Cerebral Arteriovenous Malformation Causing Epistaxis

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Summary: We present an unusual case of epistaxis due to a right frontal arteriovenous malformation and underscore the need to follow a strict protocol in the angiographic work-up for possible embolization of epistaxis.

Index terms: Arteriovenous malformations, cerebral; Epistaxis; Cerebral angiography, indications

Epistaxis can be life threatening and may not respond to conventional conservative measures such as packing. In our institution, between 1984 and 1991, we have treated 65 patients with embolization who had epistaxis that did not stop with nasal packing, with a success rate of 90%. This case report illustrates an unusual example of epistaxis due to a right frontal arteriovenous malformation (AVM).

Case Report

A 53-year-old man had a 4-week history of intermittent epistaxis. Despite anterior and posterior nasal packing for 4 days, he continued to bleed. The patient required transfusion with four units of blood. He had no history of previous nose bleeds and he was not being given anticoagulants. There was no family history of epistaxis and he had no skin telangiectasias. A left carotid angiogram showed a micro AVM (nidus less than 1 cm) (Figs. 1A, 1B, and 1C) on the undersurface of the right gyrus rectus. It was fed by an orbitofrontal branch of the right anterior cerebral artery and drained into two cortical veins that emptied into the superior sagittal sinus. In addition, a venous network was also opacified in the nasal fossa. Superselective angiography was attempted to further define the morphology of the AVM. However, we were unable to catheterize selectively the feeding artery. The AVM was resected surgically without complication through a right frontal craniotomy. Since the removal of the AVM, the patient has not experienced any further epistaxis. A control angiogram showed no residual AVM and disappearance of the nasal venous network (Fig. 1D).

Discussion

Nasal packing is the first line of treatment in epistaxis; however, during the past decade, embolization has become the treatment of choice in those that do not respond to packing. Between 1981–1988, we treated 40 patients by embolization. Most cases were idiopathic and occurred in patients over 50 years of age, probably related to hypertension. The other causes included coagulopathy, juvenile nasopharyngeal angiofibroma, Rendu-Osler-Weber syndrome, vascular malformation of the nasal cavity, and trauma (1). Epistaxis can also result from a biopsy of an unknown vascular nasal mass or a rupture of a cavernous aneurysm (2) or a carotid cavernous fistula leaking into the sphenoid sinus (3). A case report of a petrous internal carotid aneurysm causing epistaxis by rupturing into the adjacent eustachian tube reminds one to also consider the middle ear as a source of such bleeding (4).

We know of only one other report of a brain AVM resulting in epistaxis (5). In this previous report, the malformation was located in the right parietal lobe and drained partially to the face through orbital and nasoethmoidal veins via the cavernous sinus. Epistaxis occurs in 85% of patients with Rendu-Osler-Weber syndrome (6). Twenty-eight percent of these patients have associated brain AVMs (7, 8). However, in Rendu-Osler-Weber syndrome the epistaxis is caused by nasal telangiectasias, not the brain AVM.

Venous communication between cortical veins of the frontal lobe and the nasal mucosa could be the cause of the nasal bleeding in our patient. Such a communication is possible through the foramen caecum. This ostium, located in the midline between the frontal bone and crista galli, can transmit a small emissary vein that connects...
the veins in the nose with the rostral end of the
superior sagittal sinus (9). The anterior aspect of
the superior sagittal sinus is an inconstant struc­
ture. The small emissary vein previously de­
scribed is more commonly present when the
rostral aspect of the superior sagittal sinus is well
formed. The existence of this connection has,
however, been questioned by others (10).

This case report illustrates the need to follow
a strict protocol in the angiographic work-up for
possible embolization of epistaxis. Our protocol
includes the study of both internal carotid ar­
ter­ies, both internal maxillary arteries, and both
facial arteries. Adherence to this protocol will
avoid overlooking unusual causes of epistaxis.

References

1. Marotta T, et al. Presented at the Annual Meeting of the Canadian
Association of Radiologists, Quebec City, October 1989.
2. Gelbert F, Reizine D, Stecken J, Ruffennacht D, Haffont J, Merland
JJ. Epistaxis grave par rupture de la carotide interne dans le sinus
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3. Wilson CB, Markesbery W. Traumatic carotidocavernous fistula with
fatal epistaxis: report of a case. J Neurosurg 1966;24:111–113
carotid aneurysm causing epistaxis: balloon embolization with pres­
ervation of the parent vessel. Neuroradiology 1987;29:570–572
5. Frayne TA, Boldt DW. Massive epistaxis caused by cerebral arterio­
of hereditary hemorrhagic telangiectasia (Rendu-Osler-Weber dis­
ease): report of two cases and review of the literature. Ann Neurol
1978;4:130–144
8. Willinsky R, Lasjaunias P, TerBrugge K, Burrows P. Multiple cerebral
arteriovenous malformations (AVM's): review of our experience from
203 patients with cerebral vascular lesions. Neuroradiology
1990;32:207–210
Orlando, FL: Grune & Stratton, 1984:3–23
10. Kaplan H, Browder A, Browder J. Nasal venous drainage and the
foramen cecum. Laryngoscope 1973;83:327–329