Dural Arteriovenous Fistula Arising after Intracranial Surgery in Posterior Fossa of Nondominant Sinus: Two Cases and Literature Review

Abstract
The results of recent clinical and experimental studies suggest that the most important factor associated with the pathogenesis of dural arteriovenous fistula (AVF) is sinus thrombosis and subsequent venous or intrasinus hypertension. Here, we describe two patients who each developed a dural AVF after a posterior fossa craniotomy on the side of the nondominant or hypoplastic transverse (TS)-sigmoid (SS) sinuses. A 63-year-old female underwent surgical resection of a meningioma in the left cerebellopontine angle. Preoperative subtraction digital angiography (DSA) revealed a hypoplastic, ipsilateral left TS-SS and the sinus occlusion was revealed after surgery. Sixteen months later, she presented with a progressive left retroauricular, pulse-synchronous bruit. An AVF in the left TS-SS region was diagnosed by DSA and treated with transvenous coil embolization. The patient recovered without neurological deterioration. A 56-year-old female underwent surgical removal of an epidermoid tumor in the right cerebellopontine angle. Preoperative DSA revealed severe, ipsilateral right TS stenosis and the sinus occlusion was revealed after surgery. Two years later, she presented with the progressive right retroauricular, pulse-synchronous bruit, which was diagnosed by DSA as dural AVF in the right TS-SS region. She was treated with transvenous coil embolization and recovered without neurological deterioration. Sinus manipulation during intracranial surgery carries a potential risk of dural AVF development and this should be carefully considered, even when the ipsilateral TS-SS is nondominant or appears hypoplastic.

Keywords: Development hypoplastic sinus, dural arteriovenous fistulae, postoperative outcome

Introduction
Intracranial dural arteriovenous fistulae (AVF) are defined as abnormal arteriovenous shunts or connections that occur within the dura mater. Unlike arteriovenous malformations, the development of dural AVF seems more similar to that of acquired lesions.[1] Sinus thrombosis after trauma, infection, and surgery can cause venous outflow obstruction and venous hypertension and frequently arises in patients with dural AVF.[1-4] However, the pathophysiology of dural AVF development has not been elucidated. This report describes two patients who developed a dural AVF in the nondominant or hypoplastic transverse (TS)-sigmoid (SS) sinuses after the sinus occlusion during intracranial surgery. Both patients were evaluated by angiography just before the original surgery and when a dural AVF was diagnosed. We discuss the pathogenesis of dural AVF after the sinus occlusion during intracranial surgery in the posterior fossa based on our experience with these patients.

Case Reports
Case 1
A benign meningioma was surgically resected from the left cerebellopontine angle of a 63-year-old female using a left lateral suboccipital approach [Figure 1a and b]. Preoperative DSA revealed that the ipsilateral left TS-SS was hypoplastic, and the contralateral right TS-SS was dominant [Figure 1c]. The sinus occlusion was revealed after surgery and 6 months after craniotomy, three-dimensional enhanced computed tomography angiography (3D-CTA) showed venous reflux in the left TS-SS region [Figure 1d and e]. Sixteen months after craniotomy, she presented with the progressive left retroauricular pulse-synchronous bruit. Recurrent meningioma was not found using...
magnetic resonance imaging (MRI) and angiography, but 3D-CTA showed worsening venous reflux [Figure 1f]. Dural AVF in the left TS-SS fed by the left ascending pharyngeal and occipital (OA) arteries with retrograde drainage to the left inferior petrosal sinus with SS occlusion was confirmed by DSA [Figure 2a]. She was treated with transvenous coil embolization through a transconfluence approach, and she recovered without neurological deterioration [Figure 2b-d]. The patient has remained free of recurrent dural AVF for 5 years.

**Case 2**

An epidermoid tumor in the right cerebellopontine angle of a 56-year-old female was surgically resected through a right presigmoid, transpetrosal, and retrosigmoid approach [Figure 3a and b]. Preoperative 3D-CTA and DSA revealed severe ipsilateral right TS stenosis and dominant contralateral left TS-SS, [Figure 3c and d] and the sinus occlusion was revealed after surgery. Two years after intracranial surgery, she presented with the progressive right retroauricular pulse-synchronous bruit. Dural AVF in the right TS-SS region fed by the right OA was diagnosed by DSA, and the isolated sinus appearance with retrograde drainage to the right vein of Labbe and an inferior temporal vein was shown [Figure 4a]. She underwent transvenous coil embolization through a transconfluence approach, without neurological deterioration [Figure 4b-d]. The patient has remained free of recurrent dural AVF for 5 years.

**Discussion**

The etiology of dural AVF in the posterior fossa remains uncertain. Clinical observations and serial angiographic findings have indicated that many dural AVF are acquired and that sinus thrombosis,\(^1\) sinus hypertension,\(^1\) intracranial abscess,\(^2\) and trauma\(^3\) might contribute to their formation. The notion that this condition is acquired is supported by the finding that they frequently develop after trauma, craniotomy, and sinus thrombosis. Clinical observations indicate that pathological changes in the intracranial sinus, including sinus thrombosis, can lead to dural AVF formation.\(^4\) Notably, 39%–80% of dural AVF are associated with intracranial sinus thrombosis, and 78% of patients without thrombosis have sinus dysplasia, stenosis, septation, or distortion.\(^5-11\)

Table 1 shows published case series of dural AVF developing after intracranial surgery.\(^12-15\) Most of them occurred at the site of a prior intracranial surgery, regardless of the dominance or stenosis of venous or occluded sinuses before intracranial surgery. These authors speculated that venous thrombosis after sinus occlusion due to intracranial surgery might cause a fistula by opening the physiological shunts in the dura mater, which would consequently redirect arterial blood into the cortical veins, followed by the eventual development of a dural AVF at the site of the original intracranial surgery. On the other hand, experimental studies that have investigated the mechanisms or pathogenesis of dural AVF have found that they can be induced by venous hypertension without sinus or venous
Therefore, an exposure of the vessels to venous hypertension and subsequent local hypoperfusion might cause vessel dilation and the loss of sphincter control in arterioles. Increased intraluminal pressure in the vessels stimulates angiogenesis, and the subsequent formation of direct connections to a sinus or vein results in the development of dural AVF. Shin et al. suggested that venous sinus hypertension increases the production of vascular endothelial growth factor (VEGF), which might help to increase the angiogenesis and lead to an AVF. Matrix metallopeptidase-9 (MMP-9) is closely associated with neovascularization and might synergistically work with VEGF. Furthermore, cerebral venous sinus thrombosis induces MMP-9 upregulation and inflammatory activation. Dural AVF developed in our patients

<table>
<thead>
<tr>
<th>Authors</th>
<th>Age</th>
<th>Sex</th>
<th>Disease for intracranial surgery</th>
<th>Precraniotomy sinus findings of DSA</th>
<th>Craniotomy side</th>
<th>Development duration for DAVF</th>
<th>Location of DAVF</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sakaki et al.</td>
<td>50</td>
<td>Male</td>
<td>Right mandible ameloblastoma</td>
<td>No abnormality</td>
<td>Right</td>
<td>6 months</td>
<td>Right TS-SS</td>
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<tr>
<td></td>
<td>59</td>
<td>Female</td>
<td>Left retromastoid meningioma</td>
<td>Left SS occlusion by the tumor</td>
<td>Left</td>
<td>42 months</td>
<td>Left TS-SS</td>
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<tr>
<td></td>
<td>65</td>
<td>Male</td>
<td>Right neurinoma meningioma</td>
<td>No abnormality</td>
<td>Right</td>
<td>60 months</td>
<td>Right TS-SS</td>
</tr>
<tr>
<td></td>
<td>56</td>
<td>Male</td>
<td>Right hypoglossal neurinoma</td>
<td>Right SS occlusion by the tumor</td>
<td>Left</td>
<td>54 months</td>
<td>Right TS-SS</td>
</tr>
<tr>
<td>Converse et al.</td>
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<td>Female</td>
<td>Left glomus jugulare tumor</td>
<td>No abnormality</td>
<td>Right</td>
<td>22 months</td>
<td>Left TS-SS</td>
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<td>N/A</td>
<td>Right tentorial meningioma</td>
<td>SSS and right TS occlusion</td>
<td>Right</td>
<td>4 months</td>
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<tr>
<td>Nabors et al.</td>
<td>70</td>
<td>Female</td>
<td>Right trigeminal neuralgia</td>
<td>Not mentioned</td>
<td>Right</td>
<td>4 months</td>
<td>Right TS-SS</td>
</tr>
<tr>
<td></td>
<td>62</td>
<td>Male</td>
<td>Left hemifacial spasm</td>
<td>Not mentioned</td>
<td>Left</td>
<td>24 months</td>
<td>Right TS-SS</td>
</tr>
<tr>
<td>Sasaki et al.</td>
<td>58</td>
<td>Male</td>
<td>Right trigeminal neurinoma</td>
<td>No abnormality</td>
<td>Right</td>
<td>24 months</td>
<td>Right TS-SS</td>
</tr>
<tr>
<td>Present cases</td>
<td>63</td>
<td>Female</td>
<td>Left CP angle meningioma</td>
<td>Left hypoplastic TS-SS</td>
<td>Left</td>
<td>6 months</td>
<td>Left TS-SS</td>
</tr>
<tr>
<td></td>
<td>56</td>
<td>Female</td>
<td>Right CP angle epidermoid tumor</td>
<td>Right TS severe stenosis</td>
<td>Right</td>
<td>24 months</td>
<td>Right TS-SS</td>
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</tbody>
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DAVF – Dural arteriovenous fistulae; CP – Cerebellopontine; TS – Transverse sinus; SS – Sigmoid sinus; SSS – Superior sagittal sinus; DSA – Digital subtraction angiography; N/A – Not applicable
The sigmoid and transverse sinuses can be occluded during presigmoid, transpetrosal posterior fossa craniotomy. Patients were discovered asymptomatic sinus occlusion after surgery. Jean et al. reported that 10% of patients were discovered asymptomatic sinus occlusion after posterior fossa craniotomy. The sigmoid and transverse sinuses can be occluded during presigmoid, transpetrosal approaches to the skull base. There are scant data available on the incidence of venous sinus occlusion after surgery. Jean et al. reported that 10% of patients were discovered asymptomatic sinus occlusion after posterior fossa craniotomy. The sigmoid and transverse sinuses can be occluded during presigmoid, transpetrosal approaches to the skull base tumor. It is important to follow-up MRI because there is a possibility of occurrence of dural AVF if asymptomatic sinus occlusion is observed after posterior fossa surgery.

Transfemoral embolization of the feeding arteries, embolization of the affected sinus or complete surgical excision is recommended to treat dural AVF. We believe that all of these procedures, including surgical excision, are valuable in the appropriate setting. Transvenous embolization of venous channels appears to be the first choice because to penetrate a fistulous point using glue injected via transarterial embolization is difficult. Shunts in our patients were completely occluded after transvenous sinus embolization through a transvenous approach because these affected sinuses were not associated with normal cerebral venous drainage. Embolization using Onyx (eV3 Neurovascular, Inc., Irvine, CA, USA), which penetrates better and allows for robust injections, is a new and promising modality for treating dural AVF. However, Onyx has the same disadvantage as other embolic materials in that its ability to penetrate small dangerous anastomoses to cranial nerves or vessels supplying the brain cannot be controlled. Findings from long-term follow-up are awaited to establish the value of embolization with Onyx for dural AVF.

Conclusion

Sinus manipulation during intracranial surgery of the posterior fossa carries a risk of sinus occlusion and developing postoperative dural AVF. This should be considered even if the ipsilateral TS-SS is nondominant or appears hypoplastic.

Informed consent

Informed consent was obtained from all individual participants included in the study.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent forms. In the form, the patient has given her consent for her images and other clinical information to be reported in the journal. The patient understands that name and initials will not be published and due efforts will be made to conceal identity, but anonymity cannot be guaranteed.

Financial support and sponsorship

Nil.

Conflicts of interest

There are no conflicts of interest.

References


