Cutaneous Metastasis of Chondrosarcoma: A Case Report With Literature Review

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The metastasis to respiratory system secondary to chondrosarcoma is a common finding; however, metastasis to other organs such as skin or bones is much less common. In the current report, we described a case with the history of chondrosarcoma of the mandible that recently referred with the metastatic lesions in her scalp skin. Our case is a female patient that its secondary metastasis occurred only 4 months after her initial tumor diagnosis. More interestingly, among all baseline laboratory parameters, only inflammatory biomarkers increased as the nonspecific diagnostic indices. In other words, the accurate diagnosis of metastasis to bone following chondrosarcoma may be delayed and even masked with early inflammatory reactions. On the whole, in all patients who suffer from chondrosarcoma, early metastasis to skin or bones should be considered, especially when inflammatory indices are high.

ABSTRACT

The metastasis to respiratory system secondary to chondrosarcoma is a common finding; however, metastasis to other organs such as skin or bones is much less common. In the current report, we described a case with the history of chondrosarcoma of the mandible that recently referred with the metastatic lesions in her scalp skin. Our case is a female patient that its secondary metastasis occurred only 4 months after her initial tumor diagnosis. More interestingly, among all baseline laboratory parameters, only inflammatory biomarkers increased as the nonspecific diagnostic indices. In other words, the accurate diagnosis of metastasis to bone following chondrosarcoma may be delayed and even masked with early inflammatory reactions. On the whole, in all patients who suffer from chondrosarcoma, early metastasis to skin or bones should be considered, especially when inflammatory indices are high.

Introduction

Chondrosarcoma is one of the most frequent malignant tumors of the skeletal system accounting for about 20% of all types of sarcomas [1]. This tumor frequently appears in the early fourth decade of life with the male predominance. The pattern of the tumor progression, metastasis, is directly related to tumor grading so that metastasis of chondrosarcoma is infrequent in low-grade category (grades I to II). While half of the patients with grade III and almost all those with a dedifferentiated form of cancer suffer from metastatic lesions [2].

In the patients with high-grade chondrosarcoma, the metastasis to respiratory system is a common finding, and less so in the regional lymph nodes and the liver [3].

However, metastasis to other tissues such as skin or bones is much less common. In the current report, we presented

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a case with the history of chondrosarcoma of the mandible that recently referred with the scalp metastatic lesions.

**Case Presentation**

Our case is a 39-year-old woman with a history of mandibular chondrosarcoma four months before admission. In her pathological report on the time of tumor diagnosis, mandibular lytic bone lesions were macroscopically appeared. In the microscopic examination, foci with marked cellularity, pleomorphism, and bizarre tumor giant cells with mitotic pattern were seen confirming conventional mandibular chondrosarcoma grade II. The patient was referred to our hospital four months after the first assessment with the bony pain, especially at the scalp point. In laboratory analysis on admission, in-

<table>
<thead>
<tr>
<th>Author (y) Reference</th>
<th>Age (y) / Gender</th>
<th>Primary Tumor</th>
<th>Metastatic Region(s)</th>
<th>Interval Period*</th>
<th>Survival Length**/ Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cruickshank (1945) [8]</td>
<td>66/M</td>
<td>Middle pharynx</td>
<td>Face, neck, trunk</td>
<td>26 mon</td>
<td>Unknown</td>
</tr>
<tr>
<td>Froimson (1967) [9]</td>
<td>30/M</td>
<td>Fibular head</td>
<td>Lung, right thumb, left ring finger</td>
<td>NS</td>
<td>3 mon</td>
</tr>
<tr>
<td>King (1978) [10]</td>
<td>31/F</td>
<td>Scapula</td>
<td>Vulva, right thumb, right middle toe, scalp, heel, gingiva</td>
<td>30 mon</td>
<td>2 mon/DOD</td>
</tr>
<tr>
<td>Sherr (1986) [12]</td>
<td>47/F</td>
<td>Left ischium</td>
<td>Nose, supraorbital region, buttock, hemipelvectomy scar</td>
<td>103 mon</td>
<td>5 mon</td>
</tr>
<tr>
<td>Karabela-Bouropoulou (1986) [13]</td>
<td>41/F</td>
<td>Right thumb (proximal phalanx)</td>
<td>Abdomen, chest, frontal</td>
<td>68 m</td>
<td>2 mon/DOD</td>
</tr>
<tr>
<td>Amadio (1987) [14]</td>
<td>32/M; 59/M</td>
<td>Humerus; Femur</td>
<td>Left thumb; Left thumb</td>
<td>NS</td>
<td>5 mon/NS</td>
</tr>
<tr>
<td>Leal-Khouri (1990) [16]</td>
<td>85/M</td>
<td>Left shoulder</td>
<td>Both lungs, nose</td>
<td>115 mon</td>
<td>8 mon/DOD</td>
</tr>
<tr>
<td>Lambert (1992) [17]</td>
<td>36/M</td>
<td>Right femur</td>
<td>Lung, bilateral ring finger</td>
<td>19 mon</td>
<td>5 mon/DOD</td>
</tr>
<tr>
<td>Malek (1994) [7]</td>
<td>66/F</td>
<td>Right tibia</td>
<td>Forearm</td>
<td>84 mon</td>
<td>2 wk/alive</td>
</tr>
<tr>
<td>Damron (1995) [18]</td>
<td>76/F</td>
<td>The fifth metacarpal bone</td>
<td>Forearm</td>
<td>6 mon</td>
<td>6 mon/alive</td>
</tr>
<tr>
<td>Aramburu-Gonzalez (1999) [19]</td>
<td>64/M</td>
<td>Forearm soft tissue</td>
<td>Face</td>
<td>15 mon</td>
<td>28 mon/alive</td>
</tr>
<tr>
<td>Arce (2000) [20]</td>
<td>69/M</td>
<td>Right humerus</td>
<td>Right inguinal, infra mammary</td>
<td>Initial present with metastasis</td>
<td>21 d/DOD</td>
</tr>
<tr>
<td>Ozcanli (2006) [21]</td>
<td>47/M</td>
<td>Proximal part of humerus</td>
<td>Both hands</td>
<td>24 mon</td>
<td>18 mon/DOD</td>
</tr>
<tr>
<td>Stanisz (2012) [22]</td>
<td>51/M</td>
<td>Left os ileum</td>
<td>Lung, left atrium, back, right upper leg</td>
<td>8 mon</td>
<td>&lt;12 mon/DOD</td>
</tr>
<tr>
<td>Dominguez-Durán (2014) [23]</td>
<td>60/M</td>
<td>Larynx</td>
<td>Chin, nose</td>
<td>30 mon</td>
<td>35 mon/DOD</td>
</tr>
<tr>
<td>Our case (2017) 07/01</td>
<td>30/F</td>
<td>Mandible</td>
<td>Scalp</td>
<td>4 mon</td>
<td>24 mon/alive</td>
</tr>
</tbody>
</table>

* Period between primary tumor and metastasis to the skin; **Survival length with skin metastasis. DOD: Died of disease; NS: Not stated.
creased ESR (50mm/h) and positive CRP (3+) were the only finding of the recent manifestation.

Other laboratory parameters were normal. Hence, the patient was candidate for pathological assessment of suspected metastatic lesions of the primary tumor. In gross pathology, frontal lesion, including a piece of irregular skin tissue measuring 0.8×0.3×0.3cm and scalp mass consisting of two pieces of pale-tan brownish tissue measuring 5×3×1cm were revealed. Histopathologic examination revealed well-circumscribed cartilaginous lobules in deep dermis composed of atypical chondrocytes with mild to moderate pleomorphism and irregular hyperchromatic nuclei located in the lacunae within a bluish chondroid matrix (Figure 1). Microscopically, metastatic lesions of chondrosarcoma to the skin in the frontal zone and anterior scalp margin were confirmed. Thus the patient was scheduled for surgical resection with clear margins in combination with chemotherapy and radiotherapy approaches.

Discussion

Metastasis to the lungs due to chondrosarcoma is a common finding, but its metastasis to other organs such as skin or skeletal system rarely occurs. Distant metastasis also develops mostly secondary to primary chondrosarcoma than to secondary type. Moreover, the rate of distant metastasis is higher in those with local recurrence than in those without local recurrence [4]. In conventional chondrosarcoma, the risk of metastasis is directly dependent on the tumor grade so that grade I tumor does not metastasize, while about two-thirds of grade III tumors do. In the dedifferentiated type of tumor, hematogenous metastasis occurs in most patients with a considerable low long-term survival [4].

In some types of tumor such as clear cell chondrosarcoma, distant metastasis can occur even in a low-grade tumor. Reviewing the literature shows rare reports of bone and cutaneous metastatic lesions secondary to chondrosarcoma [4]. In other words, although cutaneous lesions have been recently reported, bone metastasis is very rare. Similarly, Ragsdale et al. reported a 22-year-old male with osteosarcoma of the pelvis and metastasis to the lungs and chest wall who developed a clinically unsuspected solitary cutaneous metastasis in the scalp whose incisional biopsy disclosed a solid tan nodule of chondro-osseous sarcoma [5].

In another report by Shah et al. a 60-year-old man had chondrosarcoma of the lung with two local recurrences, including recurrent cutaneous metastases and skeletal metastasis [6]. Malek et al. reported a case of a solitary subcutaneous metastasis from chondrosarcoma that occurred 7 years after excision of the primary neoplasm [7]. Table 1 presents the published case reports on metastatic chondrosarcoma to skin up until now. Most patients are men (F/M: 6/20), and the primary tumors were mostly located in the peripheral bones.

The interval period between primary tumor and metastasis to skin varies from two weeks to 115 months, and the length of survival varies from two weeks to 66 months. Four of 20 cases were extra-skeletal chondrosarcoma (arisi-
ing from heart, lung, larynx, and soft tissue) and one patient initially presents with metastasis to skin.

Contrary to previous reports and as a more rare case, our case is a young woman whose primary tumor confined to a central bone (mandible), and its secondary metastasis occurred only four months after initial tumor diagnosis. More interestingly, among all baseline laboratory parameters, only inflammatory biomarkers were high as the nonspecific diagnostic indices. In other words, the accurate diagnosis of metastasis following chondrosarcoma may be delayed and even masked with early inflammatory reactions. On the whole, in all patients who suffer from chondrosarcoma, early metastasis to skin or bones should be considered, especially when inflammatory indices are high.

Ethical Considerations

Compliance with ethical guidelines

All ethical principles were considered in this article.

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Conflict of interest

There was no ethical considerations to be considered in this research.

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References


