Introduction

We present an interesting case of a 41-year-old lady who works as an administrative assistant in a local high school.

She reports that while lifting heavy boxes some six months ago, she developed sudden and severe interscapular pain. Her symptoms had persisted and worsened over the period in spite of the best efforts of her local doctors. She denied any pain or weakness in any of her limbs. Her past medical history was unremarkable for any trauma or significant medical conditions. Clinically, she was in severe pain, and walked in a stooped fashion. She had exquisite tenderness in the upper dorsal spine area, but no deformity. Neurological examination was unremarkable.

X-rays of the dorsal spine were negative; however, a magnetic resonance imaging (MRI) scan of the area demonstrated the presence of two distinct, fluid filled dorsal extradural lesions at the T3 and T4 level (Figures 1 and 2).

The differential diagnoses being considered by the radiologists included parasitic disease, neoplasms and infections, none of which seemed to tally with the history as given by the patient.

In theatre, two distinct masses containing cerebrospinal fluid (CSF) were found, enclosed within a fibrous capsule. On opening the capsules, there was CSF egress and dural defects were visible at both sites which were repaired in the standard fashion. There were no neural elements herniating into the sacs, and the walls were sent for histology which demonstrated arachnoid enmeshed in a fibrous capsule.

A diagnosis of thoracic pseudomeningocoele was hence established.

Post-operatively she did very well indeed with immediate resolution of her pain.

Discussion

Pseudomeningocoeles are so called to differentiate them from true meningoceles.

In the former, CSF escapes from the subarachnoid space to lie in the extradural space, and in time becomes covered by a false capsule or in some instances an intact arachnoid layer, while in the latter all layers of the meninges are present in the sac.

Pseudomeningocoeles usually form in the setting of trauma or as a complication of intentional or accidental durotomy at surgery, while true meningoceles occur in the setting of spinal dysraphism.

It is reported that the diagnosis of pseudomeningocoeles is not always straightforward and a number of them are asymptomatic.
A history of spinal trauma or else surgery to the spine is helpful in increasing the index of suspicion for the condition, and this should always be considered in the appropriate clinical setting.1,2,4-7

Where symptoms do occur, these have been ascribed to the sheer size that some of these lesions can achieve, or else neurological deficits occasioned by herniation into or compression of nerve roots and the spinal cord into the sac.2,4-7

The management of symptomatic pseudomeningocoeles of the spine is generally by direct exposure of the lesion, removal of the cyst wall and repair of the dural defect.1,2 Some large ones have been managed indirectly by CSF diversion shunts,8 while there are reports of spontaneous resolution of pseudomeningocoeles.8,9

This case report is unique in that there is hardly any mention in the literature of this condition occurring in the setting of physical exertion. While sudden spinal pain following physical exertion is common in clinical practice, it is mainly encountered in the lumbar spine and where sprains and strains have been excluded, it is usually due to annular tears of intervertebral discs or else, in a few cases, disc prolapses or fractures of osteoporotic vertebra in our experience. While pseudomeningocoeles have been reported to occur in conditions such as neurofibromatosis and Marfan’s syndrome, in fact the majority of these have been found to be true meningoceles.10 This lady had no such conditions.

There are case reports in the literature of intracranial hypotension occurring in the setting of spontaneous spinal CSF leaks. In most cases, the patients are females, and present with orthostatic headaches, as well as nausea, vomiting and even abducens cranial nerve palsies.11,12 It is interesting to note that this patient did not have any of these symptoms.

This publication alerts spine surgeons as to this rare condition, which should be included in the differential diagnosis of acute and severe spinal pain following physical exertion.

As demonstrated in our case report, while the pain and discomfort were excruciating, the management was relatively straightforward and the outcome rewarding for patient and surgeon alike.

**Teaching points**

1. Spinal surgeons must be aware of spinal pseudomeningocoeles presenting in unusual fashions and should have a high index of suspicion for these lesions in appropriate circumstances.
2. Careful consideration of investigations such as MRI scans is essential for timeous diagnosis of pathology.
3. This condition lends itself to effective treatment once diagnosed correctly.

**REFERENCES**

4. Johnson DB, McGrath FP. Case report: post-traumatic thoracolumbar pseudomeningocoele—an unusual cause of upper

---

**Figures 1 and 2: Sagittal T1 and T2 Magnetic resonance imaging (MRI) scans respectively showing dorsal, cyst like lesions at the T3 and T4 levels**

---

VOL. 56 NO. 3 SEPTEMBER 2018 SAJS 37


