Case Report

Sphenoid wing dural arteriovenous fistula: A case report and literature review

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ABSTRACT

Background: Sphenoid wing dural arteriovenous fistula (SWDAVF) is rare that is typically fed by middle meningeal artery feeders and that drain through the sphenoparietal sinus or middle cerebral vein. Here, we report a case of SWDAVF treated by coils placed in the venous aneurysm through the contralateral cavernous sinus (CS).

Case Description: A 37-year-old woman was admitted to our hospital with headache and bilateral oculomotor nerve palsy. Magnetic resonance images and an angiogram showed a venous aneurysm in the right middle cranial fossa. A DAVF, consisting of two main feeders, was diagnosed based on the angiogram findings. The fistula drained into the left inferior petrosal sinus (IPS) through the left CS and right IPS. Given the remarkable extent of venous ectasia together with the headache and right abducens nerve paralysis, endovascular treatment was initiated. A transvenous approach through the right IPS was not feasible, as it is strenuous to insert the microcatheter into the right IPS. Thus, we tried an approach through the left IPS. The venous aneurysm was embolized with coils. The postoperative course was uneventful, and postoperative cerebral angiography confirmed disappearance of the fistula.

Conclusion: A SWDAVF is extremely rare. In our case, since the AVF drained into the contralateral CS, contralateral ocular symptoms occurred. Endovascular occlusion of the venous aneurysm and fistula was achieved through a transvenous approach.

Keywords: Cavernous sinus, Dural arteriovenous fistula, Endovascular, Venous aneurysm

INTRODUCTION

Intracranial dural arteriovenous fistula (DAVF) is sporadic, pathologic arteriovenous connections that most commonly involve the wall of a major dural venous sinus. The sphenoid bone forms the foundation of the anterior and middle cranial fossae. It is a centrally placed bone, through which critical neurovascular connections are transmitted through vital foramina. DAVF of the lesser sphenoid wing is often considered to be abnormal connections between the middle meningeal artery (MMA) and the sphenoparietal sinus.[5,18] Most sphenoid wing DAVF (SWDAVF) are fed by the MMA and drain into the sphenoparietal sinus and the superficial middle cerebral vein (SMCV).[11] In our case, the DAVF drained into the cavernous sinus (CS), mimicking CS DAVF. Endovascular occlusion of fistulous connections by transarterial or transvenous approaches is the primary therapeutic strategy to cure these potentially dangerous lesions. Here, we describe a case of DAVF in the sphenoid wing accompanied with ocular symptoms; we successfully treated this patient with transvenous embolization.
CASE PRESENTATION

A 37-year-old woman presented with headache, bilateral ptosis, and right proptosis. She denied a history of head trauma and hypertension, and her medical history was unremarkable. Her laboratory data were all normal. Physical examination revealed bilateral oculomotor nerve palsy. Magnetic resonance (MR) imaging demonstrated varix in the right middle cranial fossa [Figure 1a and b]. Cerebral angiography revealed a high-flow, multichannel fistula between the right MMA and a venous aneurysm wall at the right greater sphenoid wing, which drained into the left CS superior orbital vein (SOV) and inferior petrosal sinus (IPS). The petrosal outlets of the right CS were occluded on the way. On the venous side, retrograde venous drainage was observed from the right SWL to the right facial vein through a right SOV with venous ectasia. Second, the venous drainage from the DMCV drained into the right CS and then the left CS, through the inter-CS (ICS), and into the left IPS [Figure 2a-c]. No connection between the SWL and the basal vein was observed. Given the remarkable ectasia of the draining vein, accompanied by the presence of headache and right abducens nerve paralysis, endovascular treatment was initiated.

Our approach was through the left IPS since the right IPS was occluded. A 6 Fr FUBUKI (ASAHI INTECC, Nagoya, Japan) guiding catheter was placed in the left internal jugular vein. A 4.2 Fr FUBUKI distal access catheter (ASAHI INTECC, Nagoya, Japan) was introduced through the left IPS and then placed in the right CS at the orifice of the sphenoparietal sinus. Excelsior SL-10STR (Stryker, Kalamazoo, MI, USA) and ASAHI CHIKAI (ASAHI INTECC, Nagoya, Japan) were advanced into the venous aneurysm of the SWL. Following the introduction of the microcatheter into the SWL, superselective angiography suggested a high-flow, multichannel fistula between the right petrosal branch of the MMA and a venous aneurysm wall [Figure 2c]. We assigned this area as the fistulous point, and the venous aneurysm was occluded using a Target 360 soft coil (Stryker Neurovascular) followed by the application of additional down sized coils. Complete obliteration of the venous aneurysm was achieved after the insertion of 10 coils [Figure 3a]. A final right external carotid angiography showed disappearance of the fistula [Figure 3b and c]. The postoperative course was uneventful. After the procedure, the symptoms of headache and diplopia improved.

Postoperative cerebral angiography and time-of-flight MR image on the 3rd month showed disappearance of the venous aneurysm and fistula [Figure 4a and b].

DISCUSSION

DAVF within the greater and lesser sphenoid wings has vascular features distinctive from the CS. The sphenoidal group of cerebral veins formed by the terminal ends of the superficial Sylvian, and occasionally, the deep Sylvian veins drain into the sphenoparietal or CS. It less commonly drains into the sphenobasal or sphenopetrosal sinuses that course on the inner surface of the sphenoid bone. The incidence rate of SWDAVF was reported to be 1.54% of all
Intracranial DAVF, and it is suggested that a DAVF diagnosis might be difficult because it could easily be confused with a CS DAVF. In our case, venous drainage from the right CS to the left CS was shown in the early venous phase. Arteriovenous fistulae cause an increase pressure of the left CS and also right CS; hence, contralateral ocular symptoms occurred. Clinical manifestations include ptosis, global amnesia, headache, visual field defects, and dizziness. Although few patients exhibit aggressive presentation, such as intracranial hemorrhage or progressive neurological deficits, treatment is strongly recommended due to the presence of varix similar to our case. The treatment options for SWDAVF have been described in previous reports and include TAE, TVE, and surgical obliteration. However, TVE is the primary therapeutic strategy for the curative treatment of DAVF. If it is impossible to approach the affected sinus through the IPS, it is necessary to consider another venous access route, such as the facial vein or the vein of Galen. In the current era of treating DAVF with curative embolization using Onyx, TAE of fistulous connections is the primary treatment technique. Flow control using coils, acrylics, or temporary balloon occlusion is useful for high-flow shunts to decrease the likelihood of distal embolic migration and to improve the penetration of embolic material into the arteriovenous connections.

We summarized the clinical data of SWDAVF [Table 1]. SWDAVF is rare, and only 15 cases have been reported in the literature, including our case. The median patient age was 52.9 years (range, 27–71 years), and the patients were predominantly male (80%). Nine out of 15 patients had varix. The four cases who exhibited cerebral symptoms had venous drainage related to reflux into the SMCV or through the petrosal veins into the perimesencephalic and cerebellar veins. In these patients, however, cerebral symptoms and SAH were always concomitantly present with CS symptoms, such as proptosis and retinal hemorrhage. In addition to the SMCV, the CS and SOV were seen during angiography. Five cases presented with orbito-ocular or CS symptoms and had antegrade (through the IPS) and/or retrograde (through SOV) venous drainage routes.

Shi et al. reported two cases treated by successful endovascular treatment. Embolization through a transarterial or transvenous approach is the primary therapeutic strategy for these lesions. Our patient was fortunately cured by only a transvenous approach.

Endovascular treatment has been favored because the draining vein just proximal to the fistula point at the lesser sphenoid wing is easy to access through the IPS and CS. Six patients underwent endovascular treatment only, four patients underwent surgery only, and five underwent both [Table 1]. Although arterial embolization for SWDAVF has only been performed to reduce the risk of the surgical approach, successful endovascular occlusion for SWDAVF has been reported recently. Fukuda et al. reported two cases with successful endovascular treatment and concluded that SWDAVF could be cured safely by endovascular treatment with proper strategy and instruments.

CONCLUSION

Here, we report the rare patient of SWDAVF presenting with ocular symptoms. TVE was performed, and the patient had
a favorable outcome. To the best of our knowledge, this is the first report of a patient presenting with bilateral ocular symptoms due to SWDAVF with progressive bilateral ocular symptoms related to cranial nerve compression is warranted surgical intervention.

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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