Case Report
Lymphangioma of the dorso-lateral tongue in an adult male: A rare case report

Kafil Akhtar1,*, Fatima Meraj1, Fauzia Talat1, Sadaf Haiyat1
1 Dept. of Pathology, Jawaharlal Nehru Medical College, AMU, Aligarh, Uttar Pradesh, India

ABSTRACT
Lymphangioma is a relatively uncommon benign tumour occurring due to malformation of the lymphatic vessels. Lymphangioma in adults is a rare occurrence. Intraoral lymphangiomas occur most frequently on the dorsal tongue and rarely on the palate, buccal mucosa, gingiva and lips. Clinically, lymphangiomas of the oral cavity usually have a plaque formed of small thin-walled vesicles and similar in surface to frog eggs. We present a case of lymphangioma of the tongue in a 34-year-old male who presented with complaints of painless swelling in his tongue with difficulty in speaking for the last 25 days. CT scan revealed a lobulated sharp-contoured soft tissue communicating with the superficial planes with minimum vascularity in colour doppler studies. En masse excision of the firm, solid erythematous mass was performed which microscopically showed multiple dilated endothelial lined vessels filled with a proteinaceous fluid and lymphocytes. A diagnosis of lymphangioma of the tongue was given.

1. Introduction
Lymphangioma, described by Redenbacher in 1828, is a relatively uncommon benign tumour occurring due to malformation of the lymphatic vessels.1,2 These tumours comprise 4% of all the vascular tumours and 25% of all benign vascular tumours in children. No racial predominance or sexual predilection has been reported in them.3 They are also called lymphatic hamartomas and are usually detected at birth (50%) or in early childhood, and the majority (90%) of congenital cases develop before two years of age.4 Lymphangioma in adults is a rare occurrence.4

The head and neck region is frequently involved (50%–75%) in lymphangiomas; however, the oral cavity is rarely affected. Other sites include shoulder, armpit, abdomen, neck, pharynx, eyelids, and conjunctiva.5 Intraoral lymphangiomas occur most frequently on the dorsal tongue and rarely on the palate, buccal mucosa, gingiva and lips.6 Tongue lymphangiomas represent 6% of these tumours in the body.7,8 The most common site for intraoral lymphangioma leading to macroglossia is the anterior two-thirds of the dorsal surface of the tongue.9

2. Case Report
A 34-year-old male reported to the otolaryngology Clinics of Jawaharlal Nehru Medical College Hospital with chief complaint of a painless swelling on his tongue for 25 days. The patient also had difficulty in speaking and accidental biting of tongue while eating. Intraoral examination revealed a solitary inflamed fleshy swelling on the left dorso-lateral part of the tongue, approximately 3x4 cm in size, with numerous tiny papillary projections. On palpation, the lesion was soft and tender with adherence to the superficial and deep layers. There was no bleeding on palpation, as well as no evidence of any lymph nodes. No other intraoral lesions were appreciated. The cranial nerve examination was within normal limits. The patients’ vital signs were stable.

Routine blood investigations showed all parameters within the normal range. Indirect laryngoscopy showed no extension to the vocal cords and larynx. CT scan revealed

* Corresponding author.
E-mail address: drkafilakhtar@gmail.com (K. Akhtar).
a lobulated sharp-contoured soft tissue communicating with
the superficial planes. Minimum vascularity was present in
colour doppler studies.

En masse excision was done using local anesthesia
with a margin of 1 cm from the lesion. The excision
mass was firm, solid erythematous and measured 3.5 x 4.2 cm. Histopathological examination of the lesion
showed multiple dilated endothelial lined vessels just
adjacent to squamous epithelium. These vessels contained
proteinaceous fluid and lymphocytes (Figures 1 and 2). A
diagnosis of lymphangioma of the tongues was given. Our
patient is doing well after 12 months of follow up period.

Fig. 1: Histopathological examination of the lesion showed
multiple dilated endothelial lined vessels just adjacent to squamous
epithelium, filled with proteinaceous fluid and lymphocytes.
Hematoxylin and Eosin x10X.

Fig. 2: High power of Figure 1.

3. Discussion

The etiology of lymphangioma is not fully understood, and
two major theories are proposed. The first theory explains
that the lymphatic channels develop from 5 primitive
sacs arising from the venous system and endothelial
out-pouching from the jugular sac forms the lymphatic
system in the head and neck region. The second theory
proposes that the mesenchymal clefts in the venous plexus
reticulum extend towards the center of the jugular sac
and form the lymphatic system. Congenital obstruction
or sequestration of these enlarged primitive lymphatics
leads to lymphangioma. On the other hand, the etiology
of lymphangioma acquired in adulthood is different and
includes trauma, inflammation, and lymphatic obstruction.

Lymphangioma is typically diagnosed clinically. The
extension and size of lesion affect the clinical appearance
of lymphangioma. Superficial lesions are pink or yellowish
in color and consist of elevated nodules. The deeper
lesions are soft and diffuse masses with mild variation
in color. Clinically, lymphangiomas of the oral cavity
present as pinkish small thin-walled vesicular lesions. The
vesicles contain both clear fluid (lymph) and blood content
suggesting the co-existence of lymphatic and vascular
anomalies.

De Serres LM in 1995 proposed a classification for
lymphangiomas of head and neck based on anatomical
extent, which can be used to predict prognosis and
outcome of surgical intervention. This system branched
lymphangiomas into unilateral infrapharyngeal lesions (stage I), unilateral
suprahyoid lesions (stage II), unilateral suprathyroid and infrathyroid lesions (stage III), bilateral
suprathyroid lesions (stage IV) and bilateral suprathyroid and infrathyroid lesions (stage V).

Macroglossia, with an irregular surface displaying gray
and pink projections, is a key feature of lymphangioma
of the tongue. In children, lymphangioma is a common
cause of macroglossia and is associated with difficulty in
swallowing and mastication, speech disturbances, airway
obstruction, mandibular prognathism, bleeding, and other
deformities of maxillofacial complex. Majority of cases of
macroglossia in children due to intraoral lymphangioma are
located on the anterior two-thirds of the dorsal tongue.

Lymphangiomas are classified histopathologically
into lymphangioma simplex (capillary lymphangioma),
cavernous lymphangioma and cystic lymphangioma (cystic
hygroma).

Due to their characteristic histology and classic clinical
appearance, lymphangiomas of the tongue are easy to
diagnose. The histologic features are characteristic and the
clinical appearance often classic. Generally, any “bump of
the tongue” is included under the clinical differentials of
lymphangiomas. Hematoxylin and eosin stained sections
are best for differentiating them from other differentials.

In the present case, the dorsolateral site may suggest
entities such as granular cell tumor, lingual thyroid,
and mesenchymal tumors, but are not limited to these.
Vascular lesions like hemangiomas, venous malformations
and arteriovenous malformations are the main mesenchymal
tumors to be considered.

Among the hemangiomas, the infantile hemangiomas
show the most resemblance to lymphangiomas. These
are mostly congenital and occur frequently in the head and neck. However, they involute after some years whereas tongue lymphangiomas do not regress. The classic frog egg clinical appearance is absent in hemangiomas. Infantile hemangiomas may show marked endothelial proliferation which is missing in lymphangiomas. There is no close connection to the superficial epidermis as in lymphangiomas, and they are very reactive (95%) to GLUT-1. Lymphangioma can be distinguished from the vascular lesions by the increased expression of D2-40 in lymphatic endothelium.5

Venous malformation also tend to be present at birth and persist like lymphangiomas. Clinically, they may be red or blue, having nodules or blebs very similar to a lymphangioma. However, venous malformations are typically compressible and may demonstrate a thrill or bruit, whereas lymphangiomas will not. Histologically, venous malformations are composed of aberrant vessels often times dilated, filled with red blood cells. Endothelial cell proliferation is generally not a feature of vascular malformations.6 Arteriovenous malformations are less common and can have excessive bleeding on biopsy. Tissue may show variable sized vessels, thrombosis and calcification.11

Pyogenic granuloma is a localized reaction found mostly in the gingiva, but also frequent on the tongue. Clinically, pyogenic granulomas may be sessile or pedunculated red to pink mass. Histologically, they may resemble granulation tissue with a vague lobular architecture of the vessels. Pyogenic granulomas may be very inflamed, demonstrating both acute and chronic inflammation with or without ulceration.3

Early recognition and appropriate treatment is necessary to achieve optimal therapeutic results and avoid complications.12,13 Lymphangioma is managed on the basis of their type, size, anatomical structures involved and infiltration to the adjacent tissues.13,14 The main purpose of therapy is to relieve pain, edema, lymph, and blood leak, as well as super infections. Aesthetic improvement is also important for patients psychological well-being. Surgical excision is the treatment of choice, but it causes scar formation and is not feasible in all cases. Cryotherapy, radiation therapy, steroid administration, sclerotherapy, electro-cautery, embolization, ligation, laser surgery, and radiofrequency tissue ablation are the other modalities used in such cases. It is essential to include a surrounding border of normal tissue without damaging vital structures for minimal recurrence and successful outcome.15,16

4. Source of Funding
No financial support was received for the work within this manuscript.

5. Conflict of Interest
The authors declare they have no conflict of interest.

References