

Extradural arachnoid cyst with bony erosion: a rare case report

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Abstract: The authors present the case of an otherwise healthy 38-year-old female with an atypical extradural arachnoid cyst with multi-level involvement in the lumbar spine leading to left quadriceps weakness and dysesthesia. Upon presentation, a lumbar spine MRI with contrast and plain radiographs revealed extensive L4 bony erosion. An MR angiogram and cervical spine MRI with contrast were then obtained in order to rule out any aortic root or cervical spine pathology. With no other apparent clinically relevant pathology revealed by these additional tests, an L3-5 posterior decompression and fusion procedure was performed. Her preoperative symptoms were successfully resolved following the procedure, with no resultant surgical complications. The cyst is atypical not only due to its size and location, but also due of the significant bony erosion of the left L4 pedicle and vertebral body. To the authors' knowledge, this is the first reported case of an extradural arachnoid cyst in the lumbar spine with bony erosion of the pedicle and vertebral body. In cases such as this, a CT myelogram may be useful in planning the operative approach through visualization of the exact communication between cyst and dura. This approach may also aid in diagnosing and identifying atypical cyst presentations such as the one presented here.

Keywords: Arachnoid; cyst; lumbar vertebrae; case report

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Introduction

Arachnoid cysts are collections of cerebrospinal fluid lined by the arachnoid mater within the cranium and spine. A minority of arachnoid cysts occur within the spine (1,2) and most are asymptomatic, not requiring intervention (3,4). Spinal arachnoid cysts tend to occur within the thoracic and cervical regions, with lumbar arachnoid cysts rarely described in the literature (1,5-7). These are usually small in size (8). It is theorized that a valve like communication with the sub-arachnoid space may be an important factor in their enlargement (9,10). They usually present in patients 30-50 years old and can be extradural, intradural, or perineural (11). Their etiology is proposed to be congenital, traumatic, iatrogenic, and infectious (12,13). For specific cases, the pathogenesis is typically undetermined (14). Surgical intervention is rarely required (5). Indications for surgery are neurologic impairment due to mass effect and

asymptomatic cyst expansion.

The authors report a case of an unusually large, symptomatic, extradural lumbar arachnoid cyst causing significant bony erosion of the pedicle and vertebral body in an otherwise healthy 38-year-old female. This is the first report detailing a cyst with the above-mentioned characteristics. We present the following article/case in accordance with the CARE reporting checklist (available at http://dx.doi.org/10.21037/jss-20-590).

Case presentation

A 38-year-old woman presented with 5 years of progressive weakness in her left leg (grade 4/5 quadriceps) with dysesthesias extending from her thigh to her left foot. Her symptoms had grown noticeably worse in the immediately preceding 6 months. Her medical history included

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appendectomy, cholecystectomy, cleft palate repair, gastric bypass, and thyroid disease. She had no history of trauma, infection, previous spine conditions, connective tissue disease, neurofibromatosis, or other cyst formation. In addition, the patient never had an epidural catheter placed, epidural or spinal injection, or CT myelogram performed.

A lumbar MRI without contrast revealed a large intraspinal fluid collection extending from L3 to L5, causing severe cauda equina compression with near complete erosion of the left L4 pedicle and the posterior superior portion of the L4 vertebral body (*Figure 1*). Plain radiographs demonstrate a benign appearing spine on lateral imaging, with a loss of the left L4 pedicle on the AP view (*Figure 1*).

With the atypical imaging findings suggestive of dural ectasia, an MR angiogram and cervical MRI were performed to rule out disorders of the aorta and cervical spine. The aortic root appeared healthy (*Figure 1*). The cervical MRI revealed borderline Chiari malformation with no other significant cervical pathology (*Figure 1*).

Due to the likelihood for further worsening of symptoms, surgical intervention was recommended. The goals of surgery were to decompress the cauda equina and to preserve spinal stability. An L3–L5 posterior decompression and fusion procedure was thus recommended. A complete timeline of this case report can be found in *Figure 2*. All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee(s) and with the Helsinki Declaration (as revised in 2013). Written informed consent was obtained from the patient.

Surgical approach

A CSF drain was placed percutaneously at the L1–2 level prior to spine exposure.

Following a posterior midline exposure, pedicle screws were placed bilaterally at L3, on the right at L4, and bilaterally at L5. Titanium rods were placed with mild distraction to provide indirect neurologic compression and stabilization of the segment. A laminectomy of L3 and L4 was performed. A posterior fusion procedure was indicated due to the concern about potential for iatrogenic instability after a wide decompression in the presence of an incompetent L4 pedicle/facet complex

Once the cyst was visualized, a plane was developed between the thin-walled cyst and the dura using a blunt nerve probe. The distal end of the cyst was fully mobilized relative to the dura and affixed with a suture to aid in elevation of the cyst from distal to proximal (*Figure 3*). During the final stages of dissection, the site of communication between the cyst and the intra-dural space was encountered. Separation of the cyst from the dural stalk led to the abrupt deflation of the cyst and partial deflation of the dura. A nerve root was observed to herniate through the 2mm wide by 5mm long dura stalk opening and obstruct further CSF leakage. (*Figure 3*). The dural stalk opening was midline at the level of the L3 foramen. The defect was repaired with three 4-0 Neurolon sutures, with subsequent fibrin glue placement (*Figure 3*).

Operative time was 3 hours 19 minutes. Estimated blood loss was 200 ml. The patient tolerated the procedure well without complications. Due to concern about the potential for CSF leakage, the patient was kept on bedrest for 48 hours as the CSF drain was adjusted to produce 10ml per hour. After two days, the CSF drain was stopped, and the patient allowed to ambulate. Absent any symptoms or signs of CSF leakage, the drain was removed, and the patient was discharged to home on postoperative day 4.

The excised cyst was sent to pathology due to its unusual presentation and was determined to be consistent with an arachnoid cyst with immunostaining for EMA highlighting the cyst lining meningothelial cells.

The patient experienced complete resolution of her preoperative radiculopathy and returned to normal function without activity restrictions by two months after surgery (*Figure 4*).

Discussion

The authors present the case of an extraordinarily large arachnoid cyst that is atypical due to its location, size, dramatic bony erosion, and lack of discernable precipitating factors. Extradural arachnoid cysts typically occur in the thoracic region, with rare occurrences found in the lumbar spine. Lumbosacral cysts account for a small minority of identified spinal cysts, with two centers reporting that over a 7- and 10-year period, 81% and 77% of all spinal cysts identified were located in the thoracic region respectively (2,5,6). In addition, out of 20 patients who underwent surgical treatment for spinal cysts over a four-year period at another institution, there were no reported cases of lumbar arachnoid cysts (1).

While arachnoid cyst size may vary based on pathogenesis and the underlying mechanism of formation, most are relatively small and localized (10,15). It has been



Figure 1 Preoperative sagittal midline (A) and axial L4 (B) MRI images reveal large cystic lesion with the as compression and near complete erosion of the left L4 pedicle. Preoperative lateral (C) and anteroposterior (D) radiograph reveal healthy appearing spinal anatomy excepting the atrophic left L4 pedicle. Preoperative sagittal midline cervical MRI (E) shows the cerebellar tonsils to extent ~1 cm below the foramen magnum. Preoperative MR angiogram (F) reveals healthy appearing aortic root.



Figure 2 The patient's complete course of treatment, including dates of additional testing and clinic visits. Initial symptoms: 5 years of left leg weakness (grade 4/5 quadriceps) with accompanying dysesthesias markedly worsening over the immediately preceding 6 months. Initial consult: lumbar MRI and plain radiographs reviewed with patient upon presentation to clinic revealing an intraspinal fluid collection and L4 pedicle and vertebral body erosion. Additional imaging: MR angiogram and cervical MRI performed revealing no clinically significant abnormalities. Operation: 3-hour and 1-minute L3-5 posterior decompression and fusion procedure performed with no surgical complications. Postoperative visit: patient demonstrated complete resolution of symptoms at the 2-month postoperative visit with no further follow-up needed. No adverse events were noted during this period.



Figure 3 (A) Exposed cyst (A1) and healthy dura (A2). (B) Distal cyst tie-off. (C) Cyst removal: adhesions between cyst and healthy dura. (D) Deflated cyst (D1) and healthy dura (D2). (E) Dural defect with nerve root protrusion. (F) Dural defect repair.

hypothesized that a ball-valve mechanism of formation, amongst others, may contribute to the enlargement in select cysts (5,6,9). The findings in this case of: a discrete connection between the dura and the cyst, the visualization of a nerve root transiently blocking CSF leakage from the thecal sac, the large cyst size, and bony erosion all support the one-way valve theory. To the authors' knowledge, the currently described case is the first extradural arachnoid cyst in the lumbar spine with subsequent bony erosion of the pedicle and vertebral body in the literature. While there are a few reports of bony erosion of the cranium with a variety of cysts, there is only one reported case of a lumbar arachnoid cyst with subsequent bony erosion; however, this was an intradural



Figure 4 Standing anteroposterior (A) and lateral (B) radiographs.

cyst with mainly vertebral body erosion (5). This was determined after a comprehensive review of the English medical literature using terms: arachnoid cyst, erosion, pedicle, vertebral body, lumbar, thoracic, contiguous, and all possible permutations and combinations of said terms. We theorize that the bony erosion resulted from the cyst expansion secondary to the one-way valve mechanism of communication. The concept is that a nerve root or dural flap located at the site of the connection between the dura and the cyst could allow egress of CSF into the cyst during times of peak intrathecal pressure, but not allow reversion of the pressure gradient. It is the theorized heightened pressure within the extra-dural cyst causes the cyst's expansion, resultant dural compression, and eventual bony erosion.

The cause of the arachnoid cyst in this case remains undetermined. She had no known dural trauma. Although the preoperative MRI shows her cerebellar tonsils to extend approximately 1 cm below the foramen magnum, she has ample space for CSF flow and an absence of typical Chiari malformation symptoms. An association between Chiari malformation and lumbar arachnoid cyst formation has not been previously described. The authors are not proposing such a connection between the Chiari malformation and arachnoid cyst in this case.

Due to the apparent bony erosion and multi-level contiguous involvement, we considered various differential diagnosis. This included dural ectasia among others. Dural ectasia is typically associated with neurofibromatosis or connective tissue disorders and has radiographic findings of dural expansion and bony erosion. The patient's lack of signs/symptoms of connective tissue disorders or neurofibromatosis, however, made such seem less likely. Concern regarding dural ectasia as the underlying disorder prompted the ordering of a thoracic MRI to rule out aortic root pathology and the existence of other foci of dural pathology within the thoracic spine.

Arachnoid cyst was also on our preoperative differential diagnosis. Such was considered less likely; however, as an extensive literature search did not identify any cases of an arachnoid cyst with bony erosion or multi-level contiguous involvement as was present in the current case. Additionally, while arachnoid cysts are a common incidental finding on intracranial imaging, their prevalence in the spine, and specifically the lumbar spine, is very rare (14,16).

This case represents a valuable addition to the literature on the abnormal presentation of arachnoid cysts, as well as to their potential methods of communication with the subarachnoid space. The current procedure involved a multilevel decompression and fusion, due to the size of the cyst, and concern for spine instability secondary to the bony erosion of the left L4 pedicle and the anticipated need for a wide decompression. Had we found more evidence in the literature supporting an arachnoid cyst presentation similar to our case, we may have rationalized the use of a preoperative myelogram CT to elicit additional information about the character of the dura-compression process, including the location of the cyst communication, without the need of the MR-angiogram or cervical spine MRI. Mishra, et al. 2017 describe the potential benefit of dynamic intraoperative MR myelography in determining the location cyst/dura communication (17). One can speculate that for the current case, were the nature of the cyst known and its stalk location well understood, a minimally invasive and focal stalk ligation and dural repair procedure might have been possible with avoidance of instrumented arthrodesis.

Conclusions

The authors report an unusually large, lumbar extradural arachnoid cyst in an otherwise healthy 38-year-old woman. This case has unique combined attributes that have not previously been reported with an arachnoid cyst: significant pedicle erosion, multi-level involvement, and location in the lumbar spine. The authors propose at CT-myelogram to be potentially useful in determining the precise location of communication between the cyst and dura. This case demonstrates the need for careful preoperative planning, and consideration for atypical presentations of cystic lesions.

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Footnote

Reporting Checklist: The authors have completed the CARE reporting checklist. Available at http://dx.doi.org/10.21037/jss-20-590

Conflicts of Interest: Both authors have completed the ICMJE uniform disclosure form (available at http://dx.doi. org/10.21037/jss-20-590). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee(s) and with the Helsinki Declaration (as revised in 2013). Written informed consent was obtained from the patient for publication of the Case report and any accompanying images.

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