Case Report

The common peroneal nerve schwannoma: A case rare case report with brief review of literature

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1. Introduction

Neurilemmomas (neurinoma-schwannoma) are benign, solitary, well-demarcated usually as a capsule, slowly growing tumours. They arise from the proliferation of active peripheral Schwann cells.¹,² Although they represent the commonest benign peripheral nerve sheath tumours, the occurrence on the lower limbs account for 1% of all cases.³ Frequently they are seen between the ages of 20 and 50 years.

Cases of common peroneal nerve schwannomas are very rare in the literature (Table 1).²,⁴ They usually have a clinically silent course. Rarely, they may present as knee pain or paraesthesia due to mechanical compression of the nerve. A thorough examination and prompt investigation helped us make a timely diagnosis and initiate appropriate treatment. Schwannomas should be included in the differential diagnosis of lumps in the vicinity of nerves. These tumours may affect the nerve conduction of the nerve as indicated by the nerve conduction study in our case. This case presentation highlights the importance of screening for the possibility of a peroneal schwannoma in patients presenting with popliteal swelling despite nonspecific findings of physical examination.

2. Case Report

A 12 years old male student patient presented to OPD with a complaint of swelling in right popliteal fossa since 5 years. It was initially of size of peanut which gradually increased over period to present size about 2x2 cm. It was insidious in onset, gradually progressive, not associated with any pain or restriction of movement or any neurological deficit. There was no h/o trauma, pin prick, fever, reduction of swelling or any other.

Examination revealed a firm mass in the proximal lateral aspect of his right leg, measuring 3cm x 3cm, located in the subcutaneous plane with little mobility in the vertical plane and good mobility in the horizontal plane (perpendicular to nerve direction). It was non tender, non-reducible, non-fluctuant, non transilluminating. Swelling disappeared on flexing the knee. No other masses were found. Motor function of the deep and superficial peroneal nerves was normal. By lightly tapping the leg lump, a positive Tinel’s sign was elicited.
Table 1: Some related case reports.

<table>
<thead>
<tr>
<th>S. No.</th>
<th>Case reports</th>
<th>Symptoms</th>
<th>Excision</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Shariq et al. (2012)</td>
<td>Knee lump</td>
<td>Required intralesional excision</td>
</tr>
<tr>
<td>2</td>
<td>Houshian et al. (1999)</td>
<td>Knee Pain</td>
<td>Successful</td>
</tr>
<tr>
<td>3</td>
<td>Mahitchi et al. (2001)</td>
<td>Knee Pain and numbness</td>
<td>Successful</td>
</tr>
<tr>
<td>4</td>
<td>Cato et al. (1995)</td>
<td>Knee web space pain</td>
<td>Successful</td>
</tr>
<tr>
<td>5</td>
<td>Birol Aktaş et al. (2017)</td>
<td>Knee lump. pain numbness</td>
<td>Successful</td>
</tr>
</tbody>
</table>

Routine laboratory diagnostic tests and x rays showed no specific findings. Therefore, an MRI scan was obtained (Image A). Nerve conduction study was done which was suggestive of decreased motor amplitude of the right common peroneal nerve at ankle and fibular head as compared to left common peroneal nerve.

The MRI revealed an ovoid solid mass at posterolateral aspect of right knee about 2.9 cm x 1.79 cm x 2.96 cm in size overlying the head of gastrocnemius and anterolateral just abutting the bicep femoris. It was slightly hyper intense than muscle on T1 weighted images and was hyper intense on T2 weighted images and at the perimeter was involving the common peroneal nerve and demonstrating nerve continuity.

During surgery, a longitudinal incision was made and the lesion was exposed by longitudinal cutaneous and subcutaneous incisions. The peroneal nerve was located and dissected secured at either ends. After longitudinal dissection of the perineurium, the lesion was excised with its roots in the common peroneal nerve. The perineural sheath was repaired with prolene 8-0 suture.

The excised material was sent for histopathological examination which showed a capsule formed in the epineurium with spindle cells arranged in short whorls, with hyper and hypo-cellular areas and numerous verocay bodies.

The patient’s complaints completely disappeared in the post-operative period without development of any neurological deficit.

3. Discussion

A Schwannoma, also known as neuroma, neurilemmoma or Schwann cell tumour, arises from the neurilemmal cells which forms myelin over peripheral nerves. Schwannomas are slow growing masses with rare malignant transformation. Schwannomas is more predilection for females in usual age group of fifth decade. Schwannomas are located in varied location with rare reports of being located in leg and foot. Schwannomas are usually clinically silent lesions with main presentation as a painless growing mass as was in our case. MRI is the investigation of choice of peripheral nerve schwannomas which reflects nature of pathology and planes with respect to adjoining anatomical structures. Nerve conduction studies and other ancillary tests may reflect subtle changes in nerve physiology where no obvious clinical symptoms of nerve compromise are evident as was there in our case with decreased NCV.

Surgical excision remains the treatment of choice with careful enucleation of schwannoma preserving nerve fascicles is the main goal of treatment. The main challenge is to preserve nerve continuity and its function in post operated period. Our case had no deficits in post operated period and is doing fine after 1 year of follow up.

The schwannomas are grossly solid or solid cystic lesions with presence of Antoni A and Antoni B pattern on microscopy. In Antoni A, there is presence of spindle cell array with collagen matrix and presence of verocay bodies while in Antoni B loose arrangement is present with a mucinous matrix.
There is usually a good prognosis after complete excision of peroneal nerve schwannomas.

4. Conclusions

Schwannoma of peroneal nerve may present as a painless palpable mass in leg and foot and should be kept in consideration while evaluating such cases specially swelling in popliteal region. Complete excision of lesion with preservation of nerve function and integrity remains the mainstay of treatment with low rates of recurrence.

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6. Conflict of Interest

None.

References


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