Lenticulostriate Artery Aneurysm Associated with Cerebral Arteriovenous Malformation: Report of Rare Association and Its Endovascular Management

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Introduction

Aneurysms involving the lenticulostriate arteries (LSA) are rare and are described in the literature in small case series or single-case reports. A recent literature review identified fewer than 60 cases reported till date, and the management of these aneurysms varied from conservative approach to endovascular occlusion and surgical excision. The literature on endovascular treatment is rare; however, recent studies increasingly report on the usefulness of minimally invasive therapies in the management of these aneurysms.

Case Description

An 11-year-old female patient with no known comorbidities presented with symptoms of sudden onset severe headache and multiple episodes of vomiting. Computed tomography (CT) scan of the brain revealed an acute small hematoma in the roof of the right lateral ventricle and mild intraventricular bleed. CT angiography was obtained 4 days later, which showed a parenchymal arteriovenous malformation (AVM) involving the corona radiata and the posterior lentiform nucleus. She was initially treated conservatively and was referred to our institution for further management. She underwent digital subtraction angiography after 2 months of clinical presentation. Angiography confirmed the presence of the AVM and identified the feeders from lateral lenticulostriate arteries of right middle cerebral artery (MCA) as well as multiple medullary arteries from the M2 and M3 segments of the MCA. In addition, there was a 5-mm aneurysm at the distal segment of the lateral LSA (LLSA) feeding the AVM. Considering the risk of rupture of the aneurysm and its associated neurologic sequelae, a targeted approach for treatment of an associated aneurysm could be considered to reduce the risk of rebleed until the AVM is obliterated.

Abstract

Keywords
► lenticulostriate artery aneurysm
► cerebral arteriovenous malformation
► endovascular embolization

Aneurysms involving the lenticulostriate arteries are rare and are described in the literature in small case series or single-case reports. In this report, we discuss a case of a young woman who presented with acute lateral ventricular bleed and was found to have an arteriovenous malformation within the corona radiata and an aneurysm involving the distal lenticulostriate artery. The aneurysm was treated successfully by endovascular embolization. In deep-seated arteriovenous malformations (AVMs) that are conventionally treated by stereotactic radiosurgery, a targeted approach for treatment of an associated aneurysm could be considered to reduce the risk of rebleed until the AVM is obliterated.

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into distal LLSA for stability. The wire slack was removed and was gently torqued to either sides to identify the position that resulted in minimal wire prolapse. While maintaining a counter torque, the microcatheter was then advanced and the LLSA beyond the cisternal segment was catheterized. The microcatheter was positioned immediately proximal to the aneurysm, and the artery along the aneurysm and a part of the nidus was obliterated using 33% n-butyl cyanoacrylate (nBCA; Histoacryl, B Braun, Melsungen, Germany) (►Fig. 1). Follow-up CT scan of the brain revealed no infarction and she was discharged neurologically intact on the second post-operative day. Follow-up angiogram after 2 months of the embolization revealed no intranidal or feeding artery aneurysms. The residual AVM was treated by stereotactic radiosurgery. The patient remained clinically asymptomatic at 12 months follow-up, and presently, she is considered for repeat angiography to assess response to radiotherapy.

**Discussion**

Aneurysms of the LSA are often located within the basal ganglia and hence these aneurysms carry worse prognosis in the event of rupture. Indeed, most of the reported aneurysms presented with parenchymal hemorrhage and the extent of recovery varied.\(^2\) The aneurysms involving the distal LSA may be associated with moyamoya disease, dissection, or hypertension. However, the majority of these aneurysms are idiopathic. The natural history of these aneurysms is unknown and some aneurysms are known to spontaneously disappear on follow-up studies. This may be true in dissecting aneurysms where the dissected arterial wall heals over a period of time, allowing spontaneous thrombosis, and hence a wait-and-watch policy has been adopted by some authors as one of the therapeutic strategies.\(^2\) Aneurysms of LSA in association with cerebral AVMs are rare, with only four reported cases so far. Aneurysms in association with cerebral AVMs are found in 3 to 58% of cases, and the presence of these extranidal aneurysms are thought to pose additional risk of future hemorrhage.\(^3,4\) Exaggerated hemodynamic stress on the feeding artery and additional underlying intrinsic vascular defects may play a crucial role in the formation of these aneurysms.\(^5\) Though conservative approach has been advocated for LSA aneurysms per se as well as for the distal feeding aneurysms coexisting with AVM, we considered definitive therapy as the aneurysm had sizeable dimension and the aneurysm was located within the basal ganglia.\(^6\) Also, a recent study reported an association between the size of the aneurysm and the risk of AVM rupture.\(^4\) Surgical approaches include exclusion of the aneurysm or parent artery sacrifice; however, these approaches are

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*Fig. 1* Diagnostic angiography reveals an arteriovenous malformation within the basal ganglia (A, B) and an aneurysm at the distal aspect of the lenticulostriate artery (LSA) (arrow in B). Three-dimensional rotational angiography depicts the tortuous recurrent course of the LSA (C) that was catheterized. Angiography through the microcatheter confirms the presence of the aneurysm (D) that was treated with occlusion of the LSA with n-butyl cyanoacrylate (E). Final angiography reveals exclusion of the aneurysm (F) and residual arteriovenous malformation.
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Santhosh Kumar Kannath contributed in concept, design, data analysis, data interpretation, manuscript preparation, and critical revision.

Conflicts of Interest
The authors have no personal or financial conflict of interest to disclose.

References