Case Report

Horner’s syndrome secondary to T1-T2 intervertebral disc prolapse

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ABSTRACT

Background: T1-T2 intervertebral disc prolapse (IVDP) is a rare clinical condition. Horner's syndrome is an extremely rare clinical finding in these patients.

Case Description: A 56-year-old man presented with the left C8 T1 radiculopathy, left hand grip weakness, and ipsilateral Horner's syndrome. Magnetic resonance imaging of the spine showed a contrast-enhancing lesion in the left T1 foramen compressing the left T1 nerve root. He underwent left T1 hemilaminectomy, upper half of left T2 hemilaminectomy and removal of the left foraminal lesion. A biopsy of the lesion was sent for histopathological diagnosis which revealed tissue consistent with disc material. Postoperatively, he had near-complete recovery with residual minimal Horner's syndrome.

Conclusion: T1-T2 IVDP should be considered in the differential diagnosis when a patient presents with C8 T1 radiculopathy and Horner's syndrome.

Keywords: Horner's syndrome, Radiculopathy, Spine, Thoracic disc prolapse

INTRODUCTION

The incidence of thoracic disc prolapse is reported to be around 1 in 1000–1 in 1,000,000.[8,28] This accounts for <1% of all intervertebral disc prolapse (IVDP). Among these, only around 4% of cases are above the level of T3.[9] Only around 24% of upper thoracic disc prolapses are found to be associated with Horner’s syndrome.[27] In this case report, we present the diagnostic challenges and management of an extremely rare case of T1-T2 IVDP associated with Horner’s syndrome.

CASE REPORT

A 56-year-old man with no previously known comorbidities presented with a 3-week history of radicular pain along the medial aspect of the left arm and forearm, numbness along the medial aspect of the left arm and forearm, and progressive left-hand grip weakness.

On examination, he had left-sided partial ptosis and miosis [Figure 1a], wasting of the left forearm muscles and small muscles of the hand [Figure 1b], weakness of the intrinsic muscles of the left hand, and decreased sensation along the medial aspect of his left arm and forearm. His reflexes were normal.
Blood counts and inflammatory markers were normal. Chest X-ray and computed tomography (CT) thorax were normal. A magnetic resonance imaging (MRI) of the cervicothoracic spine revealed T1-T2 disc prolapse on T2-weighted images [Figures 2a and b]. On injecting gadolinium, contrast-enhancement [Figures 3a and b] was seen around the T1-T2 disc space. The lesion was seen compressing the left T1 nerve root. The radiological differential diagnosis of the lesion we considered was (i) inflammatory lesions such as pyogenic discitis or Pott’s spine, (ii) benign lesions such as schwannoma, neurofibroma, or meningioma, (iii) malignant lesions such as metastasis, lymphoma, or Pancoast tumor, or (iv) cervicothoracic IVDP.

Under general anesthesia, he underwent left T1 hemilaminectomy, upper half of left T2 hemilaminectomy, and removal of the lesion through a posterior approach in prone position with electrophysiological monitoring. Intraoperatively, the foraminal lesion was seen filling the left T1 foramen and pushing the T1 nerve root superiorly and the dural tube medially. The lesion appeared fibrous and was seen extending into the T1-T2 disc space. The lesion was removed and the nerve root and dural tube were made lax. The dura and the nerve roots were not infiltrated by the lesion. There was no purulent material. Biopsy of the lesion revealed tissue consistent with disc material with no evidence of granuloma, atypical, or malignant cells. The tissue culture from the lesion was reported as sterile. Immediately after surgery, his left upper limb radicular pain improved significantly. At 3-month follow-up, there was a total improvement in his left upper limb radicular pain, numbness, and weakness. Left forearm wasting was gradually improving, and there was residual minimal Horner’s syndrome.

**DISCUSSION**

T1-T2 IVDP is an extremely rare clinical entity.\(^{2,26}\) The uncommonness of this condition is probably due to the relative stability of the vertebral column at this level.\(^{17}\) Horner’s syndrome in association with T1-T2 IVDP\(^{5,6,11,13-15,20,23,25,27,28}\) has only been rarely reported.

The oculosympathetic pathway consists of central, preganglionic, and postganglionic neurons with two relay centers, the ciliospinal center of Budge and the superior cervical ganglion.\(^{2,21}\) Interruption of any segment of the oculosympathetic pathway can cause Horner’s syndrome.\(^{2,21}\) The injury could be traumatic, neoplastic, vascular, or inflammatory etiology.\(^{2}\) Horner’s syndrome or oculosympathetic paralysis presents as a triad of ipsilateral pupillary constriction, facial anhidrosis, and ptosis caused by the relaxation of the superior tarsal muscle.\(^{1,2,21,23,27}\)

In T1-T2 IVDP, the disc can impinge on the sympathetic chain within the T1 root.\(^{27}\) This causes disruption of the ipsilateral preganglionic sympathetic pathway with unopposed parasymptathetic activity resulting in Horner’s syndrome.\(^{27}\) We believe that compression of the sympathetic chain by the T1-T2 disc at the T1 foramen could be the cause of Horner’s syndrome in our case. The exact reason why
| Reference                     | Age/Sex | Clinical features                                                                 | Investigation                                                                 | Treatment                              | Outcome                                           |
|------------------------------|---------|-----------------------------------------------------------------------------------|--------------------------------------------------------------------------------|----------------------------------------|--------------------------------------------------|
| Hammon[14] (1968)            | 33/Male | Lt T1 radiculopathy, Lt ptosis and miosis                                        | Myelogram - Lt T1-T2 defect                                                        | Surgery - Hemilaminectomy and discectomy | Total recovery                                    |
| Gelch[13] (1978)            | 40/Male | Lt upper limb radiculopathy, Lt hand grip weakness, Lt ptosis and miosis          | Myelogram - Lateral defect at T1-T2                                                | Surgery - Partial laminectomy and discectomy | Near-total recovery with mild residual miosis |
| Lloyd et al.[20] (1979)     | 50/Male | Interscapular back pain radiating to Lt chest and arm, Lt hand intrinsic muscles weakness, Lt ptosis and miosis | Myelogram - Smooth extradural defect at Lt T1-T2                                   | Surgery - Laminectomy and discectomy     | Total recovery                                    |
| Simper and Jorgensen[16] (1989) | UK     | Lt upper limb radiculopathy, Lt ptosis and miosis                                | Myelogram and CT spine - Herniation of T1-T2 disc                                  | Surgery - Laminectomy and discectomy     | Total recovery                                    |
| Caner et al.[6] (2003)      | 57/Male | Lt chest pain radiating to Lt upper limb, Lt T1 sensory loss, Lt hand intrinsic muscle weakness, Lt ptosis and partial miosis | MRI spine - Herniation of T1-T2 disc                                                | Surgery - Discectomy through anterior approach | Total recovery                                    |
| Spacey et al.[27] (2014)    | 54/Male | Periscapular pain radiating to the Rt medial forearm, Rt T1 sensory loss, Rt hand intrinsic muscle weakness, Rt ptosis and partial miosis | MRI spine - Lesion at the level of T1–2 with compression of the Rt T1 root          | Conservative                           | Near-total recovery with mild residual ptois |
| Foss-Skiftesvik et al.[11] (2015) | 60/Male | Thoracic back pain radiating to Rt upper limb, Rt T1 sensory loss, Rt hand intrinsic muscle weakness, Rt ptosis and miosis | MRI spine - Paramedian herniation of T1-T2 disc compressing Rt T1 nerve root       | Conservative                           | Near-total recovery with mild T1 paraesthesia and mild ptosis |
| Heckmann and Pauli[15] (2016) | 45/Female | Midscapular pain radiating to Lt upper limb, Lt T1 sensory loss, Lt hand grip weakness, Lt ptosis and partial miosis | MRI spine - Lt lateral disc herniation at T1-T2 level obliterating the T1-T2 intervertebral foramen | Surgery - Foraminotomy and discectomy via posterior approach | Total recovery                                    |
| Teixeira et al.[28] (2017)  | 34/Female | Neck and scapular pain radiating to Lt upper limb, Lt T1 sensory loss, Lt ptosis and miosis | MRI spine - Lt disc herniation at T1-T2 with intraforaminal compression of T1 nerve root | Surgery - Lt T1-T2 laminotomy, partial medial facetectomy and microdiscectomy. | Total recovery                                    |
| Bhandutia et al.[9] (2017)  | 29/Male | Rt upper limb radiculopathy, Rt T1 sensory loss, Rt ptosis, miosis and anhidrosis  | MRI spine - Rt paracentral disc herniation at T1-T2 level                           | Surgery - Hemilaminectomy medial facetectomy and discectomy | NK |
| Possley et al.[23] (2018)   | 29/Male | Periscapular pain radiating to Lt upper limb, Lt T1 sensory loss, Lt handgrip weakness, Lt ptosis and miosis | MRI spine - T1-T2 Lt paracentral disc prolapse with Lt lateral stenosis with compression of Lt T1 nerve root | Surgery - Lamino-foraminotomy and discectomy | Near-total recovery with mild ptosis and miosis |
| Our case                    | 56/Male | Lt C8 T1 radiculopathy, Lt T1 sensory loss, Lt handgrip weakness, Lt ptosis and miosis | MRI spine - Contrast-enhancing lesion in the Lt T1 foramen with compression of the Lt T1 nerve root | Surgery - Complete T1 and partial T2 Lt hemilaminectomy and discectomy | Near-total recovery with mild ptosis and miosis |

MRI: Magnetic resonance imaging, Lt: left, Rt: Right, NK: Not known, CT: Computed tomography
anhidrosis was not seen in our case is unknown. This could be due to incomplete disruption of the sympathetic pathway at the T1 foramen.

There are multiple causes for the upper limb radiculopathy associated with Horner’s syndrome. Pancoast tumor is reported to be the most common cause of T1 nerve root compression presenting with Horner’s syndrome.[28] Other causes include (i) T1-T2 IVDP,[5,6,11,15,20,23,25,27,28] as seen in our case, (ii) traumatic spinal cord injury at the cervicothoracic junction,[7,30] (iii) infective spine lesions such as osteomyelitis[18] or Pott’s spine, and (iv) neoplastic lesions such as vertebral metastasis,[4] neurofibroma,[19,20] or schwannoma.[16]

Careful evaluation of the history and proper clinical examination can help localize the lesion.[21] MRI of the spine is the investigative procedure most commonly used to confirm the diagnosis of thoracic disc prolapse.[21] Contrast-enhanced MRI is indicated when inflammatory or neoplastic lesions are suspected. On contrast MRI, gadolinium enhancement is rarely seen in IVDP[1,10,29] as seen in our case. Due to its atypical presentation on contrast MRI, we initially suspected the lesion to be inflammatory or neoplastic. However, the biopsy sent for histopathological diagnosis revealed tissue consistent with disc material. Our case is novel in that it is the first reported case of T1-T2 disc prolapse showing contrast-enhancement on MRI.

With technological advancement, intraoperative ultrasound and neuronavigation are being increasingly used for image-guided spine surgery.[12] These modalities allow for minimally invasive spine surgeries and permit maximal resection of spinal lesions while preserving vital neural structures.[12] Hence, liberal use of intraoperative imaging should be done to ensure complete resection of spinal lesions without damage to the spinal cord and surrounding neural structures.

In 2019, Rahimizadeh et al.[24] identified previously reported 32 cases of T1-T2 IVDP. In addition to this, they analyzed and reported four new cases from their own experience. We were able to identify six more previously reported cases[5,11,13,15,23,25] in addition to the 36 cases reviewed by Rahimizadeh et al. This makes our case the 43rd reported case of T1-T2 IVDP published in the literature. Among these reported cases of T1-T2 IVDP, we were able to identify Horner’s syndrome in only 11 cases (26.2%).[5,6,11,13,15,20,23,25,27,28]

Table 1 summarizes all the 11 cases of T1-T2 IVDP associated with Horner’s syndrome published in the literature. The age of these patients ranged from 29 to 60 years. There was a male preponderance. Scapular pain,[15,20,23,27,28] thoracic back pain,[11] upper limb radiculopathy,[5,6,11,13,15,20,23,25,27,28] sensory loss,[5,6,11,13,15,20,23,27,28] and weakness of intrinsic muscles of the hand[5,6,11,13,15,20,23,17] were the presenting symptoms. In most cases, an MRI spine with or without contrast was done to confirm the diagnosis.[5,6,11,15,23,27,28] For cases from the pre-MRI era, myelogram and CT were used as investigative modalities.[13,14,20,25] T1-T2 IVDP is managed either surgically or conservatively.[27] A small percentage of patients improve with conservative management[11,27] while most require surgical treatment.[5,6,11,15,20,23,25,28] Surgically, T1-T2 IVDP was approached either through an anterior[6] or a posterior[5,13,15,23,25] approach. Posterior approach to T1-T2 IVDP was done through a facetectomy,[25,28] hemilaminectomy,[14] or foraminotomy.[15,23] Only two cases were reported where the patients were managed conservatively.[11,27] All patients showed improvement in their symptoms and they had good outcomes.

CONCLUSION

T1-T2 IVDP should be considered in the differential diagnosis when a patient presents with C8 T1 radiculopathy and ipsilateral Horner’s syndrome. A careful neurological examination needs to be done to look for Horner’s syndrome in these patients. A contrast-enhanced MRI spine should be the investigation of choice in these patients as it helps to rule out inflammatory and neoplastic causes of C8 T1 radiculopathy. The MRI should always extend to the upper thoracic vertebral levels in these cases. The majority of these patients require a discectomy. A small subgroup of patients with T1-T2 IVDP improves with conservative treatment. These patients generally have good outcomes.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent.

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Conflicts of interest

There are no conflicts of interest.

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