Embolization of An Intracranial arteriovenous Malformation Followed By immediate Surgical Resection: A Case Report

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Arteriovenous malformations (AVMs) consist of tangles of vessels of different wall thickness and diameter associated with arteriovenous shunting. In AVMs, arterioles and venules communicate directly without the interposition of a capillary bed. Ninety percent of intracranial AVMs are supratentorial. They often have associated aneurysms. Hemorrhage is the most dreaded presentation. Treatment and complete cure is challenging, especially for Spetzler grade IV and V. Glue embolization offers cure in some cases but most of the time it is adjuvant to surgery or gamma knife therapy. The following case report shows a comprehensive multidisciplinary approach to treat AVM in which embolization was immediately followed by surgical resection under the same anesthesia.

CASE REPORT

A 22-year-old man presented with headache, right sided weakness and slurred speech for one week. CT scan showed hemorrhage in the left parietal lobe (Figure 1). Cerebral angiography showed Spetzler Grade II AVM in the left parietal region. He was referred to our facility for elective embolization of AVM.

During embolization, the initial angiographic run revealed an approximately 1.2 cm-sized AVM nidus supplied by two
branches of the left middle cerebral artery (MCA) that was draining into a superficial cortical vein (Figure 2). With the help of a flow-guided microcatheter, the nidus was occluded via the major feeder with Histoacryl glue. Reflux of glue was noted into the draining vein (Figure 3).

A subsequent angiographic run revealed part of the nidus persistently filling up by a tiny proximal branch from the left MCA (Figure 4). It was not possible to manipulate the catheter through this tiny vessel despite multiple attempts. Due to reflux of glue into the draining vein, the risk of re-hemorrhage was significantly increased and immediate surgery was planned. The patient was shifted to the operating room where left fronto-parietal craniotomy was performed under general anesthesia and successful excision of the left parietal AVM was done (Figure 5), along with evacuation of left parietal hematoma.

Following surgery, the patient was hemodynamically stable. Although he continued to have right-sided weakness and slurred speech, he was following commands (pre-procedural state). No endovascular or surgical complication was observed.
DISCUSSION

Arteriovenous malformations (AVMs) are the most common intracranial vascular malformation. Incidence of AVMs is one-seventh to one-tenth that of aneurysms, and AVMs occur in about 0.02% to 0.05% of the population. Varying degrees of thrombosis and calcification may be seen. Atrophy of the surrounding parenchyma caused by steal can be seen.

The usual presenting symptoms are seizures, progressive neurologic deficit, intractable headache, intracranial hemorrhage and hydrocephalus. About 64% of cases present by 40 years of age; 30% to 50% present secondary to hemorrhage. The risk of death from bleeding in a patient with an AVM is about 29%; an additional 20% to 30% of patients suffer neurologic deficits. Each bleeding episode carries a mortality rate of 10% to 15%. Hemorrhage from an AVM is usually related to an associated aneurysm, outflow restriction, or pure deep venous drainage.

For treatment of AVMs, the nidus of the lesion must be removed or obliterated. Surgical resection is the mainstay of treatment. Only when the surgical approach or actual removal of the AVM is risky is endovascular therapy contemplated. Because the endovascular cure rate for AVMs is only 5% to 10%, endovascular therapy usually is used for preoperative assistance, or to shrink the mass as much as possible before radiosurgery, which is another therapeutic option.

Embolization is a good adjuvant to surgery. Although a very large range of materials have been used to try and embolize intracranial AVMs, it was the introduction of acrylate adhesive agents that have had the most impact. Acrylate-based glues are now the mainstay of intracranial AVM embolization.

Endovascular therapy for intracranial AVMs is now employed widely with a high degree of success and relatively infrequent complications. Recently, the more predictable embolic agent onyx has been introduced which allows targeted delivery to the nidus. Risks associated with the use of these agents include stroke or cranial nerve dysfunction due to unintended branch occlusions, hemorrhage, obstruction of venous outflow due to reflux in the draining vein, and cementing of catheter in the vessel.

In our case, reflux in the draining vein occurred which occluded outflow while inflow was still present by the other tiny arterial feeder. The patient’s blood pressures were kept low during the procedure and there was a high probability that as soon as blood pressures returned to normal, hemorrhage would recur. Therefore it was imperative that immediate surgery was planned for the resection of the AVM. Through this report, we also wish to highlight the multidisciplinary approach that must underpin any effective clinical approach to AVMs.

REFERENCES