CASE REPORT
GIANT HAEMANGIOMA OF NASOPHARYNX: A RARE CASE

Haissan Iftikhar, Moghira Iqbaluddin Siddiqui
Department of Otolaryngology / Head and neck surgery, Aga Khan University, Karachi-Pakistan

A 60-year-old female presented to us with a 5-year history of progressive right sided nasal obstruction and recurrent epistaxis. On examination lesion was seen in the right nostril which was firm and bled on probing. CT-scan paranasal sinuses showed a right sided lesion of nose and naso-pharynx obstructing posterior nasal choanae. Dimensions were reported to be 9×4.1×3.8 cm. A punch biopsy was taken in operating room under general anaesthesia which resulted in profuse bleeding. Suction cautery was used to control bleeding and the nose was packed. The mass was firm to hard and provided resistance during the time of biopsy. On second post-operative day, the mass visibly changed its colour and became blackish in appearance. Patient had an episode of cough in the evening and expelled out the entire mass orally.

**Keywords:** Hemangioma; Head and neck; Otolaryngology

INTRODUCTION

Although the hemangiomas can arise anywhere in the head and neck region from the skin to deeper structures such as muscles and bone, hemangiomas arising in the nose are rare. 80% arise from the kiesselbach’s plexus and 15% from the lateral wall of the nasal cavity. Nasal bleeding with obstruction can be associated with benign and malignant lesions. The exact origin of haemangiomas is still debatable. Some have proposed embryonic unipotent cells to be involved while others advocate vascular elongation. Infantile haemangiomas however ultimately represent endothelial cell proliferation.

Haemangiomas are classified on the basis of the vessels they contain: Infantile haemangioma (HOI), (cavernous, capillary) or mixed. Capillary haemangioma are common and are seen in childhood which spontaneously regress and do not need any intervention unless they are symptomatic. However infantile hemangioma (HOI) is usually seen in adulthood and warrants an intervention if symptomatic.

In our case the differentials were nasopharyngeal carcinoma, inverted papilloma and HOI.

CASE REPORT

A 60-year-old female presented to our outpatient clinic with a 5-year history of progressive right sided nasal obstruction and epistaxis which was recurrent and profuse. On nasal examination, the lesion was seen in the right nostril which was firm and bled on probing, oral examination showed bulge in soft palate and a mass in nasopharynx. Blood counts were done which revealed no haematological disturbance and normal haemoglobin. CT-scan head and neck with intravenous contrast was done which revealed a lesion occupying the right side of nose, nasopharynx and extending up to the oropharynx (Figure-1) with the dimensions of 9×4.1×3.8 cm. Patient was electively taken for examination under anaesthesia and biopsy. A punch biopsy was taken in operating room under general anaesthesia which resulted in profuse bleeding. Suction cautery was used to control bleeding and the nose was packed. The mass was firm to hard and provided resistance during the time of biopsy. On post-operative day 2 patients had an episode of cough which led to expelling a large lesion out of her mouth (Figure-2). Biopsy was consistent with haemangioma of infancy. The patient was then symptom free and was subsequently discharged the next day.
DISCUSSION

Haemangioma of infancy are slow growing lesion and when symptomatic cause epistaxis, nasal obstruction and at times haemoptysis. The first two symptoms are consistent with our patient. The types of haemangioma vary histologically but have no clinical significance of their histological variation. Mean age of presentation of HOI (previously named cavernous haemangioma) is in 40s whereas the sex predilection appears to be equal. The underlying bone is otherwise normal but might be re-modelled due to pressure effect. MRI images show low intensity signal on T1-weighted images and very high intensity signals on T2-weighted images. Angiography is another modality that not only offers intensity signals on T2-weighted images.

The main stay of treatment for haemangioma of Sino nasal anterior skull base remains surgery which might include arterio-embolization preoperatively. Angioembolization minimizes blood loss intra-operatively. In our case angioembolization was not considered as we planned a biopsy initially. The first successful case of endoscopic removal of Sino nasal infantile haemangioma was in 2003. There is now growing evidence and positive outcomes for treatment of Sino nasal tumours: benign and malignant lesions endoscopically.

What makes our case special is its size and auto-expulsion post biopsy. In an attempt to control bleeding intra-operatively we probably were able to cauterize the main blood supply of the mass which resulted in necrosis and the tumour dropped down into the oropharynx.to the best of our knowledge we report the first ever case of such expulsion of sino nasal haemangioma post-operatively.

AUTHORS’ CONTRIBUTION

HI had substantial contributions to the conception, design of the work and the acquisition, analysis, and interpretation of data for the work. MIS had critically revised the manuscript for important intellectual content and final approval of the version to be published.

REFERENCES


Address for Correspondence:
Dr. Moghira Iqbaluddin Siddiqui, Department of Surgery, Section of Otolaryngology, Aga Khan University Hospital, Karachi-Pakistan
Cell: +92 306 2463146
Email: moghira.siddiqui@aku.edu

Received: 4 January, 2017
Revised: --
Accepted: 12 February, 2017