Hairy polyp of the rhinopharynx – a rare and unrecognized entity

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ABSTRACT

Hairy polyps are tumors that have been described in the literature as congenital masses usually found in the nasopharynx. Clinical symptoms are a consequence of disease dimensions and anatomical positioning. Although rare, this kind of tumors can be mistaken for neurologic impairment of an infant. Histological examination is the base for classification of this malformation. Earlier management of the disease could prevent possible complications (of the airway) and allow a better long-term growth of the patient.

Keywords: hairy polyps, rhinopharynx, tumor

INTRODUCTION

Hairy polyps are tumors that have been described in the literature as congenital masses usually found in the nasopharynx [1]. Their incidence is low [2,3]. Histologically, they contain both ectodermal and mesodermal tissues [2]. The point of insertion into the pharynx can vary: most of the tumors are linked to the soft palate [2], but some authors published palatopharyngeus muscle as the origin point [4].

Clinical symptoms are a consequence of disease dimensions and anatomical positioning. Disordered breathing and dysphagia can be found [5]. Diagnosis can be suspected at clinical examination of the patient and completing the evaluation with imagistics: computed tomography scanning (CT) or magnetic resonance imaging (MRI). Surgical removal is the therapeutic approach and pathologic examination confirms the diagnosis.

There is no clear origin of the tumor but, most of the authors hypothesize the branchial origin of this malformation [6]. Some even presumed these tumors to be the equivalent of accessory auricles trapped into the nasopharynx [7]. In this hypothesis, some authors consider hairy polyps origins to be located in the lower third of the Eustachian tube [8].

CASE PRESENTATION

We present a case of a 1-year-old girl, referred to our department for feeding problems, with intermittent choking, lasting from birth but worsened during the last month. At clinical examination we noted a tumor originating from the posterior face of the soft palate. A computed tomography (CT) scanning showed the tumor advancing into the proximal esophagus for at least 3 cm (figures 1-2).
Under general anesthesia (GA), we found a pedunculated tumor, apparently covered with skin and originating from the posterior left face of the soft palate. It could be surgically removed and its origin cauterized (figure 3) after soft palate retraction, under direct vision. Histology showed keratinizing squamous epithelium covering fat tissue and cartilage. No complications were noted postoperatively. The child recovered quickly and no similar symptoms could be elicited at 1-month follow-up.

DISCUSSION

Although rare, this kind of tumors can be mistaken for neurologic impairment of an infant. Its congenital nature does not bring much light into the differential diagnosis, regarding deglutition troubles after birth.

Good clinical examination in a child can be difficult due to high position of the hyoid-base of tongue complex or the presence of an endotracheal tube [3]. Detailed examination can sometimes need nasopharyngeal fibroscopy and to a much lesser extent general anesthesia.

Tumors noted in the naso-pharyngeal region are usually benign or congenital. Teratomas, foreign bodies, glioma, meningo-encephalocele, lingual thyroid gland, vallecular cysts must all be differentiated from hairy polyps. Histological examination is the base for classification of this malformation.

Also, pedicle insertion point could bring clues to its origin. Our case demonstrated a long insertion pedicle into the left rhinopharynx.

Theoretically, these esophageal polyps, given the long pedicle, could eventually become dislodged from their esophageal position and threaten the airway at the level of the larynx [9,10]. We did not get a history of such symptoms, but only feeding difficulties as the only complaint.

CONCLUSIONS

Although rare, this malformation should be reminded whenever it is present from birth and has a typical appearance (hairy), especially if located posterior to the soft palate. Treatment is by surgical excision. Prognosis is good, with almost no recurrences reported. We think that earlier management of the disease could prevent possible complications (of the airway) and allow a better long-term growth of the patient.
REFERENCES


