

The endovascular treatment of bilateral infarction of middle cerebellar peduncles

Etiology and endovascular treatment analysis

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ABSTRACT

يظهر احتشاء الدماغ السويقتين الأوسط عادة من الأمراض الوعائية الدماغية . وقد وصف ادامز في بداية عام 1943م متلازمة (AICA) الشريان المخيخي الأمامي التحتي، ولكن فشلت الدراسات السريرية للتصوير تتبع هذا لإلقاء المزيد من الضوء حتى ظهور التصوير بالرنين المغناطيسي وتحسن بذلك فهم AICA بشكل ملحوظ، وتعد أمراض احتشاء للدماغ السويقتين الوسطى (MCP) غير شائعة، وعلاج هذا المرض يعد نادر . نستعرض في هذا التقرير 4 مرضى باحتشاء للدماغ الثنائي في وقت واحد، ومنهم 2 مريضين تم علاج الأوعية الدموية داخل الجمجمة، وأظهرت الأعراض تحسنا . وتشير النتائج لنا إلى أن المرضى الذين يعانون من ضيق الشريان القاعدي مع احتمال احتشاء للدماغ الثنائي يمكن أن يستفيدون من العلاج داخل الأوعية .

Infarction of the symmetrical middle cerebellar peduncles is often induced by ischemic cerebrovascular disease. Adams described the anterior inferior cerebellar artery (AICA) syndrome as early as 1943, but clinical and imaging studies following this failed to shed more light regarding the condition until the advent of magnetic resonance imaging that comprehension regarding AICA improved significantly. Infarction of the middle cerebellar peduncles (MCP) is uncommon and the endovascular treatment of this condition is even more rare. We studied 4 patients with simultaneous bilateral cerebellar infarction of whom 2 received intracranial vascular therapy and demonstrated improvement in symptoms. Our findings suggest that patients with vertebral basilar artery stenosis with potential bilateral cerebellar infarction may benefit from endovascular treatment.

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The middle cerebellar peduncle (MCP) is supplied mainly by the anterior inferior cerebellar artery (AICA) and partly by the superior cerebellar artery. Bilateral infarction of the MCP is rare, and the therapy report of the stent is even more rare. There was only one case report of endovascular stent treatment for bilateral infarction of the MCP. We described 4 patients with bilateral infarction of the MCP, and 2 patients were treated with endovascular stenting and received impressive results.

Case Report. Case 1. A 50-year-old male patient with long-standing hypertension and episodes of dizziness for 2 years, presented with slurred speech, right side numbness and instability while standing. Neurologic examination revealed right side hearing loss, weakness of right side limbs, right side ataxia, and hyperalgesia. He had recurring transient left eye abduction dysfunction and diplopia following admission. He underwent left vertebral artery angioplasty on January 13, 2013 (Figure 1) and the symptoms improved significantly after stenting. There was significant relief in right side weakness and dysarthria while nystagmus and dyskinesia recovered significantly.

Case 2. A 61-year-old male patient presented with slurred speech, unsteady gait, dizziness and standing instability for 20 days. Following 3 days of

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work, he presented with limited abilities in left limb, dysarthria, dysphagia, aspiration, bilateral hearing loss and somnolence. Physical examination revealed that the pupil size of both eyes was not equal. Left pupil diameter was approximately 3 mm and the right was 4mm. Both eyes showed horizontal nystagmus, hearing significantly decreased, muscle strength of all four limbs was reduced and Babinski's sign was elicited on the left side. On presentation at the hospital, he was started on dual antiplatelet medication (acetylsalicylic acid and clopidogrel). The patient's physical activity significantly improved when following stent implantation after admission and hearing was restored. However, serious limbs and trunk ataxia persisted when he was discharged (Figure 2).

Case 3. A 68-year-old male patient presented with recurrent attacks of nausea, vomiting, vertigo and severe ataxia for 3 years, sudden slurring of speech and unsteady gait. He was admitted to a local hospital. He had transient dysarthria, somnolence, hearing loss, unsteady gait after vertigo, vomiting and nausea. Physical examination

revealed bilateral horizontal nystagmus and ataxia of limbs and trunk. He was unable to complete both heel-knee-tibia test and finger-nose test. Diffusion weighted image (DWI) examination is shown in Figure 3. After 14 days of hospitalization conservative treatment there was a slight improvement in dysarthria, but there were significant ataxia and bilateral deafness.

Case 4. A 51-year-old male patient presented with dizziness that had persisted for more than 5 hours, accompanied by nausea, vomiting, slurred speech, diplopia dysphagia, tinnitus, weakness and unsteady gait. He was admitted. Physical examination revealed that alternating hand movement test and heel-knee-tibia test were mildly abnormal. Following conservative therapy, the patient still had slurred speech and gait instability. Diffusion Weighted Image examination is shown in Figure 4.

Discussion. Image analysis of 4 cases demonstrated that the infarction lesion was located in the middle cerebellar peduncles and other AICA supply area with or

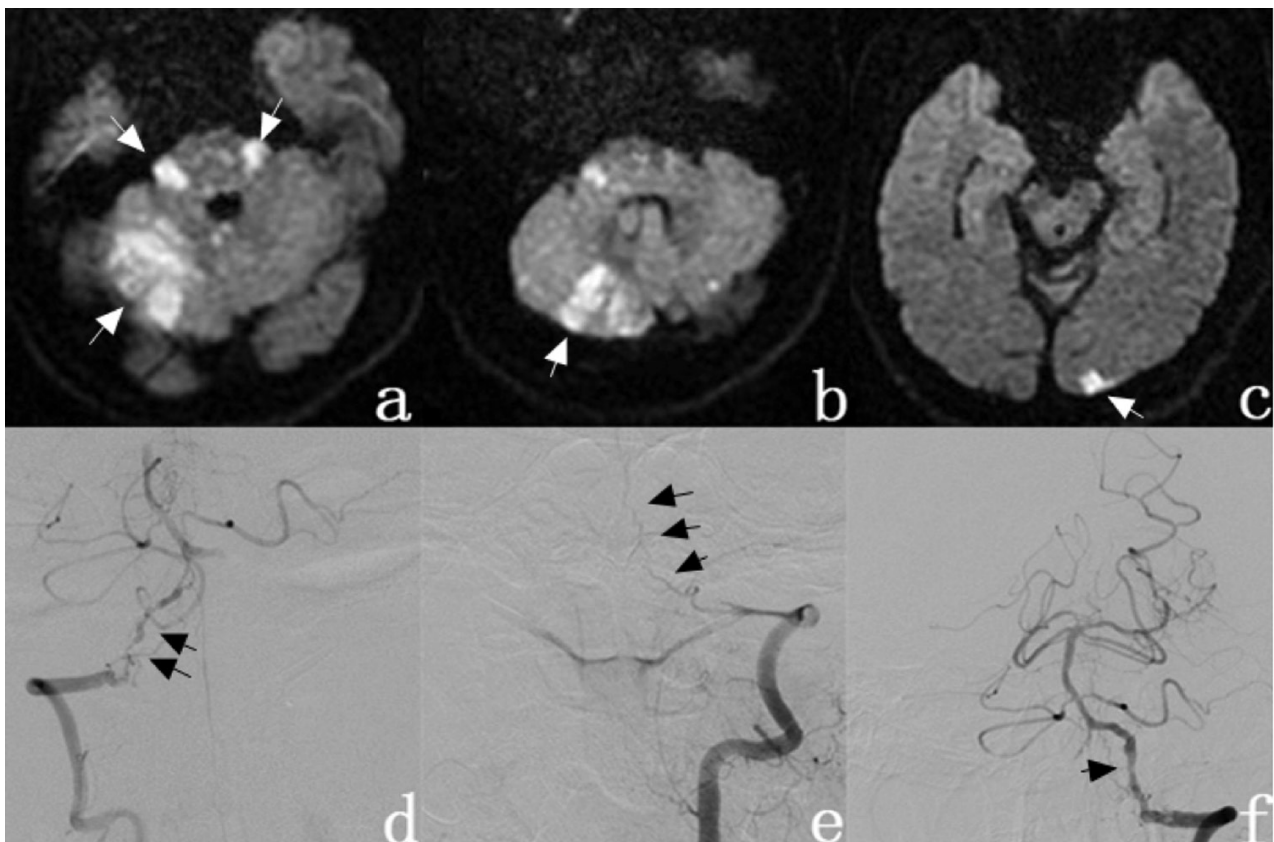


Figure 1 - Diffusion Weighted Image showing A & B) spotty high signal in bilateral cerebellar hemispheres and vermis of the cerebellum, pons, right thalamus (arrow); C) with left occipital high-intensity (arrow). D) Digital subtraction angiography showed right vertebral artery and basilar artery stenosis (arrow) and E) left vertebral artery distal was totally occluded. F) Left vertebral artery, basilar artery, bilateral inferior cerebellar artery, and bilateral superior cerebellar artery were visible after left vertebral artery recanalization.

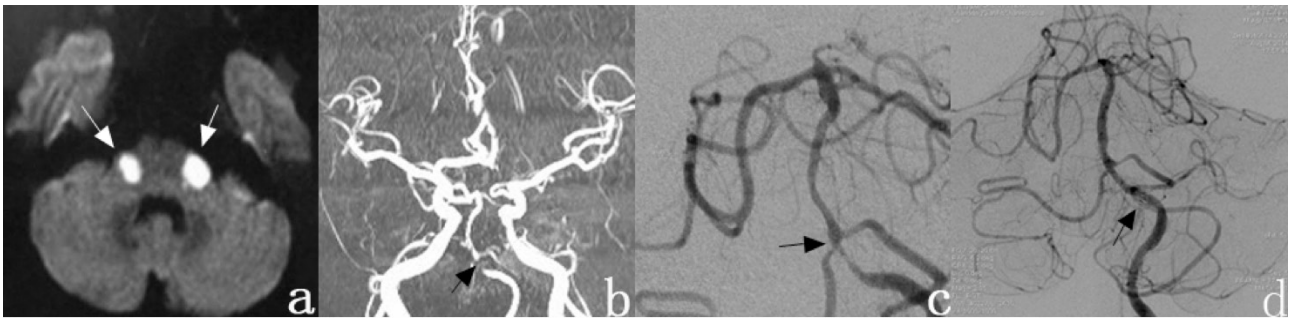


Figure 2 - Diffusion Weighted Image shows A) high-intensity bilateral symmetric round infarctions (arrow). B) MRA shows that the right vertebral artery is invisible, severe stenosis of the segment from the left vertebral artery distal to the basilar artery or occlusion (lower arrow). C) Cerebral angiography shows diffuse severe stenosis from the V4 segment of the left vertebral artery to the proximal end of the basilar artery and moderate stenosis at the origin of the right AICA (arrow). D) The stenosis was relieved following stent implantation (arrow) in the inferior segment of the basilar artery and the stenosis of the right vertebral artery. artery stenosis (arrow).

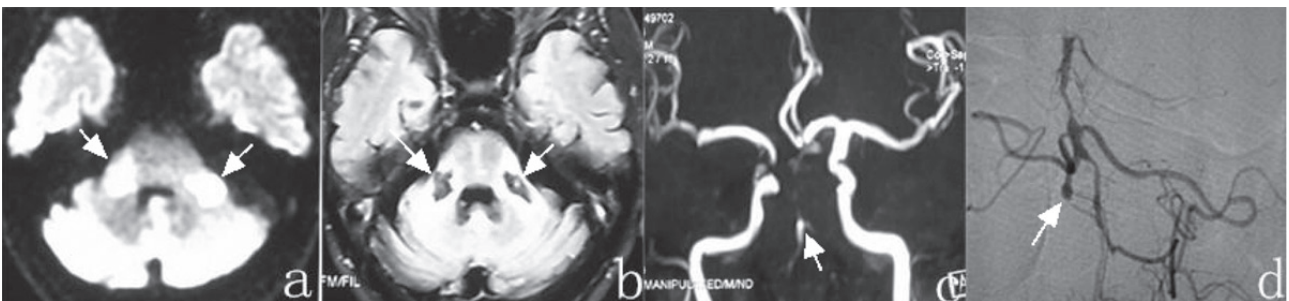


Figure 3 - Diffusion Weighted Image shows A) high-intensity areas in the bilateral middle cerebellar peduncles (arrow). B) Bilateral middle cerebellar peduncle shows symmetrical hypodensity areas on T1-weighted images (arrow). C) MRA shows that bilateral vertebral artery and the basilar artery at the end of the vertebral artery can be visualized (arrow). Bilateral anterior inferior cerebellar artery, superior cerebellar artery and posterior cerebellar artery are not invisible. D) Cerebral angiography showed occlusion of the V5 segment (arrow) of the right vertebral artery, moderate to severe stenosis of the left vertebral artery, basal segment of the basilar artery shows swelling, bilateral superior cerebellar artery visible but obviously thin, the right posterior cerebellar artery is invisible and moderate to severe stenosis of the left posterior cerebellar artery.

without PICA or involvement of SCA supply area. We observed an infarct in the SCA area in case 1. Separate AICA infarction can be considered in cases 2, 3 and 4. The treatment consisted mainly of anticoagulation, antiplatelet therapy. Application of stent at the site of stenosis can significantly improve the symptoms in 2 patients.

Infarction of the MCP is a rare type of AICA syndrome. The incidence of anterior inferior cerebellar artery infarction in acute stroke is only 0.9% and represents 5.2% of patients affected the vertebrobasilar territory. Patients with infarction localized in the bilateral middle cerebellar peduncles were rare.¹ Adams¹ noted rotational dizziness, vomiting, tinnitus and dysarthria on the side of the infarction as well as dysmetria, facial palsy, trigeminal sensory impairment, Horner syndrome and loss of pain and temperature on the opposite side.² The clinical picture described by Adams¹ was called the classic AICA syndrome. The presence of vertigo, ataxia, peripheral facial palsy and ipsilateral hypoacusia means a “complete AICA infarct”.¹

The anterior inferior cerebellar artery is an almost constant artery. The artery usually arises from the basilar artery, from the lower third in 75% of the cases. Sometimes from the middle third, and occasionally from its lower limit.³ The AICA has 2 separate areas of supply, the proximal branch, and the lateral branch. The proximal branch supplies the lateral area of the pons. The lateral branch, after crossing the 8th nerve, divides into 2 branches. One sends a constant trunk along the medial surface of the cerebellar hemisphere to anastomose with a cerebellar branch of the posterior inferior cerebellar artery. The MCP and an area of variable size in the lower two-thirds of the pons are supplied by the small arteries from the other branch.^{3,4} The MCP is the core of the affected territory and may be considered as the radiological hallmark of this syndrome.¹ The distribution of classical AICA is primarily lateral pontine extending to include a small portion of the anterior, medial, and inferior region of the cerebellar hemisphere.⁵ It frequently involves the

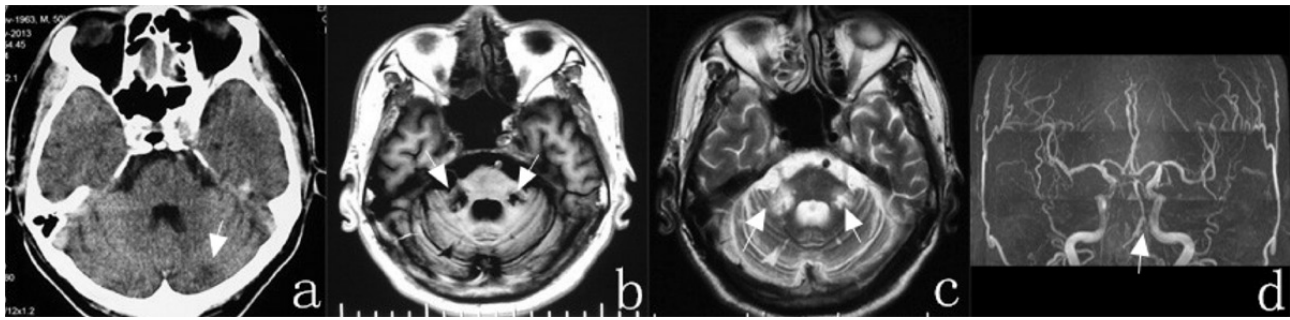


Figure 4 - CT scan showed A) patchy low-density foci in the left cerebellar hemisphere (arrow). B & C) MR shows low signal intensity from bilateral middle cerebellar (arrow). D) MRA shows bilateral vertebral artery stenosis (arrow).

nuclei and roots of the fifth, seventh and eighth nerves which are characteristics of the clinical syndrome.¹

Deafness and tinnitus caused by the eighth nerve damage are common symptoms of AICA syndrome since AICA is the source of the internal auditory artery (IAA), which exclusively supply the inner ear. Since deafness and tinnitus are characteristic symptoms of AICA, these symptoms can be used to differentiate between AICA and PICA infarct. Internal auditory artery occlusion is often accompanied by labyrinthine dysfunction. Labyrinthine dysfunction might have been masked by severe ataxia, but it can explain the horizontal nystagmus in cases 2 and 3. Some of the branches of the AICA may constitute the upper and posterior branches of the lateral medulla, supplying the fibers of the ninth and tenth nerves and the superior part of the nucleus ambiguus.³ The involvement of these sites can lead to dysarthria and dysphagia. Our 3 patients presented with obvious dysarthria suggest the possibility of upper medulla involvement. The AICA also supplies the lateral part of the abducens nucleus. Case 3 had recurrent left eye abduction restriction, which may be due to transient ischemia of the left side of the abducens nucleus or abducent nerve. Motor weakness is the common concomitant symptom of AICA, associated with corticospinal tract involvement. From the clinical features of our 4 patients, they have several obvious common features, such as vertigo, ataxia, slurring speech, unsteady gait, and hearing loss (Table 1). Middle cerebellar peduncle is largely supplied by the AICA, but it also receives blood supply from the SCA which anastomoses with the terminal branches of the AICA. As such, the MCP is a watershed area for these 2 arteries.^{3,6}

Large artery occlusive disease was more frequently observed in cerebellar watershed infarction.⁶ The common etiology of the AICA infarct is thrombosis superimposed on atheromatous stenosis which is

located in the AICA or the basilar artery lower part.² Symmetrical cerebellar infarction is mostly caused by lesions on one side of the vertebral artery and basilar artery, or bilateral vertebral artery lesions, but the occlusion rarely occurs in AICA. Less frequently the occlusion occurs on the distal part of both vertebral arteries or in the distal part of the ipsilateral vertebral artery.

We speculate that occlusion of both vertebral arteries may lead to hypoperfusion in the watershed area between the AICA and the superior cerebellar arteries, resulting in infarction in the bilateral middle cerebellar peduncles. John described bilateral cerebellar infarction cases, although bilateral AICAs were invisible, but also speculated that the possible mechanism of bilateral cerebellar infarction to be due to arterial stenosis or occlusion, based on blood flow reduction leading to AICA involvement.⁷

In our study, cerebral angiography showed severe stenosis or occlusion of the vertebral basilar artery system, except in case 2 where there was mild stenosis at the initiation site of the right AICA, while the other cases showed that AICA was not. However, all of our 4 cases demonstrated severe stenosis or occlusion of the vertebral basilar artery. The pathogenic mechanism of MCP infarcts may be the result of hypoperfusion based on arterial stenosis leading to decreased blood flow to the AICA, which could not be completely compensated by the collateral circulation. From the point of risk factors, the 4 patients have long history of hypertension and hyperlipidemia (Table 1). We consider the hyperlipidemia and hypertension lead to atherosclerosis when suddenly decreased blood pressure leads to the hypoperfusion for cerebral infarction. The history of atrial fibrillation cannot be excluded as the cause of cardioembolic infarcts resulting in MCP of case 2. Unfortunately, the patient did not undergo an ultrasound examination. In case 1, the left cerebellum

Table 1 - Clinical manifestation from the clinical features of our 4 patients, they have several obvious common features, such as vertigo, ataxia, slurring speech, unsteady gait, and hearing loss.

Cases	Age (years)/gender	Vascular risk factors	Symptoms/signs	Diagnose	Treatment	Outcome
1	50/M	Hypertension, hypercholesterolemia, smoking and drinking history	Vertigo, slurred speech, right side numbness and weakness, unsteady gait, right hearing loss, right ataxia, right hemihypoesthesia. left eye abduction dysfunction, and diplopia	Infarction of MCP	Angioplasty-stent placed, oral dual antiplatelet (acetylsalicylic acid and clopidogrel)	Relief in right side weakness and dysarthria, nystagmus and dyskinesia recovered significantly
2	61/M	Hypertension, hypercholesterolemia, smoking and drinking history, atrial fibrillation	Vertigo, slurred speech, unsteady, standing instability, dysarthria, dysphagia, aspiration, bilateral hearing loss, and somnolence	Infarction of MCP	Angioplasty-stent placed, oral dual antiplatelet (acetylsalicylic acid and clopidogrel)	Physical activity significantly improved, hearing was restored. Ataxia persisted
3	68/M	Hypertension, hypercholesterolemia, diabetes	Vertigo and severe ataxia, slurring of speech and unsteady gait, dysarthria, somnolence, hearing loss, unsteady gait, vomiting and nausea. bilateral horizontal nystagmus	Infarction of MCP	Oral dual antiplatelet (acetylsalicylic acid and clopidogrel)	Slight improvement in dysarthria, but there were significant ataxia and bilateral deafness
4	51/M	Hypertension, hypercholesterolemia, diabetes	Vertigo, severe ataxia, slurring speech, unsteady gait, dysarthria, somnolence, hearing loss, bilateral horizontal nystagmus, ataxia of limbs and trunk	Infarction of MCP	Oral dual antiplatelet (acetylsalicylic acid and clopidogrel)	Still had ataxia, slurred speech, and gait instability

MCP - Middle Cerebellar Peduncles, M - Male

showed high signal intensity and the right superior cerebellum showed patchily or spotted high signal intensity, suggesting that the superior cerebellar artery or anterior inferior cerebellar artery perforating branch occlusion may be associated with hypoperfusion. Case 3 showed basilar artery dilatations distal to the stenosis suggesting arterial dissection. According to the case 4 cerebral angiography, clinical symptoms, the time of onset, and an obvious responsibility lesion in imaging, we excluded the multiple system atrophy.

Anterior inferior cerebellar artery syndrome occurs mostly due to vertebral basilar artery stenosis. The treatment mainly comprises of anticoagulation and antiplatelet therapy. Stent implantation in the stenosed artery has become an important therapeutic avenue. Huo et al⁸ and Gao et al⁹ had shown that the combined use of mechanical thrombectomy with angioplasty and stent placement in the basilar artery, with severe underlying intracranial atherosclerotic stenosis, is technically feasible and safe. A previous single case report suggests a possible stent therapy for bilateral cerebellar infarction.¹⁰ In this study, we found that cases 1 and 2 had significant improvement following stent placement. We found that the overall prognosis of stenting was better than that without

stent implantation. Patients with AICA infarction who have mild symptoms have a good prognosis and fewer sequelae. However, patients with severe stenosis of the vertebral basilar artery system, if not treated in a timely fashion, have an increased risk of large artery infarction that hinders posterior circulation and threatens their lives.

We collected 16 relevant case reports, most of which were treated conservatively in 1991 to 2016, with only one case referred to the stent (Table 2). It can be seen that the endovascular treatment in the bilateral infarction of middle cerebellar peduncles is very rare. We analyzed that the previous treatment for bilateral infarction of middle cerebellar peduncles was a more conservative treatment rather than endovascular treatment for the reason that the development of intravascular treatment technology was relatively backward. But with the development endovascular treatment techniques in recent years, endovascular techniques in the treatment of bilateral infarction of middle cerebellar peduncles will become a new and effective method.

In conclusion, cerebellar infarction is the core area involved in AICA infarction and the main clinical symptoms are vertigo and ataxia. The cause of AICA infarction in almost all cases is atherosclerosis or

Table 2 - Previous case reports of bilateral cerebellar infarction.

Year	Number of cases	Therapy	Author + Reference
1991	1	—	Tsukamoto T et al ¹¹
1994	1	—	Ichikawa H et al ¹²
1998	1	—	Sato K et al ¹³
1998	1	—	Roquer J et al ¹⁴
2000	1	—	Akiyama K et al ¹⁵
2001	1	Anticoagulation	Lee H, et al ¹⁶
2005	1	Oral antiplatelet	Sunami E, et al ¹⁷
2007	1	Anticoagulation	Iwanami H, et al ¹⁸
2009	3	—	Lee H, et al ²
2010	1	—	Renard D et al ¹⁹
2011	1	Intravenous ozagrel and oral ticlopidine	Kataoka H et al ⁶
2013	1	Dual antiplatelets (acetylsalicylic acid and clopidogrel)	John S et al ⁷
2013	1	Stent	Kattah JC et al ¹⁰
2016	1	Anti-platelet and symptomatic treatment	Özkan A et al ²⁰

“—” means the treatment was not explicitly mentioned in the literature

thrombosis superimposed on the basilar artery stenosis. Patients with severe stenosis of the vertebral basilar artery system, if not treated in a timely fashion, have an increased risk of large artery infarction that hinders posterior circulation and threatens their lives. In addition to the traditional drug therapy, interventional therapy can play a significant role.

At present, the evidence of the conventional basilar artery angioplasty/ stenting of infarction of the middle cerebellar peduncles is lacking. However, there is a growing of literature that shows that angioplasty stenting as a generally acceptable choice in patients with the rapidly deteriorating neurological status from the advanced vertebrobasilar stenosis.

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References

- Adams RD. Occlusion of the anterior inferior cerebellar artery. *Arch Neurol Psychiatry* 1943; 49: 765-770.
- Lee H, Kim JS, Chung EJ, Yi HA, Chung IS, Lee SR, et al. Infarction in the territory of anterior inferior cerebellar artery: spectrum of audiovestibular loss. *Stroke* 2009; 40: 3745-3751.
- Lee H. Audiovestibular loss in anterior inferior cerebellar artery territory infarction: a window to early detection? *J Neurol Sci* 2012; 313: 153-159.
- Duvernoy HM, editor. The human brain: surface, three-dimensional sectional anatomy with MRI, and blood supply. Berlin (Germany): Springer Science & Business Media; 2012.
- Neugebauer H, Witsch J, Zweckberger K, Jüttler E. Space-occupying cerebellar infarction: complications, treatment, and outcome. *Neurosurgical focus* 2013; 34: E8.
- Kataoka H, Izumi T, Kinoshita S, Kawahara M, Sugie K, Ueno S. Infarction limited to both middle cerebellar peduncles. *J Neuroimaging* 2011; 21: e171-e172.
- John S, Hegazy M, Cheng Ching E, Katzan I. Isolated bilateral middle cerebellar peduncle infarcts. *J Stroke Cerebrovasc Dis* 2013; 22: e645-e646.
- Huo X, Gao F, Sun X, Ma N, Song L, Mo D, et al. Endovascular mechanical thrombectomy with the solitaire device for the treatment of acute basilar artery occlusion. *World Neurosurg* 2016; 89: 301-308.
- Gao F, Lo W, Sun X, Mo D, Ma N, Miao Z. Combined use of mechanical thrombectomy with angioplasty and stenting for acute basilar occlusions with underlying severe intracranial vertebrobasilar stenosis: preliminary experience from a single Chinese center. *AJNR Am J Neuroradiol*. 2015; 36: 1947-1952.
- Kattah JC, Nair D, Talkad A, Wang DZ, Fraser K. A case of vestibular and oculomotor pathology from bilateral AICA watershed infarcts treated with basilar artery stenting. *Clin Neurol Neurosurg* 2013; 115: 1098-1101.
- Tsukamoto T, Seki H, Saitoh J, Watanabe K. A case of bilateral cerebellar peduncle infarction. *Jpn J Med* 1991; 30: 376-378.
- Ichikawa H, Fukui T, Negishi A, Sugita K, Kanda M. [A case of bilateral cerebellar peduncle infarction with bilateral hearing impairment of a sudden onset]. *Rinsho Shinkeigaku* 1994; 34: 569-576. Japanese
- Sato K. [Bilateral middle cerebellar peduncle infarction]. *No To Shinkei* 1998; 50: 1036-1037.
- Roquer J, Lorenzo JL, Pou A. The anterior inferior cerebellar artery infarcts: a clinical-magnetic resonance imaging study. *Acta Neurol Scand* 1998; 97: 225-230.
- Akiyama K, Takizawa S, Tokuoaka K, Ohnuki Y, Kobayashi N, Shinohara Y. Bilateral middle cerebellar peduncle infarction caused by traumatic vertebral artery dissection. *Neurology* 2001; 56: 693-694.
- Lee H, Whitman GT, Lim JG, Lee SD, Park YC. Bilateral sudden deafness as a prodrome of anterior inferior cerebellar artery infarction. *Arch Neurol* 2001; 58: 1287-1289.
- Sunami E, Nagayama H, Yamazaki M, Katsumata T, Katayama Y. [A case of infarction in brainstem and cerebellum as a initial symptom with bilateral hearing loss]. *No To Shinkei* 2006; 58: 791-795. Japanese
- Iwanami H, Odaka M, Hirata K. [Bilateral cerebellar infarction caused by intracranial dissection of the vertebral artery after long periods of “Shiatsu”]. *Brain Nerve* 2007; 59: 169-171.
- Renard D, Taieb G, Castelnovo G, Ghanem S, Ayrygnac X, Labauge P. Isolated bilateral anterior inferior cerebellar artery infarction. *Arch Neurol* 2010; 67: 766-767.
- Özkan A, Balci Ç, Şen HM, Adam G, Özişik Karaman HI. Bilateral anterior inferior cerebellar infarction presenting severe vertigo and hearing loss. *Turkish Journal of Cerebrovascular Diseases* 2016; 22: 41-44.