Schwannoma of the intermediate nerve: A rare type of cerebellopontine angle tumor

Schwannoma del nervio intermedio: Un tipo raro de tumor del ángulo pontocerebeloso

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Resumen

Antecedentes: El schwannoma del nervio intermedio es poco común. La mayoría de las lesiones del ángulo pontocerebeloso (APC) son schwannomas vestibulares. Descripción del caso: Mujer de 50 años que consulta por dolor de oído, parálisis facial e hipoacusia. La resonancia magnética reveló un tumor APC. Se realizó abordaje retrosigmoideo izquierdo y microcirugía y el origen tumoral fue del nervio intermedio. Se utilizó monitorización neurofisiológica intraoperatoria (MNIO) durante la cirugía. Conclusión: Los schwannomas del nervio intermedio son indistinguibles de los schwannomas vestibulares en la imagen preoperatoria, sin embargo, es posible identificar el origen del tumor durante la disección microquirúrgica con la ayuda de MNIO y el origen del nervio intermedio es una realidad.

Palabras clave: Nervio intermedio, Schwannoma, tumor APC.

Abstract

Background: Schwannoma of the Intermediate Nerve is rare. The most majority of cerebellopontine angle (CPA) lesions are vestibular Schwannomas. Case description: A 50-year-old woman presented with intense ear pain, facial palsy and hypoacusia. MRI revealed a CPA tumor. A left retrosigmoid approach and microsurgery was performed and the tumor origin was from the intermediate nerve. Intraoperative neurophysiologic monitoring (MNIO) was used during the surgery. Conclusion: Schwannomas of the intermediate nerve is indistinguishable from vestibular schwannomas on preoperatory image, however it’s possible to identify tumor origin during microsurgery dissection with aid of MNIO and origin from intermediate nerve is a reality.

Key words: Intermediate nerve, Schwannoma, CPA tumor.

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Introduction

The intermediate nerve (IN) is described as a root containing sensory and parasympathetic fibers of the facial nerve and its fibers are found in close relation to the motor part of the facial nerve in the cerebellopontine cistern. The first person who identified the IN was Bartholomeu Eustachius in 1563 and the first one to describe was Heinrich Albert Wirsberg in 1777 due to this is often referred as Wrisberg nerve. It is responsible for the sense of taste in the anterior two-thirds of the tongue, floor of the mouth, palate and sensory information from the skin of the external auditory meatus, as well as the lacrimal and mucous glands of the nasopharynx and nose. The IN originates at the brainstem between the facial nerve and the vestibulocochlear nerve in the lateral medullopontine sulcus; the IN accompanies the facial nerve or the vestibulocochlear nerve and with high-resolution 3T MR imaging allows reliable depiction of the IN in some cases.

About 10% of intracranial tumors occur in the internal auditory canal (IAC) and cerebellopontine angle (CPA). The vast majority (95%) of these tumors are vestibular schwannomas, these types of tumors originate from myelin schwann cells of the vestibular portion of the vestibulocochlear nerve. Schwannoma of the IN is rare and there are only 3 cases reported in the literature. Although, 3 other cases with origin from IN were found in other series.

Case Description

A 50-year-old woman presented with intense ear pain, facial palsy classified as House-Brackmann (HB) V/VI and hypoacusis on the left. MRI revealed an enhanced CPA lesion in the left (Figure 1).

Patient underwent microsurgical resection via left retrosigmoid approach, in semi-sitting position and intraoperative neuropsychologic monitoring (MNIO) was used during the surgery. MNIO consisted of motor evoked potentials for corticospinal tract with anodal C3 or C4 stimulation and recording with a pair of needles in right and left muscles abductor pollicis brevis, anterior tibialis and abductor hallucis. Somatosensory evoked potentials after right and left median and tibial nerves stimulation resulted in normal N20 and P40 potentials. Motor evoked potentials for corticobulbar tract used C6-Cz assembly and recordings were realized in muscles of the V, VI, VII, IX, X, XI and XII nerves. Consistent muscle potentials were obtained for V, VII, IX and X nerves, which remained unchanged throughout the procedure. Free run electromyography of the same cranial muscles showed intermittent short duration type A neurotonic activity at VII and X innervated muscles, promptly stopped with warm saline irrigation. Direct nerve stimulation with a bipolar probe at intensities of 0,2 and 0,4 mA, with pulse duration of 200 uS, identified V, VII and IX in different sites. Of remark, in the internal auditory canal, VII nerve was identified and was not the origin of the tumor.

Tumor origin was identified from the intermediate nerve (Figure 2a) and MNIO confirmed that origin was not from motor root of facial nerve. A gross total resection of tumor was achieved preserving IN, the motor portion of facial and vestibulocochlear nerves. Histopathological examination confirmed schwannoma (Figure 2b, Figure 3).

The patient presented a good postoperative recovery with complete improvement in ear pain after surgery and only mild...
also schwannoma of the intermediate nerve 11. Hemangiomas, facial motor schwannomas, meningiomas and the IAC such as peripheral nerve sheath tumors, cavernous reported rare tumors that mimic vestibular schwannomas in sign of tumor presence at IAC. When small sized tumors as the Hitselberger sign, can occur. It is considered an early the posterior aspect of the external auditory canal, known or compression, loss of lacrimation, decreased sensation in IAC are being detected. In case of intermediate nerve section asing number of asymptomatic smaller tumors localized to the keep in mind.

With advances in MRI neuroimaging technology, an increasing number of asymptomatic smaller tumors localized to the IAC are being detected. In case of intermediate nerve section or compression, loss of lacrimation, decreased sensation in the posterior aspect of the external auditory canal, known as the Hitselberger sign, can occur. It is considered an early sign of tumor presence at IAC 1. When small sized tumors grow and become more symptomatic, they are often treated with radiosurgery without histologic confirmation. However, if tumors grow and symptoms progress enough, microsurgery is required 11.

During surgery, the facial motor, intermediate nerve, vestibulocochlear and other cranial nerves can be identified by direct nerve stimulation 3,7,9. The stimulation of the intermediate nerve gives no potential, since it is a sensory nerve. On the other hand, the facial nerve stimulation results in a muscle action potential. This simple technique is used to help identify them both, like in our case.

The operative approaches that can be used are retrosigmoid, trans labyrinthine, suboccipital or subtemporal. In cases of schwannoma of the NI found in the literature, the retrosigmoid approach 3,11 and the suboccipital craniotomy were performed 3,7. In our patient, we performed the retrosigmoid approach with good route and total resection.

Figure 3. Post-operative axial MRI showing complete removal of the tumor.

Discussion

Intermediate nerve tumor is indistinguishable from vestibular or facial motor schwannoma on MRI 11. Watanabe et al., reported rare tumors that mimic vestibular schwannomas in the IAC such as peripheral nerve sheath tumors, cavernous hemangiomas, facial motor schwannomas, meningiomas and also schwannoma of the intermediate nerve 11 and they also cited glioblastoma multiforme, metastases, primary adenocarcinoma, arachnoid cysts, lipomas, lipochoristoma, melanoma from another series 11 which is important to neurosurgeon to keep in mind.

Conclusion

Schwannoma of the intermediate nerve is rare. When there is a tumor arising from this nerve it cannot be differentiated from vestibular or facial schwannomas by imaging. However, the identification of the origin of tumor from IN is possible during microsurgery dissection with direct stimulation and, also, symptoms can suggest the diagnosis. It is worthy to keep in mind that schwannoma of the IN is a possibility.

Conflict of Interest: None.

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