Coil Embolization of a Trigeminal-Cavernous Fistula

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Summary: A 53-year-old woman spontaneously incurred a right trigeminal artery–cavernous sinus fistula, manifested by an intracranial bruit and right sixth nerve palsy. This lesion was successfully managed by coil embolization via the transvenous and transarterial routes.

The trigeminal artery is a persistent, embryonic, vascular anastomotic connection between the cavernous internal carotid artery (ICA) and the basilar artery. Its rate of occurrence is between 0.1% and 0.3% (1, 2). Generally, flow within the trigeminal artery is from the ICA to the basilar artery. We describe a case of a spontaneous fistula from the trigeminal artery to the cavernous sinus.

Case Report

A previously healthy right-handed 53-year-old woman was gardening when she experienced the sudden onset of a pulsatile bruit in her right ear and double vision. Upon examination in an emergency department 10 days after the onset of symptoms, she was found to have a right orbital bruit and a right sixth nerve palsy. An arteriogram showed the presence of a Saltzman type 2 persistent trigeminal artery (3) associated with a right-sided trigeminal-cavernous fistula (Fig 1A) as well as a carotid-cavernous fistula (Fig 1B). No discrete aneurysm or cortical venous filling was seen. The direction of filling of the trigeminal artery was from the basilar artery to the right cavernous sinus.

One week after the initial angiogram, elective endovascular treatment of the lesion was undertaken. Via the right transfemoral approach, using systemic heparinization, and with the patient under general endotracheal anesthesia, a 6F guiding catheter was directed into the right ICA. Through this guiding catheter, a microcatheter distally mounted with an Interventionsal Therapeutics Corp. detachable silicone balloon was directed under digital roadmap imaging into the distal right ICA. The microcatheter with its detachable silicone balloon could not be flow-directed across the rent in the cavernous right ICA. Next, using the left transfemoral venous approach, a 7F Bernstein catheter was directed into the right internal jugular vein. A coaxial microcatheter was directed over a 0.016-inch guidewire through the right inferior petrosal sinus, into the right cavernous sinus, and finally into the right superior ophthalmic vein. Ninety fibered platinum microembolization coils were deposited. A repeat vertebral arteriogram showed complete occlusion of the fistula (Fig 1E). The patient was awakened from general endotracheal anesthesia and was noted to have sustained mild left hemiparesis and dysarthria, believed to be the result of a right hemispheric stroke. Immediately after the procedure, the patient noted that her intracranial bruit had resolved. The hemiparesis and dysarthria gradually resolved over the next several weeks with physical and speech therapy. The right sixth nerve palsy and diplopia resolved over the next 3 months. At 12 months after embolization, the patient had had no recurrence of intracranial bruit or diplopia.

Discussion

The blood supply to the developing hindbrain is derived from a series of channels connecting the developing carotid arteries and the longitudinal neural arteries. These connections include the posterior communicating, trigeminal, otic, hypoglossal, proatlantal, intersegmental, and persistent cervical intersegmental arteries. All of these normally regress, except for the posterior communicating artery (4, 5). The trigeminal artery is the most common persistent connection. It is associated with vascular steal, ocular palsies, aneurysmal formation, vascular rupture, subarachnoid hemorrhage, trigeminal neuralgia, and arteriovenous fistula (1, 2, 6–9). In our case, an aneurysm was not found, but the spontaneous development of a trigeminal-cavernous fistula suggests that one may have been present. There is a 14% chance of an intracranial aneurysm being found in association with a trigeminal artery, but aneurysms of the trigeminal artery itself have a significantly lower rate of occurrence of (10). An alternative explanation for the fistula in our patient is spontaneous vascular rupture. Some authors have suggested that structural abnormalities may exist in the wall of the trigeminal artery that predispose toward vascular rupture (11, 12).

The goal of therapy in repairing carotid-cavernous fistulas is to preserve vision, eliminate intracranial bruit, and avoid ischemic and hemorrhagic complications. Emergency treatment of these lesions is neces-

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sary when there is progressive vision loss and the development of cortical and/or deep cerebral venous drainage (13, 14). Treatment is generally by transarterial delivery of a detachable balloon within the affected cavernous sinus (2, 7, 8, 15, 16). Carotid-cavernous fistulas have also been treated by transvenous embolization using balloons, coils, and liquid adhesive agents (13, 17).

Conclusion

This is an unusual case of a trigeminal-cavernous fistula that was occluded by coil embolization of the cavernous sinus and the trigeminal artery via the transvenous and transarterial routes, respectively.

References


Fig 1. 53-year-old woman with sudden onset of a pulsatile bruit in the right ear and double vision.

A, Lateral left vertebral digital arteriogram shows a persistent trigeminal artery (arrowhead) with flow of contrast material into right cavernous sinus, indicating a trigeminal-cavernous fistula.

B, Lateral right ICA digital arteriogram shows a large carotid-cavernous fistula.

C, Right ICA lateral digital arteriogram after coil embolization of the posterior right superior ophthalmic vein and right cavernous sinus shows minimal residual flow through the carotid-cavernous fistula to the right inferior petrosal sinus.

D, Repeat lateral left vertebral digital arteriogram shows continued early venous opacification (arrow), the result of residual flow through the trigeminal-cavernous fistula.

E, Postembolization lateral left vertebral digital arteriogram shows no further filling of the trigeminal artery and obliteration of the trigeminal cavernous fistula.