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The primary aim of World Journal of Clinical Cases (WJCC, World J Clin Cases) is to provide scholars and readers from various fields of clinical medicine with a platform to publish high-quality clinical research articles and communicate their research findings online.

WJCC mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

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RESPONSIBLE EDITORS FOR THIS ISSUE
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Eustachian tube teratoma: A case report

Jin-Ye Li, Li-Xin Sun, Na Hu, Ge-Sheng Song, Wei-Qiang Dou, Ruo-Zhen Gong, Chuan-Ting Li

Abstract

BACKGROUND
Mature teratoma composed of all three basic germ cell layers of the head and neck is a rare disease. Teratomas involving the temporal bone are particularly scarce.

CASE SUMMARY
A 48-year-old male patient with a history of chronic otitis of the left ear from infancy, for which he had been operated on twice, was referred to our hospital for chronic otitis, cholesteatoma and a middle ear mass. Computed tomography (CT) scan and magnetic resonance imaging (MRI) revealed an eustachian tube teratoma, in which the anterior lower part and posterior upper part were connected by a thin membranous tissue. The mass was removed completely under general anesthesia by mastoidectomy. As of last follow-up (2 years post-surgery), the disease had not relapsed.

CONCLUSION
Pre-operative CT and MRI are necessary for eustachian tube teratoma. Complete surgical resection provided excellent prognosis.

Key Words: Eustachian tube; Teratoma; Chronic otitis; Computed tomography; Magnetic resonance imaging; Case report

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Core Tip: Mature teratoma of the head and neck is a rare disease. We present a case of eustachian tube teratoma, in which the anterior lower part and posterior upper part were connected by a thin membranaceous tissue, in a 48-year-old male patient who had a history of chronic otitis of the left ear from infancy and had already been operated on twice. When a long history of chronic otitis is encountered, combined with polyps in the tympanum and/or external auditory canal, a combination of computed tomography and magnetic resonance imaging is necessary pre-operation.


INTRODUCTION
Teratomas of the head and neck account for 5%-15% of all teratomas and principally involve the nasopharynx and neck[1]. Reports of teratomas involving the temporal bone are scarce[2,3]. Herein, we present a case of mature teratoma of the eustachian tube (ET) in an adult male with a history of chronic otitis of the left ear from infancy.

CASE PRESENTATION
Chief complaints
A 48-year-old man was referred to our hospital for chronic otitis, cholesteatoma and a middle ear mass.

History of present illness
The patient had experienced chronic otitis of the left ear from infancy and underwent surgery for cholesteatoma in the tympanum at another hospital. However, his clinical symptoms had persisted. His conscious hearing was poor, and he suffered from earache, ear boredom, headache and dizziness.

History of past illness
The patient had no significant past medical history.

Personal and family history
The patient’s family history was unremarkable.

Physical examination
Otoscopy examination demonstrated a large amount of pus in the left external auditory canal, a fleshy polyp present at a deeper site, and mucosal edema in the previously operated area (Figure 1). Audiometric assessment confirmed a severe conductive hearing loss; hearing threshold was 80 dB and auditory brainstem response was 70 dB on the affected side. Nasopharyngoscopy showed that the orifice of the left ET opened well.

Laboratory examinations
The laboratory examination was otherwise unremarkable. The laboratory assessment included routine blood tests. Tests for C-reactive proteins and viral hepatitis markers were negative. Glucose and serum insulin levels were normal.

Imaging examinations
The unenhanced computed tomography (CT) of the temporal bone showed some changes, including a well-circumscribed, mixed density tumor with a fat density area in the ET; the lesion extended down to the left part of the tympanum and external auditory canal (Figure 2), without ossicular chain, which had resulted from the mastoidectomy. The T1- and T2-weighted magnetic resonance imaging (MRI) in the
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Figure 1 Endoscopic appearance and intraoperative appearance of the patient’s teratoma. A: Otoscopic examination demonstrated a large amount of pus in the left external auditory canal (white arrowhead) and a fleshy polyp at a deeper site (black arrowhead); B: The part of the mass in the tympanum and external auditory canal appeared as a fleshy polyp (arrowhead); C: “Hairs” were present on the surface of the mass and cartilage was surrounded by the mass in part of the eustachian tube (arrowhead).

Figure 2 High-resolution computed tomography scan of the patient’s temporal bone. Computed tomographic imaging showed a well-circumscribed, mixed density tumor (white arrowhead) with a fat density area (black arrowhead) located in the eustachian tube.

Transverse plane showed a 3.2 cm × 1.3 cm × 2.0 cm, well-defined, homogeneous lesion with high signal intensity along the left ET. The mass showed signal intensity similar to that of the fat on all sequences and with little cartilage signal (Figure 3). The lesion extended down to the left part of the tympanum and external auditory canal, where the signal was slightly higher than the part in the ET. On fat-saturated T1- and T2-weighted sequences, the part of the mass in the ET demonstrated a decreased signal intensity, indicating that the mass was consistent with macroscopic fat. The mass was surrounded by a smooth, thick, hypointense capsule, which was enhanced slightly after contrast administration.

Pathological results
The ET diameter was expanded, to about 0.8 cm. The part of the mass in the tympanum and external auditory canal was classified as a fleshy polyp. However, “hairs” were visible on the surface of the mass and cartilage surrounded by the mass could be seen in the ET area (Figure 1). The mass, in which the anterior lower part and posterior upper part were connected by a thin membranaceous tissue, was in the ET. It was tightly bonded to the former ET wall and could not be removed in toto, so serial partial excision was performed. The resulting cavity (composed of the tympanum and external auditory canal) was then resolved by filling with abdominal fat.

On gross examination, the resected mass appeared as a cluster of irregular, soft, grey-colored tissue. Microscopically, a photomicrograph of the mass revealed characteristics of keratinized squamous epithelium, adipose, sweat gland and mature skeletal muscle tissues. Photomicrographs of the areas including the tympanum and external auditory canal revealed squamous epithelial mucosal polyps (Figure 4).
Figure 3 Magnetic resonance imaging of the patient’s head and neck. A, D, G: Three-dimensional (3D) T2 weighted image (WI); B, E, H: Fat suppression (FS) 3D T1WI; C, F, I: FS 3D T1WI with contrast (C+); A-C: Magnetic resonance (MR) images in the transverse plane showed the part of the mass which was a homogeneous lesion with slightly higher signal intensity and with enhancement (white arrowheads) in the tympanum and external auditory canal; D-F, G-I: MR images in the transverse plane showed the part of the mass which was a well-defined, homogeneous lesion with high signal intensity along the left eustachian tube (white arrowheads), and on FS 3D T1WI, a lesion with decreased signal intensity consistent with macroscopic fat and with a contrast-enhancing rim (black arrowhead) was seen.

FINAL DIAGNOSIS
Mature teratomas.

TREATMENT
The mass was removed completely by mastoidectomy, with the patient under general anesthesia.

OUTCOME AND FOLLOW-UP
After follow-ups at 7 mo and 2 years, the patient showed no signs of disease relapse.
DISCUSSION

Mature teratoma is a true neoplasm composed of all three basic germ cell layers (ectoderm, mesoderm, and endoderm), which differs from dermoids and epidermois [4]. Most cases involve a midline or paraxial location, and the most common site is in the sacrococcygeal region (40%-60% of cases)[5]. Only 2%-10% of cases have involved the head and neck regions[5], especially the cervical and nasopharyngeal regions. We used the PubMed database to search for relevant publications on (keywords) “mature teratoma” and “eustachian tube”. We found only eight relevant publications among the English-language literature, excluding reports on dermoid cysts, which represent a special category of teratomas. According to the histologic classification of teratomas in the head and neck region that is most commonly used today, the term ‘dermoid’ (as it was proposed by Arnold in 1888) is most appropriate[6]. This implies an origin from epidermal and mesodermal elements, which differentiates them histologically from teratomas composed of elements from all three germinal layers and from cholesteatomas, which are only of ectodermal origin[7]. Of these 9 total cases reported to date (including this case), 4 were male and 5 female; the near 1:1 ratio indicates that there is no sex preponderance for ET teratoma. The age of the 9 patients ranged from 1 d to 48 years, with a median of 10-mo-old. The cases had various clinical manifestations, as follow: foul-smelling left otorrhea[8], peripheral palsy of the seventh cranial nerve[5], a tongue-like structure which protruded when the patient cried and retracted into the mouth when swallowing[9], a history of refractory otitis media and a nasopharyngeal mass[3], a discharging left ear since infancy[10], respiratory distress in a premature infant[11] with recurrent otitis media and chronic otorrhea of the left ear[12], and a history of chronic otitis of the right ear from infancy[13]. However, all these manifestations were due to obstruction of the channel or oppression of the surrounding structures.

The case presented here is the first reported for an ET teratoma with the anterior lower part and posterior upper part connected by a thin membranaceous tissue, accompanied by a fleshy polyp in the tympanum and external auditory canal. The patient, who had a history of chronic otitis of the left ear from infancy, was older than the other 8 patients reported for previous cases. It is, therefore, possible that the history of chronic otitis of the left ear and previous operations he undergone led to the formation of a fleshy polyp in the tympanum and external auditory canal. Of note, in
patients with a history of chronic otitis that exhibit polyps in the tympanum and/or external auditory canal, it is important to avoid the assumption that these are merely a result of inflammation. Equally important is to avoid making an immediate call for surgery. Both CT and MRI examinations must be performed to appraise the condition of the ET and identify possible neoplasms. For our patient, the only examination made before his previous surgery was a CT scan. This led to the failure of identifying the ET tumor. Had a preoperative MRI examination been conducted, the ET tumor would likely have been detected and the following operation would have addressed both the inflammation and the tumor.

Ultrasound is an ideal initial imaging modality to investigate neck masses, as it reveals the solid or cystic nature in most cases and localizes the lesion in relation to surrounding structures[14], but it has some limitations when applied to the temporal bone region. A combination of CT and MRI scans provides valuable clinical information to exclude alternative cranial base pathologies and assess the extent of the lesion. The CT scan allows for better evaluation of the bone architecture, whereas the MRI scan can better demonstrate the relationship between the mass and the carotid artery beneath the temporal bone[7]. The typical clinical presentation of ET teratomas includes recurrent episodes of otitis media and chronic otorrhea[15,16], caused by the obstruction of the ET by the teratomas. Ultimately, ET teratoma requires surgical removal and has excellent prognosis upon complete resection[12].

CONCLUSION

In summary, this case highlights the necessity of examination via a combination of CT and MRI pre-operation for ET teratoma. The surgical removal itself provides an excellent prognosis upon complete resection.

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