Medullary Hemorrhage Caused by Foramen Magnum Dural Arteriovenous Fistula Successfully Obliterated using Combination of Endovascular and Surgical Treatments: A Case Report and Literature Review

Abstract
The authors describe an extremely rare case of foramen magnum dural arteriovenous fistula (DAVF), Cognard type V, presented with medullary hemorrhage caused by venous varix on the lateral medullary draining vein embedded into the medulla oblongata. Following mild myelopathy for 3 days, a 20-year-old male developed dyspnea, generalized seizures, loss of consciousness, and finally cardiac arrest. After successful resuscitation, computed tomography scan (CT) of the brain was obtained and showed acute medullary hemorrhage. Subsequent magnetic resonance imaging of the brain revealed diffuse venous congestion or edema of the medulla with multiple dilated flow voids surrounding the medulla, more prominent on the left side, with venous varix embedded into the left-sided of the lower medulla. He was sent to the emergency department of the local hospital and intubated promptly. A few minutes later, the patient had a cardiac arrest. Digital subtraction angiography (DSA) demonstrated DAVF of the foramen magnum supplied mainly by dural branches of bilateral hypertrophic posterior inferior cerebellar arteries (PICAs), slightly by the posterior meningeal branch of the left vertebral artery, and the jugular branch of the left ascending pharyngeal artery (APA) originating from the occipital artery. Transarterial embolization through the bilateral dural branches of the PICAs was successfully performed using N-butyl-2-cyanoacrylate (NBCA), resulting in complete obliteration. The patient had excellent recovery and lost to annual follow-up. Seven years later, he had a recurrent of the fistula presented with occipital headache. DSA with angiographic CT in three-dimensional reconstruction and maximum intensity projection reformatted images clearly demonstrated the exact location of the DAVFs at the posterior rim of the foramen magnum, mainly recruited by the hypertrophic jugular branch of the APA originating from the occipital artery. The fistula was successfully treated surgically following transarterial embolization through the jugular branch of the APA using NBCA. Follow-up DSA confirmed complete obliteration of the DAVF. The patient has remained clinically asymptomatic 2 years after the operation.

Keywords: Brainstem congestion, dural arteriovenous fistula, foramen magnum, medullary hemorrhage, pial supply, venous varix

Introduction
Intracranial dural arteriovenous fistulas (DAVFs) are abnormal arteriovenous connections located within the leaflets of the dura matter, commonly near the venous sinuses. Arterial supply usually arises from dural or meningeal arteries. Dural branches from the pial arteries are rare. Venous drainage may occur into the dural sinuses, dural, osteodural, and/or leptomeningeal veins.\(^1\)\(^2\) Symptoms of the patients usually relate to the location of the fistulas and venous drainage patterns.\(^3\) Based on the pattern of venous drainage by Cognard et al.,\(^4\) intracranial DAVFs were classified into five types, including Type I: Fistulas drain into the main sinus with antegrade flow; Type IIa: Retrograde flow into the sinus (es) only; Type IIb: Retrograde flow to cortical vein (s) only; Type III: Direct cortical venous drainage without venous ectasia; Type IV: direct cortical venous drainage with venous ectasia; and Type V: Into the spinal perimedullary veins. Patients harboring DAVFs manifest with intracranial hemorrhage at initial diagnosis in approximately 18%–24% of all intracranial DAVFs.\(^5,6\) Foramen magnum DAVFs are rare and account for 1.5%–2.3% of all location of intracranial DAVFs.\(^4,6,8\) DAVFs of this location usually occur in a middle-aged patient with a...
strong male predominance. They commonly manifest with progressive myelopathy and posterior fossa intracranial hemorrhage (i.e., subarachnoid hemorrhage [SAH]).

We describe a patient with foramen magnum DAVF, Cognard type V, presenting with respiratory insufficiency following by seizures, unconsciousness, and developing of cardiac arrest caused by medullary hemorrhage from ruptured venous varix on the draining vein invaginated into the medulla oblongata. We also review the literature of patients having foramen magnum DAVFs.

**Case Report**

A 20-year-old smoker male experienced mild weakness and paresthesia of lower extremities for 3 days. There was no underlying diseases or history of trauma. He went to the clinic and took home some medicine. Three days later, the patient developed dyspnea, generalized seizures, and subsequent loss of consciousness. He was sent to the emergency department of the local hospital and intubated promptly. A few minutes later, the patient had a cardiac arrest. Immediate cardiopulmonary resuscitation was performed nine times until the presence of vital signs. Computed tomography (CT) scan of the brain was obtained and showed a hyperdense lesion in the dorsal region of the left-sided of the medulla oblongata, corresponding to medullary hemorrhage [Figure 1]. The patient was sent to the intensive care unit for close observation until clinical stable. Ten days later, the patient was sent to another larger local hospital. Following day, magnetic resonance imaging (MRI) of the brain revealed diffuse hyperintensity of the medulla oblongata on fluid-attenuated inversion recovery and T2-weighted sequences, corresponding to venous congestion or edema of the medulla. There were multiple dilated flow voids surrounding the medulla, more prominent on the left side, with venous varix embedded into the left-sided of the lower medulla. T1-weighted images demonstrate a round mixed iso and crescent hyperintense lesion in the dorsal region of the enlarged upper medulla, representing acute to subacute hematoma. The trajectory of hemorrhage projected superiorly from the left-sided of the lower medulla into the mid-dorsal region of the upper medulla [Figure 2]. Tracheostomy was performed due to prolonged intubation and recurrent pneumonia. The disabled patient gradually improved and could follow command in the next 2 weeks with quadripareysis status (power Grade 1–2/5). Two weeks later, the patient was transferred to Prasat Neurological Institute (PNI) for further investigation and proper

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**Figure 1:** Computed tomography scan of the brain, obtained after initial aggressive symptoms, shows a hyperdense lesion in the dorsal region of the left-sided of the medulla oblongata (arrowhead), corresponding to acute medullary hemorrhage.

**Figure 2:** Magnetic resonance imaging of the brain obtained 11 days after initial aggressive symptoms. (a) Axial fluid-attenuated inversion recovery image shows diffuse edema of the medulla with a venous varix embedded into the left-sided of the medulla. (b) Axial T2-weighted image also reveals diffuse edema with prominent dilated flow-voids surrounding the lower medulla. (c) Sagittal T1-weighted image demonstrates a round mixed iso and crescent hyperintense lesion in the dorsal region of the enlarged upper medulla, representing acute to subacute hematoma. (d) Coronal T1-weighted image discloses the trajectory of hemorrhage from the left-sided of the lower medulla into the mid-dorsal region of the upper medulla. (e) Coronal T2*-weighted gradient-echo (GRE) image shows thin hypointense rim of hemosiderin.
Figure 3: Cerebral angiography of the brain obtained 1 month after initial symptoms. Anteroposterior (a and c) and lateral views (b and d) of bilateral vertebral arteries injections show dural arteriovenous fistulas of the foramen magnum mainly supplied by bilateral hypertrophic posterior inferior cerebellar arteries. The posterior meningeal branch of the left VA also feed the fistulas. Anteroposterior (e) and lateral (f) views of late venous phase of the right vertebral artery injection demonstrate deep venous drainage to both sides of the brainstem along cerebellomedullary and cerebellopontine cisterns. On the left side, the left basal vein of Rosenthal receives venous blood from the anterior pontomesencephalic vein (white arrowhead) and the left lateral mesencephalic vein (black arrowhead) with subsequent drainage into the great vein of Galen. On the right side, the right dilated superior hemispheric vein connects from the dilated right transverse pontine vein with further drainage into the right proximal transverse sinus via the tentorial vein (black arrow). In addition, anteroposterior (g) and lateral (h) views of the left external carotid artery injection reveal the minimal supply from the neuromeningeal trunk of the left ascending pharyngeal artery originating from the occipital artery.

Figure 4: Anteroposterior (a and c) and lateral (b and d) views of superselective catheterization of bilateral posterior inferior cerebellar arteries injections clearly reveal dural branches of both posterior inferior cerebellar arteries supplying the fistulas. During embolization with two injections via the right posterior inferior cerebellar artery, lateral views (e and f) demonstrate glue cast penetrating into the proximal draining vein. Postembolization, anteroposterior (g) and lateral (h) views of the right vertebral artery confirm complete obliteration of the fistulas.
management. The neurological examination revealed fully consciousness, evidence of spastic quadripareisis (power grade 2–3/5), hyperreflexia of upper and lower extremities, and the presence of Babinski’s sign. Digital subtraction angiography (DSA) demonstrated DAVF of the foramen magnum supplied mainly by dural branches of bilateral hypertrophic posterior inferior cerebellar arteries (PICAs), slightly by the posterior meningeal branch of the left
vertebral artery (VA), and the neuromeningeal trunk of the left ascending pharyngeal artery (APA) originating from the occipital artery. The venous drainage was split in two directions: superiorly along cerebellomedullary and cerebellopontine cisterns into both petrosal veins. On the left side, the
left basal vein of Rosenthal receives venous blood from the anterior pontomesencephalic vein and the left lateral mesencephalic vein with subsequent drainage into the great vein of Galen. On another side, the right dilated superior hemispheric vein connects from the dilated right transverse pontine vein with further drainage into the right proximal transverse sinus through the tentorial sinus [Figure 3]. Transarterial embolization through the bilateral dural branches of the PICAs was successfully performed using N-butyl-2-cyanoacrylate (NBCA). Postembolization angiography confirmed complete obliteration of the fistulas [Figure 4]. MRI of the brain obtained 1 week after endovascular treatment revealed a large round mixed central isointense and peripheral hypointense mass with hyperintensity in a venous aneurysm, probably representing a thrombosed venous aneurysm with resolving hematoma. There was the disappearance of previously seen venous congestion of the medulla and multiple dilated flow-voids surrounding the medulla [Figure 5]. The patient gradually improved, and tracheostomy tube was removed before discharging home 2 weeks after treatment. Six months after endovascular treatment, the patient showed no residual neurological deficits. Routine annual follow-up was scheduled for him, but the patient was lost to follow-up without any reasons.

Seven years later, the patient experienced occipital headache without the stiffness of the neck for 7 days. He went to the local hospital and was sent back to PNI again due to the previous history of DAVFs of the foramen magnum. There were no neurological deficits on neurological examination. Follow-up MRI of the brain showed multiple dilated flow-voids, more prominent on the right side, along both cerebellomedullary and cerebellopontine cisterns, representing recurrent DAVFs of the foramen magnum. There was the disappearance of a previously seen thrombosed venous aneurysm and complete resolution of the hematoma in the medulla with the small residual...
hypointense area of hemosiderin stain in the dorsal region of the medulla [Figure 6]. DSA with angiographic CT in three-dimensional reconstruction and maximum intensity projection (MIP) reformatted images of the cranio-cervical junction clearly demonstrated the exact location of the DAVFs at the posterior rim of the foramen magnum, mainly supplied by the hypertrophic jugular branch of the APA originating from the occipital artery. Without supplying from the right PICA, the left PICA and posterior meningeal branch of the left VA partially fed the fistulas. The venous drainage drained to both sides of the medulla and run superiorly along cerebello-medullary and cerebello-pontine cisterns into both petrosal veins. On the left side, the fistulas drain superiorly into the great vein of Galen through the left dilated vein of cerebellomesencephalic fissure. On the right side, the fistulas still drained into the right proximal transverse sinus through the dilated superior hemispheric vein and tentorial sinus, respectively. Inferiorly, it also drains into anterior medullary vein connecting to anterior spinal vein [Figure 7]. Endovascular treatment was performed through the jugular branch of the left APA using NBCA [Figure 8]. DSA after embolization revealed residual fistulas. Due to incomplete obliteration of the fistulas, the patient then was informed about surgical option and accepted this option.

Following suboccipital craniotomy without C1 laminectomy, resecting the dural leaflets and disconnecting leptomeningeal medullary draining veins were successfully performed using indocyanine green fluorescence imaging during operation [Figure 9]. In addition, watertight duraplasty was done with synthetic dural graft. The postoperative course was uneventful. Follow-up DSA, obtained 1 month after surgery, confirmed complete obliteration of the fistula [Figure 10]. The patient has remained clinically asymptomatic 2 years after the operation.

Discussion

The common etiologies of the medullary hemorrhages are cavernous malformation or hemorrhagic transformation following infarction.[10] To the best of our knowledge, there was only one previously reported case of medullary hemorrhage caused by DAVF at the cranio-cervical junction, supplied by the meningeal branch of the VA with draining superiorly into the anterior medullary vein in elderly woman.[11] She was complicated by acute neurogenic pulmonary edema, resulting in acute respiratory distress syndrome. In the present study, our case was the first case report of foramen magnum DAVF presented with acute medullary hemorrhage, leading to respiratory dysfunction, subsequent seizure, and finally developing cardiac arrest.

Involuntary convulsive-like movements, for example, jerky, tonic-clonic, intermittent shaking, or decerebrate postures, may occur in patients with brainstem stroke.[12] Without electroencephalography, we could not conclude that our case had real seizures. However, foramen magnum DAVF may present with epilepsy due to venous drainage superiorly to the temporal lobe, leading to temporal venous congestion.[13]

According to Cognard classification,[4] a foramen magnum DAVF with perimedullary and spinal venous drainage in our case is classified as Type V. The patients with Cognard type V intracranial DAVFs may experience bulbar palsy, and/or respiratory failure due to medullary venous congestion.[14] Without descriptions in details, Cognard et al.[4] disclosed that all three cases of foramen magnum DAVFs, classified as Cognard type IV and V, presented with hemorrhages. Based on the study of the relation between clinical presentation and venous drainage of intracranial DAVFs with spinal venous drainage by Brunereau et al.,[15] they found that the patients presenting with hemorrhage had venous drainage limited to the cervical cord, whereas those with myelopathy had extensive spinal venous drainage descending toward the conus medullaris.

The common types of hemorrhage in intracranial DAVFs are SAH and intracerebral hemorrhage.[3,16] Brainstem hemorrhage is extremely rare. Nakajima et al.[17] reported an elderly man with DAVFs of the sinus of the lesser sphenoid wing presented with pontine hemorrhage
<table>
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<tr>
<th>Authors</th>
<th>Gender/age</th>
<th>Symptoms and signs/images findings</th>
<th>Arterial supply</th>
<th>Location of draining veins</th>
<th>Associating venous varix</th>
<th>Treatment</th>
<th>Neurological outcome</th>
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<td>Embolization with Onyx</td>
<td>Right hemiparesis (power grade 4/5)</td>
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caused by venous varix on lateral mesencephalic vein invaginated into the pons. Another case report by Lasjaunias et al., they demonstrated small midbrain hemorrhage probably caused by ruptured perimesencephalic draining vein from a foramen magnum DAVF in middle-aged man.

The presence of retrograde cortical venous drainage or leptomeningeal venous drainage in DAVFs is the most imperative risk factor of intracranial DAVFs associated with hemorrhage.\[6,18,19\] In multivariate logistic regression analysis by Singh et al., male gender, age older than 50, and posterior fossa location were also found to be independently associated with hemorrhagic manifestation. In addition, smoking was more common in the hemorrhagic group. In our case, there were many risk factors, including retrograde leptomeningeal venous drainage, male gender, posterior fossa location, and history of smoking, except young age.

We reviewed the published case reports which have sufficient clinical description and clearly demonstrated figures of foramen magnum DAVFs.\[3,9,13,16,20-35\] The collected data in this review include demographic data (i.e., gender and age of patient), presenting symptoms, the findings of image studies, arterial feeders of the fistula, location of the draining veins, the presence of venous varix, treatment of the fistula, and neurological outcomes following treatment [Table 1]. From the literature review, there were 27 cases, including our case, with 27 foramen magnum DAVFs. All patients except one were male (96.3%) with a median age 49, range 20–69 years. Of 27 cases, 15 (55.6%) were hemorrhagic presentation, including SAH, intraventricular hemorrhage, cerebellar hemorrhage, and medullary hemorrhage (our case). The most common type of hemorrhage was SAH (80%). Another 12 cases were non-hemorrhagic presentation, including progressive myelopathy, epilepsy, tinnitus, floating sensation. The median age in hemorrhagic group was 46 (range 20–58 years), whereas nonhemorrhagic group was 51.5 (range 38–69 years). Only 2 cases, including our cases, presented with myelopathy and hemorrhage. Most patients with hemorrhagic manifestation had a venous varix being the source of hemorrhage. The arterial supply included APA, VA, OA, and/or PICA. Twenty-two foramen magnum DAVFs (81.5%) supplied by the branch of APA, including hypoglossal and/or jugular branches. Four cases had blood supply only from VA. Only our case had additional supply from PICA. All patients with progressive myelopathy had venous drainage into the spinal cord, i.e., Cognard Type V. Four cases, including our cases, had respiratory insufficiency resulting from venous drainage to the medulla, leading to medullary congestion. Eighteen fistulas were treated with endovascular treatment alone, five with surgery alone, three with surgery following embolization, and one left...
untreated. Most embolic material used for treatment was liquid embolic material, including glue and Onyx. Most patients had good neurological outcome after treatment.

Retrograde leptomeningeal venous drainage, aneurysmal venous dilatation, and galenic drainage are factors predisposing to aggressive neurological presentation, i.e., intracranial hemorrhage or progressive neurological deficit.[2] Undoubtedly, leptomeningeal draining pathway with venous varix embedded into the medulla is the source of hemorrhage, leading to catastrophic neurological behavior in our case. Furthermore, we previously reviewed a rare condition of patients suffering from intramedullary hemorrhage caused by spinal DAVFs.[3] In most of the cases, venous varices on draining veins were the source of intramedullary hemorrhage. Similar to our case, Kai et al.[24] demonstrated a false aneurysm protruding into the medulla. However, this case presented with SAH. In addition, Motebejane and Choi[9] revealed that all patients harboring foramen magnum DAVFs with SAH in their series had intracranial venous drainage with venous aneurysms. Based on our review, hemorrhage in foramen magnum DAVFs usually resulted from ruptured venous varix.

Foramen magnum DAVFs can be treated by endovascular treatment, surgery, or both depend on institutions preference.[9,16,27,30] However, best management decisions should be approached by a multidisciplinary team.[3] DAVFs in the region of the foramen magnum tend to have leptomeningeal venous reflux and a high risk for intracranial hemorrhage. Therefore, aggressive treatment should be achieved.[24] The goal of treatment is to obtain the anatomical cure.[16]

Interestingly, the foramen magnum DAVF in our case mainly supplied by bilateral PICAs, i.e., pial arteries. Only 5% of all intracranial DAVFs supplied by both meningeal and pial arteries.[3] DAVFs fed by the pial arteries are associated with a higher risk of stroke or major complications either during endovascular or surgical treatment.[37] Osada and Kring[38] investigated the characteristics of pial arterial supplies in intracranial DAVFs and classified them into dilated pre-existing dural branches of pial arteries and a pure pial supply. Younger age, DAVFs within tentorium, and the presence of venous dilatation (i.e., a sign of venous hypertension) were independent predictors of a pial arterial supply. In their series, they also reported dilated dural branch of pial arteries from PCA in DAVFs of tentorium, torcular herophili, and foramen magnum DAVFs. Recognition of additional pial supply is imperative for avoiding inadvertent complication related to the reflux of liquid embolic materials into pial vessels supplying normal brain tissue. Dural vessels arising from PCA are the posterior meningeal artery and the artery of the falx cerebelli.[39] Onyx embolization in DAVFs through pial arterial supply could increase the risk of procedure-related complications, i.e., periprocedure hemorrhage.[40] In our case, the posterior meningeal arteries from PICAs pierced the dura at the posterior rim of the foramen magnum. Fortunately, the fistula was successfully treated using NBCA via these branches without any complications.

Medullary congestion can be either due to intracranial DAVFs with caudally drainage into the anterior spinal vein through the medullary or cervical DAVFs with rostrally drainage into medullary veins.[4,41] According to a systemic review by El Asri et al.[42] the poor outcomes were correlated to the presence of brainstem signal abnormalities on MRI. Unlike primary intracerebral hemorrhage, the patient suffering from extensive hemorrhagic venous infarction even caused by high-flow AVF may dramatically recover following prompt endovascular treatment.[43] In our case, the patient had regained full motor strength without residual neurological deficits after marked reduction of the flow of the fistula by transarterial embolization using NBCA through pial supply of bilateral PICAs.

Incomplete embolization, using non-permanent embolic, or partial surgical of the fistula may result in recurrent fistula, probably leading to fatal rebleeding.[16] Even though using transarterial embolization with Onyx, the recurrence of the fistula can occur.[13] Therefore, long-term follow-up with radiographical images is mandatory to early detect the recurrent fistula. Recanalization of embolized vessels occurred in our case several years later with the manifestation of occipital headache. The fistula also recruited other feeding artery, i.e., the APA.

Motebejane and Choi[9] demonstrated that all foramen magnum DAVFs in their series were supplied by the hypoglossal and/or jugular branches of the APA, commonly by hypoglossal branch. The APA may arise from the proximal occipital artery.[24] With angiographic CT in three-dimensional reconstruction and MIP reformatted images, in our case clearly demonstrated the jugular branch of the APA originating from the occipital artery. Given delay in treatment and probable subsequent poor outcomes, DAVFs at the foramen magnum may not be identified using standard 4 vessels cerebral angiography.[20,21,24] Due to common feeder by the neuromeningeal trunk of the APA, selective ascending pharyngeal angiography should be performed. Endovascular treatment through this artery may carry the risks of inadvertent embolization due to potential risk of liquid embolic materials leakage through extracranial-intracranial anastomoses or injury to vasa nervorum of cranial nerves.[1] To prevent major reflux into the main trunk, the tip of the microcatheter should be navigated close enough to the fistula.[35] In addition, balloon-augmented liquid embolic material embolization through neuromeningeal trunk of APA was used to prevent the risks of injury to the lower cranial nerves.[9,31,33] Transarterial embolization alone is likely to be ineffective due to the extensive collateral network and high vascularity of the dura.[24] Predictably, there was residual fistula
following transarterial embolization through the APA using NBCA in our case.

MRI and cerebral angiography may not depict the exact location of the fistulas at the skull base.[21] We agree with Pop et al.[23] that rotational angiographic CT is useful in delineating the complex anatomy of the foramen magnum DAVFs. In our case, angiographic CT in three-dimensional reconstructed image and MIP reformatted images can identify the exact location of fistula at the posterior border of the foramen magnum and complex venous drainages.

According to the literature review in Cognard type V intracranial DAVFs by El Asri et al.,[42] the surgical treatment appears to be more effective than endovascular treatment. Microsurgical treatment of foramen magnum DAVFs probably is an effective and more reliable method of treatment.[27] Many surgeons preferred suboccipital approach with partial or hemilaminectomy for foramen magnum DAVFs.[3,30,34] In our case, suboccipital approach without laminectomy of C1 was enough for resecting the dural leaflets and disconnecting leptomeningeal veins due to knowing the exact location of the fistula from angiographic CT.

Conclusion

We illustrated an extremely rare case of foramen magnum DAVF, Cognard type V, presented with catastrophic neurological behavior, caused by ruptured venous varix, leading to medullary hemorrhage. The patient had a complete recovery after endovascular treatment via bilateral PICAs with complete obliteration of the fistula. Seven years later, he had a recurrent DAVF presented with occipital headache. Finally, we could achieve a cure of the DAVF by surgical removal of the dural leaflets and disconnection of the fistula following endovascular treatment via the APA with excellent outcome. Understanding the angioarchitecture of foramen magnum DAVFs, including arterial feeders, venous drainage pattern, associated venous pouch, and localization of the fistulous zone, is the key to successful management. In addition, the exact location of DAVFs may influence the surgical approach.

Consent

The patient has given consent to be enrolled and has her data published.

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.

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Nil.

Conflicts of interest

There are no conflicts of interest.

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