INTRODUCTION

Osteosarcoma is the most common malignant primary bone tumour in children and adolescents. It is the sixth leading cancer in children >15 years of age. There are approximately 400 new cases each year in the US. In 1805, the French surgeon Alexis Boyer (personal surgeon to Napoleon) first used the term osteosarcoma. Boyer realized that osteosarcoma is a distinct entity from other bone lesions, such as osteochondromas (exostoses). Osteosarcoma affects males more frequently than females, with a ratio of 1.6:1. It occurs early in females due to the earlier onset of growth spurts. The most frequent sites of origin are the metaphyseal regions of the distal femur, proximal tibia and proximal humerus, although the tumor can develop in any bone. Several histological subtypes of osteosarcoma exist. These include osteoblastic (the most common), chondroblastic, fibroblastic, telangiectatic, small cell, parosteal, periosteal, high-grade surface and secondary osteosarcoma. Pain and swelling are the major symptoms of osteosarcoma. It is estimated that about 80% of patients have micrometastatic disease at the time of diagnosis, though in only 10-20% can this be initially detected by standard imaging modalities. Chest CT is more sensitive in detecting pulmonary metastases and has become the imaging procedure of choice. Time of identification of lung metastasis is an important prognostic factor. Factors that seem to negatively impact prognosis are site (axial locations fare worse), larger tumor size, poor response to chemotherapy and presence of metastatic disease. The most consistent and clinically relevant of these is presence of detectable metastases. The 5-year event-free survival rate of patients with localized disease is up to 70%, but the prognosis for patients who present with metastatic disease is still poor, with a 5-year survival rate of approximately 30%. Deaths from osteosarcoma is usually the result of progressive pulmonary metastasis with respiratory failure due to widespread disease.
This study was designed to determine the incidence of pulmonary metastasis detected at the time of diagnosis of primary osteosarcoma distal femur.

**SUBJECTS AND METHODS**

A total of nine patients of both gender and all ages with suspected primary osteosarcoma distal femur (pain, swelling) were admitted through Out Patient Department (OPD) of Orthopaedic Surgery Unit, Mardan Medical Complex Teaching Hospital, Bacha Khan Medical College, Mardan, KPK, Pakistan from March 2011 to September 2012. All patients with secondary osteosarcoma, and those receiving chemotherapy or any surgical intervention were excluded from the study. The study protocol was approved by the Ethics Committee of the hospital. Informed written consent was obtained from parents of all the patients participating in the study. In all the included subjects complete history and clinical examination were done. X-ray distal femur, MRI, bone scan and CT chest was done to stage the tumour\(^{21,22}\). The presence of lung metastasis was assessed by radiologists using CT with a standard cutting distance of 5 mm. The frequency and patterns of pulmonary metastasis on CT chest was recorded. Incisional biopsy was taken under general anaesthesia. The histological types of osteosarcoma were noted and patients were referred to neo adjuvant chemotherapy before any definite surgical procedure. The data was analyzed with SPSS version 11 and represented in a table where necessary.

**RESULTS**

Nine patients including 7 males (77.7%) and 2 females (22.2%) with mean age 12.4 years (Range 8 to 19 years) were included in our study. Majority (n=6, 66.6%) of the patients had right sided distal femur tumour while three (33.3%) patients had left sided involvement. Symptoms referable to the primary osteosarcoma were present for an average of 3 months (range, 3 week to 7 months) prior to diagnosis. 66.6 % (n=6, 5 males, 1 female) had pulmonary metastasis on CT chest at initial presentation while 33 % (n=3, 2 males, 1 female) had no pulmonary metastases on CT chest. The frequency and distribution of pulmonary metastasis on CT chest is shown in Table-I. The lung nodules were almost always located in the peripheral lungs or subpleural regions; in cases of multiple pulmonary nodules, intraparenchymal nodules occurred in 3 cases. The distribution of histologic subtypes of primary tumors comprised osteoblastic (n=5, 55.5%), chondroblastic (n=2, 22.2%), telangiectatic (n=1, 11.1%) and fibroblastic (n=1, 11.1%). Metastatic disease at diagnosis was significantly associated with histologic subtype of osteosarcoma. Of the cases with metastases at diagnosis, 55.5% had osteoblastic histology compared with 33.3% of those with non metastatic disease. Majority (n=7, 77.7%) of the patients had tumour size of more than 10 cm (Five patients had pulmonary metastasis) while in only two (22.2%) patients tumour size was less than 10 cm (One patient had pulmonary metastasis). Three patients (33.3%) had palpable inguinal lymph nodes. One (11.1%) patient had pathological fracture at initial presentation.

<table>
<thead>
<tr>
<th>Pulmonary Metastasis</th>
<th>No. of patients</th>
<th>%age</th>
</tr>
</thead>
<tbody>
<tr>
<td>Right upper lobe</td>
<td>1</td>
<td>16.6%</td>
</tr>
<tr>
<td>Right lower lobe</td>
<td>2</td>
<td>33.3%</td>
</tr>
<tr>
<td>Left upper lobe</td>
<td>1</td>
<td>16.6%</td>
</tr>
<tr>
<td>Both lungs (All lobes)</td>
<td>2</td>
<td>33.3%</td>
</tr>
<tr>
<td>Solitary pulmonary nodule</td>
<td>2</td>
<td>33.3%</td>
</tr>
<tr>
<td>Multiple pulmonary nodules</td>
<td>4</td>
<td>66.6%</td>
</tr>
<tr>
<td>Nodular calcifications</td>
<td>4</td>
<td>66.6%</td>
</tr>
</tbody>
</table>

**DISCUSSION**

Osteosarcoma is the most common primary malignant bone tumor in children and adolescents\(^{23}\).
The most frequent site for metastatic presentation is the lung\textsuperscript{23}. The development of pulmonary metastases is a factor of poor prognosis and most of untreated patients died within 6 months of diagnosis\textsuperscript{25}. In our study 66.6\% (n=6, 5 males, 1 female) had pulmonary metastasis on CT chest at initial presentation (one patient had simultaneous bone and pulmonary metastases) while 33\% (n=3, 2 males, 1 female) had no pulmonary metastases on CT chest. Of 1,765 patients with newly diagnosed, previously untreated high-grade osteosarcomas of bone registered in the neo adjuvant Cooperative Osteosarcoma Study Group studies before 1999, 202 patients (11.4\%) had proven metastases at diagnosis\textsuperscript{26}. This study also noted that only multiple metastases at diagnosis (relative hazard rate [RHR] = 2.3) and macroscopically incomplete surgical resection (RHR = 2.4) remained significantly associated with inferior outcomes. Another study on 259 patients with primary osteosarcoma referred to St. Jude Children's Research Hospital from March 1, 1962 through December 31, 1989, 45 (17\%) had metastatic disease at diagnosis. The 31 patients with lung metastases at diagnosis of osteosarcoma ranged in age from 6 to 21 years (median, 15 years). There was a preponderance of male patients (ratio, 2:1); 24 patients were white, and 7 were black\textsuperscript{13}.

Similar to our study the Brazilian Osteosarcoma Treatment Group reported a very high incidence rate (n=37, 78.7\%) of pulmonary metastases at diagnosis while six (12.8\%) had simultaneous bone and pulmonary metastases\textsuperscript{27}. Munajat and Zulmi reported that of the 70 patients with osteosarcoma, 33 (47\%) had evidence of lung metastasis, whereas 37 (53\%) did not. An increase in tumour volume represented an increase in the chance of lung metastasis with a positive predictive value of 69\%. The rates of lung metastasis were 34\% and 69\% in patients with the tumour volume of <371 cm\textsuperscript{3} and \geq 371 cm\textsuperscript{3}, respectively. They concluded that larger tumours are more likely to correlate with lung metastasis\textsuperscript{28}. This is similar to our study findings as majority (n=7, 77.7\%) of our patients had tumour size of more than 10 cm (Five patients had pulmonary metastasis) while in only two (22.2\%) patients tumour size was less than 10 cm (One patient had pulmonary metastasis). Hiroyuki and Yoshimitsu\textsuperscript{14} identified 46(16.4 \%) patients with lung metastasis at initial presentation in their study while Kuei Wu and Chen\textsuperscript{29} reported an incidence rate of 10.2 \% of pulmonary metastasis at diagnosis with 5-year survival rate significantly worse in patient with more than one lobe involved (27.0, P=0.006) and more than three pulmonary nodule metastases (21.3\%, P = 0.002).

Metastatic disease at diagnosis was significantly associated with histologic subtype of osteosarcoma. In our study, 55.5\% patients had osteoblastic histology compared with 33.3\% of those with non metastatic disease. Kaste and Pratt documented that cases with metastases at diagnosis, 53\% had osteoblastic histology of osteosarcoma compared with 34\% of those with non metastatic disease (P=0.049)\textsuperscript{30}.

The rate of incidence of computed tomography detected pulmonary metastases was found to be 14\% (31 of 215 patients) at diagnosis in this study.

Symptoms referable to the primary osteosarcoma were present for an average of 3 months (Range, 3 week to 7 months) prior to diagnosis in our study and this delayed presentation might be responsible for such a higher incidence of pulmonary metastasis detection (66.6 \%) in our study.

Despite the strengths of our study, a few limitations deserve mention.

Our sample size may not be large enough. As very little research data on this topic is available so far in this country. The need for more research cannot be over emphasized.

**CONCLUSIONS**
Majority of osteosarcoma distal femur presented with pulmonary metastasis at initial presentation. The absence of signs of malignancy on the first plain film radiographs and the later development of the osteosarcoma emphasize the importance of a detailed history and further imaging. It is necessary to reduce the interval between the onset of signs and symptoms and the diagnosis of cases of osteosarcoma thus increasing the chances of survival for these patients. For this, it is essential that orthopaedic surgeons be aware of the characteristic early signs and symptoms, such as pain, increase of local volume, fever, and in some cases, the original pathological fracture.

Copyright © 15 Jan, 2013.

REFERENCES


AUTHOR(S):

1. DR. FAAIZ ALI SHAH
FCPS (Orthopaedics)
Senior Registrar, Orthopaedic Unit
Mardan Medical Complex Teaching Hospital
Bacha Khan Medical College, Mardan KPK

2. DR. ABDUL AZIZ ZIA
DMRD (Radiology)
Associate Professor/Incharge Department of Radiology
Mardan Medical Complex Teaching Hospital
Bacha Khan Medical College, Mardan KPK

3. DR. ZAHIR KHAN
FCPS (Orthopaedics)
Medical Officer Orthopaedic Unit
Mardan Medical Complex Teaching Hospital
Bacha Khan Medical College, Mardan KPK

4. Dr. Kifayatullah
FCPS (Orthopaedics)
Associate Professor Orthopaedic Unit
Mardan Medical Complex Teaching Hospital
Bacha Khan Medical College, Mardan KPK

Correspondence Address:
Dr. Faaiz Ali Shah
faaizalishah@yahoo.com

Article received on: 02/10/2012
Accepted for Publication: 15/01/2013
Received after proof reading: 01/02/2013