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Keratosis Obturans

IMAGE

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A 13-year-old male patient presented with pain and hearing loss in the right ear for 5 days. Examination revealed a brownish-white flaky mass

surrounded by granulation tissue in the right external auditory canal. A provisional diagnosis of keratosis obturans was made, and the mass was removed under general anesthesia (Fig. 1a). Postremoval examination revealed erosion and granulation in the external auditory canal (Fig. 1b), and postoperative biopsy confirmed the diagnosis (Fig. 1c). The patient was started on antibiotics and steroid drops for 7 days and was asymptomatic 3 months postoperatively. Keratosis obturans is characterized by abnormal accumulation of desquamated keratin in the external auditory canal, leading to the occlusion and expansion of the bony portion of the canal.1 It is common in young patients and can be bilateral in 50% of the cases. The etiology may be linked to abnormal shedding of the epithelium and failure of cell

migration from the surface of the

tympanic membrane, leading to the

accumulation of a mass (1, 2). The

condition is more common in pa-

b c

Figure 1. (a) Whitish flaky mass removed from the external auditory canal. (b) Postoperative picture showing granulation tissue and erosion of the external auditory canal. (c) Histopathological image revealing laminated flakes of keratin with chronic inflammatory infiltrate $[HE, 40\times]$

tients with bronchiectasis and chronic sinusitis and in smokers. The presentation is usually of severe ear pain and conductive deafness (2). Computer tomography typically demonstrates soft tissue plug in the external ear canal with an expansion of the osseous part. The differential diagnosis includes external auditory canal cholesteatoma and impacted cerumen (2, 3). Softening of the mass using sodium bicarbonate and glycerine solution followed by removal under anesthesia is the treatment of choice (3). Widening of the external auditory canal followed by skin grafting is rarely required. Prognosis is good; however, because recurrence is common, close follow-up of the patients is advised (1).

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