Free non-vascularized fibular graft reconstruction after resection of metacarpal unicameral bone cyst: A case report

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Abstract. Unicameral bone cyst (UBC) or simple bone cyst (SBC) is a benign cystic lesion commonly found in the humerus and femur that is mainly encountered during childhood. UBCs typically remain asymptomatic and represent ~3% of all primary bone lesions (1). UBC is a serous or serosanguineous fluid-filled cavity, which is typically inactive and remains asymptomatic until a pathological fracture occurs. The clinical presentation, location and radiographic features are rather characteristic and the diagnosis may be readily established. In 86% of the cases, UCB occurs within the first two decades of life and the proximal humerus is the most common site, whereas it is rarely observed in the hand. Very few cases of UBC have been described in the hand to date, including the metacarpals, phalanges, hamate and lunate bones. Men are twice as likely to be affected compared with women (2).

The location of UBC may be classified as active (immediately juxtaposed to the growth plate, not to be confused with Enneking stage 2 tumors) or inactive (growth plate not adjacent to the cyst) (3). On clinical examination, the lesion appears as a well-defined fluid-filled cyst in the metaphysis and diaphysis of the affected bone that may cause pain and swelling. Radiological examination is crucial for establishing the diagnosis, based on the following characteristics: Well-demarcated osteolytic lesion without marked marginal sclerosis, central location, thinning of the cortex and intralesional fracture fragment (‘fallen leaf’ or ‘fallen fragment’ sign) (3). The treatment for UBC of the hand commonly involves curettage, bone grafting, partial resection with or without grafting, multiple drilling, fracture immobilization and observation alone, and steroid injection (4).

Introduction

Unicameral bone cyst (UBC) or simple bone cyst (SBC) is a benign cystic lesion commonly found in the humerus and femur that is mainly encountered during childhood. UBCs typically remain asymptomatic and represent ~3% of all primary bone lesions (1). UBC is a serous or serosanguineous fluid-filled cavity, which is typically inactive and remains asymptomatic until a pathological fracture occurs. The clinical presentation, location and radiographic features are rather characteristic and the diagnosis may be readily established. In 86% of the cases, UCB occurs within the first two decades of life and the proximal humerus is the most common site, whereas it is rarely observed in the hand. Very few cases of UBC have been described in the hand to date, including the metacarpals, phalanges, hamate and lunate bones. Men are twice as likely to be affected compared with women (2).

The location of UBC may be classified as active (immediately juxtaposed to the growth plate, not to be confused with Enneking stage 2 tumors) or inactive (growth plate not adjacent to the cyst) (3). On clinical examination, the lesion appears as a well-defined fluid-filled cyst in the metaphysis and diaphysis of the affected bone that may cause pain and swelling. Radiological examination is crucial for establishing the diagnosis, based on the following characteristics: Well-demarcated osteolytic lesion without marked marginal sclerosis, central location, thinning of the cortex and intralesional fracture fragment (‘fallen leaf’ or ‘fallen fragment’ sign) (3). The treatment for UBC of the hand commonly involves curettage, bone grafting, partial resection with or without grafting, multiple drilling, fracture immobilization and observation alone, and steroid injection (4).

Case report

A 9-year-old right-handed female patient presented in February 2017 to the Department of Orthopaedics and Traumatology of Saiful Anwar Hospital (Malang, Indonesia) with a chief complaint of a large, fast-growing lump over the first metacarpal of the left hand. The swelling was first observed 6 months earlier. The pain was described as dull and constant, with no diurnal variation. There was no history of trauma or infection. Physical examination revealed a firm mass, ~3x4 cm in size, which was tender on palpation, but with no local rise in temperature (Fig. 1). Flexion of the first metacarpophalangeal joint was limited. There was minimal soft tissue extension and no sensory-motor deficit; the distal circulation was also normal. On imaging, plain radiographs of the left hand revealed cystic enlargement of the first metacarpal that included the base (Fig. 2).
A biopsy was performed, which revealed that the cyst was filled with serosanguinous fluid with a haemorrhagic tinge (Fig. 3). After the diagnosis was established, we decided to perform total resection of the cyst with placement of a non-vascularized fibular graft. The cyst was completely removed by resecting the shaft of the metacarpal with preservation of the epiphyseal plate (Fig. 4). A bone graft was obtained from the fibula and inserted in the gap, distally attached to the epiphyseal plate of the metacarpal and fixed with Kirshner wire proximally. A radiograph revealed solid union of the bone graft to the epiphyseal plate at the head after 7 weeks, with improving function of the thumb. No recurrence has been reported after the last follow-up appointment during January 2018 (Fig. 5).

**Discussion**

Simple bone cyst or UBC is a serous or serosanguineous fluid-filled cavity that very rarely develops in the metacarpal bone. The exact cause is unknown, but epiphyseal plate defect or venous outflow obstruction have been suggested as possible causes (3). Once diagnosed, UBC poses a dilemma for clinicians, as its natural history and management remain controversial. The recurrence rate of UBC ranges from 20 to 50% after various treatments (5). The differential diagnosis includes aneurysmal bone cyst and fibrous dysplasia. When additional examinations are required, magnetic resonance imaging most accurately delineates the central fluid collection. If a pathological fracture has occurred, a fluid level may be observed, mimicking the appearance of an aneurysmal bone cyst. There is no convincing evidence, however, that a UBC may transform to an aneurysmal bone cyst or other bone lesion (6). The gross pathology of UBC generally includes a singular fluid-filled cavity, unlike an aneurysmal bone cyst, which is composed of blood-filled cystic cavities that are lined by a thick, fleshy membrane with an endothelial-like inner layer. In some cases, multilocular lesions may be present. The bony architecture surrounding the cyst is non-reactive and otherwise unremarkable. Microscopic histological analysis shows a fibrous tissue membrane, with occasional giant cells present. The fluid within the cysts contains high levels of prostaglandins and enzymes (7). Traditionally, plain radiography has been used to diagnose and monitor UBCs, but advanced imaging has become more popular as technology has improved. On plain radiography, UBCs appear as lytic, expansile lesions within the medullary cavity of long bones. The cortex is thinned, but is not typically compromised structurally. Rarely, a UBC may develop in the diaphysial portion of long bones and is known as a ‘latent’ cyst (8).

Among the various treatment methods mentioned above, the choice of treatment may vary according to the patient’s age, the bone involved and the location of the cyst within the bone, and proper treatment should be applied to decrease the risk of recurrence (4,5). Several agents have been used for intralesional injection of UBCs; Scaglietti et al first described the use of methylprednisolone injections in patients with UBCs. Recurrence rates of 15-88% have been reported after an average of three injections (9).
A more aggressive approach, such as in the present case, is occasionally necessary to treat UBC. The patient was radically treated by complete excision and a fibular graft. Our rationale was that, according to Jaffe (1), these cysts are active, as demonstrated by complete destruction of the metacarpal bone; furthermore, the patient was aged <10 years and was statistically more prone to recurrence (10). This method may also be used in other bone lesions, such as aneurysmal bone cyst, as reported by Gundes et al (11), in 2005.

Before proceeding with treatment, a biopsy was performed to confirm the diagnosis. We decided to perform total resection of the cyst with preservation of the periosteum and placement of a non-vascularized fibular graft. The proximal epiphyseal plate was preserved and fixed with a Kirschner wire. After 7 weeks, the function of the thumb was evaluated and union of the graft was observed. There was no recurrence after 3 months of follow-up, and the Kirschner wire was removed. The only known predictor of treatment success is the age of the patient; patients aged >10 years display higher rates of healing (90%) compared with younger patients (60%), regardless of the treatment regimen (8) due to the presence of the active growth plate in younger patients.
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Availability of data and materials

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Authors' contributions

IGMOR and TECJH were involved in the drafting of the manuscript and the revision of the manuscript for important intellectual content. ARBS and SPPI made substantial contributions to the design of the case report and were involved in the surgery. III provided final approval of the version to be published. All the authors have read and approved the final version of this manuscript.

Ethics approval and consent to participate

Ethics approval was obtained from the local Ethics Committee of Saiful Anwar Hospital.

Patient consent for publication

The legal guardians of patients included in the present study provided written, informed consent for the publication of case details and associated photographs in this manuscript.

Competing interests

The authors declare that they have no competing interests.

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