Osteoma of Internal Auditory Canal - A Rare Pathology

Bhushita Nilesh Guru1, Bhushan Narayan Lakhkar2

1Department of Radiology, Datta Meghe Institute of Medical Sciences, Sawangi (Meghe), Wardha, Maharashtra, India. 
2Department of Radiology, Datta Meghe Institute of Medical Sciences, Sawangi (Meghe), Wardha, Maharashtra, India.

PRESENTATION OF CASE

A 27-year-old female patient visited the Department of Radiology with complaints of right sided facial palsy and sensory-neural hearing loss from past 10 years. Otologic examination revealed both tympanic membranes to be normal. Audiometry revealed right sided sensory neural hearing loss. The patient was also having multiple facial spasms.

HRCT temporal bone of the patient was done, and it showed a well-defined round to oval bony out-pouching arising from posterior wall of right internal auditory canal causing severe stenosis of porus acusticus with only 7 mm patency. (Figure 1) The lesion was noted to be over the vestibulo-cochlear and the facial nerves. The cortex of the lesion was continuous with that of the parent bone. (Figure 2). The left internal auditory canal was normal.

DISCUSSION

Osteomas are one of the common benign bone pathologies. They are often confused with exostosis; hence, the diagnosis of osteoma and exostosis go hand in hand. The most common sites for osteomas include the frontal sinus, ethmoid sinus, the bony calvarias and external auditory canal.1

Osteomas are benign slow growing tumours of the bone. It is one of the commonest pathologies within the external auditory canal, however they are one of the rarest pathologies of the internal auditory canal. In the external ear they occur as a result of repeated trauma, infection or irritating stimuli; but its pathogenesis within the internal auditory canal is still unknown. Till now only 21 cases of the same have been reported in literature. 2

The most common differential diagnosis of osteomas is exostosis; however, exostosis generally appears bilaterally.3 Biopsy of the lesions of internal auditory canal is not possible; hence, the radiographic appearance plays an important role in differentiating exostosis from osteoma or any other bony lesion. Exostoses are usually multiple and bilateral having a smooth border, have a broad based and grow towards internal auditory canal; whereas, osteomas are solitary, stalked lesion with its narrow continuous with that of the parent bone.4

There are a number of causes that can compromise the lumen of internal auditory canal, like Paget’s disease, otosclerosis or fibrous dysplasia. They may both lead to expansion of the bones of canal or new bone formation within the canal and cause its narrowing. However, typical appearance of an osteoma helps to differentiate it from rest of the causes.5

Such patients usually present with sensory neural hearing loss, tinnitus, or facial palsy and spasm. It might be due to either direct compression or the lesion acting like a fulcrum causing repeated indentations over the nerves with movement. An osteoma will narrow the caliber of the canal and compress the 7th and 8th cranial nerve.

Treatment is still controversial and is determined by the severity of the symptoms. Asymptomatic patients are usually observed every 6 months. Patients with mild symptoms are advised serial audiograms and CT scans. Patients with severe symptoms, such as those with progressive hearing loss or vertigo and near complete occlusion of the IAC may require surgical excision and nerve decompression before irreversible damage occurs.2
CONCLUSIONS

Bony osteoma is one of the rarest pathologies of internal auditory canal. Patients presenting with unilateral hearing loss or palsy should be investigated for it and should be treated according to the severity of symptoms.

Financial or Other Competing Interests: None.

REFERENCES