Facial nerve palsy secondary to middle-ear lipoma

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Abstract

Objective: We present the first reported case of a middle-ear lipoma presenting with facial nerve palsy. We review the available literature on middle-ear lipomas and alert the surgeon to the possibility of a lipoma occurring in this location.

Case report: A 33-year-old man presented to our unit with a right-sided, House-Brackmann grade two, lower motor neurone facial palsy. A computed tomography scan revealed abnormal soft tissue in the epitympanic recess, extending to the region of the geniculate ganglion. At middle-ear exploration, a lump of fatty tissue was found filling the anterior middle-ear cleft, juxtaposed to the horizontal portion of the facial nerve. The patient's facial palsy resolved within a few weeks of surgery.

Conclusion: Lipomas are a rare but real differential diagnosis of a mass in the middle ear. Early imaging is advised.

Key words: Middle Ear; Facial Paralysis; Lipoma

Introduction

Facial nerve palsy is a relatively common presentation to an ENT department. The majority are idiopathic. ¹ Lipomas in the head and neck are common, benign neoplasms. They are mesenchymal in origin and are typically found in the subcutaneous tissues. They are less commonly encountered in internal organs and are usually asymptomatic. Only five previous cases of middle-ear lipomas have been described in the English literature. To our knowledge, none presented with facial palsy.

Case report

A 33-year-old man presented to the ENT emergency clinic with a one-day history of right-sided, lower motor neurone facial palsy. The preceding week, he had suffered from an upper respiratory tract infection, with some right otalgia and clear otorrhoea. As a child, he had undergone insertion of ventilation tubes for bilateral otitis media with effusion, and had also undergone a right modified radical mastoidectomy for cholesteatoma. He had been asymptomatic for 20 years.

On presentation to our unit, the patient had a grade two House-Brackmann palsy and a mildly congested right tympanic membrane. The rest of the ENT examination was normal.

A diagnosis of acute otitis media and right facial palsy was made. The patient was treated with a seven-day course of oral prednisolone 60 mg once daily, ciprofloxacin 500 mg twice daily, acyclovir 800 mg five times daily and Gentisone HC ear drops (Roche, Basel, Switzerland) thrice daily.

At one-week review, the patient's facial palsy had deteriorated to House-Brackmann grade three. The tympanic membrane remained congested and was noted to be retracted with an underlying effusion. The pure tone

audiogram revealed a moderate right conductive hearing loss.

At four-week review, the patient's facial palsy had improved (to House-Brackmann grade two). However, the apparent middle-ear effusion persisted. A computed tomography (CT) scan revealed abnormal soft tissue in the epitympanic recess, extending to the region of the geniculate ganglion (Figure 1).

In view of the patient's history of mastoidectomy for cholesteatoma, recurrence was suspected, and it was decided to undertake a middle-ear exploration. This revealed an encapsulated lump of fatty tissue filling the anterior middle-ear cleft, juxtaposed to the horizontal portion of the facial nerve.

Histological analysis confirmed the presence of a lipoma, with evidence of chronic inflammation within the benign tumour (Figure 2).

Post-operatively, the patient made a good recovery. His facial palsy resolved within a few weeks of surgery.

Discussion

Lipomas in the middle ear are rare, and only five cases have been previously reported. We performed a literature review by searching PubMed and Ovid, using the key words 'lipoma', 'middle ear' and 'mastoid'.

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Luetje et al.² found that 17 per cent of 240 temporal bones examined at the House Ear Institute demonstrated varying amounts of fat juxtaposed to the mastoid vertical portion of the facial nerve. Essentially, two theories exist as to the origin of fat in the middle ear. Luetje et al. postulated that, embryologically, fat marrow which has persisted in the mastoid could lead to the growth of ectopic fat in the middle ear. Their other hypothesis was that mastoid and VIIth nerve development could have dragged fat cell precursors into the middle ear, which could then lead to lipoma development.²

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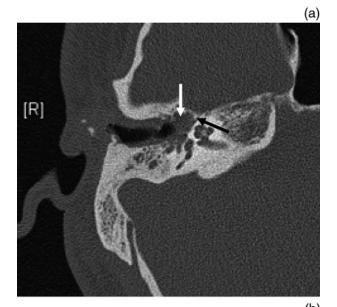




Fig. 1

(a) Axial and (b) coronal computed tomography images demonstrating a soft tissue mass (white arrow) adjacent to the horizontal portion of the facial nerve in the fallopian canal (black arrow).

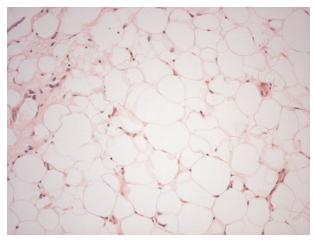


Fig. 2

Histological section of the mass found in the right middle ear, demonstrating adipose tissue and scanty lymphocytes in a fibrous stroma (H&E; \times 100).

Stegehuis *et al.*³ described a most unusual case of a middle-ear lipoma that extended down the eustachian tube into the oropharynx, presenting with airway distress in a 64-year-old woman. Selesnick *et al.*,⁴ Abdullah *et al.*⁵ and Edmonds *et al.*⁶ all reported middle-ear lipomas in children presenting with middle-ear effusions and conductive hearing loss. Abdullah *et al.* described bilateral middle-ear lipomas which were found to be obstructing the eustachian tube and causing a middle-ear effusion in a five-year-old child.⁵ Luetje *et al.* reported the case of a 16-year-old adolescent with bilateral lipomas who presented with an incidental, unilateral, high frequency sensorineural hearing loss.² None of the previously reported cases of middle-ear lipoma has been associated with facial palsy. Agarwal *et al.*, ⁷ however, described a case of unilateral facial nerve palsy caused by a primary liposarcoma in a four-year-old boy.

Our patient's lipoma had presumably developed since his childhood mastoidectomy, as there was a considerable quiescent period after this procedure. It is possible that the mastoidectomy disturbed fat precursor cells from persisting bone marrow fat, or triggered a growth response in the adipose cells already present in the middle ear, by the mechanisms proposed above. Possible aetiologies for the VIIth nerve palsy include direct nerve compression from the lipoma, or, more likely, the effect of the acute otitis media due to eustachian tube obstruction secondary to the lipoma. This latter theory is supported by the fact that the patient's facial nerve palsy showed signs of improvement prior to surgery.

This case report reminds us to be thorough in our evaluation of patients with facial nerve palsies, especially when there is a history of previous mastoid surgery. Recurrent cholesteatoma is the most likely aetiology, and surgical exploration and decompression would be advised. Computed tomography is the first-line investigation; in this case, magnetic resonance imaging may have shown a mass with similar intensity to fat.

Conclusion

Lipomas are a rare but real differential diagnosis of a mass in the middle ear. When removed, the surgeon should expect any pre-existing facial nerve palsy to recover. To our knowledge, the present case represents the first published report of a facial nerve palsy arising secondary to a middle-ear lipoma.

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CLINICAL RECORD 3

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