A primary vertebral epithelioid osteosarcoma in middle age adult

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ABSTRACT

Primary vertebral epithelioid osteosarcoma is an extremely rare malignancy, characterized by predominant epithelioid cells and occurs mainly in adolescents and young adults. To our knowledge, only one previous report described a lumbar vertebral epithelioid osteosarcoma including the clinical information, pathologic features and immunohistochemical profile. This report is a first case of primary epithelioid osteosarcoma arising in the eighth thoracic vertebrae (T8) in middle aged man. Whole body scans as well as histological and immunohistochemical examinations were performed for excluding the possibility of metastatic malignancy or secondary osteosarcoma. Metastatic lesions showing the same morphological and immunohistochemical characteristics were demonstrated at the fifth lumbar vertebra (L5) and sternum were described 1 year after excision and high dose of Methotrexate (MTX) chemotherapy.

Key Words: Epithelioid osteosarcoma, Adult, Spine

1. INTRODUCTION

Osteosarcomas are the most common sarcomas, arising in the long bones metaphysis in adolescents and young adults. These lesions have been histologically classified into subtypes, including epithelioid, chondroblastic, fibroblastic, osteoblastic types etc.[¹] Because osteosarcoma is typically a metaphyseal lesion and occurring most commonly from ages at 10 to 25 years of ages, the vertebra is considered as an unusual site for osteosarcoma. Only one previous case of middle age man’s epithelioid osteosarcoma was reported in second lumbar vertebra.[²] In this report, we describe a first primary epithelioid osteosarcoma case arising in the thoracic vertebrae in middle aged man.

2. CASE REPORT

A 50-year-old man was admitted to Kangwon National University Hospital with sudden back pain. He had no past medical, family, or history of trauma. Radiologically, there was a 3.1 cm sized osteolytic lesion at the left posterior fifth thoracic vertebral body (T8) with focal cortical destruction (see Figure 1A). Whole body PET scanning was performed to exclude the possibility of metastatic condition. A posterior and anterior total enbloc spondylectomy and entire histologic examination were performed. The specimen consisted of multiple tan colored hemorrhagic bony and soft tissue fragments (see Figure 1B). On low power view, the tumor consisted of solid sheets of densely and uniformly

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proliferated cells with destroyed bone trabeculae (see Figure 1C). On high power view, most of the tumor cells showed similar sized epithelioid appearance with a vesicular nucleus and distinct nucleoli. The cytoplasm was fairly eosinophilic and its boundary was indistinct (see Figure 1D). The tumor cells also displayed a high mitotic rate with abnormal mitotic figures. On immunohistochemical (IHC) staining, all tumor cells showed strong immuno-positivity for vimentin, and some cells expressed high molecular weight cytokeratin (HMW-CK), epithelial membrane antigen (EMA) and S-100 protein (see Figure 2A-C), but negativity for Leukocyte common antigen (LCA), Human melanoma black (HMB45) and Murine double minute (MDM2) (see Figure 2D). After surgery, high dose Methotrexate (MTX) chemotherapy was given. However, metastatic lesion with the same microscopic and immunohistochemistry were described at the fifth lumbar vertebra (L5) and sternum after 1 year later.

Figure 1. A: An osteolytic lesion in eighth vertebra (T8) is noted in computed tomogram; B: Solid sheets of tumor cells are present between bony trabeculae (H&E, ×100); C: Destructed bony structure by the tumor sheets (H&E, ×200); D: Tumor cells showing vesicular nuclei and prominent nucleoli and forming osteoid (H&E, ×600).

3. DISCUSSION
A few number of maxillary or mandibular epithelioid osteosarcoma of middle-aged men were reported. However, only one previous primary epithelioid osteosarcoma in the spine was described, presented the clinic-pathological findings. The age and sex of this previously case (49-year-old man) was similar to this case.

In the present case, the radiographic features of the affected spine showed an osteolytic lesion with cortical destruction, just similar to the conventional osteosarcoma. Cytologically, the cells were ovoid to polygonal cells with more or less a uniform size and shape with a conspicuous cytoplasm. Such cells morphologically mimic those of high grade malignancies, including carcinoma, melanoma, metastatic osteosarcoma, and large cell type lymphoma.

Although the immunophenotypic expression of epithelioid tumor cells is variable in previous literature review, the epithelioid tumor cells of the study case were strongly positive for vimentin, and focal positivity for HMW-CK, EMA and S-100 protein, but negative for LCA, HMB45 and MDM2.
The treatment of osteosarcoma is a combination of surgery, chemotherapy, and radiation therapy, but surgical excision was a mainstay of our treatment. Clear resection margins are associated with a considerably higher 5-year survival rate.\textsuperscript{9} Chemotherapy is usually tried after surgery, its effects have yet to be unequivocally demonstrated. Due to a limited number of reports, however, exact prognosis for epithelioid osteosarcoma remains unclear. Nonetheless, some previous studies have reported that osteosarcoma with epithelioid features is associated with an aggressive clinical behavior.\textsuperscript{10, 11} In this case, tumor cells also showed aggressive behavior with multiple bony metastases.

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**CONFLICTS OF INTEREST DISCLOSURE**
The authors declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

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