Cystic lymphangiomas of the tongue: 3 rare cases and a literature review of classifications and therapeutic possibilities

Kaoutar Cherrabi1*, Naouar Ouattassi1, Anouar Titou2, Dounia Kamal2 and Mohamed Noureddine El Alami1

Abstract

Background: Cystic lymphangiomas of the tongue are rare. The treatment consists of complete resection and management is generally interdisciplinary due to vital, functional, and esthetic comorbidities.

This work is a report of three rare cases of cystic lymphangioma of the tongue, with curious mode of presentation and good outcome, as well as a literature review of clinical features, classifications, and therapeutic possibilities.

Case presentation: This work presents three patients. The first one presented with a lymphangioma in the junction between base and body of tongue; the clinical aspect was first similar to hemangioma, and then the diagnosis was re-established after slight enlargement of the mass. The patient was treated with sclerotherapy with great outcome.

The second patient presented with a large lymphangioma which presented a clinical resemblance with a tumor of the tongue, the diagnosis was established upon radiological aspect and confirmed upon histological findings.

The third patient presented with a macroglossia which appeared 3 years after excision of the cystic hygroma of the neck; the patient was surgically treated with excellent outcome.

Conclusion: Cystic lymphangiomas are rare hamartomatous lymphatic malformations, and they have a predilection for head and neck but the lingual localization is quite rare. Treatment is complete excision. Management is generally interdisciplinary. The main objective is to prevent alteration to function, to treat esthetic comorbidities, and prevent speech impediment especially in very young children.

This is the cases of three young children with a rare lingual cystic lymphangioma, with adapted treatment and satisfactory outcome, with a literature review of clinical presentation, classifications, and therapeutic approaches.

Keywords: Cystic lymphangiomas, Head and neck, Anatomy, Radiology, Tongue, Congenital malformation, Macroglossia

Background

Lymphangiomas are rare congenital lymphatic hamartomatous proliferations that were first described by Virchow in 1845 [1–5]. They have predilection for head and neck regions [2–4]. Growth is generally slow, but can be accelerated due to infectious or hemorrhage episodes [6].

Case presentation

The first patient was 1-year-old when he first presented with a lesion: a small blood-filled vesicular lesions in the intersection between the body and the base of the tongue. It appeared in the first year of life and was slightly painful during feeding. The lesion was first diagnosed as a hemangioma based on its clinical aspect, and a course

*Correspondence: Kaoutarcherrabi.bz@gmail.com

1 ENT and Cervico-Facial Surgery Department, Faculty of Medicine and Pharmacy, Sidi Mohamed Ben Abdellah University, Fez, Morocco

Full list of author information is available at the end of the article
of treatment of beta-blockers was started in pediatric department. A stabilization of lesion’s size was observed. After 4 years, the patient presents a slight increase in size, with a slight speech impediment, and persistent pain. The clinical exam showed a vesicular lesion, with translucent and blood-filled vesicles, which indicated that it was in fact a lymphangioma. The patient had a cervico-facial MRI which showed a cystic lesion of the junction between the body and the base of the tongue, hyperintense on T2, isointense to muscle in T1 (Fig. 1).

Patient was started on sclerotherapy with bleomycin, with good outcome after three sessions. Follow up of 3 months was satisfactory.

The second patient was 8 years old. She presented with a large tumor of the tip of the tongue with a sub-acute increase in size for the last 2 months with difficulties in alimentation, speech, breathing in supine position, as well as impossibility to close her mouth.

The patient underwent cervico-facial MRI that showed a cystic mass attached to the tip of the tongue, without any cervical lesions associated (Fig. 2).

The third patient was 4 years old when he first presented with cystic hygroma to his neck, floor of mouth, parotid region, and masseteric region, for which he was admitted to surgery for excision. After 3 years, the patient presented with macroglossia. The lingual mass was large and it has been increasing in size in the last 3 months, articulation was impossible, alimentation very difficult, as well as respiratory distress. The clinical examination found no recurrence in cervical region, no other lesion was found in the oral cavity.

The patient had cervico-facial CT-scan which showed a heterogeneous mass of the tongue, taking on the totality of its body and base, with bulging in oropharynx. The patient underwent reductive surgery, through keyhole glossectomy technique with good follow up for 6 months, during which the patient was admitted in speech therapy.
for slight speech impediment, with satisfactory evolution (Figs. 3 and 4).

Discussion
Fifty percent of cystic lymphangiomas are noticed at birth, and 90% are discovered by 2 years of age, generally in the first decade of life [1–3, 7]. The most frequent site is the posterior triangle of the neck between the posterior border of the Sterno-cleido-mastoid muscle, the middle part of the clavicle, and the anterior edge of the trapezius [2]. They can also exist in the anterior triangle of the neck, in the parotid region, as well as the sub-mandibular region [1, 2, 7]. Cases of oral cavity lymphangioma are rare. They are generally in the tongue, palate, gingiva, lip, and alveolar ridge of mandibule [1, 2, 5]. In the tongue, the most common localization is in the anterior two thirds [2].

Precise embryological origin is still unknown. However, theories are studied [3]. Lymphangiomas are formed because of congenital obstruction or sequestration of the primitive lymphatic enlargement. Two major theories have cited to explain the origin of lymphangiomas [1–3, 5, 7].

1st theory
Lymphatic system develops from 5 primitive sacs that arise from the venous system [2]. Lymphatic system constitutes of endothelial out-pouching and travels centrifugally from jugular sac [2].

2nd theory
Lymphatic system derives from mesenchymal clefts in venous plexus reticulum and spreads centripetally towards the jugular sac [2].

Diagnosis is generally established upon superficial characteristics of the lesion. Radiology is indicated in case of inexperienced clinician or atypical lesions [8]. Clinical presentation of lymphangioma depends on extension of the lesion; superficial lesions are elevated nodules, while deeper ones are soft tissue masses [2, 3]. The patient can present with a history of enlargement due to trauma or infection, recurrent infections, airway compression, feeding difficulties, interference with normal or developmental speech, and esthetic problems [8]. Early diagnosis and management are essential for good outcome [8]. Biopsy is not indicated because of the great risk of bleeding correlated to it [8].

Macroscopic aspect shows multiple small vesicles of varying sizes [9], covering the surface of the tongue, they may be either translucent or blood-filled. While localized in cases, they can be exophytic lesions with nodular surface due to the presence of vesicles in extensive cases [8]. Macroglossia of the anterior two thirds
of the tongue causes speech impediment, poor oral hygiene, and bleeding accidents with oral trauma [2].

It is important to define the association with dental problems, especially for children with macroglossia to prevent carries and gingivitis [6].

Several classifications have been described [1, 2, 4, 10]. Histological classification is not correlated with clinical presentation or therapeutic approach [1, 4, 10].

- Lymphangiomas simplex (lymphatic channels with thin walls and capillary size).
- Cavernous lymphangiomas (dilated lymphatic channels with fibrotic adventitial coats).
- Cystic lymphangiomas: composed from cysts varying from millimeters to several centimeters in diameter.
- Lymphangiosarcoma: lymphangioma associated with angiosarcoma, which a rare malignant tumor that develops after prolonged lymphatic obstruction.

De Serres Classification of lymphatic malformations is based on localization and extension [2, 8].

Class I: infra-hyoid unilateral lesions.

Class II: supra-hyoid bilateral lesions.

Class III: supra-hyoid or infra-hyoid unilateral lesion.

Class IV: supra-hyoid bilateral lesions.

Class V: infra-hyoid bilateral lesion.

Clinical classification [8]:
A classification based on clinical aspect, and based on response to sclerotherapy, is more used:

- Microcystic: do not respect tissue planes, diffuse, and are difficult to excise.
- Macrocytic: are localized and surgically easier to excise.
- Mixte.

Classification of microcystic lymphangioma is correlated with treatment option and indicates the possibility of curative treatment (Table 1).

In Wiegand et al. classification, I, IIa, IIb, and III belong to Serres et al. In stage II, they have different treatment modalities and outcomes [8].

I and IIa are easier to treat and their complete resection is possible. IIb and III are more difficult due to the
extension of the disease and the involvement of structures [8].

MRI is the exam of choice, if treatment is indicated and in cases of large lesions in order to examine extensions [8]. The aspect on MRI consist on highly increased intensity in T2-weighted images, while in T1-weighted images, lesions have the same intensity or slightly lower intensity than muscle [7, 8].

The standard objective of management of head and neck lymphangiomas is total excision whenever it is possible [3, 4]. Incomplete removal is associated with a high risk of recurrence [10]. Restoration or preservation of function, esthetic integrity, and management of lymphatic malformations have changed in the last few years [3].

Treatment depends on size and involvement of anatomical structures [2, 8]. Conservative management should only be considered for asymptomatic cases before definite treatment. The risk of acute enlargement in cases of infection or hemorrhage should be contemplated [8]. Surgery is generally delayed until 3 years old of age [2, 4, 8].

Several techniques have been described to manage lymphangiomatosus and hemolymphangiomatosus lesions of the tongue. Peripheral glossectomy are used in cases of lesions that occupy length and width of the tongue. Keyhole glossectomy, which is used when all dimensions of the tongue are involved, and central strip glossectomy in cases that involve hemihypertrophy of the tongue [11]. Jian Glossectomy technique combines marginal and central resection, and maybe used for lesions in the anterior and middle parts of the tongue [11].

The association of antibiotic therapy and steroid course should be considered for all stages in cases of enlargement [8].

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**Table 1** Table of classification of microcystic lymphangiomas [8]

| Description                                             | Treatment                                      |
|---------------------------------------------------------|------------------------------------------------|
| Stage I [8] Isolated superficial microcystic lymphatic malformations of the tongue. | IIaCO2 laser is curative with excellent outcome. (Susanne Wiegand et al.). |
| Stage II [8] Isolated superficial lymphatic malformations of the tongue with muscle involvement. | IIa CO2 laser is curative with excellent outcome. (Susanne Wiegand et al.). |
| Stage III [8] Microcystic malformation of the tongue and the floor of mouth. | IIb useful combined and staged approach (Susanne Wiegand et al.). |
| Stage IV [8] Extensive microcystic lymphatic malformation involving the tongue, floor of mouth, further to cervical structures. | IIb useful combined and staged approach (Susanne Wiegand et al.). |
Oral cephalosprin: Cefuroxim axetil 500 mg, two times a day for 7 to 10 days, or cefuroxim 80 mg/kg a day for a maximum of 4.4 g in 3 doses by IV [8].

Steroid: 5 mg/kg/day with a maximum of 250 mg a day [8].

When surgery is indicated, the best choice the radi-cal approach. The need for repeated surgery alters the underlying structures [8] (Table 2).

Cystic lymphangiomas of the tongue can be associated with several complications:

- Rapid enlargement due to upper tract infections or hemorrhage with possible airway obstruction [3].
- Functional complications [2, 3, 8].
- Esthetic complications [1–3, 8].

Recurrence rate after total excision is 0–27%, and after partial excision 50–100% [4].

**Conclusion**

Management of large lymphatic malformation of the tongue is accomplished through complete excision, and demands inter-disciplinary approach [1, 8]. Macroglos-sia and large lymphangiomas of the tongue could lead to skeletal and dental problems like prognathism and mal-oclusion [8]. Maxillo-facial surgeons and ortho-dentists should be involved [4, 8].

This work is a report of three rare cases of cystic lymphangioma of the tongue, with curious mode of presentation and good outcome, as well as a literature review of clinical features, classifications, and therapeu-tic possibilities.

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**Authors' contributions**

KC conceived and designed analysis, collected data, performed analysis, wrote the paper, and performed surgery. NO performed surgery. AT contributed in data collection. DK supervised surgery and contributed to designing analysis. MNE supervised designing analysis. All authors have read and agreed to its content.

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**Ethics approval and consent to participate**

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**Consent for publication**

The patients’ parents provided written informed consent to publish this case report, as well as for the publication of information.

**Competing interests**

The authors declare that they have no competing interests.

**Author details**

1ENT and Cervico-Facial Surgery Department, Faculty of Medicine and Pharmacy, Sidi Mohamed Ben Abdellah University, Fez, Morocco. 2Maxillo-facial Surgery Department, Faculty of Medicine and Pharmacy, Sidi Mohamed Ben Abdellah University, Fez, Morocco.
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