Gnathic Osteosarcoma: A Retrospective Analysis over a 20 Year Period
Prabhu S,Jose M,Iyengar S

ABSTRACT
Background
Osteosarcomas are very rare malignant mesenchymal neoplasms affecting jaw bones. Only few studies are reported about this lesion in this particular region.

Objective
We wanted to assess the demography and tumor behaviour and histopathological pattern, retrospectively in the available cases.

Methods
A retrospective analysis of hospital records for 20 years was carried out. Clinical features, radiographic findings, histopathological details and treatment aspect of each of the lesions was considered.

Results
Findings of this study revealed only 13 cases of gnathic osteosarcomas, usually occurring in middle age, with equal sex predilection and predominantly seen in maxilla. Histopathologically, most of them belonged to osteogenic variant.

Conclusion
Only few cases of gnathic osteosarcomas are reported in this region mainly in the young males. Interestingly many of them has history of dental extraction before the onset of disease.

KEY WORDS
Demography, osteosarcoma, retrospective analysis

Corresponding Author
Sudeendra Prabhu
Department of Oral Pathology
Yenepoya Dental College
Yenepoya University
Mangalore

1Department of Oral Pathology
Yenepoya Dental College
Yenepoya University
Mangalore

2Department of Oral Pathology
SDM College of Dental Sciences
Dharwad

Citation
INTRODUCTION

Osteosarcomas (OS) are tumors associated with bone which are characterized by the direct formation of bone/osteoid by tumor cells.\textsuperscript{1} OS of jaw bones is very rare and it comprises of only 6.5% of total OS as a whole.\textsuperscript{2} Demographics and behaviour of OS are completely different compared to classic OS of long bones.\textsuperscript{3} The demographic data of OS varies in different population.\textsuperscript{4,4} We wanted to assess the clinico-pathologic correlation of OS in Indians and correlate with other races.

METHODS

The case files of patients histologically diagnosed to have an osteosarcoma treated in SDM College of Dental Sciences, Dharwad, Karnataka, India, from 1990 to 2010, were retrieved and information about these lesions was documented. The sample involves all the cases reported to the hospital. The clinical parameters included are age, sex, location, duration, family history, associated symptoms and behavior of the lesion. The radiographic appearance, histological features, treatment and follow up data were also recorded.

RESULTS

A total of 13 cases of osteosarcomas were recorded over the last twenty years period. In general, the age ranged from 4.5 to 70 years with mean age of 36.42 years (Fig 1). The male:female ratio for these patients was 1.1:1 with slight male predilection (seven men and six women). Among all the patients, lesion reported was solitary. Most of the time, these individuals complained of a slow growing swelling of the jaws and facial asymmetry. The average duration of the swelling was 4.8 months with least being 15 days and the longest being two years. In one case, a small swelling was present since three years and it started just after extraction of a tooth. But its growth was rapid for last six months. None of these cases had any family history of similar lesions. However, interestingly four of these patients (30%) had a history of extraction of the tooth in the same location. One each of the patients had a history of retinoblastoma in the same region, anti-tubercular drug and radiotherapy.

Tumor cells were plump, round to ovoid or spindle shaped with marked cellular atypia, hyper chromatism and pleomorphism and most of the time, the nucleus was vesicular. Tumor cells were dispersed in between the osteoid, with few areas undergoing mineralisation. Areas of chondroid and fibrous tissue were also evident. Stroma was fibro-cellular, few with high vascularity. Areas of hemorrhage and necrosis were also evident in few of the lesions. The common finding in majority of them was the presence of osteoclast like giant cells. Depending upon predominant secretory material, nine of them are sub divided as osteogenic osteosarcoma (69%) and three of them as chondroblastic osteosarcoma (23%) and one belonging to fibroblastic variant (8%).

DISCUSSION

Osteosarcomas are very rare malignant mesenchymal tumors occurring in the jaws.\textsuperscript{3,4} These lesions commonly seen in fourth decade, though it is reported with a wide age range.\textsuperscript{10} OS of long bones affects mainly adolescents and young adults, where as gnathic OS seen mainly in middle age (third to fourth decade of life).\textsuperscript{11-16} However, the peculiarity in our sample was that OS is mostly seen in young adults (second decade). Similar age group was affected in an Iranian study.\textsuperscript{5}
There was slight male predilection in the present study as was evident in a Srilankan data. Most of earlier work has shown a definite male predominance. On the other hand, no sex predilection was evident in a research work conducted by Jasnau S et al. In only one analysis, female predilection was seen. According to previous results, it is clearly evident, that OS are often seen in mandible compared to that of maxilla. However, in the present sample, maxilla is frequently involved which is a very uncommon finding.

Painful swelling and paresthesia is a typical complaint of the patients in gnathic OS. However, swelling was the complaint in all the cases here, with pain in 60% of the time and paresthesia in only one patient.

Interesting finding here our patients, that many of them had a history of extraction before the onset of the lesion. Such a finding is not reported before. Radiographically, lesion can be osteolytic, osteoblastic or mixture of the two with poorly defined margins. In the present data also, most of them were mixed lesions with both lucency and opacity with irregular borders.

Microscopically, they say that most common subtype is mixed one. However, the opinion is varied in this regard. Most of the time it will be osteogenic osteosarcoma, as seen in the present population. In one of the groups, chondroblastic variant is widespread followed by fibroblastic. In this African population, only 17% of the total number of cases were showing osteogenic osteosarcoma, which is a rare phenomenon. This histological subclassification is clinically significant as chondroblastic and fibroblastic has a better prognosis compared to osteogenic OS.

OS of jaws rarely metastasizes to other parts of the body as seen in the present patients where there was no metastasis. With respect to treatment, controversy still exists regarding the combination of surgery, radiotherapy and chemotherapy. However, in all 13 cases of OS, here were treated with a combination of surgery and radiotherapy.

### Table 1. Demographic features of Osteosarcomas.

<table>
<thead>
<tr>
<th>Sl No</th>
<th>Age (years)</th>
<th>Sex</th>
<th>Site</th>
<th>Clinical features</th>
<th>Radiographic features</th>
<th>Histopathology</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>45</td>
<td>M</td>
<td>Maxilla - left</td>
<td>Swelling since 1 month</td>
<td>Unilocular radiolucency, Resorption wrt 24, 25, 26</td>
<td>Chondroblastic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>2</td>
<td>17</td>
<td>F</td>
<td>Mandible-right</td>
<td>Pain and swelling since 5 months</td>
<td>Mixed opacity and lucency, osteophytes</td>
<td>Osteogenic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>3</td>
<td>50</td>
<td>F</td>
<td>Mandible - left</td>
<td>Swelling, pain, paresthesia</td>
<td>Mixed opacity and lucency, resorption wrt 33-36</td>
<td>Osteogenic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>4</td>
<td>53</td>
<td>M</td>
<td>Maxilla - left</td>
<td>Swelling since 3 years</td>
<td>Mixed opacity and lucency, resorption wrt 23-28</td>
<td>Chondroblastic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>5</td>
<td>30</td>
<td>F</td>
<td>Mandible - right</td>
<td>Swelling and pain since 3 months</td>
<td>Mixed opacity and lucency with discontinuous lower border</td>
<td>Fibroblastic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>6</td>
<td>45</td>
<td>M</td>
<td>Maxilla - right</td>
<td>Swelling since 6-8 months</td>
<td>Mixed opacity and lucency, extension into lateral wall of the orbit, maxillary sinus and naso-ethmoidal complex</td>
<td>Osteogenic (Epitheloid)</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>7</td>
<td>27</td>
<td>M</td>
<td>Maxilla - right</td>
<td>Swelling (history of retinoblastoma in the same region)</td>
<td>Mixed opacity and lucency with obliteration of right orbit</td>
<td>Osteogenic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>8</td>
<td>70</td>
<td>M</td>
<td>Maxilla - Left</td>
<td>Swelling since 1 year</td>
<td>Diffuse radiolucency</td>
<td>Osteogenic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>9</td>
<td>30</td>
<td>M</td>
<td>Maxilla - right</td>
<td>Swelling and pain since 3-4 months</td>
<td>Diffuse radiolucency</td>
<td>Chondroblastic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>10</td>
<td>40</td>
<td>F</td>
<td>Maxilla - left</td>
<td>Swelling and pain since 1 month</td>
<td>Diffuse opacification involving right maxillary sinus</td>
<td>Osteogenic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>11</td>
<td>22</td>
<td>M</td>
<td>Mandible - left</td>
<td>Swelling and pain</td>
<td>Mixed opacity and lucency, ground glass appearance, extending till angle of the mandible</td>
<td>Osteogenic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>12</td>
<td>24</td>
<td>F</td>
<td>Maxilla - right</td>
<td>Swelling since 3 months</td>
<td>Mixed opacity and lucency</td>
<td>Osteogenic</td>
<td>Excision with radiotherapy</td>
</tr>
<tr>
<td>13</td>
<td>60</td>
<td>F</td>
<td>Maxilla - left</td>
<td>Swelling and pain since 6 months</td>
<td>Diffuse lucency extending till tuberosity</td>
<td>Osteogenic</td>
<td>Excision with radiotherapy</td>
</tr>
</tbody>
</table>
CONCLUSION

The outcome of this study suggests that a very few osteosarcomas reported in this area (0.2% of total biopsies). Most of the subtype belongs to osteogenic osteosarcoma (70%). The common age group is young age (3rd decade) with slight male predilection. Interesting thing in our study is many cases had a history of extraction before the present lesion.

REFERENCES