Case Report

Clinicopathological representation of nonmetastatic Ewing sarcoma of the scapula - a case study

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INTRODUCTION

Ewing sarcoma (ES) is characterized as a small undifferentiated round cell sarcoma which evolves most prominently in bone and rarely occurs in extra skeletal soft tissues.¹ Ewing sarcoma of the bone develops as a primary sarcoma, whereas secondary sarcoma is the one that metastasize to the bone (bone metastases) from other parts of the body. Ewing sarcoma exhibits rapid growth and metastasize on bone and other soft tissues of the body, through blood. Extra skeletal metastasis of ES primarily occurs in lungs.² Ewing sarcoma occurs in children and adults and is most prevalent among people under 30 years with a median age of 20. It occurs predominantly in males than in females.³ The epidemiology of ES has found to be related to race. It is significantly prevalent in whites and is less among Africans and Asians.⁴ Ewing sarcoma arises as a result of fusion of the oncogene, EWS (Ewing sarcoma) of chromosome 22 with one of the several members of ETS (Ewing tumors) family. The translocation {t (11; 22) (q24; q12)} is the most common which results in the fusion of EWS with FLI1 (friend leukemia integration 1) of ETS family.⁵ A differential diagnosis with a combination of techniques is required to diagnose ES. Here we present a case featuring non-metastatic primary Ewing sarcoma of the scapula.

CASE REPORT

The patient is a seven year old female with swelling on the left shoulder which gradually increases by time. The
patient doesn’t have complaint of pain or pain during shoulder movement. At the time of examination, the patient had no fever, trauma, asthma or epilepsy. The patient was also found to be negative for the clinical conditions, pallor, icterus, clubbing and edema. The patient’s appetite, bowel movement, and bladder function was normal with no history of weight loss. Absence of organomegaly and no focal neurological deficit has been reported. Local examination revealed a 5×4 cm mass over the left scapular region. The MRI scan has shown altered signals involving lateral border and body of left scapula.

**Histopathology and immunohistochemistry**

The bone marrow aspiration and biopsy (1.5 cm) tissue has been processed and examined for malignancy using standard protocols. Typical small round and oval cells with, densely packed nuclei and scanty cytoplasm has been observed microscopically after staining with hematoxylin and eosin (H&E). The sample has shown positive for CD99, vimentin, and negative for the other biomarkers (Table 1) which concludes the presence of ES.

![Figure 1: Whole body bone scan showing a mass on the left scapula.](image)

![Figure 2: H&E staining- tightly arranged tumor cells with hyperchromatic nuclei and some inconspicuous needle –like osteoid particles magnification X 40.](image)

<table>
<thead>
<tr>
<th>Biomarker</th>
<th>Result</th>
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<tbody>
<tr>
<td>Keratin</td>
<td>Negative</td>
</tr>
<tr>
<td>NSE (neuron specific enolase)</td>
<td>Negative</td>
</tr>
<tr>
<td>Synaptophysin</td>
<td>Negative</td>
</tr>
<tr>
<td>Chromogranin A</td>
<td>Negative</td>
</tr>
<tr>
<td>Vimentin</td>
<td>Positive</td>
</tr>
<tr>
<td>CD56</td>
<td>Negative</td>
</tr>
<tr>
<td>CD57</td>
<td>Negative</td>
</tr>
<tr>
<td>CD 31</td>
<td>Negative</td>
</tr>
<tr>
<td>CD 34</td>
<td>Negative</td>
</tr>
<tr>
<td>CD99</td>
<td>Positive</td>
</tr>
<tr>
<td>KI 67</td>
<td>Negative</td>
</tr>
</tbody>
</table>

![Figure 3: IHC vimentin is more distinct. magnification X 10.](image)

![Figure 4: IHC reaction with CD99 is positive. magnification X 40.](image)

**Cytogenetics**

Peripheral blood sample of the patient was collected and used for chromosomal analysis. The mitotic indices were analysed by G-banding after trypsinization using standard protocol. The peripheral blood of the patient has shown a normal karyotype (Figure 5).
DISCUSSION

About 50% of ES occurs in the femur and pelvis and ES of scapula is one of the rare cases. The diagnosis of ES is difficult due to its characteristic similarity with other malignant tumors. A gradual increase of pain is one of the major symptoms of bone sarcomas. In the present case, the patient doesn’t have a complaint of pain. This may be due to the early stage of the disease condition. A well-defined information of the tumor including its expansion, relation to the adjacent blood vessels, skin, etc., can be obtained by MRI. Our MRI reports a firm, consistent, non-tender, non-adherent tumor on the left scapula. It has been found that CD99 is expressed on the cell membranes of Ewing sarcoma tumor and is therefore used as an immunohistochemical marker for the diagnosis of Ewing tumors. Vimentin is an intermediate filament (IF) protein which is overexpressed at the time of accelerated tumor growth. In the present study a positive reaction to CD99, vimentin and characteristic small round cell morphology of H&E staining concludes the diagnosis of ES. A negative result to the markers over-expressed by other cancer cells like, CD57 (leukemia, carcinoma etc.), CD56 and chromogranin A (neuroendocrine tumors), keratin (melanoma), neuron specific enolase (NSE) (neuroblastoma, small cell carcinoma of the lung, etc.), CD31 and CD 34 (vascular tumors) Ki67 (breast cancer) synaptophysin (neuroblastoma) is also a diagnostic criteria for ES. An additional translocation [t (11; 22) (q24; q12)] has been reported in other studies as a supportive data. Due to the non-availability, chromosomal analysis of bone marrow sample is not done in our study. Chromosomal analysis of the peripheral blood is done as a routine process in our institute. A normal karyotype reveals information on the absence of other chromosomal disorders like down syndrome.

CONCLUSION

Ewing sarcoma is a rare type of bone sarcoma for which a combination of methods are required for clinical diagnosis. Histopathological examination of bone marrow biopsy sample plays a vital role in the definitive diagnosis as it is the quick and reliable method. The limitation of this case study is that the cells of the bone marrow would have been karyotyped as an additional confirmatory test. Early diagnosis of ES and distinguishing it from other small cell tumors facilitates better management of treatment strategies.

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