Fatal recurrent ischemic stroke caused by vertebral artery stump syndrome

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INTRODUCTION

Vertebral artery stump syndrome (VASS) presents with recurrent posterior circulation (PC) stroke after ipsilateral vertebral artery (VA) occlusion at its origin.1 Its pathophysiology is thought to derive from a stagnant thrombus distal to the VA occlusion caused by antegrade flow from the collaterals but effective treatment remains unestablished. Here, we report a case of basilar artery occlusion (BAO) that occurred only 13 h after intravenous recombinant tissue plasminogen activator (IV rt-PA) therapy.

CASE DESCRIPTION

A 46-year-old man arrived at our emergency room with sudden, disturbed consciousness, left homonymous hemianopsia, left hemiplegia, and left hypoesthesia. The National Institutes of Health Stroke Scale (NIHSS) score was 10. Magnetic resonance (MR) diffusion-weighted imaging revealed slightly high-intensity signals in the right occipital lobe, right thalamus, and right hippocampus [Figure 1a and b] while MR-angiography showed the right posterior cerebral
artery occlusion at the P1 segment and unclear blood flow in the left VA [Figure 1c]. We immediately started IV rt-PA and performed cerebral angiography, revealing a right P1 occlusion [Figure 2a]. We pursued mechanical thrombectomy with a stent retriever against the right P1 thrombus through the right VA. Complete first-pass perfusion was achieved and the thrombolysis in cerebral infarction (TICI) score was Grade 3 [Figure 2b]. Angiography also revealed a left VA occlusion from its origin [Figure 3a], with the left deep cervical artery anastomosed to the left VA and forming an antegrade stagnant flow to the proximal end of the left VA [Figure 3b and c]. After thrombectomy, the patient dramatically improved (NIHSS score of 1) and complained only of mild left-sided paresthesia. We assumed that the VASS had an embolic source but, given the 24-h window since rt-PA use, avoided additional anticoagulation therapy at that point. Thirteen hours after the thrombectomy, the patient developed generalized seizures then a decerebrate posture. CT showed a high-density thrombus in the BA while angiography further revealed BA occlusion, necessitating a repeat mechanical thrombectomy with a stent retriever [Figure 4a]. This thrombus was retrieved but new thrombi appeared repeatedly during the procedure. After 7 passes, we achieved TICI grade 2A partial reperfusion but blood flow to the cerebellum was absent [Figure 4b] and the left VA was still occluded from its origin. The patient died 3 h after the procedure.

DISCUSSION

In 2008, Nguyen et al. reported two cases of recurrent PC stroke due to VA origin occlusion, calling it “VASS” because of the clinical similarities to carotid stump syndrome.[11] They hypothesized that the pathophysiology of this stroke was the embolism from the stagnant thrombus distal to the occlusion whereas the stagnant flow in the VA was caused by collaterals from the deep and ascending cervical arteries or the occipital artery. This anastomosis caused antegrade flow in the proximal side of the VA whose origin was occluded although the detailed fatal mechanism was unclear.[14,15] From this hypothesis, anticoagulation therapy may be effective for the prevention of recurrent ischemic events in VASS[9] as seen in a report of 12 such cases.[8] In that report, 1.2% of PC strokes were VASS-related, indicating that VASS is not a rare cause of PC stroke, but, since it is not recognized as causative for PC stroke, scarce reports exist. Given the lack of accumulated data, effective medical treatments for VASS remain controversial.

Intravenous rt-PA is known as an effective treatment for acute ischemic stroke[6] and mechanical thrombectomy is also effective for those cases where rt-PA is ineffective. In our case, mechanical thrombectomy was effective against VASS on the first attempt. Since VASS is believed to derive from embolism, thrombectomy may be efficacious but its efficacy and safety with regard to the PC has yet to be proven[5] since it cannot prevent recurrent ischemic strokes. After rt-PA use, additional antithrombotic treatment after rt-PA carries risks of hemorrhagic complications within the first 24 h,
Yamano, et al.: Fatal ischemic stroke caused by VA stump syndrome

Surgical Neurology International • 2021 • 12(445) | 2

as seen in current AHA/ASA recommendations, but some reports maintain that early antithrombotic therapy is safe. Since our patient developed recurrent stroke within 24 h of rt-PA and symptoms dramatically improved, we avoided additional antithrombotic therapy after thrombectomy. We still believe that immediate intravenous rt-PA and mechanical thrombectomy for the distal thrombus using a stent retriever were also a candidate strategy for the "artery-to-artery" thrombus originating from VA occlusion in the present case. However, retrospectively, the early introduction of additional antithrombotic therapy might have prevented the recurrent stroke.

Endovascular treatment is reported to permanently prevent recurrent stroke, as seen when Nguyen et al. reported two cases of VASS treated with coil embolization to exclude the embolic source at the VA. The coils were placed at the distal end of the VA thrombus, slightly proximal to the collateral flow to the VA while microcatheters were inserted from the contralateral VA or the collateral artery to the VA. After coil embolism, the stagnant flow at the VA disappeared and the two patients had no recurrent strokes during follow-up. On the other hand, Chen et al. reported 23 cases of VASS treated with endovascular revascularization in which a stent was placed in the proximal VA after balloon percutaneous transluminal angioplasty under the use of a distal protection device. The rate of successful technical revascularization was 91.3% and the rate of complication was 4.3%, results that held over a 3-month follow-up. Although these reports suggested the efficiency and safety of either embolization and revascularization treatment against non-acute VASS, small sample sizes reflect the need for further research and treatment against acute VASS was not reported.

There are several reports of endovascular treatment for BAO due to VASS. Revascularization treatment (either angioplasty or stenting) is done during acute BAO, and Ecker et al. showed 100% (6 out of 6 patients) recanalization of the occluded VA prior to the acute BAO treatment. In Table 1, we summarize the data of patients who underwent endovascular treatment for VASS. In a previous report, patients with minor stroke, especially recurrent strokes, were treated electively while patients with major strokes, such as BAO, were treated radically, including revascularization of the occluded VA. In our case, the patient demonstrated right P1 occlusion, and mechanical thrombectomy was effective for the lesion. At that time, dramatic symptom improvement made us hesitate on additional acute phase treatment but the thrombus occluding the P1 segment could have been large enough for urgent VA embolization or revascularization. These treatments have some technical limitations. Coil embolization to exclude the embolic source needs access to the distal part of the occluded VA through collateral arteries or the contralateral VA. The difficulty of the procedure depends on individual anatomical features and severe arteriosclerotic changes are expected in VASS patients. Revascularization of the occluded VA is also not simple since, unlike an acute artery occlusion, inability to traverse a lesion, dissection of the VA, and stroke due to distal embolization may complicate repairs. Preparation of appropriate guiding catheter support and distal protection devices is suggested.

Figure 3: Angiography of the left subclavian artery. The left VA was occluded from its origin (a). Anterior-posterior view (b) and lateral view (c) of angiography of the left subclavian artery. The left deep cervical artery was anastomosed to the left VA and its antegrade flow in the VA was stagnant. VA: Vertebral artery.

Figure 4: Angiography before the second thrombectomy. Basilar artery occlusion was revealed (a). Angiography after thrombectomy. Blood flow to the cerebellum other than the left posterior inferior cerebellar artery was absent (b).

Figure 4: Angiography before the second thrombectomy. Basilar artery occlusion was revealed (a). Angiography after thrombectomy. Blood flow to the cerebellum other than the left posterior inferior cerebellar artery was absent (b).
but, as there are no best practice guidelines for this issue,\[^{2,4}\] we hesitated to enact urgent radical treatment in the acute phase to avoid complications.

In our case, the stagnant flow at the VA continued to produce new thrombi despite IV rt-PA use. Because the volume from the VA origin to the anastomosis with the collaterals was relatively large, the stagnant flow had enough space to produce a large thrombus, which had the potential to occlude the BA. The possibility of recurrent stroke, including fatal BA occlusion, must be considered, especially in the acute phase, if antithrombotic therapy is withheld. Even though VASS is not a rare cause of PC stroke, only a few cases have been reported. The concept of VASS must be spread for the sake of accurate diagnosis and to accumulate case numbers sufficient for evaluating the efficacy and safety of early use of antithrombotic and/or radical endovascular treatment.

**CONCLUSION**

In patients with PC ischemic stroke featuring VA occlusion, VASS, as a dangerous syndrome involving the risk of fatal recurrent stroke, must be considered as its embolic source. In the present case, fatal BA occlusion developed within 24 h of rt-PA use but administration of rt-PA restricted the early introduction of antithrombotic therapy. Vigilance against such fatal, recurrent strokes must therefore be maintained, particularly when antithrombotic therapy is restricted.

**Declaration of patient consent**

The authors certify that they have obtained all appropriate patient consent.

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**Conflicts of interest**

There are no conflicts of interest.

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