Hairy Polyp in the Nasopharynx - A Case Report

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Abstract:
Background: Hairy polyp or dermoid is a benign biregional congenital germ cell tumour. It is a rare condition in the nasopharynx. Hairy polyp in head and neck region consists less than 5% of that occurs in whole body.
Objective: To report a rare nasopharyngeal benign condition presented without obstructive symptoms.
Case: A four-year-old female child presented with a pedunculated dermoid tumour attached to the left lateral wall of nasopharynx. There was an associated unilateral cleft in soft palate. The tumour was removed by dissecting with bipolar diathermy by combined endonasal endoscopic and transoral approach.
Conclusion: Hairy polyp in nasopharynx may present as an incidental asymptomatic mass in throat and may be associated with a cleft palate.

Introduction:
Hairy polyp or dermoid is a benign biregional congenital germ cell tumour composed of ectodermal and mesodermal elements1. Less than 5% of it occurs in head and neck region2. Its occurrence in nasopharynx is rare3,4. It was first reported in 17843,5. Till 2012 a total of 170 cases in nasopharynx were reported6. Usually it presents due to its obstructive features – breathing or feeding difficulties1,7. It usually presents in the first year of life; but it may present within hours to eighth decade of life3,4,8,9. Hairy polyp is associated neither with a particular congenital syndrome nor with genetic abnormality. But rarely it may be associated with other congenital malformations such as cleft palate, absent uvula, auricular deformities, facial hemihypertrophy, ankyloglossia, osteopetrosis and atresia of carotid artery7,8,10. Hairy polyps are successfully treated by simple surgical excision. With adequate removal these lesions do not or rarely recur5,8,9. A combined transoral and nasoendoscopic surgical resection offer better outcome without any late complication as compared to blind resection11.
Case Report:
A four-year-old female child presented with the complaints of her mother of having a rounded mass in the throat for about six months. There was no breathing or feeding difficulties. The mass was grayish-white in colour, smooth, rounded and hard in consistency (Fig.-1). It was pedunculated and attached to the left lateral wall of nasopharynx. From nasopharynx it was protruding into the oropharynx. There was an associated unilateral cleft in the left side of the soft palate (Fig.-2). CT scan of nasopharynx revealed a soft tissue shadow with hyperdense area in the nasopharynx (Fig.-3). The tumour was removed by dissecting with bipolar diathermy by combined endonasal endoscopic and transoral approach retracting the soft palate. The surface of the removed mass was grayish-white and smooth. After sectioning the cut surface was variegated with different types of soft tissue within it. There were two well-formed teeth like structures and bones within it (Fig.-4). Histological examination revealed that the tumour was covered with keratinizing squamous epithelium with epidermal appendages and the stroma contained fat, fibrous tissue, cartilage, bone and tooth like structures. There was no evidence of any endodermal element.

Fig.-1: Smooth rounded mass seen in oropharynx after retracting the soft palate.

Fig.-2: A cleft in the left side of the soft palate.

Fig.-3: An isodense shadow in the nasopharynx with an irregular hyperdense area within it.

Fig.-4: Bones and well-formed tooth-like structures
Discussion:
Hairy polyp or dermoid in nasopharynx is a rare condition. It was first reported in 1784 by Ford as quoted by Brown-kelly. Arnold in 1870 classified congenital germ cell tumours into 4 categories: Teratomas, Teratoids, Epignathi, and Dermoids. Teratomas consist of 3 germ layers (ectoderm, mesoderm, and endoderm). The cells and tissues are differentiated so that organ can be recognized histologically. Teratoids are also trigeminal but are composed of poorly differentiated tissues. Epignathi demonstrate highly differentiated trigeminal layers in which well-formed organs and limbs of a parasitic fetus (fetus in fetu) are present. Dermoids are derived from only two germ layers (ectoderm and mesoderm). These tumours can present at any age, but mostly present in the first year of life. Females are more frequently affected (F:M = 6:1). Most of reported cases were on the left side what is seen in the present case as well. The reason for predilection for females and left side is unknown. The present case is a four-year-old female child who presented with an incidental finding of a mass in the throat. But usual presentation is feeding or breathing difficulties and very rarely needs tracheostomy. Although it is associated neither with a particular congenital syndrome nor with genetic abnormality, rarely it may be associated with other congenital malformations such as cleft palate, absent uvula, auricular deformities, facial hemihypertrophy, ankyloglossia, osteopetrosis, and atresia of carotid artery. The present case has a small unilateral cleft in the left side of the soft palate. Hairy polyps are successfully treated by simple surgical excision. With adequate removal these lesions do not or rarely recur. In the present case the tumour could be easily and completely removed by bipolar diathermy dissection using combined endonasal endoscopic and transoral approaches by retracting the soft palate. No peroperative or postoperative consequences were observed.

Conclusion:
Hairy polyp in nasopharynx may present as an incidental asymptomatic mass in throat and may be associated with a cleft palate.

References:


