



Angioleiomyoma of the external auditory canal: a rare diagnosis—case report and review of the literature

William Howden^{1,2}, Timothy Do¹, Tony Kuo^{1,2,3}

¹Department of Otolaryngology, Gosford Hospital, Gosford, NSW, Australia; ²Department of Medicine, University of Newcastle, Newcastle, NSW, Australia; ³Department of Otolaryngology, Macquarie University Hospital, Sydney, NSW, Australia

Correspondence to: William Howden, MD, BSc. Department of Otolaryngology, Gosford Hospital, Holden St, Gosford, 2250 NSW, Australia.

Email: William.howden@newcastle.edu.au.

Abstract: Angioleiomyomas (ALMs) usually arise from the alimentary tract or uterus, and seldom in the soft tissues of the head and neck. It is particularly rare for an ALM to develop in the external ear canal. To date, including our case—only 5 of these have been described in the literature. A total of 7 cases of the broader classification of leiomyoma (LM) have been reported in the external ear canal, which are reviewed in this report. This case presents a 59-year-old woman with 30-year history of constant, bilateral, non-pulsatile tinnitus, and unilateral high-frequency conductive hearing loss. The report provides a detailed timeline including preoperative investigations with imaging, audiological assessment, intraoperative transcanal endoscopic images, detailed histopathological findings and postoperative functional and objective outcomes. The article discusses current understanding of histopathology and includes a summary and comparison of all published cases of LM of the external auditory canal (EAC) to date.

Keywords: Otolaryngology; endoscopic ear surgery; benign tumor; leiomyoma (LM); angioleiomyoma (ALM)

Received: 12 June 2019; Accepted: 13 January 2020; Published: 28 February 2020.

doi: 10.21037/ajo.2020.01.01

View this article at: <http://dx.doi.org/10.21037/ajo.2020.01.01>

Introduction

Angioleiomyomas (ALMs) are benign tumors arising from smooth muscle cells, which typically occur in the alimentary tract, uterus or lower limb in the 3rd to 6th decade of life. Tumors of smooth muscle are uncommon in the head and neck region. It is very uncommon for a leiomyoma (LM) to occur in the external auditory canal (EAC), and ALMs even more so. Only 5 other cases of ALM (1), from a total of seven cases of LM of the EAC have been reported in the literature (2-6).

In this report, we present a case of a 59-year woman with a 30-year history of symptoms. A review of the literature is also presented.

Case presentation

A 59-year-old female presented with hearing loss initially thought to be secondary to secondary to wax impaction.

She had an associated 30-year history of bilateral non-pulsatile and constant tinnitus. The patient also described a slight aural fullness in her left ear and felt her hearing may have subjectively decreased. There was no history of otalgia, otorrhea or vertigo. Her past medical history was notable only for medically-managed hypertension. She had no relevant family medical history.

Physical examination demonstrated a mildly tender, opaque and mildly vascular soft tissue mass located at the superior aspect of the left external ear canal, obstructing 30% to 40% of the ear canal lumen. The right ear was normal. The remainder of the examination including cranial nerves was unremarkable.

Magnetic resonance imaging (MRI) of her brain and posterior fossa demonstrated a small lesion in the bony portion of her external ear canal (*Figure 1*). The lesion was isointense to brain parenchyma on both T1- and T2-weighted sequences. There were no features to suggest demyelination. Computed tomography (CT) of petrous

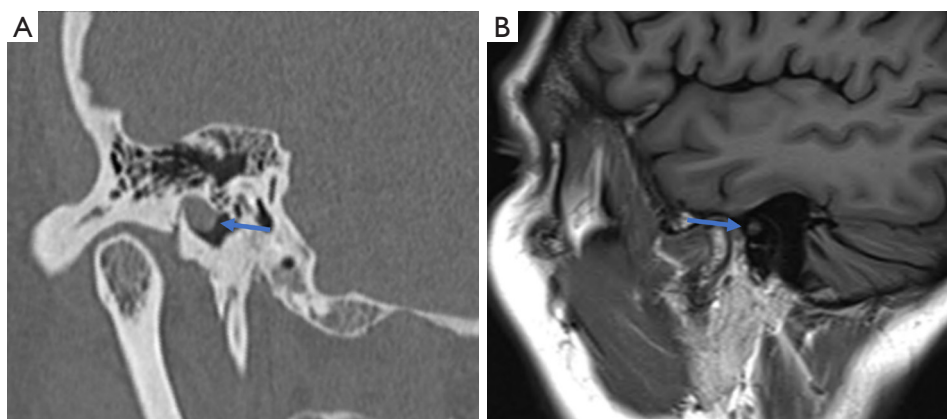


Figure 1 Imaging. (A) CT demonstrating a soft tissue lesion on the superior aspect of the bony ear canal; (B) sagittal MRI image demonstrating a lesion on the superior aspect of the bony ear canal, isointense to brain parenchyma on T1-weighted sequence. CT, computed tomography; MRI, magnetic resonance imaging.

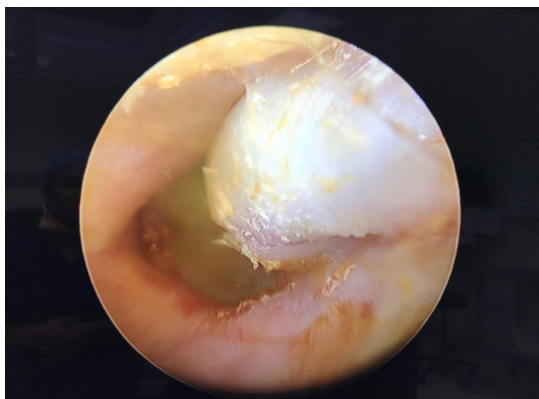


Figure 2 Intraoperative image of the superiorly-based, opaque EAC lesion. EAC, external auditory canal.

temporal bones demonstrated an asymmetric soft tissue thickening measuring 6.5 mm in diameter in the bony portion of the left EAC (*Figure 1*). The lesion was abutting the tympanic membrane. There was no associated bony erosion.

An audiogram demonstrated mild hearing loss of air conduction at 8,000 Hz in the left ear, with all other frequencies normal bilaterally.

The patient subsequently underwent endoscopic assisted excision of the lesion under general anesthetic. Intraoperatively, the superiorly based well-circumscribed opaque lesion was visualized arising from the bony-cartilaginous junction, abutting the tympanic membrane (*Figure 2*). It was partially adherent to the tympanic membrane immediately adjacent to the lateral process

of the malleus and the attic. The mass was peeled off the pars flaccida, and a subsequent small attic perforation was repaired with Gelfoam.

The macroscopic specimen was irregular and opaque, measuring 10×5×5 mm. There was no cystic structure identified. Histopathological analysis (*Figure 3*) demonstrated a mass with closely packed bundles of plump eosinophilic cells. Immunohistochemistry shows diffuse positive staining for smooth muscle actin (SMA) and negative staining for S100 and desmin. Staining for CD34 highlights the numerous small vessels within the lesion. There was no evidence of significant nuclear pleomorphism or increased mitotic activity. The findings are consistent with ALM.

The patient recovered well postoperatively with complete re-epithelization achieved within 4 weeks of surgery. The Attic repair was successful with subjective reduction in tinnitus. Her postoperative audiogram showed no significant change.

Discussion

Benign tumors of the EAC are rare with more common benign lesions being osteoma, melanoma or polyps (6). Our case represents the 8th case overall of LM, and 6th of ALM subtype of the EAC reported in the literature to date. An ALM (ICD-O code 8894/0) is a benign soft tissue tumor of vascular smooth muscle origin. Although relatively common in the lower extremities (accounting for 4–5% of all soft tissue tumors), they are infrequently found on the

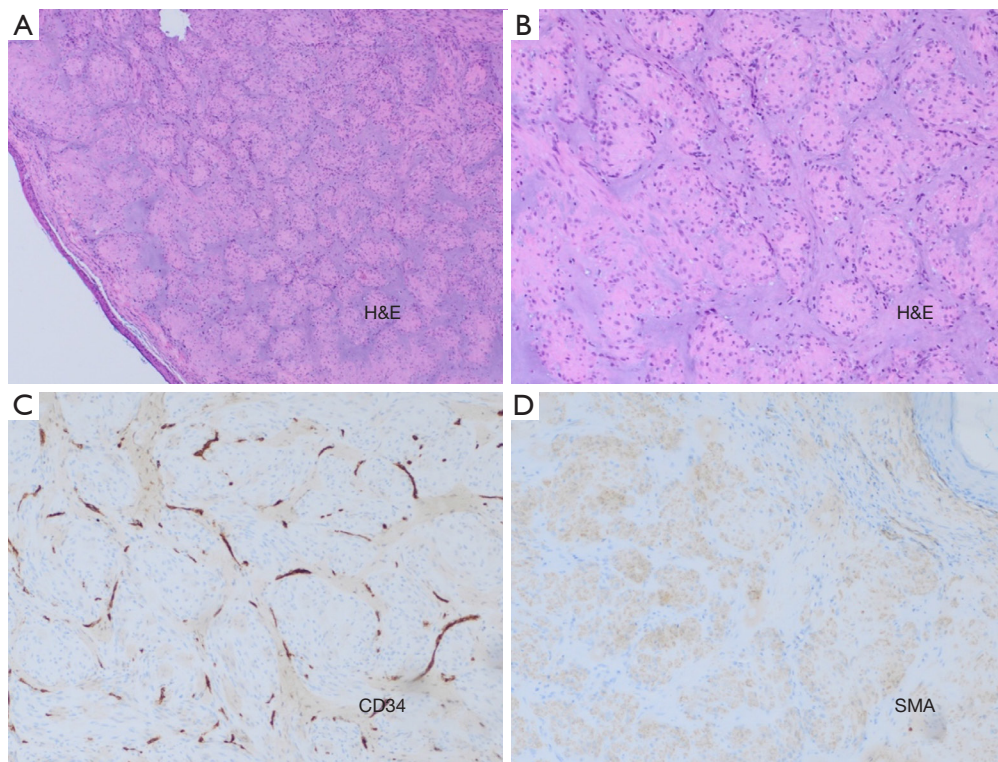


Figure 3 Microscopic images. (A) Low power view of HE staining shows a polypoid mass partly covered by attenuated squamous mucosa; (B) higher power views of the mass consists of closely packed bundles of plump eosinophilic cells with scattered small vessels present throughout; (C) staining for CD34 highlights numerous small vessels within the lesion; (D) further immunohistochemistry demonstrates diffuse positive staining for SMA with negative staining for S100 and desmin. HE, hematoxylin and eosin; SMA, smooth muscle actin.

head and neck. The mean age of diagnosis of EAC tumors is 42.5 (*Table 1*). Their incidence in females is nearly twice that of males when arising in the lower extremity, although EAC LMs are 4 times more common in males than females (*Table 1*). While the pathophysiology of LMs is not fully understood, both trauma and venous stasis have been hypothesized as contributing factors (2,7).

LMs exist on a morphological spectrum with the vascular tumor myopericytoma and are divided into solid, venous and cavernous subtypes. No cavernous LMs of the EAC have been reported to date.

ALMs arise from the smooth muscle cells of the muscular layer of vascular structures, while the solid LM (or pileleiomyoma) have been postulated to have arisen from the erector pili muscles of the skin (6,8).

Microscopically, ALMs contain well-differentiated, closely packed smooth muscle cells, arranged in many vascular channels. Immunohistochemically, the lesion stains positive for smooth muscle actin (SMA), desmin, myosin,

trichrome, HHHF35, calponin and h-caldesmon (with some variability) (9).

Further understanding of the nature of EAC LMs relies on accumulated experience through case reports and literature reviews. *Table 1* demonstrates all accounts of EAC LMs in a literature search of Medline and PubMed. The lesion tends to have a male predominance in the EAC (compared to females, which are more common outside the head and neck region). Although able to arise from anywhere along the EACs extent, the lesion seems to be more common in the bony portion.

Bony erosion on CT or MRI in the EAC would raise suspicion of either cholesteatoma or other malignant processes. Interestingly, it has been demonstrated that both cases of LM arising from erector pili muscles (6,8) have undergone a degree of bony erosion associated with the lesion (*Table 1*), perhaps suggesting a more invasive nature of the tumor compared to its vascular counterpart. This finding makes it an important differential diagnosis in the

Table 1 A summary of reported cases of ECA LMs

Case	Age	Sex	Trauma	Site	Bony erosion	Treatment	Histology	Source
1	47	M	+	Posterior bony canal wall	-	Excision + myringoplasty and mastoidectomy	ALM	Petschenik <i>et al.</i> [1996]
2	56	M	-	Anterior bony canal wall	-	Endaural excision	ALM	Iguchi <i>et al.</i> [1997]
3	47	M	-	Posterior cartilaginous canal wall	-	Retroauricular excision	ALM	Yeh <i>et al.</i> [2005]
4	5	M	-	Posterosuperior/inferior cartilaginous canal wall	-	Retroauricular excision	ALM	Wootten <i>et al.</i> [2007]
5	37	F	-	Posterosuperior bony canal wall	-	Retroauricular excision	ALM	Picciotti <i>et al.</i> [2007]
6	37	M	-	Posteroinferior bony canal wall	+	Endaural excision with endoscopy	LM	Ikeda <i>et al.</i> [2016]
7	52	M	-	Posterior bony canal wall	+	Retroauricular excision	LM	George <i>et al.</i> [2016]
8	59	F	-	Superior bony canal adhered to pars flaccida	-	Transcanal endoscopic excision with myringoplasty	ALM	Howden <i>et al.</i> [2020]

EAC, external auditory canal; LM, leiomyoma; ALM, angioleiomyoma.

investigation of suspected invasive disease.

In our case, the initial imaging investigation was MRI. This was originally reported as normal, despite the request detailing a visualized canal mass. The lesion was only visualized on MRI retrospective to CT and excision. Its presence was confirmed with the reporting radiologist. Although all previously published cases underwent CT, none detail investigation with MRI alone. The mass is certainly more discernible on CT than it is on MRI, and this may suggest a greater sensitivity of CT in the diagnosis of LMs of the EAC.

All cases published cases were managed surgically. While classically an endaural or retroaural microscopic approach had been favored, our case is the second case described of the excision of a LM solely via a transcanal endoscopic approach without extra aural incision as outlined by Ikeda *et al.* (6) in 2016. Enhanced visualization and direct vision of concealed areas are advantages of endoscopic ear surgery over the traditional microscopic techniques. Although transcanal endoscopic ear surgery has been a somewhat controversial topic in the past, its popularity is increasing among head and neck surgeons (10). The use of endoscopic ear surgery is still in its infancy and requires further high-quality data on clinical outcomes, particularly comparative

against traditional direct-vision techniques.

Conclusions

LM of the EAC is a rare diagnosis. The tumor is an important differential in the diagnosis of tumors of the EAC. It occurs mainly in adult males, in their fourth decade of life. The treatment of choice is complete excision, with a transcanal endoscopic approach an effective technique in appropriate patients.

Acknowledgments

The authors thank Dr. Robert W. Laing (MbChB, FRCPath, FRCPA) for assistance with histopathology interpretation and the associated images.

Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related

to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The patient consented for case report and publication of her condition.

References

1. Picciotti PM, Cantore I, La Greca C, et al. Angioleiomyoma of the external auditory canal. *Am J Otolaryngol* 2007;28:235-7.
2. Petschenik AJ, Linstrom CJ, McCormick SA. Leiomyoma of the external auditory canal. *Am J Otol* 1996;17:133-6.
3. Iguchi H, Nakai Y, Yamane H, et al. Smooth muscle tumor of the external auditory canal. *Otolaryngol Head Neck Surg* 1997;116:231-3.
4. Yeh SY, Chao TK. Leiomyoma of the external auditory canal--a case report and brief review of the literature. *Eur Arch Otorhinolaryngol* 2005;262:397-9.
5. Wootten CT, Fasig J, Kaylie D. External auditory canal leiomyoma: a pediatric case report. *Int J Pediatr Otorhinolaryngol Extra* 2007;2:51-3.
6. Ikeda R, Tateda M, Okoshi A, et al. Transcanal endoscopic resection for leiomyoma of the external auditory canal and a review of the literature. *Auris Nasus Larynx* 2016;43:97-9.
7. Ramesh P, Annapureddy SR, Khan F, et al. Angioleiomyoma: a clinical, pathological and radiological review. *Int J Clin Pract* 2004;58:587-91.
8. George MV, Puthiyapurayil J. Leiomyoma of external auditory canal. *Indian J Otolaryngol Head Neck Surg* 2016;68:377-9.
9. Fletcher CDM, Bridge JA, Hogendoorn P, et al. WHO classification of tumours of soft tissue and bone. 4th ed. Lyon: International Agency for Research on Cancer, 2013.
10. Kozin ED, Gulati S, Kaplan AB, et al. Systematic review of outcomes following observational and operative endoscopic middle ear surgery. *Laryngoscope* 2015;125:1205-14.

doi: 10.21037/ajo.2020.01.01

Cite this article as: Howden W, Do T, Kuo T. Angioleiomyoma of the external auditory canal: a rare diagnosis—case report and review of the literature. *Aust J Otolaryngol* 2020;3:2.