Subungal osteochondroma: A case report

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Abstract
Osteochondromas are among the most common benign bone tumors of the foot. However their occurrence is rare in subungal region. Clinically they present as slow growing mass causing deformity of the overlying nail. Here we present a case of subungal osteochondroma in an 11 year male kid with characteristic clinical, radiological & histopathological findings. The lesion was successfully treated by excision.

Keywords: Osteochondroma, exostosis, subungal

Introduction
Osteochondroma is a benign bone tumor. Subungually located osteochondromas are rare and are more common in toes than in fingers. Clinically these lesions may present as slow-growing masses, usually on the dorsum of the distal phalanx. This lesion will further cause deformity of the overlying nail. Main symptom includes pain further leads to ulceration due to pressure on the surrounding structures by the mass. Differential diagnosis includes subungal exostosis, verruca vulgaris, glomus tumor, enchondroma, pyogenic granuloma, osteosarcoma, or malignant melanoma. Early diagnosis and treatment of these lesions with their distinctive clinical, radiologic, and histopathologic features is critical in preventing overaggressive, sometimes mutilating therapy.

Case Report
An 11year old boy presented with pain in the right great toe since. Patient also complains of ulceration over dorsum of great toe medial to nail plate. There was no history trauma. On physical examination, a firm, reddish white nodule was noted medial to the distorted nail. Radiological examination revealed well circumscribed, pedunculated outgrowth over the dorsum of distal phalanx. Growth was radio opaque with sclerotic borders.

Under general anaesthesia, first the deformed nail was removed. Dissection was done till the tumor. Tumor measuring 1 x 0.8 cm was excised & sent to histopathological examination. Curretage of base was done till the medullary cavity. Dead space created by the lesion was sutured using monocryl 3-0. Mupirocin cream was applied. Nail bed dressing was done using Bactigras. A firm bandage was wrapped around the great toe, and the limb was elevated.

Wound examination was done once in 2 days. Nail bed began to heal on 14th day. On histopathological examination of the lesion, a characteristic trabecular bone pattern covered with a hyaline cartilage cap was seen, confirming the diagnosis of osteochondroma.

Discussion
Osteochondroma is the most common benign skeletal neoplasm and accounts for 10% to 15% of all bone tumors and 20% to 50% of benign bone tumors and usually is solitary (90%). Subungually located osteochondromas are rare and are more common in toes than in fingers. Osteochondromas are more frequently observed in adolescents and young adults, with a male to female ratio of 2:1. Etiology is not fully understood. Although they are thought to be congenital in origin, they usually remain asymptomatic in childhood and begin enlarging after puberty. Clinically the lesions appear as firm, shiny, smooth-surfaced, white yellow nodules, which is characteristic for the entity.
Radiographic appearance is diagnostic, exhibiting a well-defined, circumscribed, pedunculated bone growth, projecting from the dorsum of the distal phalanx. Histopathology examination has typical findings of a well-defined trabecular bone topped with a hyaline cartilaginous cap.

Osteochondromas are easily misdiagnosed. Differential diagnosis included benign conditions like verruca vulgaris, subungual fibroma, glomus tumor, pyogenic granuloma, subungual digital mucous cyst, subungual exostosis and enchondroma, and malignant lesions such as osteosarcoma, squamous cell carcinoma, and malignant melanoma. Verruca vulgaris, squamous cell carcinoma, and malignant melanoma can be easily diagnosed by histopathological examination. Subungual fibroma and glomus tumors have characteristic clinical findings.

Enchondromas are tumoral lesions composed of cartilage. They arise from the medullar cavity of the bones and they are frequently located on the dorsum of the distal phalanx. Typically they cause expansion of the underlying phalanx as they enlarge. Radiologically they may be seen as radiolucent areas surrounded by a calcified cartilage localized on the dorsum of the expanded phalanx.

Osteochondromas often are misdiagnosed as mature subungual exostosis because of their radiographic similarities. Subungual exostosis shows trabeculated bone arising from the dorsal or dorsomedial aspect of the tuft and extends distally and away from the epiphysis. This may be broad based, tapered, and upwardly extending or narrow-based with progressive widening. Ossification varies with lesion maturity, and trabeculae are commonly observed at the bases of mature lesions. Destructive changes of the phalanx and periosteal reaction are absent.

Subungual osteochondroma presents as an outpocketing of trabecular bone from the juxtaepiphyseal region of the distal phalanx. These growths usually slant away from the adjacent interphalangeal joint of the phalanx. The cortical and medullary bone of osteochondroma curves smoothly into the distal phalanx, and the lesion bases may be circumscribed, sessile, or pedunculated. Because subungual osteochondroma grows by enchondral ossification, the tumor may increase slowly until puberty and then show an accelerated growth rate throughout puberty and adolescence, until closure of the physeal plates.

The major histologic difference between subungual exostosis and subungual osteochondroma is the composition of the cartilaginous cap. In subungual exostosis, this is composed of fibrocartilage, and the distal phalangeal tuft normally develops through enchondral ossification of fibrocartilaginous anlage. Exostosis begins as a fibrocartilaginous nest just below or adjacent to the nail bed. Microscopic examination of the cap exhibits hypercellularity with plump nuclei and chondrocyte multinucleation. Therefore, a limited biopsy may create the false impression of a malignant cartilage tumor. The thicknesses of fibrocartilage caps vary and largely depend on lesion maturity; moreover, as lesions mature the thickness of cartilage rims reduce as they are replaced by bone. The cartilaginous cap of an osteochondroma is composed of hyaline cartilage, with a cell configuration similar to that of a normal growing epiphysis. This hyaline cartilage may grow to become much larger than in an exostosis, which generally measures less than 1 cm and contains a large number of chondrocytes. Active enchondral ossification also was evident at interfaces between hyaline cartilage and underlying trabecular bone, and trabecular and cortical bone were found to be continuous with the distal phalanx.

Moreover, histologic examinations of subungual osteochondromas do not suggest the possibility of a traumatic etiology.

X-Rays

Intra-Operative Images

Microscopy Images

Conclusion

Subungual exostosis and subungual osteochondroma are benign but distinct osseous pathologies, which differ clinically, radiographically, histologically, and how they develop. When a diagnosis has been made and symptoms persist, complete bony lesion excision with curettage of its base and nail bed reconstruction is the treatment of choice in both conditions.

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