Learning Point of the Article:
How to differentiate the potential differential diagnostic conditions before a diagnosis of chondromyxoid fibroma is made which is essentially a diagnosis of exclusion.

Abstract

Introduction: Chondromyxoid fibroma (CMF) is a benign rare bone tumor of slow-growing nature arising from chondroblastic derivation. CMF in most of the cases is a diagnosis of exclusion, and in this case report, we differentiate the histological and radiological findings of CMF and difficulties in diagnosis of CMF from potential differential diagnosis.

Case Report: A 38-year-old female patient presented with a history of limping for 5 months and on evaluation revealed an expansile osteolytic lesion in fibular head with septations and soft tissue component. Excision biopsy was done. Histological examination revealed a cellular neoplasm arranged as vague nodules in chondroid background with occasional mitotic figures and giant cells in periphery without any calcification. To rule out chondroblastoma, S-100 and epithelial markers were done which was negative establishing diagnosis of CMF by exclusion.

Conclusion: CMF is often misdiagnosed being a radiological and pathological mimicker. Histology remains key to diagnosis. En bloc resection remains the mainstay of management in expendable bone-like fibula.

Keywords: Chondromyxoid fibroma, benign bone tumor, en block resection, fibula.

Introduction

Chondromyxoid fibroma (CMF) is a benign rare bone tumor of slow-growing nature arising from chondroblastic derivation [1, 2, 3, 4]. Jaffe and Lichtenstein described the condition at first in 1943 [5]. In their work, they differentiated CMF from chondrosarcoma, which is a much more common, but with malignant nature. Before their description in 1943, CMF has been considered as a giant cell variant or a benign cartilaginous tumor. A myxoid element in the tumor is the defining characteristic feature.

Around 500 CMF cases have been described in the entire literature [6]. The diagnostic frequency seems to be in decline. CMF mainly affects the second and third decade of young adults. Around 80% of patients are <36 years. The tumor is not gender specific and both males and females are affected equally; however, some series showed a slight male predominance [7]. The aim of our study is to evaluate the histological and radiological findings of CMF and difficulties in diagnosis from potential differential diagnosis of CMF. Consent was obtained for publication of the case details without revealing the identity of the patient.

Case Report
We present a case report of a 38-year-old female who presented with difficulty in walking for the past 5 months associated with pain in the knee for the past 3 months. The pain was dull aching in nature, mild intensity. There was no diurnal variation. Physical examination showed mild tenderness. Overlying skin was normal with restricted range of movement due to pain. Blood investigations were under normal limits.

On magnetic resonance imaging, T1-T2 hypointense and short tau inversion recovery hyperintense lesion with thinning and endosteal scalloping of the cortex with focal break noted in the proximal fibula without any periosteal reaction as shown in (Fig. 2).

Excision biopsy was planned and after obtaining consent from the patient, an en block excision was done. The resected specimen was sent to histopathological examination.

Gross examination showed a grayish-white ill-defined lesion at the epi-metaphyseal region measuring 2.5 × 1.5 × 1 cm which is focally extending into the adjacent soft tissue. On serial microscopic sectioning and examination as shown in (Fig. 3) revealed cellular neoplasm which was well demarcated from the adjacent osseous tissue arranged as vague nodules in chondroid background. The cells were round to oval with vesicular nuclei and abundant eosinophilic cytoplasm and displayed well-delineated cell borders. Mitosis is occasional 1–2/10 hpf.

Figure 1: The X-ray and computed tomography localization of the tumor which revealed multilobulated osteolytic lesion with scalping, thinning, and expansion cortex seen in fibular head with narrow zone of transition without any matrix mineralization.

Figure 2: Magnetic resonance imaging study with T1-T2 hypointense and short tau inversion recovery hyperintense lesion with thinning and endosteal scalloping of the cortex with focal break noted in the proximal fibula without any periosteal reaction.
Giant cells are noted in the periphery of the lesion. There was no evidence of calcification. With a suspicion for chondroblastoma, S-100 and epithelial markers were done which all came out to be negative establishing the diagnosis of CMF by exclusion. Although myxoid element was not much evident on histology considering the tumor to be in the earlier stages of evolution diagnosed, early diagnosis of CMF was made.

Postoperatively, the patient was put on partial weight-bearing for initial 2 weeks and later resumed full weight-bearing by 4 weeks. The case was followed up for 2 years without any recurrence till date as shown in (Fig. 4)

Discussion

The common location of CMF is metaphysis adjacent to the growth plate, which strengthens the theory that lesion is of cartilaginous remnant origin [4]. Similar reports of fibular CMF have been made by Mane et al. [8] from India and Atalar et al. [9] from Turkey and Merine et al. [10] from New York who also corroborate the rarity of the tumor incidence and location in fibula.

To make a certain diagnosis, a thorough clinicoradiological and pathological examination was important as CMF is often misdiagnosed as chondroblastoma or chondrosarcoma due to some pathological similarities.

A myxoid element is the defining characteristic of this tumor, and hence, histology remains the key in diagnosing this rare entity. However, the myxoid element was not so evident in our case which made us to diagnose this pathology by exclusion of the other possibilities such as chondroblastoma by special stains and epithelial markers. Since the tumor is in initial stages of differentiation, myxoid element may not be evident in our case.

Grossly, the tumor is firm, grayish-white in color which may be lobulated or pseudolobulated. Their structure resembles that of fibrous tissue or hyaline cartilage as in tumors such as chondroblastoma or chondrosarcoma. The lesions often thin the cortex and rarely destroy trabecular bone. Rarely, the tumor has some areas of hemorrhagic degeneration and cystic degeneration and sometimes mimics aneurysmal bone cysts.

Many CMFs exhibit morphological features that show to be in different stages of chondrogenesis [11, 12]. The most characteristic pathological fracture is the myxoid element. Giant cells are mostly seen toward the...
periphery of the tumor architecture.
Low grade chondrosarcoma sometimes mimic CMF histologically, except for the lack of myxoid element and soft tissue involvement. Chondrosarcoma has peculiar demographic and radiographic features. Chondrosarcoma has peak incidence around the sixth and seventh decades of life, whereas CMF occurs in the second and third decade of life.

Radiographically, location of the chondrosarcoma is central and has rich calcifications. Both chondrosarcoma and CMF sometimes show cortical expansion. In chondrosarcomas with long-standing or higher grade type, soft tissue involvement and cortical erosion are noted [13, 14].

CMF histopathological picture shows a lobular pattern of growth with focal hyaline deposits and rare mitotic figures and even cellular pleomorphism sometimes which may be similar to that of chondrosarcoma. However, chondrosarcoma shows features such as rich mucinous material, higher nuclear atypia, and multiple pleomorphic or multinuclear cartilage cells. Moreover, it behaves in a more malignant fashion.

Patients with CMF may be misdiagnosed in the early stages as giant cell tumor (GCT) of bone, but they are much older than persons with CMF; moreover, cellular and radiographic features of GCT differ from that of CMF. GCTs are usually of metaphyseal origin with epiphyseal extension. Multinucleated giant cells with nucleus identical to the background nucleus of the stromal cells are the major defining histologic characteristic of GCT [15]. CMFs show only a few giant cells which are also located near the periphery. GCTs show neither myxoid nor chondroid elements.

CMFs are managed surgically with intralesional curettage or en bloc excision depending on the location of the lesion [17]. Although Jaffe and Lichtenstein showed in their original description of CMF that incomplete removal did not lead to recurrence and the remnant tissue shows regression [5], some series noted recurrence rates of approximately 25% with curettage and bone grafting which may be higher in young children who are in the first or second decade of life and in myxoid predominant CMFs. Aggressive curettage of the tumor in the vicinity of the physis resulted in physeal damage many cause growth arrest.

En bloc excision is usually curative in expendable bones. Recurrence is noted in cases, in which marginal excision was done. In a study by Bharma et al. of 22 consecutive cases of CMF treated by intralesional curettage [16], 2 patients (9%) showed local recurrence within 2 years after curettage. CMFs in axial skeleton may behave in an aggressive fashion after resection. Malignant conversion following en block resection is extremely rare. Consequently, no case of mortality was reported due to true CMF.

Extended curettage with phenol, methyl methacrylate, and liquid nitrogen has not decreased the recurrence. In occurrence of this tumor in expendable bones such as fibula, en block resection is the ideal treatment of choice to prevent future recurrence.

Radiotherapy may be considered an option for unresectable tumors [15]. Radiotherapy is generally avoided to prevent post-radiation sarcoma. However, malignant transformation in the absence of preceding radiotherapy has been noted which many authors believe could be a misdiagnosed chondrosarcoma.

Recurrence if at all happens mostly within 2 years, but sometimes recurrence after 19 years has also been reported [13, 16, 17, 18]. Hence, routine patients follow-up with periodic physical and radiological examination for a minimum period of 2 years is recommended [19, 20].

**Conclusion**

CMF is a rare tumor of bone which is often confused with other radiological and pathological mimickers and misdiagnosed many times. Histological evaluation remains a key in diagnosing this rare pathology which is at times a diagnosis of exclusion as in our case. En bloc resection remains the mainstay of the management of this tumor in expendable bone-like fibula.

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**Clinical Message**

CMF a rare tumor, is often confused with other radiological and pathological mimickers such as chondroblastoma or chondrosarcoma. However, myxoid element in histology is the defining characteristic feature. En bloc resection is curative for expendable bone-like fibula.
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