Ossifying Fibromyxoid Tumor of Soft Parts

Case Report

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Introduction

Ossifying fibromyxoid tumor of soft parts (OFMT) is a rare neoplasm first described in 1989 [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 [1]. In addition, occurrences in unusual sites, such as the neck and lips, have also been reported [2, 3]. Typically, OFMT presents as a slow-growing benign mass, which may recur locally or metastasize [4]. In this report, a new case of OFMT diagnosed in Farwaniya Hospital, Kuwait, with immunohistochemical study is presented.

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 that measured 7.5 cm × 5.5 cm × 4.0 cm; it was hard and difficult to cut. Hemorrhaging and necrosis were not seen (fig. 1). The specimen was X-rayed, and it showed a thin shell of bone within the capsule; no calcification was seen within the parenchyma (fig. 2). Formalin-fixed paraffin-embedded tissues were sectioned at 3 μm for hematoxylin and eosin (H&E) and immunohistochemical staining. Histological examination showed a thick fibrous capsule which extended into the tumor parenchyma dividing it into lobules. Within the capsule, there was a thin, incomplete shell of mature lamellar bone, partially rimmed by osteoblasts (fig. 3). The tumor showed variable cellularity and, in areas, contained myxoid stroma. The tumor cells had eosinophilic cytoplasm with vesicular round-to-oval nuclei. Mitotic counts were less than 2/50 per high-power field (HPF; fig. 4).

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


