

Archive ouverte UNIGE

https://archive-ouverte.unige.ch

Article scientifique

Article

2019

Published version

Open Access

This is the published version of the publication, made available in accordance with the publisher's policy.

Idiopathic bone marrow oedema with joint effusion: A differential diagnosis to infectious osteomyelitis

Uckay, Ilker; Christofilopoulos, Panayiotis

How to cite

UCKAY, Ilker, CHRISTOFILOPOULOS, Panayiotis. Idiopathic bone marrow oedema with joint effusion: A differential diagnosis to infectious osteomyelitis. In: International Journal of Infectious Diseases, 2019, vol. 84, p. 97–98. doi: 10.1016/j.ijid.2019.05.007

This publication URL: https://archive-ouverte.unige.ch/unige:132662

Publication DOI: <u>10.1016/j.ijid.2019.05.007</u>

© The author(s). This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives (CC BY-NC-ND) https://creativecommons.org/licenses/by-nc-nd/4.0



Contents lists available at ScienceDirect

International Journal of Infectious Diseases



journal homepage: www.elsevier.com/locate/ijid

Medical Imagery

Idiopathic bone marrow oedema with joint effusion: A differential diagnosis to infectious osteomyelitis



ARTICLE INFO

Article history: Received 29 April 2019 Accepted 3 May 2019

Corresponding Editor: Eskild Petersen, Aarhus, Denmark*Keywords:* Idiopathic bone marrow oedema Autoimmune Fractures
Osteomyelitis

A 45-year-old healthy man reported progressive right hip pain during walking without a history of trauma or systemic inflammation. The pain responded favourably to ibuprofen and/or rest. Magnetic resonance imaging (MRI) revealed homogeneous inflammation with effusion into the joint (Figure 1). Idiopathic bone marrow oedema (IBME) of the femoral head was diagnosed radiologically. The patient was advised to practice partial weightbearing. No biopsies were performed. The patient declined to use crutches and relied on aspirin, ibuprofen, and intranasal calcitonin for 4 weeks. Two months later, his pain gradually disappeared over the course of 4 weeks. A control MRI at 5 months showed the absence of inflammation. The patient remained pain-free for the next 4 years.

IBME is a rare, (auto)immune disease of unknown origin, affecting mostly males between 30 and 60 years of age, and usually the lower extremity bones. Diagnosis is by MRI (low signal on T1, lack of subchondral changes, and high signal on T2; Emad et al., 2012). Biopsies are not necessary. Resolution is spontaneous. Affected persons must be told to have patience. Recurrences, osteonecrosis, and fractures are possible in severe cases. Treatment

is symptomatic and anti-inflammatory. The beneficial effects of non-weight-bearing and calcitonin or osteoporotic medication are anecdotal, with uncertain evidence (Ikemura et al., 2016), while intravenous prostaglandins represent promising current research (Meizer et al., 2005).

Funding

There was no funding for this article.

Patient consent

The patient consented by signature to the publication of his history and photographs.

Conflict of interest

The authors have no conflict of interest to declare. This publication fulfils the ethical requirements of the Declaration of Helsinki.

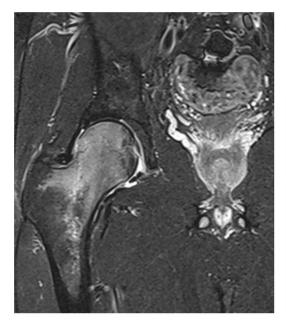




Figure 1. A 45-year-old male with idiopathic bone marrow oedema of the right femoral head and intra-articular effusions. Image obtained at diagnosis (left) and at the 5-month control (right).

Transparency declarations

None to declare. Parts of the manuscript have been presented as a poster at the Swiss National Conference of Infectious Diseases, August 30–31, 2017, Basel, Switzerland.

References

Emad Y, Ragab Y, El-Shaarawy N, Rasker JJ. Transient osteoporosis of the hip, complete resolution after treatment with alendronate as observed by MRI description of eight cases and review of the literature. Clin Rheumatol 2012;31:1641–7.

Ikemura S, Mawatari T, Matsui G, Iguchi T, Mitsuyasu H. Clinical outcomes in relation to locations of bone marrow edema lesions in patients with a subchondral insufficiency fracture of the hip: a review of fifteen cases. Br J Radiol 2016;86:20150750.

Meizer R, Radda C, Stolz G, Kotsaris S, Petje G, Krasny C, et al. MRI-controlled analysis of 104 patients with painful bone marrow edema in different joint localizations treated with the prostacyclin analogue iloprost. Wien Klin Wochenschr 2005;117:278–86.

Ilker Uçkay^{a,*} Panayiotis Christofilopoulos^b ^aInfectiology, Balgrist University Hospital, Zurich, Switzerland ^bOrthopaedic Surgery, La Tour Hospital, Meyrin, Switzerland

* Corresponding author at: Balgrist University Hospital, Forchstrasse 340, 8008 Zürich, Switzerland. E-mail address: ilker.uckay@balgrist.ch (I. Uçkay).

Corresponding Editor: Eskild Petersen, Aarhus, Denmark

Received 29 April 2019 Accepted 3 May 2019