Case Series: Long segment extra-arachnoid fluid collections: Role of dynamic CT myelography in diagnosis and treatment planning

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Abstract

We report five patients in whom spinal MRI revealed extra-arachnoid fluid collections. These spinal fluid collections most likely resulted from accumulation of cerebrospinal fluid (CSF) from a dural leak. The patients presented with either compressive myelopathy due to the cyst or superficial siderosis (SS). All of these fluid collections were long segment, and MRI demonstrated the fluid collections but not the exact site of leak. Dynamic CT myelogram demonstrated the site of leak and helped in the management of these complicated cases. Moreover, we also found that the epicenter of the fluid collection on MRI was different from the location of the leak on a dynamic CT myelogram. Knowledge of these associations can be helpful when selecting the imaging studies to facilitate diagnosis and treatment.

Key words: Long segment; extra-arachnoid fluid collections; spine; dynamic CT myelography

Introduction

Spinal extra-arachnoid fluid collections most likely result from accumulation of cerebrospinal fluid (CSF) due to a dural leak. In most patients, the site of leak can be localized using conventional myelography with delayed CT imaging. However, in some patients with spinal CSF leaks and long spinal fluid collections, there are multiple leaks or large dural tears and the time delay during transfer between myelography and CT scan allows the extrathecal contrast to spread over multiple levels, thus making identification of their source impossible. MRI in all our cases demonstrated the spinal fluid collections but not the exact site of leak. Our study demonstrates the utility of dynamic CT myelogram in identifying the site of leak and its role in the management of these complicated cases.

Case Reports

Case 1

A 56-year-old presented with a two-year history of progressive bilateral weakness, incoordination of the lower extremities, and progressive hearing loss. Physical examination revealed bilateral lower limb dysmetria, with exaggerated lower limb deep tendon reflexes and Babinski sign bilaterally. He had decreased vibration sensation and proprioception in the lower extremities, walked with a wide-based gait, and had a positive Romberg test. In addition, he had bilateral asymmetric high-frequency sensorineural hearing loss. Cerebrospinal fluid (CSF) and cytological analyses revealed xanthochromia and hemosiderin-laden macrophages.

Typical findings of superficial siderosis (SS) were seen in the brain [Figure 1A] and the spinal cord on MRI. There was long-segment spinal cord T2 signal abnormality, which
extended from the medulla to the D₂ spinal segment, with no cord expansion [Figure 1B]. Also noted was a long-segment, nonenhancing, extra-arachnoid fluid collection along the ventral aspect of the thecal sac, extending from C₆ to the D₁₂ spinal levels, with minimal mass effect on the thecal sac [Figure 1B]. On contrast-enhanced scans there was no abnormal intramedullary enhancement, but vascular enhancement was seen along the anterior and posterior aspects of the cord and lower brainstem, in the midline. The patient subsequently underwent two brain and spine angiograms for the abnormal vascular enhancement seen on the MRI and suspicion of spinal dural arteriovenous fistula, but the investigations were normal.

Dynamic CT myelography was performed, which also showed the ventral extra arachnoid fluid collection, indicating communication between the collection and the thecal sac. There was similar contrast density in the ventral fluid collection and the thecal sac at the C₅–C₆ level on the immediate dynamic CT myelogram, suggesting site of leak [Figure 1C and D]. The patient was managed conservatively with steroids, and follow-up MRI and repeat dynamic myelography done 1 year later showed spontaneous resolution of the ventral extra arachnoid fluid collection with persistent superficial siderosis. Patient had persistent ataxia, hearing loss and myelopathy which were presumed to be due to superficial siderosis.

Case 2
A 70-year-old male reported with imbalance, ascending numbness and tingling in the lower extremities and progressive hearing loss since 1 year. Sensory exam demonstrated decreased vibration sense in the hands and feet, with decreased perception of pinprick and cold sensation below the ankle. Hoffmann and Babinski signs were present bilaterally. MRI of the brain and spine demonstrated the typical findings of SS in the brain [Figure 2A-D] and spinal cord [Figure 2E and F]. No abnormal flow voids were noted on the brain and spine MRI to suggest an arteriovenous malformation. Also demonstrated was a nonenhancing, extra-arachnoid, ventral fluid collection extending from C₃ to D₃ causing minimal indentation of the anterior thecal sac [Figure 2E and F]. Dynamic CT myelogram was done, which identified a communication between the thecal sac and the ventral fluid collection at the D₁ level [Figure 2G and H]. At surgery, there was a communication between the ventral fluid collection and the thecal sac on the right side above the right D₁ pedicle, with multiple abnormal blood vessels in the right D₁ nerve root sleeve. One of them was potentially the source of the bleeding responsible for the SS and was surgically clipped and the CSF leak was repaired. After surgery, patient developed postoperative complications with pseudomeningocele formation which required re-exploration. After the second surgery patient had stable symptoms of superficial siderosis with improvement of upper extremity signs.

Case 3
A 58-year-old lady presented with history of a motor vehicle accident followed by a 10-year history of progressive painless wasting and weakness of the left upper extremity. On examination, there was decreased bulk in the intrinsic hand and distal forearm muscles on the left. Hoffman sign and finger flexor jerk were absent on the left. MRI of the spine showed an extra-arachnoid fluid collection extending anteriorly from C₆ to D₁ spinal level, with mild indentation of the thecal sac; the fluid was seen tracking to the left posterolateral epidural space from C₇ to the D₁ levels and becoming ventral in the upper to mid thoracic spine [Figure 3A-D]. Cord T2 signal changes were noted from C₄ to D₁ along with cord atrophy [Figure 3A-D].
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A dynamic immediate CT myelogram in the left lateral decubitus showed differential opacification between the thecal sac and the collection [Figure 3E] and communication between the fluid collection and thecal sac, with asymmetric accumulation of contrast in the left C₈ perineural sheath, suggestive of a traumatic pseudomeningocele [Figure 3F]. On the delayed images there was progressively increasing accumulation of contrast within the fluid collection, with homogenous contrast opacification [Figure 3G]. Given the lack of significant cord compression, surgery was not offered to the patient.

Follow MRI showed spontaneous improvement in the size of the fluid collection with stable weakness in the left upper extremity.

Case 4
A 22-year-old man presented with history of a motor vehicle crash and was evaluated for complaints of neck pain with tingling and numbness in both the upper extremities immediately after the crash, with progression over the next two weeks. His examination revealed brisk tendon reflexes in the upper extremities, with a positive Hoffman test bilaterally. Spinal MRI showed a long-segment, nonenhancing, ventral extra-arachnoid fluid collection extending from C₂-D₉ (thick black arrows). Also noted is a thin rim of T2 hypointensity around the surface of the cord (thin white arrows) due to superficial siderosis. Sagittal immediate dynamic CT myelogram (G) shows leakage of contrast into the ventral collection at T₁-T₂ level (notched white arrow). Axial immediate dynamic CT myelogram (H) shows a lucent line in the ventral fluid collection, which may represent a fibrous band (thin white arrow). At surgery, a connection between the ventral collection and the main dural cavity was identified on the right at the T₁-T₂ level. Also identified were multiple abnormal blood vessels near the right nerve root sleeve, one of which was an acute bleeder and was the potential source of bleeding responsible for the superficial siderosis.

Figure 2 (A-H): Axial gradient-echo MRIs from inferior to superior (A-D) shows superficial siderosis around the cerebellar folia and brainstem. Sagittal T2W (E) and axial T2W (F) spine MRIs show a ventral fluid collection extending from C₂-D₉ (thick black arrows). Also noted is a thin rim of T2 hypointensity around the surface of the cord (thin white arrows) due to superficial siderosis. Sagittal immediate dynamic CT myelogram (G) shows leakage of contrast into the ventral collection at T₁-T₂ level (notched white arrow). Axial immediate dynamic CT myelogram (H) shows a lucent line in the ventral fluid collection, which may represent a fibrous band (thin white arrow). At surgery, a connection between the ventral collection and the main dural cavity was identified on the right at the T₁-T₂ level. Also identified were multiple abnormal blood vessels near the right nerve root sleeve, one of which was an acute bleeder and was the potential source of bleeding responsible for the superficial siderosis.
with placement of cysto-subarachnoid drainage catheter, which was followed by significant clinical improvement.

Case 5
A 36-year-old man presented with a 3-year history of progressive weakness of his hands—left more than right. At 19 years of age he had suffered a stab injury to the base of the neck on the right and had developed right leg weakness that had progressively improved over the next few months and returned to baseline. On examination, there was decreased bulk in the left forearm, with minimally increased tone in the right upper and right lower extremities. There was significant weakness in the left first dorsal interosseous muscles, with diminished left handgrip. He had slightly reduced vibration sense up to the level of the ankle bilaterally, slightly worse on the left side. There was bilateral sustained clonus and bilateral Hoffman sign. Spinal MRI revealed a focal area of cystic myelomalacia within the right hemicond at the D₁-D₂ level, with focal right posterolateral tethering of the cord [Figure 5A-C]. A ventral extra-arachnoid fluid collection was seen extending from C₆ through L₂ levels [Figure 5A-C]. Dynamic immediate CT myelogram demonstrated a small area of triangular contrast extravasation along the right posterolateral aspect of the thecal sac at the D₁-D₂ level and an unopacified ventral fluid collection [Figure 5D-F]). Delayed CT myelogram showed a left posterolateral fluid collection, with differential opacification of the thoracic ventral fluid collection [Figure 5G-J]. At surgery, the right posterolateral leak was identified and repaired. Follow up MRI demonstrated stable ventral fluid collections with clinical improvement of the left upper extremity weakness.

Discussion
Spinal extra-arachnoid fluid collections are CSF–filled cavities located within the spinal canal and outside the thecal sac. These fluid-filled cavities have been variably referred to as meningoceles, pseudomeningoceles, diverticula, epidural cyst, or simply as ‘fluid collections.’ These spinal extra-arachnoid fluid collections are caused by extradural CSF leaks which can be spontaneous, caused by traumatic or iatrogenic (during surgery) dural injury or traumatic avulsion of the brachial plexus. This theory is supported by the observed communication between the extradural spinal collection and the thecal sac seen at dynamic CT myelography in our patients and also noted by several authors. Differential diagnosis for these spinal collections include extradural arachnoid cysts.
Figure 4 (A-L): Sagittal T2 W (A, B) and axial T2W (C) spine MRIs show the ventral extra-arachnoid fluid collection (thin white arrows) extending from C2 down to L3 level. The thecal sac and cervical cord is displaced posteriorly by this fluid collection (thick white arrow). MR myelogram (D) also demonstrates the ventral fluid collection (thick white arrows). Sagittal immediate dynamic CT myelogram (E) shows opacification of the ventral fluid collection, with similar density of contrast in ventral fluid collection and the thecal sac at C2–C5 and C5–C6 disc levels (white arrows). Axial immediate dynamic CT myelogram at C3–C4 (F) disc level shows contrast density in the thecal sac and the ventral fluid collection to be similar (thick black arrows), suggesting site of leak. Axial immediate dynamic CT myelogram at C6–C7 (G) disc level shows differential opacification of the ventral fluid collection and the thecal sac (thick white arrows). In addition, pseudomeningoceles were identified on the left at multiple levels, more prominent at C6–C7 (G) and C7–T1 (H) levels (notched arrows). The nerve root sleeves on the right appear to be separated from the collection, with the differential contrast opacity within the right-sided nerve root sleeves (arrowheads); however, the left-sided nerve root sleeves (thick white arrow) appear to be incorporated within the ventral fluid collection. Also, note the effacement of the CSF space surrounding the cord due to extrinsic compression by the ventral fluid collection (G and H). The epicenter of the compression (C5–C6) is lower than the site of leak (C2–C3). Sagittal oblique reconstructions (I) of early dynamic CT myelogram shows pseudomeningoceles at multiple levels (white arrows) on the left in the cervical spine. Sagittal delayed dynamic CT myelogram (J) shows inferior extension of the ventral fluid collection into the thoracic region (white arrows). Axial immediate (K) and axial delayed (L) images from dynamic CT myelogram of the thoracic spine show progressive equilibration of contrast in the ventral fluid collection in the thoracic region on the delayed scans (L) (thick black arrows).
Spinal extra-arachnoid fluid collections may present with symptoms related to compressive myelopathy from the collection, SS, or intracranial hypotension. The pathogenesis is usually considered to be an occult CSF leak through small defects in the meninges. Pulsatile CSF dynamics,\textsuperscript{[11]} presence of an osmotic gradient between the subarachnoid space and cyst,\textsuperscript{[12]} and valve-like mechanism between the fluid collection and subarachnoid space\textsuperscript{[13]} may play important roles in the enlargement of these spinal extra-arachnoid fluid collections, which can then cause cord compression and cord signal changes.

In patients with spinal fluid collections, it is essential to not only demonstrate the leak but also the site and morphology, because treatment strategy differs dependent on these characteristics. The goal of imaging is to diagnose the extra-arachnoid fluid collections and pseudomeningocele and also, specifically, to identify the dural defect; this is best characterized by MRI, MR myelography CT myelography,\textsuperscript{[2,14-16]} and digital subtraction myelography.\textsuperscript{[2]} 

All our cases demonstrated long-segment nonenhancing extra-arachnoid spinal fluid collections that were isointense to CSF on T1W and T2W images. Spinal MRI demonstrated the spinal fluid collections but did not demonstrate the actual site of CSF leak.

In most patients, the site of leak can be localized by using conventional myelography, with delayed CT scans for slower leaks. However, when there are multiple leaks or large dural tears, the time delay during transfer between the myelographic portion of the examination and CT scan allows the extrathecal contrast to spread over multiple levels, thus limiting the ability to localize the leaks.\textsuperscript{[11]} The exact site of dural defect in intraspinal collections spanning long segments can be accurately localized with dynamic CT myelography,\textsuperscript{[16,17]} which was first introduced for localizing high-flow CSF leaks.\textsuperscript{[11]}

The technique of dynamic CT myelography involves CT acquisition during the initial introduction of the myelographic contrast media into the thecal sac. Specifically, our technique involves puncturing the thecal sac in the lumbar region on the CT Table with the patient in the prone position. A short-segment CT scan over the lumbar segments confirms accurate intrathecal positioning of the needle (scanning the entire spine before injection of contrast is usually not necessary and also increases the radiation dose of the examination). After injection of 10-15 ml of myelographic contrast media, the patient is mobilized to...
displace the contrast into the spinal segment of interest, e.g., by having the patient adopt a Trendelenburg position to displace the contrast column into the thoracic and cervical spine. The immediate CT acquisition is obtained with the collection in the dependent position. A second delayed acquisition can be obtained after asking the patient to rotate 360°.

Dynamic CT myelogram in all of our cases showed contrast opacification of the extra-arachnoid fluid collection, confirming free communication with the subarachnoid space. Early dynamic CT myelography demonstrated the site of leak where there was similar contrast density within the thecal sac and the fluid collections. All patients had long-segment extra-arachnoid fluid collections, with the epicenter of the collection at a site different from the site of the dural leak.

We understand that dynamic CT myelography is more invasive, has higher radiation dose and is more time intensive. We thus, do not advocate routine use of this technique for all patients with spinal fluid collections.[11] We reserve this technique for a subset of patients with long segment spinal fluid collections with presumed high-flow spinal CSF leaks.

Spinal extra-arachnoid pseudomeningoceles are rare and can be an unrecognized source of chronic bleeding into the subarachnoid space, leading to SS.[14] SS occurs due to chronic or repeated slow hemorrhage into the subarachnoid space that leads to hemosiderin deposition over the cerebral and spinal leptomeninges.[19] Etiologies for SS include root avulsions, tumors, vascular malformations, and brain or spine surgery.[20] In a 1995 survey of the reported SS cases in the literature, a dural pathology was found in 47% of cases.[20] This included CSF cavity lesions (e.g., meningoceles, pseudomeningoceles, pseudoencephaloceles, cavity remaining after a hemispherectomy, and chronic suboccipital hematomas), or root pathology (e.g., root avulsions or epidural cysts).[20] Brachial plexus injury with associated nerve root avulsion is also a common association.[17,21-24]

The association of a spinal fluid collection that communicates with the subarachnoid space in patients with SS and history of trauma has been recently recognized by Kumar et al.,[16] and the authors stress the importance of looking for such a communication using dynamic CT-myelogram. In our series, two patients presented with SS, and in one of them extensive brain and spine angiograms did not reveal spinal arteriovenous malformations. It is important to remember that in patients with SS with long-segment extra-arachnoid fluid collections, the bleeding source is usually in the spine, and dynamic CT myelogram helps to locate the exact site of the leak, which then helps in surgical planning.

Conclusions

Our cases illustrate a complex, poorly understood, and difficult-to-clinically-manage entity, which on imaging is represented by extra-arachnoid fluid collections related to occult dural tear and CSF leak. The exact location of the leak is best identified using dynamic CT myelography, and accurate localization helps in planning surgical treatment.

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