A Rare Case of Non-Syndromic Multiple Non-Ossifying Fibromas

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Abstract: Syndromic Multifocal non-ossifying fibromas have been reported in patients with the Jaffe–Campanacci syndrome and in patients with Type 1 neurofibromatosis[1–3]. Evans and park reported a rare case of three members of a family with the existence of multiple, symmetrical, non-ossifying fibromatosis without associated neurofibromatosis [5]. Here we report a rare case of non-syndromic multiple non-ossifying fibromas. A 23-year-old male complained of bowing deformity in the Right knee for 10 years. The deformity was insidious in onset, gradually progressive in nature, and made him sit with a wide base between the two feet. Initially, the base was non-progressive with minimal distance between the two feet but now the distance has increased to the length of a brick. There was no history of pain, trauma, or difficulty to perform routine activities. On examination, Genu Valgum of Right knee was diagnosed and the disappearance of Deformity was noted during flexion of the knee. There were no Café-au-lait spots or other signs of neurofibromatosis. Radiograph of the Knee revealed lucency in the distal end of the Right femur. Differential diagnoses considered were Fibrous cortical defects and Non-ossifying Fibromas. MRI of the Right knee revealed multiple well defined bubbly lytic lesion with sclerotic margin and eccentrically along the cortex in the distal femur. A similar lesion was seen in the proximal tibia. Later bone curettage was performed and histopathology examination revealed findings consistent with non-ossifying fibroma. Here we present a rare case of non-syndromic multiple non-ossifying fibromas.

Keywords: Type I neurofibromatosis, multiple, non-ossifying fibromas, non-syndromic, lytic, Café-au-lait spots, Fibrous cortical defects, sclerotic, Genu Valgum.

INTRODUCTION

Syndromic Multifocal non-ossifying fibromas have been reported in patients with the Jaffe–Campanacci syndrome (multiple NOFs, café-au-lait spots, mental retardation, hypogonadism or cryptorchidism, ocular anomalies or cardiovascular malformations) [1–3] and in patients with Type 1 neurofibromatosis [4]. Evans and park reported a rare case of three members of a family with the existence of multiple, symmetrical, non-ossifying fibromatosis without associated neurofibromatosis. These patients had upper and lower extremity involvement and no family history of neurofibromatosis [5]. Here we report a rare case of non-syndromic multiple non ossifying fibroma.

EXPERIMENTAL SECTION

A 23-year-old male complained of bowing deformity in the Right knee for 10 years. The deformity was insidious in onset, gradually progressive in nature, and made him sit with a wide base between the two feet. Initially, the base was non-progressive with minimal distance between the two feet but now the distance has increased to the length of a brick. There was no history of pain, trauma, or difficulty to perform routine activities. On examination, Genu Valgum of the right knee was diagnosed and the disappearance of the deformity was noted during flexion of the knee. There were no Café-au-lait spots or other signs of neurofibromatosis. There was no evidence of limb length discrepancy, wasting of quadriceps or calf muscles and hyperextension of knee.

Radiograph of knee revealed lucency in distal end of Right femur (fig. 1 and 2). Differential diagnosis considered were Fibrous cortical defect and Non ossifying Fibroma. MRI of the right knee in PD TSE sequence revealed multiple well defined bubbly lytic lesion with sclerotic margin and eccentrically along the cortex in distal femur, in the lateral condyle measuring 1.8 x 3 cm (fig.3), distal metaphysis measuring 2 x 1.3 cm (fig.3), and in the medial condyle measuring 2 x 1.2 cm (fig.4). Similar lesion is seen in the proximal tibia measuring 1.7 x 0.9 cm (fig. 4).

The lesions appear intermediate to low signal on T1WI (fig. 5 and 6) with a high signal on T2 TSE (fig. 7) and T1 STIR imaging (fig 8 and 9). No significant surrounding edema noted. No cortical break or periosteal reaction seen.
RESULTS AND DISCUSSION

Later bone curettage was performed and histopathology examination revealed findings of consistent with Non-ossifying fibroma.

Diagnosis of None ossifying Fibroma in the long bones is based upon the characteristic radiographic and clinical appearance, typically the Non ossifying Fibroma appears as an asymptomatic multiloculated lesion often identified incidentally during radiographic evaluation indicated for another reason [6-8]. While the majority of Non ossifying Fibroma are asymptomatic, those that are particularly large may cause chronic pain and/or pathologic fracture in the long bones [9, 10]. When arising in a large, tubular bone, the Non ossifying Fibroma is always eccentric and ovoid and often results in thinning and expansion of the overlying cortex. There is usually no periosteal reaction [11] and no overt violation of the cortex [12]. The lesions are often multilocular in appearance and the inner boundary of the lesion is demarcated by bony sclerosis [13, 14]. While the definitive etiology of the Non ossifying Fibroma is unclear, it is considered to be a disturbance of growth or dystrophic calcification, rather than a tumour or neoplasm [15, 16].

Histologically, the Non ossifying Fibroma consists of spindle-celled fibrous tissue in a storiform pattern with a variable number of multinucleated giant cells, hemosiderin pigment within fibroblasts, and lipid-laden histiocytes (xanthoma cells) [17, 18]. In addition, occasional lymphocytes and endothelial-lined blood vessels are present [19-21]. The background stroma is composed of reticulin fibres [22, 23].

Radiographically all Non ossifying Fibroma lesions are categorized with respect to location, shape, border definition and presence of sclerotic changes according to Ritschl [24, 25]. Stage A: Eccentric lesion in the cortex near the epiphyseal endplate, which is small, oval to slightly polycyclic in shape without a sclerotic border. Stage B: Lesions with variable distance from the epiphysis with polycyclic shape and thin but clearly sclerotic borders, thin cortex with occasionally protruding above the surface like the shape of an hourglass; no periosteal reaction. Stage C: Lesions with properties similar to stage B but with also exhibit increasing sclerosis, which typically start from the diaphyseal side. Stage D: Complete homogeneous sclerosis of the lesion (D1), disappearing lesion (D2) and disappearance of the lesion (D3).

The regressive phase is observed radiographically as a “fading out” of the lesion: trabecular bone growth appears to occur from the periphery inward until the bone is reconstituted to a normal state [26, 27]. Treatment of None ossifying Fibroma of the long bones, particularly in asymptomatic cases, is observation. These lesions are typically self-limiting and spontaneous resolution at skeletal maturity is usually seen, especially in the smaller lesions of the Fibrous cortical defect [28, 29]. This spontaneous regression, usually starting at the end of adolescence, has been observed to occur over periods of 29–52 months [6].

CONCLUSION

Here we present a rare case of non-syndromic multiple Non-ossifying fibromas.
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