Ossifying Fibromyxoid Tumor of Soft Parts

Report of a Rare Tumor in Kuwait

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Key Words
Ossifying fibromyxoid tumor of soft parts · Bone tumors · Myxoid tumors

Abstract
Objective: To report a rare case of ossifying fibromyxoid tumor of soft tissue in Kuwait. Clinical Presentation and Intervention: A 60-year-old woman presented with a painless tumor, increasing in size and located in the left buttock, which had been present for an uncertain duration. The patient underwent an excisional biopsy. Pathological examination revealed a 7.5-cm well-circumscribed mass with a lobulated cut surface. Histologically, the tumor was encapsulated by an incomplete shell of lamellar bone. The tumor had variable cellularity, and, in areas, contained myxoid stroma. The tumor cells had eosinophilic cytoplasm with vesicular round-to-oval nuclei. Sparse mitoses were noted. Immunohistochemical stains demonstrated that the tumor cells expressed vimentin, S100 and neuron-specific enolase, with the latter expressed focally.

Case Report

A 60-year-old diabetic and hypertensive woman presented with a painless mass in the left buttock. The tumor was slowly increasing in size, without any overlying skin changes or regional lymphadenopathy. The physical examination revealed a hard, mobile mass that measured approximately 10 cm in its largest dimension. The lesion was excised.

The gross examination of the resection specimen revealed an encapsulated lobulated tumor, with a smooth external surface. Histologically, the tumor was encapsulated by an incomplete shell of lamellar bone. The tumor had variable cellularity, and, in areas, contained myxoid stroma. The tumor cells had eosinophilic cytoplasm with vesicular round-to-oval nuclei. Sparse mitoses were noted. Immunohistochemical stains demonstrated that the tumor cells expressed vimentin, S100 and neuron-specific enolase, with the latter expressed focally. Conclusion: To the best of our knowledge, this is the first case of ossifying fibromyxoid tumor of soft parts to be reported in Kuwait. Therefore, pathologists and clinicians should be aware of this tumor.

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Introduction

Ossifying fibromyxoid tumor of soft parts (OFMT) is a rare neoplasm first described in 1989 \cite{1}. It is a soft tissue tumor of uncertain histogenesis, commonly occurring in adults, with a higher incidence in males than females. The extremities and trunk are the most common sites affected \cite{1}. In addition, occurrences in unusual sites, such as the neck and lips, have also been reported \cite{2, 3}. Typically, OFMT presents as a slow-growing benign mass, which may recur locally or metastasize \cite{4}. In this report, a new case of OFMT diagnosed in Farwaniya Hospital, Kuwait, with immunohistochemical study is presented.

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Immunohistochemical stains against antisera antibodies for desmin (1:50, Dako), S100 (1:400, Dako), neuron-specific enolase (NSE; 1:100, Dako), vimentin (1:25, Dako) and CD34 (1:15, Biogenics) were performed at the indicated dilutions. The tumor cells demonstrated positive expression for S100, vimentin and focally for NSE, but were negative for desmin and CD34. The gross morphology, histological findings and immunophenotypical results were diagnostic of OFMT.

**Discussion**

In this case, the finding of an encapsulated lobulated tumor of eosinophilic spindly cells with vesicular nuclei is as described by previously reported cases [1, 4–6]. Although the fibromyxoid component of OFMT might suggest a fibroblastic origin, the expression of both S100 and NSE proteins by the tumor cells point instead to a neural origin [1]. Further support for a neural histogenesis is
provided by the discovery of Holck et al. [6] of an interrupted basal lamina, a Schwannian feature in some of the reported cases [1, 3, 7], and the demonstration of cytogenetic abnormalities involving chromosomes 6 and 18 also resonated with a malignant peripheral nerve sheath tumor.

Most cases of OFMT are cured by local excision, but up to 1/4 may recur or even metastasize. Statistically, significant factors that may predict recurrence include the size of the tumor, high cellularity, high nuclear grade and mitotic activity. Using these criteria Folpe and Weiss [4] proposed 3 classifications of OFMT: (1) typical OFMT with a low nuclear grade, low cellularity and a mitotic rate <2/50 HPF; (2) malignant OFMT with a high nuclear grade, high cellularity and a mitotic rate >2/50 HPF; (3) atypical OFMT with an intermediate grade between 1 and 2. Our case falls into the typical OFMT class.

**Conclusion**

This is the first report of an OFMT in Kuwait. An OFMT is a slow-growing tumor of uncertain, probably neuronal, origin. This tumor commonly behaves in a benign manner. Some of the tumors with high cellularity, high nuclear grade and high levels of mitotic activity tend to have local recurrence and distant metastasis.

**References**