

Epithelioid Sarcoma Presenting as Recurrent Thumb Ulcer: A Lesson to Learn

Abstract

Epithelioid sarcoma (ES) first described by Enzinger in 1970,^[1] is a rare soft-tissue sarcoma typically presenting as a subcutaneous or deep dermal mass in distal portions of the extremities of adolescents and young adults. They are frequently mistaken for ulcers, abscesses, or infected warts that fail medical management. Patients often develop multiple local recurrences of long duration, with subsequent metastases in 30%–50% of cases.^[2] We here report a case of left thumb ES that presented as an ulcer and subsequently metastasized to the forearm, arm, axillary lymph nodes, and lungs.

Keywords: Distal type sarcoma, epithelioid sarcoma, hand sarcoma, thumb ulcer

Introduction

Epithelioid sarcoma (ES) is a rare slow-growing soft-tissue sarcoma typically involving the distal extremity of young adults, with a strong propensity for local recurrence and ultimately metastasis. Microscopically, most tumors are characterized by a granuloma-like pattern: nodules of spindle and epithelioid cells circumscribe areas of central hyalinization and necrosis. In 1997, Guillou *et al.* described a different type of ES, called proximal-type ES, which is found mostly in the pelvic and perineal regions and genital tracts of young to middle-aged adults, characterized by a proliferation of epithelioid-like cells with rhabdoid features and absence of a granuloma-like pattern.^[3] These have a worse prognosis than those arising in distal locations.

Case Report

A 32-year-old male presented with complaints of swelling in his left thumb for 2 years duration. The swelling was nontender and progressively increased in size to involve the distal portion of the left thumb. The swelling underwent spontaneous rupture to form an ulcer 2 months before his hospital visit. The ulcer persisted despite multiple debridements and administration of several antibiotics. In November 2015, he underwent excision of the left thumb.

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Histopathological examination revealed a spindle cell neoplasm with epithelioid features. It also showed underlying bone and marrow infiltration by the tumor cells. Immunohistochemistry (IHC) shows diffuse positivity for epithelial membrane antigen and CD34 and focal positivity for cytokeratin and smooth muscle actin. Neoplastic cells were negative for S100. Ki-67 was around 25% [Figure 1].

The patient had not received further treatment following amputation.

In February 2016, he presented with complaint of multiple, hard mobile swellings in his left forearm and arm, with enlarged left axillary lymph nodes [Figure 2]. Excisional biopsy of one of the forearm nodules confirmed a recurrence with the same histology with positivity for same IHC markers as seen in left thumb-amputated specimen. Computed tomography (CT) thorax revealed multiple pulmonary metastases. Left axilla showed an enlarged lymph node of size 4.2×4 cm.

The patient was started on chemotherapy consisting of doxorubicin and ifosfamide. Reassessment CT scan performed after 3 cycles of chemotherapy showed progressive disease. Second-line chemotherapy (gemcitabine and docetaxel) was started but the patient did not tolerate well and hence best supportive care was offered.

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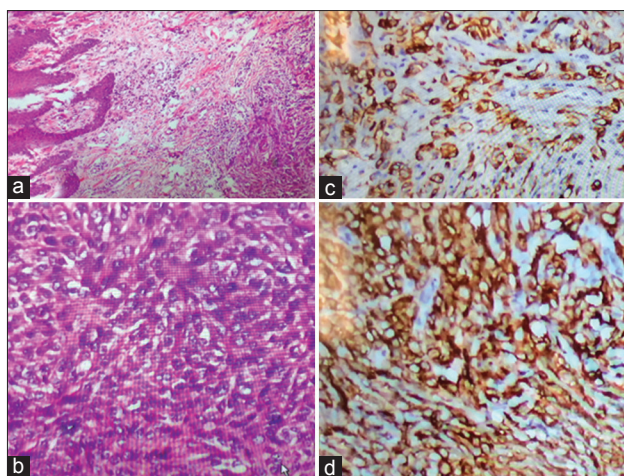


Figure 1: (a) Tumor involving the reticular dermis and subcutaneous tissue (H and E, $\times 10$). (b) Tumor cells are predominantly epithelioid and spindle shaped (H and E, $\times 40$). (c) Immunohistochemistry, tumor cells are cytokeratin positive. (d) Tumor cells are strongly positive for epithelial membrane antigen

Discussion

ES is a rare high-grade malignancy that represents the most common primary soft-tissue sarcoma of the hand. Because of its epithelial and mesenchymal differentiation, this tumor is often mistaken for chronic inflammatory conditions, necrotizing granulomas, and various fibrohistiocytic tumors.^[4] ES often presents in young men, but its significance is great because of an inherent propensity for local recurrence in addition to regional lymph node involvement and distant metastases.^[5] Males are disproportionately affected with a ratio approaching 2:1. ES initially appears as a single nodule. However, at the time of diagnosis, multiple nodules representing local disease spread may be present^[6] as seen in our patient. Periosteal bone invasion can occur.^[1]

Radical tumor excision is the primary treatment for patients with ES. Negative tumor margins must be obtained, but optimal function, particularly of the upper limb, is also to be considered.^[7] Therapeutic lymph node dissection is indicated when lymph node metastases are present. A decreased risk of local recurrence has been reported with the addition of preoperative or postoperative local radiation therapy when combined with radical surgery.^[6] Radiation can also be considered for patients with marginal primary resection, local regional recurrence, or palliative treatment.

Single agent or combination chemotherapy is the treatment offered in patients with Stage IV disease, as in our patient with lung metastasis. The most commonly administered chemotherapy regimens are single-agent anthracycline therapy or the combination of anthracycline with ifosfamide.^[8] A single group reported activity of a regimen combining gemcitabine with docetaxel, but the experience is limited to a small number of patients.^[9]



Figure 2: Clinical picture of the patient showing the partially amputated thumb and multiple cutaneous metastatic nodules over hand, forearm, and arm (arrows)

Despite the administration of palliative chemotherapy, patients with metastasis have poor prognosis. The reported median survival is around 52 weeks and the 1 and 5 year survival rates are 46% and 0%, respectively.^[8] Therefore, it's an unmet need to improve the medical management of ES patients by introducing novel agents.

Male gender, depth of invasion, and vascular invasion are associated with poor prognosis. Anatomic location of ES appears to play a role in prognostication. The overall survival and metastases-free survival are worse in lesions proximal to the elbow or knee.^[10] Close, long-term follow-up is necessary because recurrence and metastases may occur long after definitive treatment.

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Conflicts of interest

There are no conflicts of interest.

References

1. Enzinger FM. Epithelioid sarcoma. A sarcoma simulating a granuloma or a carcinoma. *Cancer* 1970;26:1029-41.
2. Chase DR, Enzinger FM. Epithelioid sarcoma. Diagnosis, prognostic indicators, and treatment. *Am J Surg Pathol* 1985;9:241-63.
3. Guillou L, Wadden C, Coindre JM, Krausz T, Fletcher CD. "Proximal-type" epithelioid sarcoma, a distinctive aggressive neoplasm showing rhabdoid features. Clinicopathologic, immunohistochemical, and ultrastructural study of a series. *Am J Surg Pathol* 1997;21:130-46.
4. Fisher C. Epithelioid sarcoma: The spectrum of ultrastructural differentiation in seven immunohistochemically defined cases. *Hum Pathol* 1988;19:265-75.
5. Spillane AJ, Thomas JM, Fisher C. Epithelioid sarcoma: The clinicopathological complexities of this rare soft tissue sarcoma. *Ann Surg Oncol* 2000;7:218-25.
6. Callister MD, Ballo MT, Pisters PW, Patel SR, Feig BW, Pollock RE, *et al.* Epithelioid sarcoma: Results of conservative surgery and radiotherapy. *Int J Radiat Oncol Biol Phys* 2001;51:384-91.
7. de Visscher SA, van Ginkel RJ, Wobbes T, Veth RP, Ten Heuvel SE, Suurmeijer AJ, *et al.* Epithelioid sarcoma: Still an only surgically curable disease. *Cancer* 2006;107:606-12.
8. Jones RL, Constantinidou A, Olmos D, Thway K, Fisher C, Al-Muderis O, *et al.* Role of palliative chemotherapy in advanced epithelioid sarcoma. *Am J Clin Oncol*

- 2012;35:351-7.
9. Pink D, Richter S, Gerdes S, Andreou D, Tunn PU, Busemann C, *et al.* Gemcitabine and docetaxel for epithelioid sarcoma: Results from a retrospective, multi-institutional analysis. *Oncology* 2014;87:95-103.
 10. Guillou L, Wadden C, Coindre JM, Krausz T, Fletcher CD. "Proximal-type" epithelioid sarcoma, a distinctive aggressive neoplasm showing rhabdoid features. Clinicopathologic, immunohistochemical, and ultrastructural study of a series. *Am J Surg Pathol* 1997;21:130-46.