CASE REPORT

Pedunculated haemangioma of the palate

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SUMMARY
Haemangiomas are the most common tumours of vascular origin of the head and neck region and can appear anywhere in the oral cavity. They are benign, appear in early childhood and usually involute as the child grows older. The dental surgeon should manage them appropriately if they are symptomatic. This report presents a rare case of a pedunculated haemangioma in a 4-year-old child.

BACKGROUND
Intra-oral soft tissue swellings are often overlooked and provisionally considered as inflammatory lesions. Clinicians should be familiar with such unusual presentations so that they are treated appropriately.

CASE PRESENTATION
A 4-year-old girl reported to the Department of Paediatric Dentistry with a swelling on the palate. According to her parents, an initially small lesion had gradually increased in size and was painless but impaired her speech. Systemic examination revealed that the child was normally built for her age and there was no medical, dental or family history of relevance. On intra-oral examination, a single swelling was seen on the palatal midline. It was pedunculated, reddish pink in colour, lobulated, soft and measured about 3×3×2 cm in diameter (figure 1). Mild blanching was noticed on applying pressure.

INVESTIGATIONS
A routine haematological examination performed according to the institutional protocol for undergoing minor surgical procedures was normal.

DIFFERENTIAL DIAGNOSIS
In view of the colour, consistency, lobulated surface and pedunculated appearance of the swelling, a differential diagnosis of pyogenic granuloma, haemangioma and squamous papilloma was made.

TREATMENT
The lesion was excised under local anaesthesia. There was more profuse bleeding from the site than expected which was stopped using electrocautery. A periodontal pack (zinc oxide eugenol paste) was placed to prevent clot dislodgement.

OUTCOME AND FOLLOW-UP
The excised specimen underwent histopathological evaluation. The microscopic picture showed stratified squamous epithelium with associated fibrovascular connective tissue. The stroma had multiple small- and large-diameter well-developed vascular channels lined with endothelial cells, some of which were congested with red blood cells (figure 2). No inflammatory component was seen. Histopathology was suggestive of infantile haemangioma.

The post-surgery period was uneventful. At 1-week follow-up, the parents reported that the child’s speech was improving. The patient presented again after 8 months with another similar lesion, but smaller in dimension (1×1 cm) and on the antero-lateral aspect of the palate (figure 3). No action was taken as it was asymptomatic, but it was placed under observation.

DISCUSSION
Haemangiomas are relatively common benign proliferations of vascular channels that may be present at birth or arise during early childhood.1 Currently, haemangiomas are considered to be benign tumours of infancy that are characterised by a rapid growth phase with endothelial proliferation, followed by gradual involution. Most haemangiomas are not present at birth but subsequently develop during the first 8 weeks of life.2

The majority of haemangiomas involve the head and neck. They are rare in the oral cavity but may occur on the tongue, lips, buccal mucosa, gingiva and palatal mucosa.3–6 Clinically, haemangiomas appear as a smooth or lobulated soft mass, are sessile or pedunculated, and may vary in size from a few millimetres to several centimetres. They are usually deep red and
undergo capillary, mixed cavernous, or sclerosing, a variety that tends to cavity are extremely rare.7

Haemangiomas occur frequently in the extremities (15%), and are grouped into infantile haemangiomas of the head and neck region (60%), followed by the trunk (25%) and the extremities (15%), and are grouped into infantile haemangiomas of the oral cavity are extremely rare.7

Haemangiomas are benign tumours of the blood vessels and are classified according to their histopathological appearance as capillary, mixed cavernous, or sclerosing, a variety that tends to undergo fibrosis.8 Haemangiomas grow by endothelial hyperplasia and should be differentiated from vascular malformations, which are not true neoplasms but are localised defects of vascular morphogenesis caused by dysfunction in embryogenesis and vasculogenesis.4 Haemangiomas are the most common benign soft tissue tumour of infancy and childhood, occurring in 12% of all infants, and are found in greater frequency in girls, white children, premature infants, twins, and those born to mothers of higher maternal age.8 The incidence in the Indian population has not been reported.9 Haemangiomas occur frequently in the head and neck region (60%), followed by the trunk (25%) and the extremities (15%), and are grouped into infantile haemangioma and congenital haemangioma. They rarely occur on the palatal mucosa.10

The pathogenesis and origin are not clearly understood. A variety of other lesions resemble haemangioma such as pyogenic granuloma, chronic inflammatory hyperplasia, epulis granulomatosa, telangiectasia and angiosarcoma.9

The management of haemangioma depends on a number of factors and most true haemangiomas require no intervention. However, 10–20% of cases require treatment because of their size, location, stage of growth, behaviour and functional compromise.5 The range of treatment includes surgery, flashlamp-pulsed laser, intra-lesional injection of fibrosing agent, treatment with interferon α, electrocoagulation and radiation.11 12

In our patient, since the tumour was causing difficulty in swallowing and was impairing speech, surgical excision was carried out.

Learning points

- Intra-oral haemangiomas can have a varying clinical appearance and may mimic inflammatory lesions such as pyogenic granuloma.
- Intra-operative complications such as bleeding should be anticipated and precautions taken during excision of even smaller benign-appearing lesions.
- A decision to observe the lesion may be taken if the diagnosis is haemangioma and it does not interfere with function or aesthetics, as 90% of them involute by 9 years of age.

Competing interests None.
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Figure 2 Histopathology (×10 magnification) showing vascular channels lined with endothelial cells.

Figure 3 Recurrence of a similar swelling at an anterior site.
Unusual presentation of more common disease/injury

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