**Hand Bones Epithelioid Hemangioendothelioma**

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**INTRODUCTION:**
Epithelioid Hemangioendothelioma is a rare tumor of vascular origin with borderline malignancy and pursues a clinical course intermediate between hemangioma and an angiosarcoma. Epithelioid Hemangioendothelioma has been reported more commonly in soft tissues and viscera involving lungs and liver but less so in bones. The tumor is usually multicentric with a predilection for bones of a particular anatomical area. We describe a case of multicentric Epithelioid Hemangioendothelioma of right hand. Roentgenographs revealed multiple lytic lesions involving the metacarpals and phalanges with impending pathological fracture. Histologically the tumor exhibited cords of epithelioid-appearing cells in a myxoid stroma and inflammatory cells with few mitotic figures. The patient was treated with primary surgical resection, curettage, bone grafting and K-wire fixation. Follow-up x-rays revealed good consolidation of bone graft with no signs of local recurrence.

**CLINICAL PRESENTATION:**
An apparently healthy 41 year male presented with complaints of pain and multiple swellings in his right hand for two months. Pain was progressive and dull aching type. Swelling was of gradual onset and increasing in size for the last one month. It interfered with his daily activities as he was unable to hold and lift weight. There was no history of trauma, constitutional symptoms or pain in any other bones or joints. Patient gave no history suggestive of tuberculosis. General examination was normal with no lymphadenopathy or general erythema. There was no complaint and with no signs of local recurrence.

**RADIOGRAPHICS:** Plain radiographs revealed multiple osteolytic lesions in 1st metacarpal, base of 3rd metacarpal and proximal phalanx of ring finger with periarticular osteoporosis. The cortex was expansile and thinned out but intact with impending pathological fracture of the 1st metacarpal.

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**Histopathology:**
Examination of the resected tissue exhibited thin trabeculae with cords and nests of plump epithelioid-appearing cells around blood vessels. The cells embedded in a myxoid-hyaline stroma possessed abundant acidophilic cytoplasm with characteristic intracytoplasmic vacuoles and a large vesicular nucleus. Focal infiltration of inflammatory cells with many eosinophils and few mitotic figures were noted. Histology was consistent with diagnosis of epithelioid hemangioendothelioma of bone.

**Follow-up:** The patient was then screened for similar lesions with x-ray chest, pelvis, left hand and sonography of abdomen and pelvis which did not detect any abnormality. Post operatively the patient had an uneventful recovery and follow-up x-rays revealed good consolidation of the bone graft. At 3-month follow-up the patient had a good functional recovery without any complaint and with no signs of local recurrence.

**DISCUSSION:** The term “Epithelioid Hemangioendothelioma” was first used in 1982 by Weiss and Enzinger1 to soft tissue vascular tumor of borderline malignancy pursuing a clinical course intermediate between hemangioma and an angiosar-
These tumors are well recognized in soft tissues, lungs, liver, skin but less so in bone. In the bone, it was first reported by Tsuneyoshi in 1986 and involvement of the bones like ilium, vertebra, skull and femur have been well-documented. One of its variants, the epithelioid and spindle cell hemangioma is described to occur in small bones of hands and feet.

Epithelioid endothelial cell tumors have been a source of controversy because of their unusual morphology, poorly understood histogenesis and unpredictable biologic behavior. Currently these tumors are classified by WHO as – epithelioid hemangio-oma (angioymphoid hyperplasia with eosinophilia), epithelioid hemangioendothelioma and epithelioid angiosarcoma.

Epithelioid hemangioendothelioma of bone usually presents as pain in the involved region, and gradually increasing swelling with pathological fracture may be seen in few cases. 50% of tumors are multicentric with a predilection for bones of a particular anatomical area. Our patient had this classical presentation.

Radiological findings of lytic lesion with thinning of cortex are similar as seen in cartilaginous tumors like enchondroma. Histopathological examination of the tumor clinches the diagnosis. Epithelioid endothelial cells possessing intracytoplasmic vacuoles and abundant eosinophilic cytoplasm are arranged in cords and form rudimentary vascular spaces within a myxoid stroma. Ultrastructure reveals specific Weibel-Palade bodies. Immunohistochemical reactivity to Factor VIII-related antigen supports the vascular nature of these neoplasms. Q-bend 10 (CD 34) i.e., hematopoietic progenitor cell antigen detected in endothelial cell is positive.

Surgical resection is the primary treatment for these tumors and radiotherapy may be useful for the surgically inaccessible ones. In few cases intensive combination chemotherapyp5 (with vincristine, adriamycin, cyclophosphamide and etoposide) was tried but proved ineffective to halt the progression of the disease. Metastasis if occurs is usually to the lungs and liver where it destroys the involved organ, resulting in death by respiratory compromise and hepatic failure respectively. Mortality is 35% after liver metastasis and 65% in lung metastasis.

CONCLUSION:
The clinical course and outcome of epithelioid hemangioendothelioma varies greatly and may not be predicted by its histological features since few reports demonstrate the aggressive behaviour and metastatic potential of histologically low-grade tumor. Local recurrence is possible. Hence a thorough clinical and radiological evaluation of all patients with epithelioid hemangioendothelioma of bone is advocated at presentation and follow-up to screen for local recurrence and metastatic disease.

Conflict of Interest: None

REFERENCE