are usually present at birth or diagnosed within the first two years of age. Almost 75% of lymphangiomas occur in the head and neck region. Its occurrence in the maxillary antrum is one of the rarest locations. Very few cases have been reported in the literature. We report a case of maxillary sinus’ lymphangioma in a 73 year old female patient which presented for a 2 years evolving 3 cm right cheek swelling. Complete surgical excision was done without later recurrence.

Keywords: Lymphangiomas; Hamartoma; Maxillary sinus.

Introduction

Lymphangiomas are developmental malformations that arise from the proliferation of the sequestered lymphatic tissues during embryogenesis. It is a benign congenital lymphatic malformation [1]. It is usually diagnosed in children under 2 years old [2]. In adults, cystic lymphangioma (CL) is a fairly rare disease [3]. Its etiology remains unclear, but it is supposed to be congenital or to result from the lymph fluid pathway obstruction and its retention in the developing lymphatic vessels. Head and neck is the main region affected probably due to the rich lymphatic network in this area. Usually lymphangiomas are asymptomatic, painless and slow growing. Associated symptoms result rather from the nearby structures local compression [4,5]. Treatment options of these lymphangiomas include surgical excision when possible, cryotherapy, sclerotherapy, laser photocoagulation, chemotherapy and radiotherapy [6].

Case presentation

A 73-year-old female without past medical history presented to otolaryngology department for a right-sided painless cheek swelling, hypoesthesia and anosmia. She does not complaint of associated symptoms such epistaxis or nasal obstruction. This symptomatology dated back to 2 years before her first consultation.

Physical examination showed the right cheek mass with an intact overlying skin. It was tender in consistency and there was no associated lymph nodes on the neck palpation. A decreased sensitivity in the V2 nerve territory was noted without any facial nerve deficiency. Nasal endoscopy showed a polypoid mass of the right middle meatus. On CT scan, this mass was described as a right maxillary tumoral process with osteolysis, extension to the anterior subcutaneous tissues, the infra-temporal fossa and to the adjoining nasal fossa (Figure 1). These findings were supported by

Figure 1: Face bones CT scan (axial section): hypodense tumor formation of the right maxillary sinus.
Magnetic resonance imaging (MRI) showing a 4x4x3 cm tumor located in the right maxillary sinus, which was in hypersignal on T1 sequences and iso-signal on T2 sequences with slight enhancement after gadolinium injection (Figure 2).

The patient underwent surgical excision under general anesthesia using Caldwell Luc approach. Intraoperatively, the tumor adhered to cheek muscles, to the orbit bony floor and extends to the infratemporal fossa through a large lysis of the maxillary sinus’ posterior wall. The surgical procedure was completed by a reaming of the different sinus walls and opening of the inter-sinuso-nasal septum. The entire surgical specimen was sent for histopathological examination.

Macroscopically, cysts bigger than 3cm were observed. Microscopically, the mass was composed by a variety of dilated lymph vessels involved in a fibrovascular stroma. The diagnosis of cystic lymphangioma was, then, retained. At the date, 5 years follow-up after surgery, no relapses were noticed.

**Discussion**

Lymphangiomas are vascular malformations that usually occur in neonates and children below 2 years with no sex predilection. According to some authors [3], several hypotheses can explain the origin of this malformation: the first hypothesis suppose a blockage or arrest of the normal primitive lymphatic network growth during embryogenesis. The second hypothesis, suppose that the proliferation of the primitive lymphatic sac is insufficient to reach the venous system. The third one and probably not the last suppose a wrong proliferation of lymphatic tissues in wrong sites during embryogenesis. The incidence of lymphangiomas ranges from 1,2 to 2,8 cases per 1000 live births and account for about 30% of all the lesions that arise from the lymphatic vessels [6]. Head and neck has a marked predilection for the lesion, in which case, it is more localized in the posterior triangle of the neck, submandibular and parotid regions. In the oral cavity, the tongue is the usually affected site followed by the palate, buccal mucosa, gingiva and the lips [7].

Based on the histopathology, lymphangiomas are categorized into simplex (small lymphatic vessels with thin walled endothelium), cavernous (dilated lymphatic vessels with surrounding adventitia) and cystic lymphangiomas (huge lymphatic vessels with surrounding fibrovascular tissues and smooth muscle) [2]. In the indexed literature, only one case of cystic lymphangioma located in the maxillary sinus was reported [9].

**Conclusion**

Lymphangiomas’ maxillary localization of is exceptional. Treatment in this case does not differ from other locations and surgery is preferred when possible. However, recurrence after surgery was reported in 40% of cases, all locations combined. Thus, early diagnosis allows appropriate treatment and reduces recurrence situations. Nevertheless, long-term follow-up is constantly required.
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References


