Coexisting sublingual dermoid cyst and heterotopic gastrointestinal cyst: Case report

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Abstract
A heterotopic oral gastrointestinal cyst coexisting with a lingual dermoid cyst is a rare condition, with only 3 reported cases in the English-language literature. Clinically, both gastrointestinal and dermoid cysts tend to manifest as an asymptomatic swelling. Cysts above the mylohyoid often present with sublingual swelling, whereas those below the mylohyoid present with submental swelling. Increased salivation, altered speech, dyspnea and difficulties in eating, swallowing and respiration may be present, depending on the size of the cyst. This paper describes the case of a 4-year-old girl presenting with swelling of the tongue and floor of the mouth that interfered with normal speech. Two cystic lesions were surgically excised using an intraoral approach, and recovery was uneventful. Histopathological examination of the specimens taken during surgery revealed the first, anterior cyst to be a dermoid cyst and the second, posterior cyst to be a heterotopic oral gastrointestinal cyst (HGIC). Although this situation is very rare, it should be included in the differential diagnosis of swellings in the submandibular region in the pediatric population.

Key words: Dermoid cyst, heterotopic gastrointestinal cyst, children, tongue.

Introduction
The term ‘heterotopia’, also referred to as ‘choristoma’ or ‘aberrant rest’, is used to describe tissue or organs displaced to an abnormal location (1). A heterotopic gastrointestinal cyst of the oral cavity – also known as a ‘lingual foregut duplication cyst’, ‘enterocystoma’, ‘choriomatic cyst’, ‘ciliated epithelial cyst’, or ‘heterotopic large bowel cyst’(2), has been observed in the duodenum, gallbladder, jejunum, pancreas, ileum, rectum, anus and the umbilicus as well as in the lips, larynx, lung, submandibular glands, epiglottis and anterior neck (1). Dermoid cysts are rare, benign, developmental lesions that occur primarily in the testes and ovaries, although they can be found anywhere in the body at an embryological fusion point, including the head and neck areas. Approximately 7% of all dermoid cysts form in the head and neck region, and of those, 23% develop in the floor...
of the mouth (3-6). However, dermoid cysts represent only 0.01% of all oral lesions (7). When dermoid cysts do occur in the oral cavity, they are usually found on the floor of the mouth in close association with the mylohyoid muscle, or, rarely, in the tongue (6). If a dermoid cyst is left untreated for a long period of time, carcinomatous change is possible, although this is very rare (8). Clinically, both gastrointestinal and dermoid cysts tend to manifest as an asymptomatic swelling. Cysts above the mylohyoid often present with sublingual swelling, whereas those below the mylohyoid present with submental swelling. Increased salivation, altered speech, dyspnea and difficulties in eating, swallowing and respiration may be present, depending on the size of the cyst (2,4,6,9). Dyspnea and respiratory distress can also be present, and tracheostomy may occasionally be required (9).

The coexistence of a dermoid cyst and heterotopic oral gastrointestinal cyst (HGIC) is extremely rare. To date, only three cases have been reported in the English-language literature. The following report describes such a case, including the pathogenesis (10).

Case Report
A 4-year-old girl presented at the Akdeniz University Dental Faculty Department of Oral and Maxillofacial Surgery with a painless sublingual swelling. Extraoral examination revealed no facial abnormalities or swellings. Intraoral examination revealed a large, compressible cystic mass in the midline of the anterior floor of the mouth, causing significant elevation of the tongue. The mucosa covering the mass was normal in color, while the mass itself was smooth, firm and fluctuant at bi-manual palpation, with no evidence of inflammation (Fig. 1). The patient’s parents were uncertain how long the swelling had been present, but stated that their child’s speech was below the level of her peers. The child’s medical history was unremarkable. No regional lymphadenopathy was noted. Magnetic resonance imaging (MRI) revealed contiguous, round, T2-hyperintense cysts with regular margins in the central and anterior parts of the floor of the mouth that were 22x22 mm and 25x21 mm, respectively, with T1-hyperintense areas observed in the anterior cystic lesion (Figs. 2a, b). Preoperative differential diagnoses included epidermoid or dermoid cysts, cystic hygroma, ranula and lymphangioma.

Under nasotracheal intubation, using an intraoral approach, a semi-lunar incision was made on the ventral surface of the tongue from left to right. Using blunt dissection, the first lesion was separated from the surrounding tissue, and the genioglossus muscle was separated from the lesion. The first lesion was then gently elevated from the mylohyoid muscle and removed in one intact piece (Fig. 3a). Next, the second lesion was accessed through a blunt dissection in the muscles of the floor of the mouth (Fig. 3b) and excised in the same manner as the first lesion. Hemostasis was established, the wound ...

Fig. 1: The patient’s preoperative clinical appearance.

Fig. 2: a: Sagittal (Magnetic resonance imaging) MRI scan showing the relationship between the two lesions in the tongue and on the floor of the mouth. b: Coronal Magnetic resonance imaging (MRI) scanned image of the two lesions.
edges were closed, and a 0.25 inch Penrose drain was inserted. The drain was removed on the postoperative 3rd day, and the region was closed primarily via suture. Histopathological examination of the specimens taken during surgery revealed the first, anterior cyst to be a dermoid cyst and the second, posterior cyst to be a heterotopic oral gastrointestinal cyst (HGIC) (Figs. 4a, b). Postoperative healing was uneventful. In line with clinical protocols, the patient presented for regular follow-up visits.

Discussion

Differential diagnosis of swellings on the floor of the mouth should include cystic lesions such as mucocele, ranula, thyroglossal duct cysts, lymphoepithelial cysts, cystic teratoma, salivary gland tumors and mesenchymal tumors as well as sialadenitis, hemangioma/lymphangioma and abscesses (3-6). The diagnostic imaging for these types of lesions includes computed tomography (CT), ultrasonography (US) and MRI (3,4). Definitive diagnosis depends on histological investigation following clinical and radiological examination (4).

The term “dermoid” is used as a general term to refer to all three histological types of dermoid cysts identified by Meyer (11) as well as to one of the specific histological categories: epidermoid, dermoid and teratoid. Epidermoid cysts have only a stratified squamous epithelial lining with no adnexal structures. In addition to a typical, squamous epithelial lining, dermoid cysts also exhibit dermal appendages, like sebaceous glands, hair follicles and sweat glands. Teratoid cysts have both squamous and respiratory epithelium, dermal appendages and distinct mesodermal components (11). Surgical excision is the treatment of choice for both HGIC and dermoid cysts. Depending upon the clinical relationship between the cyst and the mylohyoid and geniohyoid muscles, either intraoral or extraoral methods, or sometimes even a combined approach, may be appropriate (3,4,9,12). Prognosis is generally good, with no
reports of recurrence after complete surgical excision (3). However, epithelial tracts leading to the mandible or hyoid bone have been documented and require excision to avoid recurrence (12).

In conclusion, although coexisting sublingual dermoid and heterotopic gastrointestinal cysts are very rare, they should be especially included in the differential diagnosis of swellings in the submandibular region in the pediatric population.

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Conflicts of Interest
The authors have no conflicts of interest relevant to this article.