Schwannoma of the brachial plexus: a rare cause of monoparesis

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DESCRIPTION
Schwannomas, also referred as neurilemmomas, are benign, encapsulated perineural tumour of neuroectodermal derivative that originate from the schwann cells of the neural sheath of motor and sensory peripheral nerves. About 25% of schwannomas occur in the head and neck region,1 usually involving cranial nerves and sympathetic chain; however, brachial plexus schwannomas are uncommon, occurring in 0.3–0.4/100 000 person per year.2–4 Their malignant potential is low, they can be locally destructive if allowed to progress. Primary tumours of the brachial plexus are unusual. Patients commonly seek medical attention...
because of pain and loss of function of the affected part of the body. We report a case of schwannoma arising from left brachial plexus and presented as monoparesis of left upper limb.

A young 16-year-old girl presented to our hospital with sharp shooting pain on medial aspect left upper limb radiating up to medial aspect of hand since 10 months. Over the last 2 months, she noticed progressive weakness in left hand along with the inability to perform fine tasks (difficulty in gripping objects, breaking morsels). There was no history of trauma, fever, change in voice and weight loss.

Examination revealed normal vitals and no swelling, pigmentation (freckles or café-au-lait spots) or localised tenderness around left supra-clavicular region, axilla or anywhere in the body. Neurological assessment showed normal higher mental functions and cranial nerve examination. Hypotonia was noted in the left upper limb along with wasting of biceps, triceps, deltoid, supraspinatus, infraspinatus and the intrinsic muscles of the hands. Weakness was noted in these muscles (distal involvement more than proximal) with grade 3 to 4 power as per medical research council grading. Sensory modalities for touch, temperature and pin prick were lost in left upper arm. Rest of the examination was normal (figures 1 and figures 2).

On investigation, complete blood count, random blood sugar, liver and kidney function tests were normal. x-Ray cervical spine and left shoulder region did not reveal any obvious bony abnormality, MRI coronal T2 STIR sequence reveals a well-defined cone-shaped hyperintense lesion in left axilla; it also reveals the narrow tapering medial part of the lesion approaching C7/D1 neural foramina along with its widening, suggesting nerve sheath tumour (figures 3A,B).

A core biopsy was obtained through CT guidance and the histological finding of the tissue revealed benign spindle cell neoplasm arranged in fascicles, intermixed with hypocellular and hypercellular areas. The hypercellular areas showed a palisading pattern with formation of Veroay bodies (small groups of fibrils surrounded by rows of palisaded nuclei), while the hypocellular area showed dispersed spindle cells in the myxoid stroma. Immuno histochemistry examination showed that the sample was S100 antigen-positive and desmin-negative suggestive of schwannoma (figures 4 and 5). The patient was examined by a neurosurgeon and complete excision of the tumour is being planned.

Learning points
- Schwannoma may present as unrecognised slow-growing tumour.
- Benign classification underestimates their potential size and the damage they can cause to surrounding structures.
- Timely diagnosis and appropriate management may prevent such complications.

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REFERENCES