Common Peroneal Nerve Splitting in Proximal Fibular Osteochondroma: A Rare Presentation

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Learning Point of the Article:
In cases with proximal fibular osteochondroma, care should be taken to identify the entire nerve before removal of osteochondroma as failure to identify the anatomy of the nerve by the unwary surgeon may result in neurologic injury.

Abstract

Introduction: Osteochondroma is the most common benign tumor of bone. Tumors are metaphyseal in origin and commonly involve distal femur, proximal tibia, and proximal fibula in the lower extremity [1]. Osteochondromas increase in size throughout the growth period but stop when the epiphyseal plates close [2]. Osteochondroma located at proximal fibula is very close to the neurovascular bundle which can lead to compressive neuropathy of peroneal nerve [2]. It is usually a painless mass without associated symptoms. Symptoms when present may be due to impingement of contiguous tendons, major blood vessels or nerves, contusions, or fractures in rare instances. Osteochondroma located at proximal fibula can change the normal path of nerves and it may lead to the compression of vessels or peroneal nerve, leading to paralysis [3]. Here, we are describing a rare case of proximal fibular osteochondroma in an 18-year-old female, in which osteochondroma is growing through the midsubstance of common peroneal nerve without causing any neurological deficit.

Case Report

An 18-year-old female presented with pain and swelling over anterolateral aspect of the right proximal leg. Initially, it was small and painless, but gradually, over a period of 3 years, it increased in size and started causing pain. Pain was dull aching in nature,
which was aggravated on movement and relieved on rest. There was no history of trauma or fever. On examination, we noted a sizeable swelling of size approximately $6 \times 4 \times 3$ cm on proximal one-third right leg over its anterolateral aspect. The swelling was irregular, bony hard in consistency, and fixed to the underlying bone. Tenderness was absent. Knee range of motion (ROM) was comparable to the normal side, and neurovascular function was intact. X-ray and magnetic resonance imaging (MRI) were done as a part of pre-operative evaluation which revealed an eccentric large cauliflower-like growth arising from proximal fibula (Fig. 1). Indication for surgery was painful ROM around knee joint. After a thorough assessment, the patient was planned for en bloc excision of osteochondroma. In supine position, a longitudinal incision was made along the posterior border of fibula. Common peroneal nerve was explored and isolated and when traced distally, it was found to be split from midsubstance by tumor into two limbs (Fig. 2). Both the limbs of nerve were mobilized and were brought anteriorly using a suction catheter. Tumor was excised en-bloc (Fig. 3). Excised tumor was sent for histopathological examination which confirmed it to be osteochondroma, and there was no evidence of a malignant transformation. Post-operatively, distal neurovascular status was intact, and the patient was put on weight-bearing as per pain tolerance. The patient continued to follow-up till 1 year after surgery during which there was no evidence of any recurrence.

**Discussion**

Osteochondroma was initially reported by Sir Astley Cooper, in 1818 [4]. Osteochondroma is seen in 2–3% of the general population and represents approximately 36–41% of all benign bone tumors [5, 6]. Incidence of primary bone tumors involving fibula is 2.5% [7]. In our case, osteochondroma was located in proximal fibula. In 90% of the cases, osteochondromas are solitary, and these can be sessile or pedunculated [2]. In our case also, it was solitary and sessile. Osteochondromas located at proximal fibula can not only cause compression of neurovascular structures but can also distort their normal anatomical course, thus increasing the chances of their injury [4]. Furthermore, continuous increase in size of the tumor may create mass effect on adjacent tendons. In our case, main concern was growing size of the tumor along with its abnormal location which was creating mass effect, leading to pain; however, distal neurovascular function was unharmed.

MRI is the best radiologic imaging method evaluating hyaline cartilage cap [8]. It is also important for visualizing the effect of the lesion on surrounding structures and shows cortical and medullary continuity between the parent bone and osteochondroma [8]. In our study also, MRI was done, and cartilage cap thickness was found to be <1 cm. Normally, common peroneal nerve bifurcates into the superficial and deep peroneal nerves. However, this bifurcation can take place distal to the fibular neck, proximal to the joint line or distal to the joint line, and proximal to the fibular neck [9]. In our case, nerve was bifurcating distal to fibular neck. Furthermore, before the bifurcation could occur, we were surprised during our dissections to find the tumors actually growing through the peroneal nerve. Gray et al. in 2004 reported two such cases of an osteochondroma of the proximal fibula that was noted at...
surgery to grow through the common peroneal nerve, splitting it into two limbs [10]. Wankhade et al. in 2016 also reported a case of proximal fibular osteochondroma causing splitting of common peroneal nerve, leading to neuropathy in an adult [2]. Incidence of nerve compression caused by osteochondroma is <1% [11]. Moreover, it is even more rare to find nerve palsy due to osteochondroma growing through midsubstance of common peroneal nerve as was also the finding in our case [2]. Surgery for proximal fibular osteochondromas varies from simple debulking to the complete excision of proximal fibula (Malawer type I resection) [12]. In our case, tumor was excised from its base rather than removing the proximal fibula. Although the recurrence of osteochondroma after its surgical excision is rare, it may occur if a lesion which is in continuation with the physis in a growing child is not completely removed or if there is incomplete removal of the cartilaginous cap [4]. In our case, follow-up was done for a period of 1-year duration, and no recurrence was noted. Clinical features suspicious for malignant transformation comprise new onset of pain in a previously stable lesion, rapid or new growth, growth after skeletal maturity, and/or large lesions [13, 14]. In our case, there was no clinical indication of any malignant transformation and histopathology report also confirmed our clinical finding.

Conclusion

Thus, we conclude that, by turning up such a rare case, we strive to make the surgeons aware that when removing osteochondroma located at proximal fibula, care should be taken to identify the entire nerve before removal as one may encounter such a situation where tumor is extending through the common peroneal nerve and procedure done in a hurry in such a case can cause irreversible damage to the patient.

Declaration of patient consent: The authors certify that they have obtained all appropriate patient consent forms. In the form, the patient has given the consent for his/ her images and other clinical information to be reported in the journal. The patient understands that his/ her names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

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References


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