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Radiographic presentation of a multi-compartmental atypical giant lipoma of the lower extremity

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Keywords: Atypical lipoma; Giant lipoma; Soft tissue mass; Multi-compartmental mass.

Description

A 61-year-old female with no significant past medical history presented with painless progressive left lower extremity weakness and foot drop resulting in altered gait over the course of two years prior to seeking care. Examination revealed a hypertrophied left lower extremity, steppage gait, and 1 out of 5 strength with left ankle dorsiflexion. Initial x-ray of the affected ankle was unremarkable.

Further imaging of the affected extremity was performed via Magnetic Resonance Imaging (MRI), which revealed a large multi-compartmental fat containing mass occupying the anterior, lateral, and deep posterior compartments from the knee joint to just above the ankle joint (Figure 1A-C). Given the extensive spread of the mass and resulting gait alteration, surgical intervention was pursued with complete resection of the tumor. Postoperative pathology revealed the mass to be consistent with an atypical lipomatous tumor with generally mature adipocytes and occasional cytologic atypia. Following resection, the patient's gait improved with physical therapy and repeat MRI after six months revealed normal postoperative changes without signs of tumor recurrence (Figure 1D). Lipomas are the most common soft-tissue tumors and can be difficult to distinguish from rarer, malignant masses given their wide variety of presentations [1]. Lipomas can be characterized by their location, size, and underlying pathological features. In this image, we present a multi-compartmental giant lipoma extending from the knee to the ankle, with the term giant referring to lipomas greater than 10 cm in diameter or 1000 g in mass [2]. Tissue examination from this mass revealed an atypical pathology, which is the most common subtype of lipomas and commonly presents as a painless, slow-growing mass. Additional common subtypes include pleomorphic and myxoid lipomas, among many additional rarer cytologies including liposarcoma [3].

The differential diagnosis for lower extremity soft-tissue masses found on radiographic imaging is broad, adding to the challenges of identifying these tumors. Alternative diagnoses may include masses of muscle etiology (muscle hematomas, myositis ossificans, muscle strains), vascular etiology (hemangiomas, deep vein thrombosis), and other connective tissue structures (Baker's cyst, fascial tear) [4].

Giant lipomas, especially those in multiple compartments of

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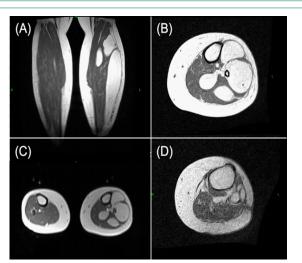


Figure 1: (A) Longitudinal view of lipoma extending from below the knee to above ankle, with surrounding muscular atrophy. (B) Multicompartmental invasion seen with cross-sectional view demonstrating multiple lobules extending into deep tissue. (C) Comparison image demonstrating normal findings of right lower extremity. (D) Postoperative imaging demonstrating resected mass with subsequent post-surgical scar tissue and greatly reduced compression of surrounding structures.

an extremity, can often cause symptoms due to compression of surrounding structures. In this case, the patient's foot drop was presumed to be due to peroneal nerve compression and leg muscle atrophy. Despite their benign nature, giant lipomas also have a risk of malignant transformation to liposarcoma and warrant at minimum a biopsy for further evaluation [5]. For patients willing to undergo surgery, wide margin excision is a preferred treatment option [5]. However, with giant lipomas, the risk of recurrence remains, and patients should undergo continued surveillance following surgical intervention to assess for relapse [2,3]. Overall, when presented with a soft-tissue mass, clinicians should utilize radiographic and pathological findings to assist with identifying and categorizing lipomas, allowing for more informed treatment and surveillance strategies.

Declarations

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