CASE REPORT

Refactory otitis media with effusion due to eustachian tube sialolipoma inform consistent with eustachian tube inflammatory polyp: rare case presentation and literature review

Khalid Al Zaabi* and Mohamed M. K. Badr-El-Dine

Abstract
Background: Unilateral otitis media with effusion in adult patient is considered alarming finding needing a detailed examination of the nasopharynx. The lesions in this area ranged from benign to more destructive malignant pathologies. Lipoma as a cause of obstruction is an extremely rare entity. Muzzi et al., in his systematic review of primary tumors and tumor-like lesions of eustachian tube (ET), searched the literature from 1888 till 2011 with no single reported lipoma case [1]. We are reporting a case of ET lipoma presented with refractory otitis media with effusion. This is an extremely rare location of such entity and to the best of our knowledge this is the third case reported in the world literature of eustachian tube lipoma after 2 cases published in 2011 and 2016 respectively [2, 3]. However, this is the first case reported in the literature of ET sialolipoma.

Case presentation: We are reporting a 27-year-old male who presented to our outpatient department with a complaint of unilateral ear block, hearing loss, and intermittent tinnitus. Clinical evaluation revealed unilateral otitis media with effusion that was confirmed by pure tone audiogram and tympanometry. Computed tomography and magnetic resonance imaging were obtained showed a lesion originating in the eustachian tube area protruding in the nasopharynx. Surgical removal was successful with complete recovery.

Conclusions: Unilateral otitis media with effusion necessitate nasal endoscopic examination to rule-out lesions at the nasopharynx. This is the first case of sialolipoma in eustachian tube in the literature. Endoscopic removal is usually successful with uneventful short recovery period.

Keywords: Otitis media with effusion, Eustachian tube dysfunction, Unilateral conductive hearing loss
Case presentation

Our patient is a 27-year-old male who was roaming around between primary care facilities and otolaryngology specialists for few months before he showed up in our clinic. His main complaint was left severe ear block sensation associated with hearing loss and intermittent tinnitus. Conservative management with watchful waiting and intranasal corticosteroid sprays were offered to him outside our facility with no successful response. He also had one trial of office-based myringotomy without ventilation tube insertion with fast recurrence of symptoms. Clinical evaluation by otoscope revealed evidence of unilateral left otitis media with effusion. Rinne test showed bone conduction (BC) > air conduction (AC) in the left ear while air conduction (AC) > bone conduction (BC) in the right side. Weber test lateralized to the left ear. Endoscopic nasal examination using 0° rigid endoscope showed deviated nasal septum to the left with a polyp originating from the left ET orifice (Fig. 1A). Pure tone Audiogram with acoustic immittance showed moderate conductive hearing loss in the left ear with normal hearing threshold in the right side. Tympanometry showed type B flat curve in the left side while type A in the right side (Fig. 2A, B).

Computed tomography (CT) scan and magnetic resonance imaging (MRI) were obtained to delineate the nature and extent of the pathology prior to any surgical intervention. Computed tomography (CT) scan and magnetic resonance imaging (MRI) were suggestive of benign lesion noted at the nasopharyngeal orifice of the ET measuring 10 × 8 mm with consequent left middle ear effusion. The most likely differential diagnosis was lipoma (Fig. 3A–F). A written consent was obtained from the patient before surgical intervention. The procedure was performed under general anesthesia via transnasal endoscopic approach. At the start of surgery, left myringotomy with insertion of a soft T-tube was performed. Then, because of the presence of evident left septal spur, endoscopic septoplasty was performed with removal of the bony-cartilaginous spur. To acquire a better exposure of the ET pharyngeal orifice, excision of the posterior bulky tail of the left inferior turbinate was done. The polyp was seen bulging and totally blocking the ET orifice. The polyp was grabbed and using sharp dissection complete excision of the polyp from its origin inside the ET was executed. Electrocautery was not used in the area of the ET orifice (Fig. 1A–D). After checking hemostasis, no nasal packing was

Fig. 1  A Transnasal intraoperative view of a polyp-like lesion protruding through the left ET orifice measuring (1 × 1 × 0.5 cm). B View of the polyp after removing the bulky tail of the left inferior turbinate to improve exposure. C Grabbing the polyp during surgery visualizing its origin from inside the lumen of the ET. D After complete excision of the polyp and the ET orifice looks widely open.
used. The patient had uneventful postoperative recovery. Histopathology report confirmed the diagnosis of sialolipoma (Fig. 4A, B). T-tube was removed 3 months after the surgery. The choice of soft T-tube enabled easy and painless removal of the tube in the office. Complete healing of the tympanic membrane occurred within the next 2 weeks and patient reported total disappearance of symptoms. Follow-up extended for 3 years after the surgery with no recurrence of symptoms or any signs of effusion.

**Discussion**

Finding a tumor or a lesion mimicking tumors originating from eustachian tube (ET) area is extremely rare [1–3]. The symptoms arising from an obstruction at the level of ET range from unilateral ear pressure or fullness to the extreme of hearing loss and referred otalgia. Such presentation warrants a comprehensive head and neck clinical examination including nasal endoscopy. Imaging are very helpful in evaluation of any suspicious fullness or identified lesion in the same area. Furthermore, it can provide
an important information about the nature and extent of such pathology. The primary pathology of isolated ET lesion might be benign like teratoma, dermoid, leio-
myoma, chondroma, and lipoma, or a malignant entity such as squamous cell carcinoma, lymphoma, and sar-
coma [1, 4]. Head and neck lipoma as an entity constitute about 13–17% of soft tissue tumors [5, 6]. The incidence of lipoma in ET area particularly is not reported in the literature [3]. Sialolipoma is a very rare variant of salivary gland lipomatous lesions characterized histologically by entrapment of ducts and acini originated from salivary gland tissues into adipocytes [7, 8]. Nagao et al. were the first to use the term “Sialolipoma” by reporting 7 cases in 2001 [9]. In 2009, Yang et al. reported that the mean age of sialolipoma is 52 years with equal ratio between both genders [10]. Our patient was only 27 years old when he presented to our clinic. The picture that he presented with is one that makes any otolaryngologist want to rule out a mass in the nasopharynx. This is because one should always stress and keep in mind that malignancy is to be ruled out in any adult patient with unilateral OME. The differential diagnosis in mind would be of an antro-
choanal polyp, inverted papilloma, or nasopharyngeal carcinoma. It is unlikely to see a pathology originating inside the ET causing OME in daily clinical practice. Our clinical examination, however, revealed a polyp that was originating in the left ET explaining the symptoms of the patient. Hearing evaluation in the clinic by audiometry and tympanometry confirmed a conductive hearing loss supporting the clinical findings. Imaging modality of the choice in this case would be a high-resolution CT scan with contrast extending from base of skull to the thoracic inlet evaluating the nature and extent of the lesion. MRI scan can be of a great help as well in evaluating the soft tissue. The findings on the CT scan were suggestive of a localized smooth well circumscribed lesion in the left ET area with no bony erosion around it. This finding is usually suggestive of benign lesion. MRI supported the diagnosis of ET lipoma. Surgical excision is the manage-
ment of such lesion. The ideal approach is endoscopic transnasal approach under general anesthesia. We were able to excise the lesion in-toto with minimal bleeding. Histopathology examination revealed the unusual rare variant of lipoma in the ET. The recovery was fast after the procedure and as originally anticipated, the T-tube was removed 3 months postoperative. The choice of soft T-tube made its office removal easier and painless compared to the standard grommet’s tube. Follow-up showed complete resolution of symptoms with no recurrence up to 3 years after the surgery.

Conclusions

Unilateral otitis media with effusion as a complaint in adult patient necessitate detailed and thorough clinical examination including rigid nasal endoscopy to rule-out lesions at the level of nasopharynx. This presentation should always raise concern of malignancy especially in older candidates. However, there are few reported cases of benign lesions in ET area. To the best of our knowl-
edge, this is the first case of sialolipoma reported in this location originating from inside the ET orifice in the liter-
ature. Endoscopic removal of these lesions is usually successful and straight forward with uneventful short recovery period. One should keep this diagnosis in mind when evaluating an adult patient with such presentation.

Abbreviations

ET: Eustachian tube; OME: Otitis media with effusion; AC: Air conduction; BC: Bone conduction; CT: Computed tomography; MRI: Magnetic resonance imaging.
Acknowledgements
Not applicable for this manuscript

Authors' contributions
KZ is the first author who contributed to the manuscript via collecting data, performing literature review and writing up the manuscript. MB is the senior author who supervised the whole scheme of work and revised the final manuscript. The author(s) read and approved the final manuscript.

Funding
Authors did not receive any funding for this work.

Availability of data and materials
Not applicable for this manuscript.

Declarations

Ethics approval and consent to participate
Ethical approval is not applicable for this manuscript. A written informed consent was obtained from the patient to use his data in this manuscript.

Consent for publication
A written informed consent was obtained from the patient to publish his data including graphs and investigations images in this manuscript

Competing interests
All authors declare that they have no competing interests.

Received: 8 December 2021 Accepted: 24 January 2022
Published online: 21 February 2022

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