CASE REPORT

Schwannoma of T12 vertebra: case report and review of literature

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Abstract
We report a case of schwannoma of the twelfth thoracic vertebra that presented with paraparesis. The tumour was excised, and posterior and anterior stabilisation was performed. Eighteen months following this procedure, the patient has solid bony union, satisfactory neurological improvement and no recurrence.

Key words: schwannoma, vertebra, thoracic

Introduction
Neurilemmomas or schwannomas are neoplasms that arise from nerve sheath cells. These rarely arise from the nerves supplying the bone. The most common bones involved are mandible, followed by sacrum. The incidence of this neoplasm in bones is reported as less than 1% of primary bone tumours.1 We report a case of schwannoma of the twelfth thoracic (T12) vertebra in a patient who presented with paraparesis.

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Fig. 1. Photomicrograph of the schwannoma (centre) invading bone that can be seen on both sides of the lesion.

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A 37-year-old man presented with back pain, weakness and numbness of both lower limbs in September 1998. He had been suffering from intermittent back pain in the past few years. In August 1998, he had developed constant back pain. His pain was mainly on his upper lumbar region and he had no leg pain. In September, he had developed weakness and numbness of his lower limbs. Weakness was worse on his right side. He did not have any bladder symptoms or perineal sensory loss.

On examination, the straight leg raising test was 50 degrees bilaterally without any leg pain. Neurological examination revealed sensory impairment of L5 and S1 dermatomes bilaterally. Motor examination revealed 0/5 power of extensor hallucis longus (EHL) and ankle evors on the right side. On the left side, the power was 3/5 of EHL and 5/5 of evors. There was loss of bilateral ankle jerks. There was no evidence of saddle anaesthesia and the anal tone was normal.

An urgent magnetic resonance imaging (MRI) scan was performed that showed a large soft tissue mass occupying most of the posterior part of the body of T12. The mass was extending into the spinal canal pushing the cord posteriorly. This mass had a well-defined margin all around. There was homogeneous enhancement with gadolinium.

This patient underwent excision of this lesion with anterior and posterior stabilisation in September 1998. Post-operative recovery was slow but encouraging. He had difficulty in emptying bladder in the initial post-operative period, which has since improved. Presently, he uses his abdominal muscles to facilitate bladder emptying. Power in his legs is grade 4/5 bilaterally and he walks without any aids.

Histology revealed highly cellular spindle cell tumour exhibiting marked nuclear palisading throughout the lesion without any nuclear pleomorphism. The tumour showed strong S100 staining with significant proliferation on Ki-67 staining.

Schwannomas arise from nerve sheath cells, particularly from sensory nerves, and are usually benign. Although it can appear in any age, the usual age group is the third or fourth decade. There appears no predilection for sex. They have been reported to arise in mandible, sacrum, lumbar, thoracic and cervical vertebrae, femur, tibia, fibula, sternum, calcaneum, etc. The reason for the increased incidence in mandible is due to either the long intra-osseous course of the nerve or the increased predilection of schwannomas to the head.
Intra-osseous schwannoma of vertebrae

Schwannomas of the spinal column present according to the site of the vertebrae involved. The schwannoma of the cervical, thoracic and lumbar vertebrae usually present with pain\textsuperscript{4,6,7} and/or neurological involvement.\textsuperscript{5,8,9} The neurology differs with the level of the lesion.

Reported cases of schwannoma of cervical spine have presented with neurological involvement.\textsuperscript{8,9} In one of the reported cases of thoracic schwannoma, the patient presented with fractured L1 vertebra following trauma but had the osteolytic lesion

and neck region, which has a large supply of sensory nerves.\textsuperscript{13}
(schwannoma) in T12, predisposing to the fracture. In another case of thoracic schwannoma at T8 level, the patient presented with 15 years history of pain. In both cases, the neurological examination was normal. A reported case of lumbar schwannoma had neurological involvement during presentation. Schwannoma of the sacrum usually presents with low back pain, with or without sciatic pain. Cases have been reported to have presented as rectal mass during routine per rectal (P/R) examination and during barium enema examination. Our patient presented with back pain and neurological symptoms as already explained.

Involvement of the bone can be explained as follows: (a) an extra-osseous tumour that can erode the bone by contiguity, (b) a tumour arising from the nutrient canal, and (c) a tumour arising centrally within the bone. The differential diagnosis includes giant cell tumour, chordoma, chondrosarcoma, and metastatic deposits.

Fig. 4. Sagittal T2W showing decompression of the theca (post-operative).
Recurrence is rare following surgical excision. In our case, 1.5 years after follow-up, there is no evidence of any recurrence of the tumour. Post-operative computed tomography and MRI scans show good bony union. The clinical improvement has been stated earlier.

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*Fig. 5. Post-operative X-ray showing the position of the implants.*
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