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Case report

Bilateral lipoma arborescens – A proliferative case demonstrating progression in an adolescent male

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ABSTRACT

Case: We describe a case of extensive symptomatic bilateral lipoma arborescens of the knee in a 19-year-old man who suffered from recurrent knee effusions for many years. This patient had MRI evidence of progression of disease prior to arthroscopic intervention. He was treated with bilateral complete arthroscopic synovectomies, demonstrating no evidence of cartilage wear.

Conclusion: Lipoma arborescens is a progressive disease. Timely diagnosis and treatment may prevent disease evolution and sequela.

1. Introduction

Lipoma arborescens is a rare cause of knee joint pain and effusion. It is characterized by a frond-like, hypertrophic synovium that may cause symptomatic effusions.\textsuperscript{1-5} The term originates from “arbor,” meaning “tree”, which describes the gross appearance of the mass. Lipoma arborescens typically occurs unilaterally in the knee, although cases of bilaterality, extraarticular disease, and locations distant to the knee joint have been described.\textsuperscript{1,3,4}

Hoffa attributed the term “lipoma arborescens” to Johannes Muller in 1904, referring to adipose disturbances in the knee joint.\textsuperscript{6} In 1953, Arzimanoglu presented a case of a 19-year-old girl with bilateral symptoms, diagnosed by open excisional debridement.\textsuperscript{7} Today, it is commonly diagnosed through MRI as nodules with a frondose morphology and hyperintense signal on T1 and T2 sequences. It is confirmed by histologic analysis as fat cells beneath the synovial tissue with small amounts of lymphocytes and plasma cells surrounding capillaries.\textsuperscript{1}

Little is known about the etiology and progression of lipoma arborescens. While MRI enables both non-invasive diagnosis in symptomatic patients and incidental discovery of lipoma arborescens in asymptomatic patients, there is no study to date documenting progression of disease to prolific pan-compartmental involvement.

2. Presentation of the case

The patient was a 19-year-old male referred to our clinic following six years of recurrent knee effusions and two recent episodes of left patella dislocation events. He had congenital transposition of the great arteries, requiring an arterial switch procedure in infancy and a subsequent aortic valve replacement at age 9, resulting in long-term use of warfarin for anticoagulation. Pediatric orthopaedic and rheumatology specialists had attributed his effusions to be due to hypertrophic synovium from chronic hemarthroses when they began occurring in his early teen years. An MRI of his left knee at age 13 showed arboreal hypertrophy of the infrapatellar fat pad, though at the time this was not diagnosed as lipoma arborescens (Fig. 1).

In addition to cartilaginous injury to the medial facet of the patella from his dislocations, the new MRI of his left knee revealed diffuse fatty hypertrophy of his synovium, with villous and arboreal architecture throughout the knee joint (Fig. 2). An MRI was ordered for his contralateral knee given its non-painful large effusion showing similar findings of the synovium (Fig. 3). Due to the severity of his recurrent effusions and difficulty performing his job duties in the automotive industry, the patient elected to have bilateral diagnostic arthroscopies with synovial biopsies and complete arthroscopic synovectomies.
2.1. Procedure

The patient was admitted to the hospital for bridging therapy of his anticoagulation. On hospital day #3, he underwent sequential bilateral knee arthroscopy, beginning with the left knee, which is described in detail. Once anesthetized, a physical exam revealed stable ligamentous testing and a grade 3+ effusion. Under tourniquet use, the knee joint was entered through an anterolateral portal and approximately 150 cc straw-colored synovial fluid was immediately evacuated. Our diagnostic arthroscopy ensued, showing a medial patellar facet injury with a corresponding loose body in the lateral gutter and, most impressively, innumerous fatty-appearing villous tumors projecting from the synovium throughout each compartment of the joint (Fig. 4a and b). The cruciate ligaments were normal appearing, but both menisci were encased in fatty tumors growing from the capsule into the joint, with the medial meniscus additionally being subluxated off of the tibial plateau surface. There was no evidence of cartilage wear on the tibiofemoral articular surfaces.

Multiple biopsies were taken from the synovial projections and the capsule itself. Complete arthroscopic synovectomy and debridement was then performed, which included using an accessory posteromedial portal for instrumentation to the posterior compartment of the knee joint (Fig. 5). His post-operative recovery in the hospital was uneventful, and his Coumadin therapy was resumed. At his 3-week post-operative visit, the patient had some persistent pain and ongoing effusion, attributed to surgical hematoma. This effusion resolved slowly; unfortunately, he did suffer some recurrence of effusions.

Surgical pathology from each knee showed proliferative synovia with primarily villous architecture of adipose tissue without atypia (Fig. 6). A focal lymphoid reaction was present, and no pigmentation consistent with pigmented villonodular synovitis (PVNS) was found (Figs. 7 and 8).

3. Discussion

To our knowledge, this is the first case of lipoma arborescens with sequential MRIs demonstrating unequivocal proliferative progression of pathology prior to surgical intervention. Differentiating lipoma arborescens from chronic synovitis and PVNS is important, as a
radical synovectomy may be necessary to treat the latter while arthroscopic management of the former has been shown to limit recurrence. While pan-compartmental involvement did occur in our case, there was no extra-articular propagation, which can occur with widespread PVNS.

Sequelae of symptomatic lipoma arborescens are not known. A recent report suggests that osteoarthritis may result from delays in the definitive treatment of this pathology. While pan-compartmental involvement did occur in our case, there was no extra-articular propagation, which can occur with widespread PVNS.

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which highlights the marked variability of this poorly understood disease.\(^9\)

The diagnosis of lipoma arborescens relies on clinical, MRI, and histopathologic studies. Although the etiology and consequences of this disease are still not well known, there appears to be a wide spectrum with different subsets. Ultimately, adequately diagnosing and treating these patients may prevent progression of the disease.

**Fig. 6.** Low power view of the synovial membrane that demonstrates a villous-like proliferation of the synovium with papillary projections expanded predominantly by mature adipocytes and focal areas of stromal fibrosis, vascular congestion, and a dense chronic inflammatory infiltrate (hematoxylin-eosin, 1x).

**Fig. 7.** Medium power view showing a chronic inflammatory infiltrate characterized by diffusely scattered plasma cells and lymphocytes with focal lymphocytic aggregates (hematoxylin-eosin, 11x).

**Disclosures**

The authors have no disclosures. Each author certifies that his institution approved or waived approval for the use of human subjects for this investigation and that all investigations were conducted in conformity with ethical principles of research.
Conflict of interest

None.

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