Case Report

Spinal Myxoma of the Facet Joint: A Case Report

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Introduction

Myxoma is a benign, slow-growing soft tissue tumor of mesenchymal origin. The estimated incidence is at one case/million population/year [1]. Even if commonly found in the left atrium of the heart, myxomas can arise in many different districts, such as large muscles (Intramuscular Myxoma) and even in continuity with joints (Juxta-articular Myxoma). Myxomas appear as ovoid painless masses often containing cystic areas. Size can range from 1-17 cm but more commonly they have been found smaller than 5cm in length, thus larger size often needs further investigations, and an alternative diagnosis should be taken into account [2-4]. Symptoms can occur consequentially to the compression on adjacent structures (vascular, neurological). Neurological symptoms such as cord compressions or nerve root compression, are rare [5]. Only 5 cases of juxta-articular myxomas of paraspinal are described in literature. We report a case of a juxta-articular myxoma in the lumbar spine causing back lumbar pain. MRI of the lumbar spine showed a big multilobulated mass at L5-S1 within paraspinal muscles, hypointense in T1-weighted sequences and hyperintense in T2-weighted. The neoplasia could have been mistaken for a giant synovial cyst due to the involvement of facet joints. Since the uncommon presentation and uncharacteristically large size, a PET-scan was conducted in order to rule out the possibility of systematic disease. Intraoperative histological examination yielded the diagnosis towards a benign neof ormation so surgical excision was conducted.

Keywords:
Myxoma
intramuscular myxoma
juxta-articular myxoma
spine surgery

Case Presentation

I Patient Description and Case History

A 61-year-old man, height 180 cm, weight 75 kg, without relevant comorbidities and pharmacotherapies, had experienced low-back pain over 1 year, which worsened in last 4-month with irradiation down the right leg (L5 root).

II Physical Examination Results

Physical examination revealed a palpable mass on the right paraspinal area in the lumbar region. Neurological examination demonstrated normal strength (5/5) in all muscle groups. The patient presented Lasegue test positive on the right side. The patient presented left leg weakness, and no other localizations. Intraoperative histological analysis was consistent with Juxta-articular myxoma (JM). Treatment of choice was surgical excision, with complete relief of symptoms afterwards.

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Figure 1: Coronal CT scan of lumbosacral spine.

Figure 2: Axial CT scan.

A pelvis MRI confirmed a multilobulated mass involving the right paravertebral muscles in the lower lumbar and sacral region that appeared hypodense on T1-weighted sequences and hyperdense in T2-weighted sequences (Figures 3 & 4).

Figure 3: Coronal MR image of lumbosacral spine show hypointense paravertebral lesion (T1 signal).

Figure 4: Axial MR image of lumbosacral spine show contrast-enhanced of the lesion (T2 signal).

IV Treatment Plan

In October 2020, a core needle biopsy of the right paravertebral muscle was conducted with non-diagnostic result. In January 2021 he underwent surgical treatment to perform an extemporaneous incisional biopsy and eventual excision of the tumor. A posterior access was performed. The tumor appeared as an encapsulated gray-white mass inside paraspinal muscles (Figure 5). The freezer histological exam confirmed a benign tumor so the dorsal portion was removed through a good cleavage plan. A clear mucinous fluid flowed from the remaining ventral part of the tumor was removed.

Figure 5: Specimen after resection.

V Pathological Examination

The histological examination was consistent for an intramuscular myxoma with cystic areas degeneration. (CD34 - / +, EMA-, MUC4, S100-, SOX10-).

VI Expected Outcome of the Treatment Plan

The patient recovered easily from the procedure. Immediately postoperatively, he experienced complete relief from symptoms.

Discussion

Myxoma is a benign mesenchymal tumor. The term myxoma is used to identify a plethora of tumor which can occur in every part of the body.
The term was first used to describe lesions that mimic the structure of umbilical cord. Myxomas are composed by a paucity of stellate cells set in a loose myxoid stroma with, minimal mitotic figures [3]. It is a rare condition and its incidence is reported to be one per million individuals, most frequently diagnosed in female patients between 40-70 years [6]. Allen extensively described multiple variants of this tumor and he identified several subgroups of the so-called “mainstream soft-tissue myxomas”; Among them Intramuscular Myxoma (IMM) is the most common form of myxoma after myocardial myxoma and can be located in any skeletal muscle groups. IMM is a solitary tumor, but it could present as part of Mazabraud’s syndrome, in which multiple myxomas associate with fibrous dysplasia in one or more bones [7]. IMM is typically seen as a well-circumscribed intramuscular mass. Magnetic resonance shows low-signal intensity in T1-weighted scan, hypointense to muscle; T2-weighted MRI displays as homogeneous, hyperintense to muscle and brighter than fat; STIR (short T1 inversion recovery) shows high-signal intensity with a surrounding fat rim [8].

On PET/CT myxoma could show 18-FDG avidity [9]. A specific variant of myxoma, resembling an IMM in almost every radiological and histological aspects, is the Juxta-articular myxoma (JAM): this tumor origin in the nearby and involves periarticular tendons, ligaments and joint capsules. JAM can indeed be histologically indistinguishable from the intramuscular form, and especially when the juxta-articular myxoma invades muscle, it may be impossible to differentiate the two on the basis of the section. Macroscopically myxomas arising from joints tend to present more focal vascularity, peripheral fibroblastic proliferation and more cystic areas than IMM. Cysts of varying sizes resembling ganglion cysts are seen in 89% of cases[10]. Given the close relationship to joints and its ganglion-like cystic spaces, this condition is closely related to ganglion cysts, which are a rather rare event. IMM, as aforementioned is completely benign and no recurrences have been described in the literature. Conservative treatment is the gold standard while excision is to be considered when symptomatic. On the other hand, JAM presents a higher risk of recurrences. Meis and Enzinger reported 34% of recurrence in their study. In the case presented, the patient suffered from radicular pain due to the mechanic effect of the mass against the nerve root. Thus, we decided to remove the tumor.

In order to remove the whole mass, including the ventral portion within the psoas major muscle, a combined surgical approach should have been conducted, with higher rate of intraoperative complications. Given the benign nature of intraoperative histological exams we opted to remove only the portion causing the compression and symptoms, avoiding a longer and unnecessary procedure.

**Conflicts of Interest**

I hereby certify, to the best of my knowledge that the authors listed in the case report presented have no affiliations with or involvement in any organization or entity with any financial interest, or non-financial interest (such as personal or professional relationships, affiliations, knowledge or beliefs) in the subject matter or materials discussed in this manuscript.

**REFERENCES**