



RESEARCH ARTICLE

CHONDROSARCOMA OF GREAT TOE : CASE REPORT

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Abstract

Here is a case of grade 1 chondrosarcoma of the distal phalanx of left great toe in a 58 year old diabetic patient .Foot is an unusual site for chondrosarcoma and involvement of phalanges are extremely rare. Patient complained of gradual swelling in left great toe since 2 years which progressed to its present stage. Xray showed lytic lesion destroying the distal phalanx with soft tissue extension. Amputation of great toe done and tissue sent for histopathological examination. Microscopically examination revealed grade 1 chondrosarcoma. The patient is on one year follow up which is uneventful.

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INTRODUCTION

Chondrosarcoma is a third most common primary malignancy of bone after myeloma / plasmocytoma and osteosarcoma and accounts for 20% of malignant bone tumours(1) . It is commonly seen in pelvic bone, proximal humerus, distal femur and ribs. Small bones of hands and feet are rarely affected(<1% of all chondrosarcoma)(2). Most patients present with symptoms of pain(3). Phalangeal chondrosarcoma behaves like a locally aggressive lesion unlike other location where metastasis is uncommon

CASE REPORT

A 58/m patient complained of diffuse swelling on his left great toe since 2 years also a yellowish discoloration of great toe nail (figure 3). Patient first reported in department of general surgery owaisi hospital where he was referred to department of orthopaedics for further management. Here the patient was advised deep tissue biopsy with evaluation of fungal elements as he is suspected for deep fungal infection (figure 2). Patient is a k/c/o dm type 2 since 2 years .

On examination : soft tissue swelling around the left great toe predominantly over the plantar aspect and measures around 5.7x3.6cm. Swelling shows smooth outerwalls with internal irregular hypodense area, suggesting variable thick fluid and multiple irregular densities.

The subcutaneous fat plane over the soft tissue swelling show variable enhancement.

The proximal 3rd of distal phalanx of the great toe show small area of extrinsic erosion over the plantar aspect.

Rest of bones of foot and toes as well as surrounding soft tissue planes appear normal.

XRAY LEFT FOOT showed lytic lesion on the distal phalanx of left great toe. Cortical destruction along the soft tissue extension. Periosteal reaction was absent. It is also confirmed by 3D CT reconstruction of left foot (fig 4)

All routine haematological biochemical and xray chest were within normal limits. Surgeon decided to excise the tumour at the metatarso-phalangeal joint after discussion with the patient.

On gross examination a glistening tumor measuring 5x3 cm was found having marked myxoid changes. Microscopically chondroid neoplasm with prominent chondroid stroma in lobulated pattern. Tumour involved the surrounding skeletal muscle and soft tissue.

18 F- FDG PET CT SCAN of whole body was done which reveals post operative case of chondrosarcoma of left great toe with no other significant active disease in the whole body (fig 5)

Patient underwent 3 cycles of chemotherapy. No local and distant recurrence after 1 year follow up



PREOPERATIVE IMAGE

figure 2



INTRAOPERATIVE IMAGE

figure 3



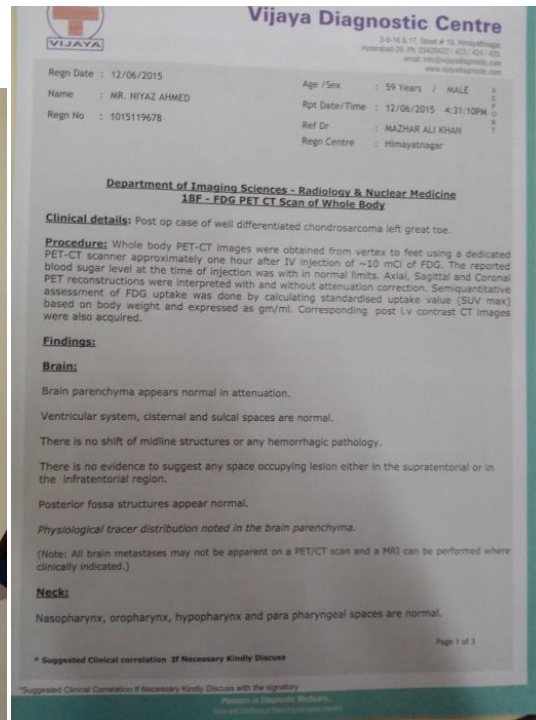
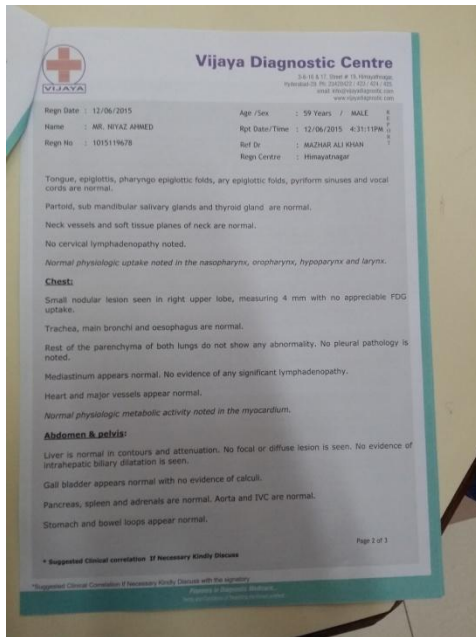
PREOPERATIVE IMAGE LATERAL VIEW

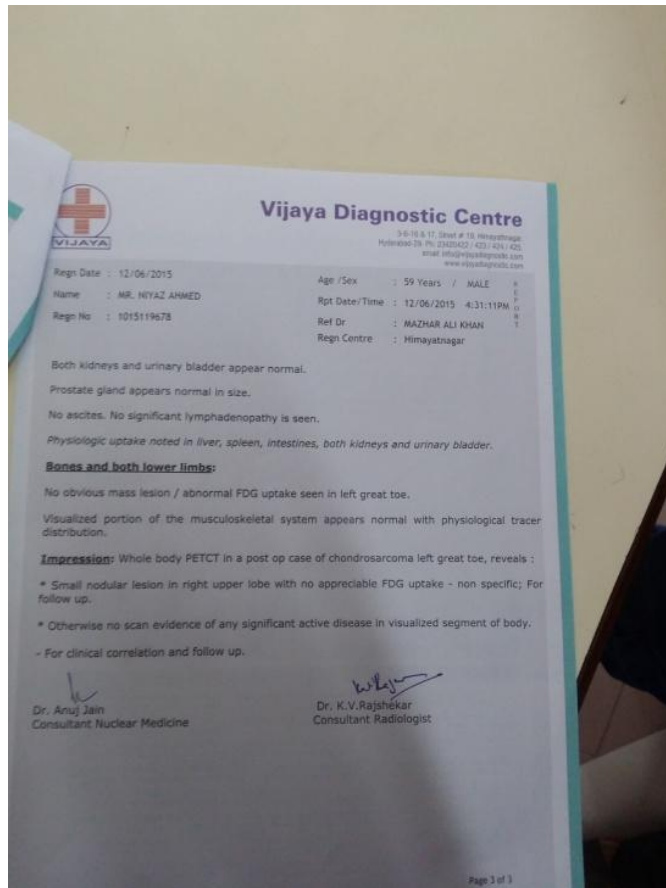
figure 4



3D CT RECONSTRUCTION IMAGE OF LEFT GREAT TOE

Figure 5





PET SCAN OF WHOLE BODY DISCUSSION

Chondrosarcoma is a malignant bone tumor containing tumor cells that produce cartilage. This tumor may form de novo as a primary lesion or from malignant transformation of pre-existing benign condition or cartilage lesion. Ogose *et al.* [4] reviewed 163 Chondrosarcoma located in the phalangeal, (meta) carpal, and (meta) tarsal bones of the hands (n=88) and feet (n=75) and suggested that these tumors have the potential to be fatal. Chondrosarcoma of the calcaneus and the talus were more likely to metastasize. [4] Kinoshita *et al.* reviewed 83 cases of bone and soft tissue tumors of the foot. They found 33 benign tumors, one primary Chondrosarcoma and 2 metastatic bone tumors. [5] For the only malignant bone tumor (Chondrosarcoma of left talus) below-knee amputation was performed but the patient (52-year old man) died because of pulmonary metastasis. [6] Patil *et al.* studied 12 Chondrosarcoma of the bones of feet, of which 4 tumors affected the tarsal bones and the rest involved the short tubular bones. The mean age of the patients was 52.3 years (range 17--83 years) and men were predominantly affected. [6]

Chondrosarcoma are graded according to the system of Evans *et al.* [7] Histologic parameters for grading include cellularity (high/moderate/low), binucleated cells (<1/1--5/>5 per high power field), cellular distribution (regular/irregular), nuclear pleomorphism (low/moderate/high), bone formation (absent/focal/diffuse), differentiation (percentage chondroid, mucoid and myxoid differentiation), calcification (absent/focal/diffuse) and cortical destruction (absent/present/unobservable). [8] In a series of 35 cases of phalangeal Chondrosarcoma, Bovee *et al.* reported 8 grade 1, 26 grade 2, and 1 grade 3 Chondrosarcoma. Median age of phalangeal Chondrosarcoma was 67 years (21--37 years) with a slight predilection for females (21 vs.14). The median duration of the disease was 18 months (range 1-480 months). [2] But the present case was a male patient of age only 37 years with a disease duration of 2 years (24 months). In this case the greatest diameter of the tumor was 3.5 cm which corroborates the median diameter of 3 cm found by Bovee *et al.* [2] Though nuclear pleomorphism, binucleation, and cellularity were not high in the present case, but cortical destruction and soft tissue extension were obvious. These findings are important to differentiate grade 1 Chondrosarcoma from enchondroma, because enchondroma of the hands and feet display greater cellularity and mild nuclear atypia than is seen elsewhere. [2],[4]

Phalangeal Chondrosarcoma occurs far more frequently in the hand compared with the foot and majority of the cases are located in the proximal phalanx.^[2] In contrast to Chondrosarcoma located elsewhere, phalangeal Chondrosarcoma appears to have female preference. Furthermore, the median age (67 years) was higher than for Chondrosarcoma in general (grade 1:44 years, grade 2:55 years).^[2]

Although for all grades and subtypes of nonmetastatic Chondrosarcoma, complete surgical treatment only offers the chance for cure, the most optimal type of surgical management is still debated. Wide, en bloc excision is the preferred surgical treatment of intermediate and high grade Chondrosarcoma.^[9] For the low-grade/grade 1 Chondrosarcoma, some prefer extensive intralesional curettage followed by local adjuvant treatment, for example, phenolization or cryosurgery (liquid nitrogen) and filling the cavity with bone graft.^[10] Below-knee amputation was also advocated by some authors. In our case excision of great toe at the metatarso-phalangeal joint was done and 1-year follow-up was uneventful.

CONCLUSION

Phalangeal Chondrosarcoma behaves as locally aggressive lesion and rarely metastasize. Curettage with local adjuvant and adequate follow-up can be tried as the first-line management, when compared to amputation which leads to functional loss.

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