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Thoracolumbar surgery for degenerative spine diseases complicated with tethered cord syndrome: A case report

Yue-Tian Wang, Guan-Zhang Mu, Hao-Lin Sun

BACKGROUND
Tethered cord syndrome (TCS) secondary to split cord malformation (SCM) is rare in adulthood. There is as yet no consensus about the optimal treatment method for adult patients with SCMs and degenerative spine diseases such as lumbar stenosis, spondylolisthesis and ossification of the ligamentum flavum (OLF). The tethered cord poses a great challenge to the decompression and fusion procedures for the intraoperative stretching of the spinal cord, which might lead to deteriorated neural deficits. Here, we report on a case to add our treatment experience to the medical literature.

CASE SUMMARY
We treated a 67-year-old female patient with type II SCM suffering from lumbar disc herniation, degenerative lumbar spondylolisthesis and thoracic OLF. The patient underwent thoracolumbar spinal fusion and decompression surgery for severe lower back pain, extensive left lower limb muscle weakness and intermittent claudication. After the thoracolumbar surgery, without stretching the tethered cord, the patient achieved complete relief of pain and lower extremity weakness at final follow-up.

CONCLUSION
For adult patients with underlying TCS secondary to SCM coupled with thoracic OLF and lumbar spondylolisthesis, a thoracolumbar fusion surgery could be safe and effective with the tethered cord untreated. It is critical to design individualized surgical protocols to reduce the stretch of the low-lying spinal cord.

Key Words: Tethered cord syndrome; Split cord malformations; Ossification of ligamentum flavum; Spondylolisthesis; Thoracolumbar surgery; Case report

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Core Tip: Tethered cord syndrome (TCS) secondary to split cord malformation (SCM) is rare in adulthood. We present a patient who underwent thoracolumbar surgery for thoracic ossification of the ligamentum flavum and lumbar spondylolisthesis complicated with TCS. A thoracolumbar fusion surgery could be safe and effective with the tethered cord untreated. It is critical to design individualized surgical protocols to reduce the stretch of the low-lying spinal cord. A literature review of SCM in adults was also performed.

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INTRODUCTION
Split cord malformation (SCM), one of the most common intraspinal malformations, is a rare disease of the spinal cord and cauda equina caused by embryonic dysplasia[1,2]. SCM refers to a spinal cord divided longitudinally into two distinct hemicords that later rejoin[1,3], which has been categorized into two types: Type I SCM refers to two hemicords, with their own dural tubes and separated by a dural-sheathed rigid osteocartilaginous median septum, and type II SCM also refers to two hemicords but housed in a single dural tube separated by a nonrigid, fibrous median septum[4]. It has been demonstrated that the presence of a bony fibrous septum anchoring the cord interferes with its normal upward ascension during growth, resulting in tethered cord syndrome (TCS)[1,3]. The typical symptoms might result from stretch-induced ischemia by traction on the cord.

SCM is commonly presented during childhood but rarely diagnosed in adults[2]. Moreover, SCM is more likely to be diagnosed in females[1,5]. Neurologic symptoms of type I SCM can progressively deteriorate if left untreated[6]. Unidentified adult SCM might also result in various degenerative spinal diseases with age, and the tethered spinal cord could pose a great challenge to the spinal fusion surgery[7]. However, there is not enough experience on spine diseases complicated by TCS secondary to SCM.

In the present study we report an adult-onset SCM patient of type II suffering from degenerative lumbar spondylolisthesis and thoracic ossification of the ligamentum flavum (OLF) who needed a spinal fusion surgery for extensive left lower limb muscle weakness.

CASE PRESENTATION
Chief complaints
A 67-year-old female was hospitalized with severe low back pain with radiating pain in the left lower limb for 5 years. Symptoms worsened in the last 1 wk. She had exhausted conservative treatments and intended to proceed with surgery.

History of present illness
Five years before hospitalization, the patient began to experience aching pain in the waist, accompanied by pain and numbness in the left lower limb ranging from the posterior thigh and the posterolateral crus to the dorsolateral foot, especially obvious at the 4th and 5th toes. There was no weakness of lower limbs and no urination difficulty. The patient was not able to walk more than 100 m without rest. The uncomfortable symptoms often occurred after overwalking and catching a cold, but could be relieved via physical therapy and NSAIDs. One week before hospitalization, the above symptoms were significantly aggravated, accompanied by progressively extensive weakness of the left lower extremity and with conservative treatment giving unsatisfactory results. Lumbar computed tomography (CT) scan demonstrated L4/5 intervertebral disc prolapse and lumbar X-ray imaging showed grade I L4 anterolisthesis from another hospital.

History of past illness
The patient was diagnosed with atrial septal defect five years before hospitalization and treated with repair surgery. She underwent operation because of lipoma located on the left trunk during the same year. She denied history of hypertension or diabetes. No history of trauma or malignant tumors were identified.

Personal and family history
There was no special history or personal history. The patient was unaware of SCM before and denied family history of SCM.
Physical examination
Examination showed normal curvature of the lumbar spine without scoliosis deformity and no foot abnormality. There was a round skin sag of 1.5 cm in diameter located in the sacrococcygeal region with chromatosis but no hair. Slight tenderness and percussion pain in the paraspinal muscles were found. The left lower extremity had slight hypalgesia and hypopselaphesia. Waist activities in different directions were somewhat limited because of pain, especially in extension. There was extensive weakness in the left lower limb, with hip flexors strength graded IV, knee flexion muscle strength graded III+, knee extension strength graded III and foot dorsiflexion strength graded III. The Lasegue sign was positive in the left lower limb. Bilateral tendon reflexes showed suspicious hyperactivity with lower limbs’ muscle tension slightly increased.

Her baseline severity of low back pain and left lower limb pain was 90 mm and 90 mm, respectively, on a 100-mm visual analog scale (VAS) when she was admitted into our hospital. We used the Oswestry Disability Index (ODI) to evaluate lumbar function and the score was 64.

Laboratory examinations
The routine blood and blood biochemical parameters of the patient were within normal limits.

Imaging examinations
Anteroposterior and lateral X-ray imaging showed a grade I L4 spondylolisthesis and the flexion-extension X-ray imaging demonstrated instability at that level. Lumbar CT scan showed the L2-S1 intervertebral disc was swollen with the dura sac compressed to varying degrees, in addition to the L4 spondylolisthesis. Thoracic CT scan showed that the left part of the ligamentum flavum was thickened and ossified at the T11-12 Level with the posterior dura sac obviously compressed and that there were block vertebrae (T8/T9) and a butterfly vertebra (T9). Lumbar magnetic resonance imaging (MRI) showed L4 spondylolisthesis and the L4-5 intervertebral disc bulge compressing bilateral nerve roots, which was more serious on the left side. Additionally, SCM was observed; the spinal was split into two hemicords from the lower aspect of T12 to the upper aspect of L2, accompanied by a low-lying conus terminating at S1 with a thickened terminal filament deposited by fatty tissue (Figure 1).

FINAL DIAGNOSIS
The clinical diagnosis was T11-12 thoracic OLF, L4 Lumbar spondylolisthesis (grade I) and TCS.

TREATMENT
Considering there was a tethered cord due to the silent SCM, neurosurgery was requested to evaluate the feasibility of concurrent transection of the filum terminale during the spine surgery. However, this was not recommended because simultaneous operation for the tethered cord required opening of the dura sac, which could lead to cerebrospinal fluid extravasation and interfere with decompression and fusion manipulations. After identifying the lesions responsible for neural symptoms, we decided to perform a thoracolumbar combined surgery to treat the thoracic OLF and lumbar spondylolisthesis but not the TCS over the same period. Posterior thoracic canal decompression through laminectomy followed by osification removal with pedicle screw fixation (T11-12) was conducted for the thoracic spinal stenosis resulting from the thoracic OLF. For the lumbar spinal stenosis due to the L4-5 intervertebral disc bulge and segmental instability caused by L4 spondylolisthesis, an L4-5 midline lumbar fusion (MIDLF) procedure was performed simultaneously to pursue bilateral L5 nerve root and spinal canal decompression, cortical bone trajectory screw fixation and intervertebral and posterior-lateral fusion. Given there was a preoperatively existing extensive decrease in left lower limb muscle strength, the intraoperative interference of the tethered cord would carry a great risk of paralysis, posing a substantial challenge for the surgeon to conduct the thoracolumbar combined operation. As a result, we adopted neural electrophysiological monitoring during the whole operation, which showed good sensory and motor conduction before and after decompression of the spinal cord (Table 1). The operation duration was 315 min and the estimated blood loss was 500 mL.

The principle of the combined operation was that the thoracic spinal decompression and pedicle screw fixation was taken as the prior task and then the bilateral compression to the T11-12 fixation was carried out to relieve and shorten the strained spinal cord for providing compensatory space. The L4-5 MIDLF was performed to avoid excessive opening of the intervertebral space, so as to reduce the stretch of the dura sac and the spinal cord. The patient's lower limbs moved well after awakening from anesthesia.
Wang YT et al. Spinal diseases complicated with tethered cord syndrome

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<td>Amplitude in μV</td>
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<td>50.3</td>
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<td>41.5</td>
<td>49.8</td>
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<tr>
<td>After lumbar decompression</td>
<td>38.8</td>
<td>49.5</td>
<td>0.6</td>
<td>38.5</td>
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</table>

The electrical stimulation value was 400 mV. MEPs: Motor evoked potentials; SEPs: Somatosensory evoked potentials.

OUTCOME AND FOLLOW-UP

The patient's symptoms were significantly alleviated postoperatively. There were no operation-related complications after surgery. By the time she was discharged, her low back pain VAS score had dropped to 3 and her leg pain VAS score to 2. She could ambulate well after leaving the bed with left lower limb muscle strength enhancement. The MRI at 1 mo after surgery showed fine decompression of the spinal canal (Figure 2). Up to the final follow-up (3 mo after surgery), her low back pain VAS score had dropped to 2 and her leg pain VAS score to 1; the ODI had dropped to 26; and the lumbar CT scan showed the intervertebral fusion status and implant position were fine (Figure 2).

DISCUSSION

SCM is described as a congenital spinal dysraphism with often a bone spur and a membranous or fibrous septum resulting in two split hemicords, with a single or separated dural layer surrounding it[4, 5,8]. SCM is often noted due to the appearance of scoliosis, skin stigmas, progressive foot deformities, calf and foot atrophy and bowel or bladder disturbances in childhood. As a result, this pathologic phenomenon is most frequently seen in childhood but rarely presented in the adult population[1,3,9]. D'Agostino et al[3] reviewed SCM in adults and from 1936 to 2018, only 25 cases of concurrent split and radiographic tethered cords were identified. Patients averaged 37 years of age at the time of diagnosis and 56% were female. A recent review summarized 146 adult SCM patients diagnosed at a mean age of 26.8 years, of which 74.6% were female[8]. SCM is often accompanied by vertebral anomalies, spina bifida occulta being the most common[9]. The female patient we presented was 67-years-old and she was not diagnosed with SCM until she was admitted to our hospital. The CT scan revealed that there were T8 and T9 block vertebrae with a T9 butterfly vertebra.

Adult SCM can occur at any level along the spine but is more common in the lumbar region, followed by lumbosacral segments and thoracic regions[2,9-11]. According to the number of dural tubes and the characteristics of the median septum, Pang et al[4] classified SCM into two types, with type I referring to
Figure 1 Preoperative radiography. A: Preoperative anteroposterior and lateral radiographs; B: Preoperative flexion–extension radiographs; C: Preoperative computed tomography scan showing block vertebrae at T8/T9 (orange arrow), butterfly vertebra at T9 (blue arrow); D: Preoperative magnetic resonance imaging showing thoracic ossification of the ligamentum flavum (red box), a split cord malformation (blue outline) with a low-lying conus and a thickened terminal filament deposited with fatty tissue, and bilateral nerve root compression at L4-5.

two hemicords, each in a separate dura sac separated by a rigid osseocartilaginous median septum, and type II involving two hemicords in a single dural tube with a nonrigid septum. A low-lying cord is usually associated with this dysraphism because the septum could prevent the spinal cord from moving upward, leading to excess strain on the spinal cord, which results in TCS. It appears that type II SCM lesions are more likely to tether than type I lesions though more data are needed to confirm this[3,5]. TCS is rarely secondary to SCM in adulthood. SCM was reported to account for 10%-38% of adult TCS diagnoses[3]. Studies revealed that adult-onset TCS is usually associated with precipitating events such as stretching of the conus medullaris, narrowing of the spinal canal or trauma[12].

The clinical presentation of SCM is variable. The most common symptom is back pain associated with the level of pathology, and radiculopathy and lower extremity weakness were also reported as common manifestations with sometimes bowel or bladder disturbance[1,13-16]. Moreover, some cases of SCM remained asymptomatic or only caused subtle symptoms until there was a factor that stretched the tethered spinal cord. The factors might be various degenerative spine diseases such as lumbar/thoracic disc herniation, spondylolisthesis, lumbar/thoracic spinal stenosis or scoliosis[17,18]. Physicians should be cautious when recommending or giving prophylactic surgery for asymptomatic SCM. Goldberg et al [19] reported 28 patients who underwent prophylactic operations of split cords. Of these patients, 10 patients had reduced lower extremity strength after treatment. For symptomatic SCM, such as pain and lower extremity weakness, if no other responsible lesions such as lumbar disc herniation or spondylolisthesis were identified on imaging, surgery for SCM or TCS such as removal of the bony diaphragm or cutting the filum terminale tended to result in good clinical outcomes[20-22].

Refractory pain and neurologic deficits in adult patients with TCS usually implicate that there might be stenosis or compression factors and that surgery is required[5]. We reviewed the reported cases of adult TCS coupled with degenerative spine abnormalities that remained asymptomatic until the compression or instability required treatment. The characteristics of these cases are summarized in the Table 2. It seems that operation on these patients with TCS was challenging because it could interfere with the existing balancing of the tethered spinal cord, causing paralysis or neurological deficits. The main principle of this kind of operation was to minimize the stretching of the spinal cord. Some surgeons argued to untether the spinal cord by filamentectomy or resection of the bony spur before treating the compression factors[5,13]. The simultaneous removal of bony spur while treating the degenerative spine lesions could be challenging and may be more suitable for type I SCM, because there is a high risk of complication of the opening of the dura sac, causing cerebrospinal fluid leakage or infection[9,13,23]. In addition, some surgeons resorted to minimally invasive procedures, such as endoscopic surgery, to remove the lesions or decompress the spinal canal in order to decrease the disturbance of the unreleasing low-lying cord[15,24]. A few cautious surgeons performed operations
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<td>Breton et al [15], 2020</td>
<td>United States</td>
<td>Case report 1</td>
<td>Female/79</td>
<td>NM</td>
<td>Leg pain; progressive gait deterioration and bilateral leg weakness</td>
<td>Lumbar spinal stenosis; spondylolisthesis</td>
<td>Sublaminoplasty for spinal cord decompression with onlay arthrodesis</td>
<td>Conduct a minimally invasive surgery with tethered cord untreated</td>
<td>NM</td>
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<td>Hui et al [26], 2014</td>
<td>China</td>
<td>Case report 1</td>
<td>Male/23</td>
<td>Type I</td>
<td>Unstable walking and progressed numbness in the lower limbs</td>
<td>Kyphoscoliosis</td>
<td>Posterior segmental pedicle screw instrumented fusion with vertebral column resection</td>
<td>Vertebral column resection above bony spur to shorten the spine and decrease the stretched power on the spinal cord.</td>
<td>No complications</td>
<td></td>
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<tr>
<td>Endo et al [17], 2014</td>
<td>Japan</td>
<td>Case report 1</td>
<td>Male/43</td>
<td>NM</td>
<td>Progressive spastic gait disturbance; numbness; muscle weakness and pyramidal tract signs in the lower limbs</td>
<td>Lumbar disc herniation</td>
<td>Herniotomy via a posterolateral approach and instrumented posterolateral fusion</td>
<td>Decompression and posterolateral fusion without intervertebral fusion</td>
<td>No complications</td>
<td></td>
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<tr>
<td>Srinivas et al[23], 2012</td>
<td>United Kingdom</td>
<td>Case report 1</td>
<td>Female/77</td>
<td>NM</td>
<td>Severe low back pain and progressive paraparesis</td>
<td>Lumbar disc herniation</td>
<td>Posterior decompression</td>
<td>Indirectly decompression by the falling back spinal cord</td>
<td>Deep wound infection</td>
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<td>Köning et al [25], 2012</td>
<td>United Kingdom</td>
<td>Case report 1</td>
<td>Female/26</td>
<td>Type II</td>
<td>Severe low back pain, and bilateral L5/S1 sciatica</td>
<td>Spondylolisthesis</td>
<td>Anterior in situ fusion coupled with pedicle screw fixation</td>
<td>Anterior fusion to minimize manipulation of neural structures</td>
<td>No complications</td>
<td></td>
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<tr>
<td>Kawamura et al[27], 2010</td>
<td>Japan</td>
<td>Case report 2</td>
<td>Male/69; Male/36</td>
<td>NM</td>
<td>1 Legs and low back pain with intermittent</td>
<td>Lumbar spinal stenosis</td>
<td>Pedicle subtraction osteotomy and</td>
<td>Pedicle subtraction osteotomy to</td>
<td>NM</td>
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Wang YT et al. Spinal diseases complicated with tethered cord syndrome

<table>
<thead>
<tr>
<th>Kramer [28], 2009</th>
<th>Canada</th>
<th>Case report</th>
<th>Female/54</th>
<th>NM</th>
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<td>claudication; 2 Numbness and severe muscle weakness in the lower limbs</td>
<td>yellow ligament resection</td>
<td>shorten the spine</td>
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<tr>
<td>Progressive pain and sensorimotor symptoms in the lower back and limbs</td>
<td>Thoracic disc herniation</td>
<td>Posterior partial vertebral body resection and decompression</td>
</tr>
<tr>
<td>Thoracic disc herniation</td>
<td>Osteotomy to shorten the spine</td>
<td>NM</td>
</tr>
</tbody>
</table>

ALIF: Anterior lumbar intervertebral fusion; NM: Not mentioned; SCM: Split spinal malformation; TCS: Tethered cord syndrome.

Figure 2 Radiography after surgery and at follow-up. A: Postoperative anteroposterior and lateral thoracolumbar radiographs; B: Postoperative thoracolumbar computed tomography (CT) scan showing the absence of thoracic ossification of the ligamentum flavum; C: Magnetic resonance imaging at 1 mo after surgery showing good decompression of the spinal canal (red and blue circle); D: Lumbar CT scan at 3 mo after surgery showing the intervertebral fusion status and good position of screws.

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avoiding stretching of the spinal cord through indirect compression operation without removing the compressive lesions, adopting an anterior approach to fusion, or pursuing posterolateral fusion without intervertebral fusion[17,23,25]. Another important method to protect the tethered cord is to shorten the spine, such as vertebral column resection or pedicle subtraction osteotomy[26-28]. As for the instability and compression from spondylolisthesis, fixation to acquire stability is fairly necessary. In order to minimize manipulation of neural structures, in situ fusion instead of reduction is a good choice[25].

To the best of our knowledge, this is the first case report of an adult patient with TCS due to SCM coupled with both thoracic OLF and lumbar spondylolisthesis who needs thoracolumbar combined surgery. Given the complexity of this case, we treated the lesions according to the following principles: (1) First, we treated the thoracic OLF by removing the ossification to decompress the thoracic spinal canal; (2) After the thoracic pedicle screw fixation was completed, bilateral fixation compression was
conducted to reduce the spinal cord strain to provide compensatory space for the following L4-5 MIDLF; (3) There was no reduction operation for the lumbar spondylolisthesis when performing the in situ MIDLF, so the split and tethered cord was not stretched during the whole operation period; and (4) Finally, a safe and uneventful intraoperative neural electrophysiological monitoring enhanced the confidence of the surgeon and improved the safety of this combined surgery. The follow-up outcomes demonstrated our treatment was successful.

This is only a case report and it remains to be confirmed that our treatment strategy is optimal through studies with larger sample sizes. Moreover, the follow-up was relatively short; longer follow-up is needed to provide information on the long-term decompression effects.

CONCLUSION

For adult patients with underlying TCS secondary to SCM coupled with thoracic OLF and lumbar spondylolisthesis, combined thoracolumbar fusion surgery could be safe and effective with the tethered cord untreated. It is critical to design individual surgical protocols to reduce the stretch of the low-lying spinal cord.

FOOTNOTES

Author contributions: Wang YT and Mu GZ reviewed the literature and contributed to manuscript drafting; Wang YT analyzed and interpreted the imaging findings; Sun HL was the patient’s spine surgeon and was responsible for the revision of the manuscript; All authors have read and approved the final manuscript.

Informed consent statement: Informed written consent was obtained from the patient for treatment and publication of this report.

Conflict-of-interest statement: All authors declare that they have no conflict of interest to disclose.

CARE Checklist (2016) statement: The authors have read the CARE Checklist (2016), and the manuscript was prepared and revised according to the CARE Checklist (2016).

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Country/Territory of origin: China

ORCID number: Hao-Lin Sun 0000-0002-4938-9198.

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