Extensive Spinal Cord Infarction after Surgical Interruption of Thoracolumbar Dural Arteriovenous Fistula Presenting with Subarachnoid Hemorrhage

Sang-Hun Lee, M.D., Ki-Tack Kim, M.D., Sung-Min Kim, M.D., Dae-Jean Jo, M.D.
Departments of Orthopaedic Surgery, Neurosurgery, Spine Center, Kyung Hee University East West Neomedical Center, Seoul, Korea

Nontraumatic intracranial subarachnoid hemorrhage (SAH) attributable to the thoracolumbar dural arteriovenous fistulas (DAVF's) has been extremely rare. A 41-year-old male patient was admitted with severe acute headache, neck stiffness, and pronounced low-back pain radiating to both legs. The T2-weighted MR imaging showed irregular signal void and enlarged, varix like pouch formation with spinal cord compression at the T11-12 level. The angiogram revealed a DAVF.
We report a DAVF case with SAH that revealed an extensive infarction from C5 to the conus medullaris after undergoing operative treatment.

KEY WORDS: Dural arteriovenous fistula · Subarachnoid hemorrhage · Spinal cord infarction.

INTRODUCTION

Dural arteriovenous fistula (DAVF, type I spinal arteriovenous malformation) is often developed at the thoracolumbar junction of middle-aged men with clinical symptoms including paraparesis, sensory deficits, and disturbance of bowel/bladder function resulted from venous hypertension and ischemic myelopathy. There are also cases presenting with subarachnoid hemorrhage (SAH) when venous pressure is high. Reports on nontraumatic intracranial SAH due to spinal DAVFs involving the cervical vertebra or the cervicothoracic junction have been made. To our knowledge, there is only one reported case of SAH occurred at the L4 level and no study has described SAH located at the thoracolumbar junction.

Spinal cord infarction is a very rare disorder, accounting for less than 1% of all strokes. Its pathogenesis and prognosis remain largely unknown. Some of the identified risk factors include reperfusion injury after aortic clamping during thoracoabdominal aortic surgery, excessive hypoxia or systemic hypotension that sustained during operation, epidural anesthesia or lumbar puncture complications, unexpected embolic episode associated with embolization of the renal artery or mesenteric artery, cardiogenic thromboembolus, and artherosclerosis. In lower frequencies, cases with vascular diseases, polycythemia, decompression sickness, hypercoagulability, and meningeval infection have been reported.

In a meta-analysis of 44 spinal cord infarctions in 1996, Chesire et al. reported 3 cases presenting with both spinal arteriovenous malformation and spinal cord infarction. However, the infarction extended from the cervical level to the thoracolumbar level was not found.

We report a DAVF case with SAH that revealed an extensive infarction from C5 to the conus medullaris after undergoing operation.

CASE REPORT

History. A 41-year-old male presented with severe acute headache, neck stiffness, and pronounced low-back pain radiating to both legs. He had intermittent both leg radiating pain during last two years. He had no history of diabetes, hypertension, smoking or coagulopathy.

Examination. Neurological examination demonstrated...
nuchal rigidity and a positive straight-leg raising sign. Laboratory study showed mild leukocytosis (14,500/μL), normal platelet count, prothrombin time (PT), and activated partial thromboplastin time (aPTT).

The T2-weighted whole spinal magnetic resonance imaging showed irregular signal void and enlarged, varix like pouch formation with spinal cord compression at the T11-12 level (Fig. 1). SAH was noted at the posterior cranial fossa and at the L5-S1 level. His cervical body to canal ratio at C5 was 0.45 and a narrow spinal canal from C3 to the thoracic spine was noted. High signal intensities were found at the C5-6 level with central disc herniation (Fig. 2). Under the impression of SAH associated with spinal DAVF, a spinal angiogram was subsequently obtained. The angiogram revealed a DAVF with arterial blood supply from the segmental artery between T12 and L1 in an early phase and venous drainage into the intrathecal venous plexus involving varicose radiculospinal veins. But, the artery of Adamkiewicz (AKA) was not found (Fig. 3).

Operation. A midline posterior approach centered on the T12 region was performed with electrophysiological monitoring of somatosensory-evoked potential and motor-evoked potential. After laminectomy and vertical midline durotomy, exposed variably sized dilated tortuous and serpentine peri-medullary veins with the hematoma compressing the spinal cord. An enlarged fistula measured 2mm in diameter was located at the ventral side of the T12 root (Fig. 4). Temporary ligations were placed at the proximal end, interrupting the venous drainage of the fistula and causing the vein to

**Fig. 1.** Right parasagittal (A) and midsagittal (B) T2 images showing irregular fluid void and enlarged, varix like pouch formation (arrows) with spinal cord compression at the level T11-12. Collection of subarachnoid hemorrhage (arrow head) is noted at the level L5-S1.

**Fig. 2.** Preoperative right parasagittal (A), midsagittal (B) and left parasagittal (C) sagittal T2 images showing narrow spinal canal from C3 to thoracic spine. High signal intensities (arrows) are found on both parasagittal images at the level C5-6 with central disc herniation. Subarachnoid hemorrhage within posterior cranial fossa is noted (arrow heads).

**Fig. 3.** Anteroposterior angiogram revealing the dural arteriovenous fistula with arterial blood supply from the segmental artery between T12 and L1 in early phase (A) and venous drainage into the intrathecal venous plexus occurs via varicose radiculospinal veins (B and C).

**Fig. 4.** Intraoperative photographs showing enlarged serpentine venous structures at the dorsal surface of spinal cord (A) and temporary ligation of fistula located just ventral side of T12 root.

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impression of epidural hematoma, we decided to perform emergent decompressive surgery. However, there was no evidence of hematoma in the epidural space, we opened the durotomy site for exploration. Severe cord swelling was seen. Thus, we carried out extended decompressive laminectomy with duroplasty using a dural substitute (Neurapatch : B. Braun Melsungen AG, Melsungen, Germany), and placed a lumbar CSF drain. The patient exhibited a complete paraplegia after the second operation.

Postoperative Course. Spinal MR imaging was obtained postoperatively, which confirmed spinal cord infarction from C5-6 to the conus medullaris (Fig. 5-1, 2). During the first 3 months after operation, neurological deficits improved gradually and reached a plateau in recovery. At the 1-year clinical follow-up examination, the patient still complained of right leg weakness (grade IV) and disturbance of bowel and bladder function. He could ambulate with support of a cane. Repeated MR imaging 8 months later showed decreased size and intensities of spinal cord infarction, but hyperintensities were still visible (Fig. 6).

**DISCUSSION**

The anterior spinal artery is the one involved with the pathogenic mechanism in spinal cord infarction. The most important feeder among them is the AKA, which branches between T9 and T12 on the left side in 65-75%. Interruption of blood supply due to embolism or injury of the artery has been cited as the main cause of spinal cord infarction.

Spinal cord infarction is also referred to as acute spinal cord ischemia syndrome (ASCIS). The general factors of ischemia include global hypoperfusion, reperfusion injury, coagulopathy, vasculitis, and decompression sickness.

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**Fig. 5-1.** Immediate postoperative right paracentral (A), midline (B) and left para central (C) sagittal T2 images showing widespread hyperintensities (arrows) with cord swelling from the level C5 to L1, consistent with spinal cord infarction. There is no residual arteriovenous fistula or varix like dilated vein.

**Fig. 5-2.** Immediate postoperative axial T2 images showing bilateral hyperintensities corresponding to the whole gray matter.

**Fig. 6.** Eight months after surgery, the size and intensities of spinal cord infarction (arrows) are decreased but still visible on right para-central (A), midline (B) and left para central (C) sagittal T2 images.
while the local factors are epidural hematoma and thromboembolic accident.

Our case showed extensive infarction reaching the cervical cord without notable risk factors after surgical interruption of thoracolumbar DAVF. However, this patient had some features to note: 1) the cervical neural canal was so narrow that preoperative MR imaging showed a nearly obliterated CSF space surrounding the spinal cord, 2) a high signal intensity resulted from the central disc herniation at C5-6 was observed on T2W sagittal MR imaging, 3) SAH extended to the posterior cranial fossa and the sacral area, and 4) a number of enlarged, dilated and tortuous epidural veins were observed during operation. These four features led us to presume that the patient had high intradural pressure within a relatively narrow neural canal.

Considering the above features, we reached the following two pathogenic mechanisms for the extensive infarction:

1) The blood vessel from which DAVF originated might have been the AKA. Preoperative angiography confirmed the feeding site on the right, but the AKA was not found on the left. Although the DAVF originated from the intervertebral foramen between T12 and L1 on the right side, the AKA may extend to anywhere between T5 and L2. About 25% of the AKA is known to extend on the right side. Hemodynamic changes or thrombosis in the spinal cord caused by the interruption of DAVF originating from the AKA might have resulted in the extensive infarction. However, this possibility alone does not clearly explain the infarction extending from the upper thoracic to the cervical cord.

2) The next possible mechanism is the rapid increase of blood flow in the spinal cord that had been under the ischemic state in an abnormal CSF dynamic pathway within a narrow and high-pressed spinal canal when DAVF was interrupted. As a result, reperfusion injury might have been sustained, leading to secondary injury and swelling of the spinal cord.

Chronic spinal cord compression is known to cause subarachnoid fibrosis, reducing collateral blood flow and the compensation mechanism for ischemic events. In fact, there are reports on the cases that developed spinal cord infarction due to compression caused by cervical myelopathy. In our case, the narrow spinal canal is considered to have played a major role in developing extensive infarction.

According to recent studies, a severely compressed cervical spinal cord due to cervical spondylopathy accompanied significant swelling after decompression surgery. There are also studies that suggest the transient root palsy observed after decompressive surgery is an incomplete cord injury caused by a reperfusion injury caused by a reperfusion injury and secondary infarction resulting from the temporarily increasing arterial pressure when the fistula shunt was blocked.

The prognosis for spinal cord infarction remains unclear, but the early neurologic symptoms and age of a patient have been cited as major factors that affect prognosis. Our patient showed gradual recovery of motor and sensory functions after operation, but partial paralysis persisted on the right lower limb. In addition, bladder and bowel functions have not recovered, showing a severe sequel.

CONCLUSION

Based on this case report, we recommend that the risk of neurologic complications be considered in advance when performing operation on a DAVF patient with a narrow spinal canal. Surgeons are also advised to check the status of blood supply to the spinal cord by confirming the AKA of the spinal cord and the feeding artery on preoperative angiography, in an effort to predict and prevent postoperative complications.

References


