



Case Report

Osteoid Osteoma of The Toe in Children: Case Report and Literature Review

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Introduction

Osteoid Osteoma (OO) constitutes 10-12% of all benign bone tumours. It has a male predilection and often occurs in younger patients. They may present with a nocturnal pain getting better by non-steroidal anti-inflammatory medication. Radiologically, a particular feature is the presence of a cortical lucent nidus surrounded by sclerotic bony reaction.

Some painless OOs of phalanx reported often have a similar circumstance of discovery: painless swelling of the toe accompanied by an enlargement of the nail (hippocratism) [1]. Closer the lesion

is to the skin, more the inflammation of the surrounding soft tissues will be visible and will manifest itself by swelling.

Case Report

We report about a 15-and-a-half-year-old boy. Since two years after a football match he complained about pain at the end of the second toe of the right foot. The additional swelling started painless with periods of night pain and some kind of painful swelling after sports. The toe showed a widening of the nail and makrodaktylie (Figure 1).



Figure 1



Figure 2

MRI (Magnetic Resonance Imaging) in fat-suppressed T2-weighted sequences revealed a hyperintense signal alteration of the distal phalanx, indicative of bone marrow edema, along with involvement of the adjacent soft tissues. In T1-weighted sequences, these structures appeared hypointense. During the contrast-enhanced phase, there was an increased contrast uptake in both the soft tissues and the bony distal phalanx, with a hypointense demarcation medially at the phalanx (Figure 2). Radiological findings suggested a high suspicion of osteomyelitis; however, no inflammatory laboratory markers were present.

Operation

Due to persistent symptoms and an enlarged distal phalanx, partial nail resection was performed, removing one-third of the nail along with adjacent soft tissues. This approach allowed the resection of the osseous defect while achieving a favorable cosmetic outcome.

Histology

Osteoblasts and osteocytes were observed at the peripheral margins, accompanied by osteoclastic giant cells. No atypical cellular components or necrosis were detected. Cartilage tissue

was not present. The findings were most consistent with an osteoid osteoma.

Literature Review and Our Case

Reviewing literature, we found 12 Case reports with an Osteoid Osteoma of the toe in children up to 16 years. Together with our boy we had 10 boys and 3 girls. The average age was 12,7 years.

Nearly all (12) had night pain more or less, 10 had macrodaktylie. In 6 cases we had the great toe-, 4 had the second-, one had the 3- and in two children the 4th toe that was affected. In 11 cases the distal phalanx had the problem. The majority (10) had a widening of the nail.

Literature	Year	age (years)	gender	toe	distal	nightpain	macro daktylie	differential diagnoses
Bordelon et al. [2]	1975	14	m	1 (first)	y	y	y	
Bellemans [3]	2021	11	m	4	y	y	y	
Haoudou et al. [4]	2021	13	m	3	y	y	y	
LaCroix et al. [5]	2001	7	m	4	y	y	n	osteomyelitis (OM) ?
Ozturk et al. [6]	2000	9	f	1	y	y	y	
Prietz et al. [7]	2009	12	f	2	y	y	y	trauma, OM ?
Rapp and Kaiser [8]	2010	14	m	1	y	n	y	
Seo et al. [9]	2023	13	f	1	MTP	y	y	OM ?
Sproude et al. [10]	2004	14	m	2	y	y	y	OM ?
Torrent et al. [11]	2017	16	m	1	MTP	y	n	enchondroma ?
Wang et al. [12]	2019	12	m	2	y	y	y	tumor ?
Yamaga et al. [13]	2015	16	m	1	y	y	n	OM ?
Our Case	2024	15	m	2	y	y	y	enchondroma, OM ?
Total		12,7	10 m/ 3f	6x1,4x2,1x3,2x4	11 vs 2	12 vs 1	10 times	

Thiemann et al. [14] reviewed 34 studies reporting a total of 37 cases of children and adults between 1975 and 2020. The mean age of these patients was 24.3 years (range: 7-53 years), with a male-to-female ratio of 1.6:1. The most frequently affected site was the great toe (n=22, 61%), while the other toes were less commonly involved (second toe: n=7, 19%; third toe: n = 2, 6%; fourth toe: n = 3, 8%; fifth toe: n=2, 6%). In the majority of cases, the distal phalanx was affected (n=31, 86%).

The most common differential diagnoses included infection (n=17, usually suspected osteomyelitis or paronychia) or a benign/malignant tumour (n=14).

This was comparable to our review with the children. But we had 10 boy's vs 3 girls affected.

When the radiological appearance is not typical because central calcification was lacking and other entities could be possible such as such as nonossifying fibromas, enchondromas, eosinophilic granulomas, tuberculosis could be possible as Noordin et al. [15] described.

In the case of osteoid osteoma of the toe in children, surgical treatment is generally considered the therapy of choice although in long tubular bones, Radiofrequency Ablation (RFA) is regarded as the gold standard of treatment [16].

But RFA is not a viable option for toe lesions. The main reasons include the frequently uncertain dignity of the lesion, the very small lesion size, the risk of expected skin necrosis due to the heat from the needle tip, and the immediate proximity of the lesion to neurovascular structures in the toes. Nevertheless, when complete resection of the nidus is achieved and surgical treatment also demonstrates high success rates, ranging from 88% to 100% [16,17].

Finally, our patient was happy without pain and a normal looking toe.

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