Melorheostosis of the Fossa Intercondylaris Femoris – A Case Report and Review of the Literature

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Learning Point of the Article: Melorheostosis is a rare benign bone condition characterized by excessive segmental sclerosis of cortical bone being reminiscent of dripping candle wax. Manifold clinical manifestations like impingement or nerve compression syndromes are possible and potentially require surgical treatment.

Abstract

Introduction: Melorheostosis is a rare benign bone condition characterized by excessive segmental sclerosis of cortical bone being reminiscent of dripping candle wax. It typically affects the long bones and can cause impingement and nerve compression syndromes that may require surgical treatment.

Case Report: We report the case of a 49-year-old male patient with a 12-month history of the left-sided knee pain and a concomitant limitation of his left knee flexion to 90 degree. Radiographic and magnetic resonance imaging revealed the typical radiographic appearance of melorheostosis with an extraosseous lesion in the fossa intercondylaris femoris being causative for the limited knee range of motion. Following the resection of the extraosseous part of the lesion through a direct open approach, the patient is pain free with a maximum of 110 degree knee flexion at 12-month follow-up.

Conclusion: Melorheostosis can present with manifold clinical manifestations that potentially require surgical treatment. Even in patients with a challenging localization of extraosseous lesions, a good to excellent functional outcome is possible.

Keywords: Melorheostosis, Léri disease, Candle wax disease, Surgery, Fossa intercondylaris femoris, Notch.

Introduction

Melorheostosis is a rare benign bone condition, first described in 1922 by Léri and Joanny [1]. Although several eponyms such as “Léri disease” and “candle wax disease” are known, its incidence is reported to be only 0.9 per 1 million population [2, 3, 4]. It typically manifests during early childhood and adolescence with 40–50% of all cases being evident by the age of 20 [2].

Case Report

We present the case of a 49-year-old male patient (1.83m, 88 kg) with a 12-month history of left-sided knee pain with concomitant limitation of the knee flexion to 90 degree. The initial conservative treatment consisted of analgesics and physiotherapy. Persistent pain led to subsequent radiographic (Fig. 1a and b) and magnetic resonance imaging (Fig. 2a-c) that revealed an excessive hyperostosis of the left femur with an extraosseous lesion in the fossa intercondylaris femoris. The patient was subsequently referred to our specialized center for musculoskeletal oncology.

Based on the clinical and radiographic findings, we established the diagnosis of melorheostosis and the patient underwent surgical resection of the symptomatic extraosseous lesion in the fossa intercondylaris through a direct open approach under meticulous protection of the cruciate ligaments. Postoperatively, the histological findings were consistent with the diagnosis of melorheostosis (Fig. 3) and the patient was...
The precise etiology of melorheostosis remains unclear [5, 6, 7]. However, more recently, Kang et al. identified somatic mosaic mutations in the regulatory domain of the MAP2K1 kinase which seem to be causative for the development of melorheostosis in some, but not all patients [6, 7]. Although this potentially opens possibilities of a targeted treatment, Wordsworth and Chan strengthen that targeting the MAP pathway will also affect not mutated cells resulting in potential off-target side effects [7].

While only around 500 cases of melorheostosis are reported in literature, Smith et al. presented the largest single-center cohort of this entity in 2016 with 24 patients aged 3–68 years [7, 8]. Although no gender predilection has previously been reported, they found an unexpected and yet inexplicable female to male ratio of 4:1 [8]. Predominant affected bones in their cohort were the leg (67%), ankle or foot (38%), and the arm (33%) with 20 patients (83%) reporting a single limb involvement and 4 patients (17%) multiple affected body regions [8]. Of the three major clinical manifestations pain, limited range of motion, and deformity, pain was the most frequent clinical appearance in their cohort with 20 patients (83%) complaining of nociceptive, neuropathic, or skeletal pain [8].

The typical radiographic appearances of melorheostosis are excessive segmental sclerosis of cortical bone being reminiscent of the eponymous dripping candle wax [7]. Although these findings are characteristic for melorheostosis, they are not consistent and differential diagnosis to other sclerosing bone conditions can be challenging [7]. Clinical manifestations can require a surgical treatment, especially in patients where lesions cause nerve compressions or impingement syndromes [7]. Several authors report the surgical treatment of melorheostosis around the knee: John et al. reported a 42-year-old female patient that underwent repeated open excision of intra-articular lesions to the patellofemoral joint [9]. Moulder and Marsh reported a 40-year-old male patient who underwent total knee arthroplasty after a 9-year long history of symptoms and two previous arthroscopic removal of loose bodies [10]. Several authors strengthen the necessity of repeated surgeries in these patients with Smith et al. reporting that 54% of patients in their cohort had to undergo a mean number of two surgeries and conclude that a progressive nature of melorheostosis is likely [8].

When classifying our presented case in the literature, we find a typical melorheostosis manifestation of the long bone but report a previously not described appearance of an extraosseous lesion in the intercondylar notch. While future revision surgery seems statistically likely in our patient, he currently remains pain free and satisfied with the post-operative functional improvements at 12-month follow-up.

Discussion

The precise etiology of melorheostosis remains unclear [5, 6, 7]. However, more recently, Kang et al. identified somatic mosaic mutations in the regulatory domain of the MAP2K1 kinase which seem to be causative for the development of melorheostosis in some, but not all patients [6, 7]. Although this potentially opens possibilities of a targeted treatment, Wordsworth and Chan strengthen that targeting the MAP pathway will also affect not mutated cells resulting in potential off-target side effects [7].

While only around 500 cases of melorheostosis are reported in
Melorheostosis can have manifold clinical appearances that potentially require surgical treatment. Although subsequent revision surgeries to address new or repeated lesions cannot be precluded, a good to excellent functional outcome is possible. Nevertheless, melorheostosis remains a diagnosis that should be treated by orthopedic surgeons specialized in musculoskeletal oncology.

Clinical Message

Melorheostosis is a rare benign bone condition characterized by excessive segmental sclerosis of cortical bone being reminiscent of dripping candle wax. Clinical manifestations vary, but in patients with pain, deformity, and/or limited range of motion, surgical treatment might be required.

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