De Novo Cerebral Arteriovenous Malformation: Case Report and Literature Review

J. Stevens, J.L. Leach, T. Abruzzo and B.V. Jones

*AJNR Am J Neuroradiol* 2009, 30 (1) 111-112
doi: https://doi.org/10.3174/ajnr.A1255
http://www.ajnr.org/content/30/1/111
De Novo Cerebral Arteriovenous Malformation: Case Report and Literature Review

**SUMMARY:** We describe a rare case of a de novo cerebral arteriovenous malformation (AVM) in a 9-year-old girl. MR imaging at 6 years of age demonstrated band heterotopia. Follow-up MR imaging 3 years later demonstrated a new 3.5-cm AVM in the left parieto-occipital region, confirmed by conventional angiography. This report, along with limited previous reports, suggests that AVMs can be acquired lesions and that AVM development is a dynamic process extending into the postnatal period.

**Case Report**
A 6-year-old girl was evaluated with MR imaging because of seizures and developmental delay. Imaging demonstrated continuous band heterotopia (Fig 1A). No vascular abnormality or abnormal signal intensity was demonstrated.

Three years later the patient presented with increased seizure activity, behavioral changes, increased ataxia, and aphasia after a minor head injury. CT demonstrated no evidence of a skull fracture or hemorrhage. MR imaging (Fig 1B) revealed a tangle of serpiginous flow voids along the margin of the left lateral ventricle at the junction of the left temporal and occipital lobes, consistent with an AVM. Cerebral conventional angiography (Fig 2) delineated a 3.5-cm AVM nidus with predominant arterial supply from distal branches of the left middle and posterior cerebral arteries. Venous drainage was through the ipsilateral thalamostriate veins and also through parenchymal veins extending into the left transverse sinus. The patient is currently undergoing radiation therapy for this lesion.

**Discussion**
We present a rare example of de novo AVM formation in a child with abnormalities of neuronal migration. A literature review identified 5 previously reported cases of de novo AVM development. Three of these cases are of patients with pre-existing vascular pathology who developed AVMs in or near pathologically altered brain. Only 2 previous reports document a cerebral AVM arising in brain parenchyma shown to be without vascular pathology on prior MR imaging. To our knowledge, no previously reported case has documented a de novo cerebral AVM developing in a patient with a neuronal migrational disorder.

The pathogenesis of AVMs is not completely understood and has largely been theorized. It has been suggested that cerebral AVMs are primarily congenital, originating at or before the 40- to 80-mm embryo length stage and may be related to a primary abnormality of primordial capillary or venous formation. However, excluding Galenic fistulas, we have found few reports of confirmed cerebral AVMs diagnosed in utero, and they are much more commonly diagnosed in adults than in children, challenging the assumption that all AVMs are congenital. Other concepts of cerebral AVM pathogenesis focus on the potential temporal vulnerability of susceptible vascular cellular elements to a physiologic trigger. This trigger could be mechanical, hormonal, thrombotic, hemodynamic, thermal, ischemic/hypoxemic, or inflammatory in nature.

Vascular endothelial growth factor (VEGF) may be 1 link between insult and AVM formation. One study demonstrated significantly higher levels of VEGF in patients with AVMs compared with a control group. Patients with hereditary hemorrhagic telangiectasia (HHT) have an increased prevalence of AVMs and also have elevated levels of VEGF. Hypoxia is known to stimulate expression of VEGF. Hemorrhage and thrombogenesis can result in the release of VEGF from platelets. Infection can lead to the release of VEGF at the source of infection. Thus, in theory, insults such as stroke, brain contusion, and encephalitis could all potentially promote the development of AVMs.

Cerebral AVMs are not static lesions: Growth and shrinkage of AVMs with time have been documented. Several cases of AVM recurrence in the same location following some form of intervention (ie, gamma knife, excision, embolization, etc) have been reported in pediatric patients and adults. Abnormalities of arterial venous, and microvascular structures have been described in relation to areas of abnormal neuronal migration. Abnormal vascular structures have been described in association with periventricular nodular heterotopia in filamin 1 gene mutation, suggesting 1 potential link between neuronal migrational disorders and vascular malformations.

**Conclusions**
This case report demonstrates postnatal development of an arteriovenous malformation in a patient with extensive band heterotopia. This case and the few previous reports of de novo AVM challenge the concept that cerebral AVMs are purely congenital lesions. AVMs appear to be dynamic lesions that...
have the potential to develop and grow with time. It is possible that acquired insults can provide the catalyst for the formation of AVMs later in life, and as suggested by this case, disorders of neuronal migration may share common genetic ground with abnormalities of endothelial proliferation.

References