Low-grade fibromyxoid sarcoma: a rare condition with high proliferation

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ABSTRACT

Low-grade fibromyxoid sarcoma (LGFM S) is a type of high metastatic potential of the fibrosarcomas. Most of the time there is a long interval between tumor presentation and metastasis. We present 2 cases of LGFM S. The first is a 31-year-old female with a mass in anterior aspect of her left thigh, and the other is a 68-year-old female with mass in posterior part of her neck. Both cases underwent operation for several times and confirmed as LGFM S histopathologically, there is no exact protocol for postoperative follow-up to detect early metastases according to the relative variety of LGFM S. So informing the patients about the long-standing metastatic potential of their disease is important.

Keywords: fibrosarcoma, low-grade fibromyxoid sarcoma, surgery

Introduction

No Low-grade fibromyxoid sarcoma (LGFM S) is a differentiated sub-type of fibrosarcoma. Evans first described this pathologic entity as a rare high metastatic potential soft-tissue tumor despite its benign histologic appearance in 1987 [1]. Usually pathologists, radiologists, and surgeons fall in problem for fundamental principle for tumor management due to long interval period between tumor presentation and metastasis behavior. It is still difficult to estimate LGFM S incidence because many cases are not diagnosed as LGFM S. Although LGFM S can be found in unusual places, like head, retroperitoneum, or the chest wall, these tumors usually occur in the proximal extremities and trunk [2, 3]. Subfascial location is the most common place for LGFM S occurrence, as subcutis or dermis may be affected in rare occasions [4]. LGFM S especially happen in young to middle-age adults, by the way a large member of cases have been reported in pediatric ages [3, 5, 6]. In this report, we presented 2 cases of LGFM S; a 31-year-old female with a mass in anterior aspect of her left thigh, and a 68-year-old female with a mass in posterior part of her neck.
Case Presentation

Case 1
A 31-year-old female presented with a recurrent large tumor mass on the anterior aspect of her left thigh operated for three times before admitted to our clinic. She had been operated due to a similar tumor presentation 16 and 14 years ago at another medical center, and 6 years ago at our clinic. The mass has been growing slowly during past 6 years, but the patient did not seek for medical treatment until the tumor size disturbed her regular life. Physical examination revealed firm, multifocal, mobile masses without tenderness, redness, or warmth (Figure 1a). Laboratory evaluations and plain radiographs were unremarkable. Magnetic resonance imaging (MRI) demonstrated a contrast enhanced tumor with myxoid and fibrous pattern (Figures 1b and 1c). The tumoral masses were excised. Due to infiltration of muscular and fatty tissues, quadriceps femoris and adductor longus muscles were excised (Figures 1d and 1e). Medial and lateral parts of femoral bone periosteum were invaded by tumor and resected. After surgery the patient experienced no major complication and was discharged 5 days after the operation. The histopathological diagnosis was LGFMS (Figure 1f).

Case 2
A 68-year-old female patient presented with large recurrent tumoral masses on posterior part of her neck, radiating to her shoulders, and upward of right shoulder. During past 25 years she has been operated 3 times because of neck masses. Physical examination revealed neck stiffness and firm multifocal mobile masses especially radiated to right shoulder. There was no impairment of sensory and motor nerves. Cervical MRI demonstrated multiple sized and firm contrast enhanced masses which spread to both shoulders (Figure 2). The detected masses were excised. Periosteal infiltration was observed at spinous

Figure 1. (a) Preoperative left leg mass; (b, c) The fibrous components were identified as hypointense area on T1- and T2-weighted MR images and slightly enhancing on T1-weighted MR images after intravenous administration. The myxoid components of the mass were recognized as hypointense on T1-weighted MR images and hyperintense on T2-weighted MR images, and enhancing on T1-weighted MR images after contrast administration; (d) Postoperative appearance of the patient’s leg; (e) The excised specimen. The surface of the tumoral mass was smooth and glistening with white-gray color. There was no lenfovascular invasion or necrotic area; (f) Histopathological appearance of the mass. Immunohistochemistry study of the tumor revolves; EMA: focal positive, SMA: focal positive, MSA: negative. P53: negative, Ki 67: 3% positive.
processes of C6 and C7 and vertebral lamina was resected together with infiltrated parts of bone. After surgery, the patient experienced no major complication and discharged 2 days after surgery. The histopathological evaluation was reported as LGFMS (Figure 2).

Discussion

TLGFM S manifestation is usually long-standing and depended on the anatomic area of the lesion. Mostly its presentation is as painless soft tissue masses with duration of over 5 years in 5% of patients [2]. Acute presentation of LGFMS is rare, and it can happen in chest wall infiltration as acute respiratory distress syndrome (ARDS) or chest pain. It may be presented with seizure when intracranial infiltration is suspected [3, 7]. In this paper we presented two cases, diagnosed at 16 years and 25 years ago.

Special MRI and CT finding for LGFMS are defined, although imaging findings are nonspecific [3, 8-10]. On CT images without contrast, the fibrous structure of these tumors interprets data density of muscular tissue, and the myxoid part was evaluated as hypodense. The fibrous structure was interpreted as hypointense on T1- and T2-weighted images, and there was a slight contrast enhancement on T1-weighted MRI images. On the other hand, the myxoid part has been found as hypointense on T1- and hyperintense on T2-weighted images, and strongly contrast enhancing on T1-weighted images. Sometimes calcification can be detected [10]. Both of our cases had contrast enhancement at radiological examination similar to literature.

LGFMS is a sub-type of fibrosarcoma, characterized by a mixture of hypercellular myxoid nodules in collagenized area with low cellularity [6]. Tumoral parts are commonly characterized by round to ovoid nuclei small cells without nucleoid and mildly eosinophilic cytoplasm. Mitosis increased at atypical region determined by hypercellularity; only in 10% of the cases nuclear hyperchromatism, and necrosis has been reported. Tumor cells are commonly determined
by absent of sparse mitotic finding, nuclear anaplasia or necrosis. It is positive for vimentin and negative for other antibodies, such as S100 protein, desmin, keratin, epithelial membrane antigen such as CD31, and CD34 at most cases. Lesions showing proliferation of spindle cell with or without fibrous component in myxoid pattern are evaluated at differential diagnosis of LGFMS [11]. Tumors with both fibrous and myxoid pattern include fibromatosis, neurofibroma, malignant peripheral sheath tumor, perineurioma, fibrous histiocytoma, or the tumors with only predominant myxoid areas without fibrosis such as myxomas, myxoid neurofibroma, angiomyxoms, myxoid liposarcoma, and low-grade myxofibrosarcoma have to be evaluated at differential diagnosis [10]. Also, desmoids tumors such as desmoplastic fibrosarcoma, and low-grade differentiated liposarcoma should be remembered in the differential diagnosis of LGFMS. If tumor has been removed completely, it is not difficult to diagnose LGFMS due to morphologic pattern and immune phenotypic features. In such cases, an excisional biopsy should be performed before surgical resection according to the fact that it is not commonly possible to diagnose with needle core biopsy or fine needle aspiration. If the diagnosis still remains unclear for myxoid, this kind of cytogenetic for rare cases of LGFMS can be beneficial [4].

Goodlad et al. [5] pointed that LGFMS were inconsistently aggressive tumor. Although all the cases were primarily diagnosed and treated as benign lesions in retrospective study, local recurrence was reported as 68%, and the death rate was reported 18% [5]. It is clear that patient selection has affected the rate of metastases and recurrence rate because most of them selected according to unexplained metastases. In another large series death rate, local recurrence, and metastasis was detected as 2%, 54%, and 6%; respectively [6]. There was no significant relation between recurrence or metastasis and the presentation of high cellularity focal area, nuclear enlargement, increased mitotic activity, and necrosis. Because of potential of late metastasis of LGFMS, as reported 45 years after primar diagnosis, these patients had to be followed for a long period of time [4]. After initial diagnosis of LGFMS the patients must be follow-up by expert oncological group. However exact interval for periodic chest imaging is unclear, as the most metastatic area is lung, several chest CT scan have to be performed during long-term follow-up of these cases.

Conclusions

The 2 cases report presented herein, enriches the literature for the best diagnosis and surgical treatment of this rare tumor. In addition, it is important to inform patients about long-time metastatic potential of the disease as there is no dedicated protocol regarding follow-up examinations for early metastatic mass diagnosis.

Informed consent

Written informed consent was obtained from the patient for the publication of this case report.

Conflict of interest

The authors declared that there are no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

References