Sigmoid Sinus Dural Arteriovenous Malformation Resulting from Jugular Foramen Schwannoma

—Case Report—

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Abstract

A 69-year-old male presented with a jugular foramen schwannoma occluding the sigmoid sinus and associated with sigmoid sinus dural arteriovenous malformation. The patient presented with dizziness and pulsatile tinnitus following an extended period of hearing loss beginning several years before. Both lesions were resected successfully after transarterial embolization of the malformation. The sequence of symptom development suggests the dural sinus thrombosis caused the dural arteriovenous malformation.

Key words: dural arteriovenous malformation, jugular foramen, schwannoma, sinus thrombosis

Introduction

Dural arteriovenous malformations (AVMs) may result from congenital factors as shown by occurrence in infancy and characteristic angiographic features, but most dural AVMs are acquired. Dural AVM may develop after trauma, surgery, and dural sinus thrombosis. Dural sinus thrombosis is regarded as the causative event in cases of acquired dural AVM. Dural AVM may develop in various intracranial sites, but two-thirds of cases involve the transverse or sigmoid sinus.

Tumors near the sigmoid sinus often infiltrate and occlude the sinus. Jugular foramen schwannomas comprise 2.9% to 4% of intracranial schwannomas. Angiography shows that one third of jugular foramen schwannomas completely occlude the jugular bulb and the sigmoid sinus, and jugular foramen schwannoma is associated with a high incidence of sinus thrombosis. We describe a case of jugular foramen schwannoma associated with a sigmoid sinus dural AVM.

Case Report

A 69-year-old male had noted loss of hearing in the right ear for many years. A recent otological examination had been unrevealing. He developed intermittent dizziness and pulsatile tinnitus several months prior to hospital admission on July 26, 1994. On admission, auscultation revealed a bruit in the right retromastoid region. Lower cranial nerve function was intact. Audiometry indicated thresholds of 45 dB on the right and 25 dB on the left. Auditory brainstem responses to right-sided stimuli detected no response except for peak I. Otological examination suggested a retrocochlear lesion. Magnetic resonance imaging revealed a dumbbell-shaped tumor near the right jugular foramen extending in the intra- and extracranial directions (Fig. 1). Angiography revealed a dural AVM supplied by the right external carotid artery and involving the right sigmoid sinus. The right jugular bulb and proximal sigmoid sinus were occluded. The dural AVM was cross-drained via the torcular sinus to the opposite transverse and sigmoid sinuses (Fig. 2).

The patient underwent transarterial embolization of the dural AVM using polyvinylalcohol particles (150 to 250 μM). His pulsatile tinnitus was ameliorated by the embolization. The jugular foramen tumor and dural AVM were then resected using a transjugular approach. The vestibulocochlear nerve in the cerebellopontine angle cistern was compressed and displaced rostrally by the intradural tumor. The tumor filled the jugular bulb and sigmoid sinus where angiography had revealed occlusion (Fig. 3). The histological diagnosis of the resected tumor was schwannoma.
Postoperative angiography showed complete obliteration of the dural AVM (Fig. 4).

**Discussion**

Previously, angiography has shown dural AVM was preceded by dural sinus thrombosis in two cases, and multiple dural AVMs evolved from sinus thrombosis induced by transvenous embolization in one case. Brain tumors near dural sinuses often infiltrate the sinus and cause thrombosis. Superior sagittal sinus occlusion by a parasagittal meningioma is fairly common. A case of meningioma involving the sigmoid sinus groove and associated with sigmoid sinus thrombosis and dural AVM involving the transverse sinus suggested that sinus thrombosis due to tumor involvement had caused development of the dural AVM.

Dural AVMs most frequently involve the transverse and sigmoid sinuses. Schwannoma is the most common of the many tumors occurring near the jugular foramen. Jugular foramen schwannomas often occlude the sigmoid sinus. The present patient had a jugular foramen schwannoma, sigmoid sinus thrombosis, and a sigmoid sinus dural AVM. The chronology of development of these lesions, presumably in the order of jugular foramen schwannoma, sinus thrombosis, then dural AVM, was not proven by serial angiography spanning the period of dural AVM formation. However, the patient developed hearing loss several years before the onset of pulsatile tinnitus. Such hearing loss is consistent with the observed compression of the
cochlear nerve\(^6\) by the jugular foramen schwannoma in the cerebellopontine angle. Pulsatile tinnitus, the most common symptom of dural AVM involving the transverse and sigmoid sinuses,\(^7,10\) occurred later in the present case and was ameliorated by embolization of the dural AVM. The link between the tumor and the dural AVM was sigmoid sinus thrombosis due to tumor involvement. Despite the absence of an angiogram preceding dural AVM formation, the symptom progression is convincing evidence of the causative sequence in this case.

References


Fig. 3 Schema showing the jugular foramen schwannoma, sigmoid sinus thrombosis, and sigmoid sinus dural arteriovenous malformation (AVM). The right dumbbell-shaped jugular foramen schwannoma has totally filled the jugular bulb and sigmoid sinus (arrow). There is an associated sigmoid sinus dural AVM (arrowhead).

Fig. 4 Right carotid angiograms after surgical resection of the schwannoma and the dural arteriovenous malformation indicating no residual lesion.

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