Aneurysmal Bone Cyst of Clavicle in a Child: A Rare Case Report

Suresh Borah¹, Hemonta Kr Dutta²* and Manasjyoti Das¹

¹Department of Orthopaedics, Assam Medical College, India
²Department of Paediatric Surgery, Assam Medical College, India

Abstract

Aneurysmal bone cyst is a benign expanding osteolytic lesion of bones and usually affects teens and young adults. Metaphyseal end of long bones are usually affected followed by spine and flat bones. Involvement of clavicle is rare. We report a seven year old child presenting with an aneurysmal bone cyst of the clavicle which was excised with a bone graft.

Keywords: Aneurysmal bone cyst; Bone tumors; Giant cell tumor

Introduction

Aneurysmal bone cyst (ABC) is a benign but locally destructive tumor of bone. There is presence of spongy or multiloculated blood-filled spaces of variable size separated by connective tissue septa containing trabeculae or osteoid tissue and osteoclast giant cells [1]. It accounts for 1% of all benign bone tumors [2]. Eighty per cent of aneurysmal bone cysts occur in skeletally immature patients who are under age 20 years. There is no sex predilection; the peak incidence is in the second decade of life [2]. Aetiology and pathogenesis of this lesion remains unclear [3]. Aneurysmal bone cysts may involve almost any bone, but the most frequent sites are long tubular bones and vertebrae [4,5]. Most common locations include the proximal humerus, distal femur, proximal tibia, and spine. Among flat bones, the pelvis and scapula are well-known locations. The clavicle is a relatively rare site for this lesion, and not many have been reported in literature [6]. We present a case of aneurysmal bone cyst of lateral end of clavicle in a seven year old girl child.

Case Presentation

A 7-year-old girl presented in the outpatient department with a swelling over the lateral end of the right clavicle for the last 6 months. The swelling increased in size gradually and attained the size of an egg at the time of examination, with smooth surface, mildly tender on palpation and there was no local rise of temperature. The skin was not adherent to the swelling. Consistency was bony hard with egg shell cracking like feeling. The margins were distinct and no local lymph nodes were found. The child had full range of motion at the shoulder without any pain. No other similar swelling was found in the body. Radiographs (Figure 1) showed a well-defined expansile osteolytic lesion with thin sclerotic margins arising from the lateral end of the right clavicle. Thin septations could be seen within the lesion. Blood investigations including ESR, CRP and serum alkaline phosphatase were within normal limits. The FNAC report was inconclusive. Incisional biopsy was performed and was sent to two different laboratories; one of which reported as aneurysmal bone cyst and the other as benign fibro-osseous lesion. Excision of the mass with a portion of normal bone and fibular transplant with k wire was done (Figure 2A and 2B). The shoulder was immobilized for three weeks followed by active movement and physiotherapy. The excision biopsy reports confirmed the case as ABC. The patient was followed up at 3 weeks, 3 months, 6 months and at 1 & 2 years. The child returned to school after 3 weeks. The patient has full range of movements with no functional problems.

The implant was removed after one year and the x-rays showed complete union and no signs of recurrence (Figure 3A and 3B). The child showed full range of movement without any difficulty (Figure 4).

Discussion

Aneurysmal bone cyst usually presents with pain and swelling over a variable period of time. Rarely may they present with any compression features of the adjacent structures. In the long
bones, they commonly involve the metaphyseal region. But they rarely cross the joints [5]. Radiographs show an expansile lytic lesion with blowout or ballooning distension of the periosteum. The mass is outlined by a thin sheet of periosteal bone. It may also show soap bubble like appearance [6,7]. CT scan shows multiloculated cystic nature of the tumor. It may also show fluid levels in the cyst. MRI depicts the expansile nature and also helps to comment regarding the nearby structures [8].

The term aneurysmal bone cyst was coined by Jaffe and Lichtenstein [9] in 1942 based upon the radiographic findings. ABC is a locally aggressive benign tumor and its pathogenesis is uncertain. It is postulated that it may arise from a local circulatory disturbance leading to an increased venous pressure and subsequent production of hemorrhage in the bone [10]. This may explain why there is low incidence of ABC in clavicle due to low venous pressure here. Recent studies have shown that most primary ABCs demonstrate at (16;17) (q22;p13) fusion of the TRE17/CDH11-USP6 oncogene. This fusion leads to increased cellular cadherin-11 activity that seems to arrest osteoblastic maturation in a more primitive state. ABC can be confused with giant cell tumor, chondromyxoid fibroma and telangiectatic osteosarcoma [11]. GCT commonly appears after physeal closure. It is less cystic and seldom grows as rapidly as an ABC. Chondromyxoid fibroma is a rare tumor that generally affects men in the second and third decade. It is slow growing and most commonly involveibia or femur. Its radiological appearance might be confusing but histological differentiated on the basis of findings of a mixture of fibrous, myxomatous and chondroid tissue. Distinction from telangiectatic osteosarcoma is difficult because the conditions have overlapping clinical and radiologic features. The differentiation is made from the histologic features. The presence of highly anaplasmatic sarcomatous cells with atypical mitoses producing osteoid is highly diagnostic of osteosarcoma.

There are numerous ways of treating an ABC like curettage, excision, saucerization, radiotherapy, cryotherapy, vascular occlusion etc. Complete excision of the tumor remains the best modality of treatment [12]. However this is associated with functional impairment. So most commonly curettage and bone grafting is performed [13]. There are reports of spontaneous resolution of ABC [14]. The overall cure rate is 90% to 95% [15,16]. A younger age, open growth plates, and a metaphyseal location all have been associated with an increased risk of recurrence. Embolization of the feeding vessel also serves as an important treatment by reducing the size of the tumor and allowing lesser exposure and dissection during surgery. Adjuvant treatment is done in inaccessible tumors and in conditions where there is potential risk to damage to nearby structures.

References


