Facial Spasm and Paroxysmal Tinnitus Associated with an Arachnoid Cyst of the Cerebellopontine Angle
—Case Report—

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Abstract

A 59-year-old female presented with a 3-year history of hemifacial spasm and paroxysmal tinnitus associated with an arachnoid cyst of the cerebellopontine angle, which was identified preoperatively by magnetic resonance imaging. Surgical decompression of the cyst and arterial decompression of the facial and acoustic nerves at their root exit zones resulted in complete resolution of the symptoms.

Key words: cerebellopontine angle, arachnoid cyst, hemifacial spasm, tinnitus

Introduction

Hemifacial spasm, sometimes associated with hyperdysfunction of the acoustic nerve such as tinnitus, is most commonly caused by compression of the facial and acoustic nerve roots by an aberrant vessel, usually a branch of the anterior inferior cerebellar artery (AICA), posterior inferior cerebellar artery, or vertebral artery. More serious but rarer causes of compression include posterior circulation aneurysms, vascular malformations, and various types of tumors. We report a rare case of a cerebellopontine (CP) angle arachnoid cyst that indirectly caused hemifacial spasm and tinnitus.

Case Report

A 59-year-old female was admitted to our hospital on April 1995, with a chief complaint of twitching of the left facial muscles and paroxysmal tinnitus on the left side for 3 years. The spasm originated in the orbicularis oculi and extended to the lower part of the face. She had been treated with tranquilizers for several months without obvious beneficial effects before referral. The spasm and paroxysmal tinnitus had become more severe and more frequent. No other neurological abnormalities including hearing deficits were detected.

Fig. 1 Computed tomography (CT) scan showing a cystic lesion in the left cerebellopontine angle and brain stem displacement (left). Metrizamide CT cisternogram showing a cystic lesion well filled with contrast medium 3 hours after injection (right).

Computed tomography (CT) demonstrated a homogeneous lesion in the left CP angle with similar density to the cerebrospinal fluid (Fig. 1 left). Metrizamide CT cisternogram revealed no filling defect in the left CP angle 3 hours after injection (Fig. 1 right) and similar clearance 48 hours after injection. Magnetic resonance (MR) imaging showed enlargement of the left CP angle cistern, which had caused deviation of the lower pons (Fig. 2 left). Coronal MR imaging demonstrated the vascular
loop impinging on the left pontomedullary junction (Fig. 2 right). Vertebral angiography disclosed elongation of the left vertebral artery and the left AICA (Fig. 3).

**Fig. 2** Axial T2-weighted magnetic resonance (MR) image showing enlargement of the left cerebellopontine angle cistern and brain stem deviation (left). Coronal T1-weighted MR image showing the vascular loop impinging on the pontomedullary junction (right).

**Fig. 3** Left vertebral angiogram showing elongation of the left vertebral artery and anterior inferior cerebellar artery.

The diagnosis was vascular cross-compression of the facial and acoustic nerves associated with a left CP angle arachnoid cyst. Microvascular decompression and cyst decompression by the suboccipital approach were performed in May 1995. Exposure of the cyst in the CP angle showed the cyst wall was clear and transparent. The cyst was located around the facial and acoustic nerves and extended from the porus acusticus to the brain stem. Excision of the outer wall of the cyst allowed clear cerebrospinal fluid to escape from the cyst. Histological examination of the cyst wall biopsy specimen indicated reactive arachnoid membrane. Two branches of the AICA were located around the facial and acoustic nerves. One large branch ran between the facial and acoustic nerves and another small branch ran along the dorsal aspect of acoustic nerve, compressing the facial nerve ventrally and pinching the acoustic nerve at the root exit zone (Fig. 4). The large branch of the AICA was separated from the facial and acoustic nerves and small pieces of Teflon felt were inserted between the artery and the facial nerve, and between the artery and the acoustic nerve. The small branch of the AICA was separated from the acoustic nerve by careful dissection of the arachnoid between the artery and the nerve so that neither artery could compress the facial and acoustic nerves directly. The Teflon felts were fixed to the dura of the petrous bone side to prevent displacement.

Immediate postoperative condition was excellent, and she was completely relieved of hemifacial spasm and tinnitus. At follow-up 2 years after the operation, she continued to be free of the symptoms and MR imaging demonstrated the size of the arachnoid cyst was reduced (Fig. 5).
Discussion

The clinical symptoms produced by a cyst in the CP angle may closely mimic those of an acoustic neuroma, with sensorineural hearing loss, impaired corneal reflex, trigeminal neuralgia, and in the late stages cerebellar signs and increased intracranial pressure. In our patient, the CP angle arachnoid cyst had caused compression of the brain stem, resulting in stretching or bending of the facial and acoustic nerves and deviation of the ipsilateral AICA. The secondary result was contact between the AICA and the root exit zone of the facial and acoustic nerves. Hyperactive dysfunction of the acoustic nerve, such as tinnitus, associated with facial spasm may be caused by cross-vascular compression of the facial and acoustic nerves at their root exit zone. A similar case of hemifacial spasm associated with CP angle arachnoid cyst was previously described. Evacuation of the arachnoid cyst alone had no beneficial effect, and the additional microvascular decompression was needed for the relief of hemifacial spasm. Another case of hemifacial spasm associated with cerebellar arachnoid cyst was treated effectively by removal of the cyst and decompression of the facial nerve. Like other CP angle tumors responsible for hemifacial spasm or other cranial nerve compression syndromes, the CP angle arachnoid cyst may be only indirectly responsible for the syndrome, acting by displacing a small artery to impinge upon the root entry zone of the affected facial and acoustic nerves. Successful treatment of hemifacial spasm associated with CP angle arachnoid cyst can be achieved by decompression of the cyst and arterial decompression at the nerve root exit zone.

References


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