Primary Myxoid Liposarcoma of the Lower Back in an Adult Female Patient: An Extremely Rare Presentation

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Abstract

Myxoid liposarcoma is the second most common type of liposarcoma that typically presents in the lower extremities and has a predilection for men. Manifestation of this tumor in the lower back is rare. We report the first known case of an adult woman presenting with a soft tissue mass in her lower back. Following excision and further workup, the mass was positive for rearrangements of the DDIT3 (CHOP) gene at the 12q13 locus, consistent with the diagnosis of a myxoid liposarcoma. The tumor was noted to be intramuscular and extended into the margins, requiring re-excision at the previous resection site. Biopsy of the re-excised area was negative for evidence of residual myxoid liposarcoma. We report this case to highlight the importance of considering myxoid liposarcoma as a differential diagnosis in soft tissue masses presenting in the lower back.

Keywords: Myxoid; Liposarcoma; Lower back; Case report

Introduction

Soft tissue sarcomas are relatively rare malignant tumors of mesenchymal origin, representing as little as 1% of all malignant neoplasms. The most common type of soft tissue sarcoma is a liposarcoma that consists of four different subtypes [1]. Myxoid liposarcomas are the second most common type of liposarcoma that frequently occur in the fourth to fifth decade of life with a propensity to present in men [1-3]. The most prevalent area for this subtype of liposarcomas to develop is in the lower extremities [1]. Liposarcomas rarely present in the back [4]. We present a case of a 29-year-old woman found to have a myxoid liposarcoma located in her right lower back.

A 29-year-old woman presented to the hospital for an elective excision on a soft, mobile mass located on her right lower back. The patient stated that the mass had been causing her discomfort and had been present for quite some time. The mass was excised and during the procedure was noted to be intramuscular. Upon biopsy results, the gross specimen was partially capsulated measuring 11.0 × 10.0 × 5.5 cm, tan-pink to brown in color with a smooth outer surface. The tumor extended into the margins of the excision. The mass consisted of mainly hypocellular, vaguely lobulated, proliferation of small ovoid to spindle cell admixed with scattered lipoblasts, in a background of abundant myxoid material and arborizing vascularity. At the periphery of the lobules, increased cellularity was noted with moderately increased cytological atypia. Mitotic activity was very rare and areas of round cell composed less than 5% of the total tumor. In many of the tumor cells, stain with p53 showed evidence of p53 overexpression. Stain with Ki-67 showed a proliferation index of 2-8%. Fluorescence in situ hybridization (FISH) analysis was positive for detection of rearrangements of the DDIT3 (CHOP) gene at the 12q13 locus, consistent with the diagnosis of myxoid liposarcoma. Following the diagnosis, the patient had a computed tomography (CT) scan of the chest, abdomen and pelvis, which revealed a resection scar along the right paraspinous musculature with no evidence of lymphadenopathy, lesions or pathology. Due to evidence of the tumor extension into the margins, the patient underwent re-excision at the previous resection site of the mass. The patient tolerated the procedure well and will be followed up for subsequent post-operative visits. The biopsy results of the re-excision revealed no evidence of residual myxoid liposarcoma.

Discussion

Liposarcomas are one of the most common types of soft tissue sarcomas. According to the World Health Organization, there are four types. These subtypes are myxoid/round cell, pleomorphic, well differentiated and dedifferentiated [3, 5]. Of the four subtypes, myxoid liposarcomas account for about 20%. Myxoid liposarcomas typically present as a painless slow growing mass [6]. They are frequently located in the intramuscular fascia, under or on muscle [4]. The most common area for myxoid/round cell liposarcomas to develop is in the lower extremities, which make up about 75-80% of cases. More specifically, about two-thirds of...
cases manifest in the thigh [1, 3]. In about 8% of cases they present in the retroperitoneum and 5% appear in the upper extremities [1]. It is atypical for a liposarcoma to present in the back [4]. To the best of our knowledge, no cases of primary myxoid liposarcomas subtype located in the lower back have been reported. We report a rare case of a myxoid liposarcoma presenting in the right lower back.

Histologically, this subtype exists on a spectrum with variable amounts of round cells. The existence of greater than 5% of round cells within the mass is associated with a poor prognosis and is considered high-grade [1, 7]. Approximately 95% of cases of myxoid/round cell liposarcoma subtype have a TLS-CHOP fusion gene on cytogenetic analysis [7, 8]. This fusion gene is specific for this subtype and is the result of t(12;16)(q13;p11) translocation. It has been suggested to be a dysfunctional transcriptional regulator that interrupts differentiation of adipocytes resulting in proliferation [8, 9]. In our case, the mass was found to have this specific fusion gene. Seen in less than 5% of cases, EWS-CHOP fusion gene is a rare variant that has also been observed [8, 9].

The standard of care for patient with local disease is wide local surgical resection with the confirmation of negative margins [10, 11]. Low-grade myxoid liposarcomas have a low local recurrence rate. At a 5-year follow-up, approximately 8% of diagnosed patients will have local disease recurrence [7]. In comparison to other types of soft tissue sarcomas, myxoid liposarcoma is more radiosensitive [11]. Adjuvant radiation is typically considered for patients who have considerable number of round cells composing the mass [7].

Myxoid/round cell liposarcoma has a tendency to metastasize to extrapulmonary sites, specifically, soft tissue and skeletal sites such as the mediastinum and the retroperitoneum [12, 13]. With the potential to metastasize to distant sites, prompt diagnosis as well as providing the best possible treatment options is critical [14].

Conclusions

This case illustrates the challenges associated with the diagnosis and management of these rare tumors. Myxoid liposarcomas should be included as a possible differential diagnosis in cases of soft tissue masses that present in the lower back. An early accurate diagnosis will not only result in appropriate treatment from the get-go, but also save both the patient and clinician valuable time and resources. Although a rare entity, considering myxoid liposarcoma in the diagnosis of soft tissue masses may prevent misdiagnosis and possibly alter the clinical approach.

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Financial Disclosure

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Conflict of Interest

None to declare.

Informed Consent

Not applicable.

Author Contributions

SQ participated in the case. ARZ wrote the case report. SQ, JZ and AZ revised and edited the case report.

Data Availability

The authors declare that data supporting the findings of this study are available within the article.

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