Ruptured partially thrombosed anterior inferior cerebellar artery aneurysms: two case reports and review of literature

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ABSTRACT

Aneurysms arising from the distal anterior inferior cerebellar artery (AICA) are very rare. When the parent artery is an AICA–posterior inferior cerebellar artery (PICA) variant, occlusion of the artery, even distal to the meatal loop, leads to a significant area of cerebellar infarction. We report two cases of ruptured partially thrombosed distal AICA aneurysms. In both cases, the parent artery was an AICA–PICA variant. The aneurysms were clipped in one case and trapped following occipital artery (OA)–AICA anastomosis in another case. It is important to keep the OA as a donor artery for revascularization in the treatment of the AICA–PICA variant aneurysms, especially when the absence of intra-aneurysmal thrombus is not confirmed preoperatively.

Key Words: anterior inferior cerebellar artery, anastomosis, aneurysm, thrombosed, revascularization

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INTRODUCTION

Anterior inferior cerebellar artery (AICA) aneurysms account for less than 1.0% of all intracranial aneurysms,1,2) and distal AICA aneurysms are even more rare and constitute only about 0.1%.3,6) Most AICA aneurysms are found after rupture,3,7) and in such situations, it is not always easy to detect the presence of thrombus in the dome preoperatively from computed tomography (CT) and digital subtraction angiography (DSA) findings. In the presence of intra-aneurysmal thrombus, neck clipping of the aneurysm sometimes becomes difficult, necessitating an alternative treatment strategy such as trapping. We report two rare cases of ruptured partially thrombosed distal AICA–posterior inferior cerebellar artery (PICA) variant aneurysms, in one of which, preservation of the occipital artery (OA) as a donor artery for revascularization was helpful in avoiding ischemic complications.
Case 1
A 40-year-old woman presented with occipital headache of sudden onset. Her Glasgow Coma Scale (GCS) score was E3V5M6 on admission. Head CT showed the presence of subarachnoid hemorrhage distributed dominantly in the posterior fossa and clot packing the fourth ventricle (Fig. 1A). DSA was performed, and left vertebral angiogram showed a hypoplastic PICA and a 2-mm saccular aneurysm on the distal portion of the left AICA, which was also supplying the distribution of the PICA (Fig. 1B). Under endotracheal general anesthesia, left lateral suboccipital craniotomy was performed following a hockey stick-shaped skin incision. After dural incision, the facial, vestibulocochlear, and lower cranial nerves were identified, and the aneurysm was exposed, which was partially thrombosed and much larger (10 mm in diameter) than noted on preoperative DSA. The neck of the aneurysm was clipped successfully, and there developed no ischemic lesion in the distribution of the parent artery (Figs. 1C, D, E). Postoperatively, she had laryngeal edema and hydrocephalus. After treatment for these complications, she was discharged without any neurological deficits.

Fig. 1  Case 1
A. Computed tomography scan (CT) on admission showing subarachnoid hemorrhage distributing dominantly in the posterior fossa associated with clot packing the fourth ventricle.
B. Left vertebral angiograms: (1) anteroposterior view, (2) lateral view, and (3) right anterior oblique view demonstrating a 2-mm saccular aneurysm (arrow) on the distal portion of the left anterior inferior cerebellar artery–posterior inferior cerebellar artery variant.
C. Intraoperative photograph showing the neck of the aneurysm (arrow).
*: left lower cranial nerves.
D. Postoperative three-dimensional computed tomographic angiography demonstrating obliteration of the aneurysm (arrow) with a patent parent artery.
E. Postoperative CT showing absence of ischemic lesion in the distribution of the parent artery.
**Case 2**

A 62-year-old man presented with occipital headache and nausea. On admission, his GCS score was E3V5M6, and he had mild right hemiparesis, mild right facial paralysis, right abducens nerve palsy, and dysarthria. Head CT showed the presence of diffuse subarachnoid hemorrhage (Fig. 2A). DSA was performed, and left vertebral angiogram showed a hypoplastic PICA and a 4-mm saccular aneurysm on the distal portion of the right AICA, which was also supplying the distribution of the PICA (Fig. 2B). Under endotracheal general anesthesia, right lateral suboccipital craniotomy was performed following a hockey stick-shaped skin incision. The aneurysm was partially thrombosed and much larger (13 mm in diameter) than noted on the preoperative DSA. Because of thrombus, the neck of the aneurysm was hard and not amenable to clipping. Therefore, an OA−AICA anastomosis was performed, and the aneurysm was trapped (Fig. 2C). Postoperative magnetic resonance imaging demonstrated the thrombosed aneurysm clearly in the right cerebellomedullary cistern and absence of ischemic lesions in the cerebellum (Fig. 2D). Postoperative DSA showed good patency of the bypass (Fig. 2E). In the subacute period, his right hemiparesis showed some improvement and his right abducens nerve palsy remained unchanged, so he was transferred to a rehabilitation hospital with his modified Rankin Scale score of 3.

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**Fig. 2** Case 2

A. Computed tomography scan on admission showing diffuse subarachnoid hemorrhage.

B. (1) Right vertebral angiogram, anteroposterior view, showing that the right anterior inferior cerebellar artery (AICA) is an AICA–posterior inferior cerebellar artery variant; (2) left vertebral angiogram, anteroposterior view, demonstrating a 4-mm saccular aneurysm (arrow) on the distal portion of the right AICA.

C. Intraoperative schema showing anatomical location and appearance of the aneurysm; the aneurysm was partially thrombosed and its diameter was about 13 mm. Occipital artery (OA)–AICA anastomosis was performed, and the aneurysm was trapped.

D. Postoperative magnetic resonance images; (1) T2-weighted image showing the real size of the thrombosed aneurysm (arrow), (2)Diffusion-weighted image showing absence of ischemic lesion.

E. Postoperative right carotid angiogram showing patency of the OA–AICA anastomosis.
DISCUSSION

AICA aneurysms account for less than 1.0% of all intracranial aneurysms.\(^1,2\) Most AICA aneurysms are located at the proximal portion, and distal AICA aneurysms are very rare, constituting only about 0.1%.\(^3,4\) To the best of our knowledge, only a few partially thrombosed distal AICA aneurysms have been reported in the English literature (Table 1).\(^4,7-13\) Most AICA aneurysms are found after rupture.\(^3,7\) In cases of non-ruptured AICA aneurysms, patients present with headache, facial hypesthesia, trigeminal neuralgia, hearing disorder, or facial paralysis,\(^7,9,11\) and an intrameatal aneurysm may be diagnosed as a tumor involving the internal auditory canal preoperatively.\(^7,9\)

| No. | Author/year | Age/sex | Presentation | Aneurysm | Treatment | Sequela |
|-----|-------------|---------|--------------|----------|-----------|---------|
| 1   | Cantore, 1982 | 35/M    | vertigo, facial pain & hypesthesia, facial paralysis, hearing disorder, nystagmus | non-ruptured distal giant saccular | trapping & thrombectomy | incomplete facial paralysis |
| 2   | Zager, 1991 | 25/F    | headache, facial numbness | ruptured distal 20 mm fusiform | trapping & aneurysm excision | facial numbness |
| 3   | Pritz, 1993 | 33/M    | headache | non-ruptured proximal giant calcified mass | decompression | cricopharyngeal achalasia |
| 4   | Ildan, 1996 | 43/F    | trigeminal neuralgia, headache, drowsiness | ruptured distal 15 mm saccular | clipping & aneurysm excision | none |
| 5   | Sarkar, 2004 | 60/M    | headache, facial paralysis, hearing loss, nystagmus | non-ruptured distal 10 mm fusiform | trapping & thrombectomy | facial paralysis, hearing disorder, ataxia, nystagmus |
| 6   | Bambakidis, 2009 | 58/F | headache, dizziness | non-ruptured proximal 10 mm saccular | clipping | none |
| 7   | Oyama, 2010 | 65/F    | nausea, occipital headache, respiratory arrest | ruptured distal large fusiform | trapping with OA–AICA anastomosis & thrombectomy | dysphagia, hemiplegia |
| 8   | Pasler, 2011 | 22/M    | headache, hearing loss, facial paralysis | non-ruptured distal N.A. saccular | aneurysm excision with end to end anastomosis | hearing loss, headache |
| 9   | Present Case 1 | 40/F | headache | ruptured distal 10 mm saccular | clipping | none |
| 10  | Present Case 2 | 62/M | headache, hemiparesis, facial paralysis, dysarthria | ruptured distal 13 mm saccular | trapping with OA–AICA anastomosis | mild hemiparesis, diplopia |

AICA: anterior inferior cerebellar artery, N.A.: not available, OA: occipital artery

AICA aneurysms have been treated by microsurgery, endovascular therapy, or a combination of the two. Occlusion of the AICA may lead to ataxia, nystagmus, dysarthria, trigeminal sensory impairment, lateral gaze palsy, facial palsy, hearing loss, and motor weakness.\(^14-19\) On the other hand, some authors have reported that therapeutic occlusion of the AICA resulted in no deficits.\(^20-26\) Factors affecting these neurological outcomes include the site of parent artery occlusion, the volume of blood supply from the collateral circulation, and involvement of nerve-related branches such as the labyrinthine artery, the recurrent perforating arteries, and the subarcuate artery. For example, if the lesion is located distal to the meatal loop or a supplementary AICA is present,\(^25,27\) hearing preservation may be possible after occlusion of the affected site. It has been reported that the size of the infarcted area due to AICA occlusion is inversely related to the sizes of the PICA and superior cerebellar artery.\(^28\) In both of our cases, ruptured aneurysms were located in the distal portion of the AICA–PICA variant, suggesting preservation of the parent artery was necessary to avoid serious cerebellar infarction.
Ruptured thrombosed AICA aneurysms

As in Case 2, it is sometimes difficult to place a clip appropriately across the neck of a partially thrombosed aneurysm. In such a case, trapping of the aneurysm with an OA–AICA anastomosis may be a safer procedure than the more challenging neck clipping after thrombectomy. Trapping of AICA aneurysms or dissection in combination with OA–AICA anastomosis has been reported by several authors.8,29-31 Another option for revascularization of the AICA is aneurysm excision with end-to-end anastomosis of the AICA.7 Depending on the situation, the former may be easier than the latter because of a more shallow site of anastomosis. We also think that building a good anastomosis may have an advantage to maintain enough distal tissue circulation in case of vasospasm in the proximal portion of the parent artery. Neurological deficits at the time of discharge in Case 2 were mostly related to the first impact of hemorrhage.

In conclusion, when treating AICA–PICA variant aneurysms surgically, it is safer to preserve enough length of the OA in a hockey stick-shaped musculocutaneous flap as a donor for OA–AICA anastomosis in case of partially thrombosed aneurysms that necessitate trapping for obliteration.

CONFLICTS OF INTEREST DISCLOSURE

The authors have no personal, financial, or institutional interest in any of the drugs, materials, or devices presented in this article. All authors who are members of the Japan Neurosurgical Society (JNS) have registered online using the self-reported COI disclosure statement forms through the website for JNS members.

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