# **Aneurysmal bone cyst arising in iliopubic chondromyxoid fibroma – A case report**

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# **Running Title:** Association between aneurysmal bone cyst and chondromyxoid fibroma in a location never described before

# Abstract:

Chondromyxoid fibroma is rare cartilaginous tumor, accounting for 0.5% of all primary bone tumors and 2% of benign bone tumors. Areas of aneurysmal bone cysts (ABC) may be found within CMF in 8.6% of cases. A 20-year-old man presents pain due to a mass on iliopubic ramus that was diagnosed as an aneurysmal bone cyst arising from a chondromyxoid fibroma. This case confirms the rare association between aneurysmal bone cyst and chondromyxoid fibroma in a location never described before. Although aneurysmal bone cyst is more frequently associated with highly vascularized tumors, it is important to consider the diagnosis in the presence of a chondromyxoid fibroma.

Key-words

Chondromyxoid fibroma; Secondary aneurysmal bone cyst; Iliopubic ramus

# Introdution

Chondromyxoid fibroma (CMF) was described by Jaffe and Lichtenstein as a distinctive entity in 1948.1 The definition given by the World Health Organization (WHO) is benign cartilaginous neoplasm, composed of lobules formed by spindle-shaped cells with myofibroblastic features at the periphery, and stellate and chondrocyte-like cells towards the center, the extracellular matrix of the lobules is fibrous at the periphery and myxoid and chondroid towards the center. 2

This rare cartilaginous tumor, accounting for 0.5% of all primary bone tumors and 2% of benign bone tumors, occurs predominantly in adolescents and young adults and more commonly in males.23 This tumor can occur at almost any osseous site, with predilection for bones of the lower extremities, usually the proximal tibia.2,3 Flat-bones account 25 to 30.3% of cases, with ilium been the most affected.2,4,5

Areas of aneurysmal bone cysts (ABC) may be found within CMF in 8.6% of cases. 4

ABCs is result of specific pathologic change, which is probably the result of trauma or a tumor-induced anomalous vascular process.6 The most common as giant cell tumor, but other like osteoblastoma, angioma, chondroblastoma, telangiectatic osteosarcoma are common to. Rarely is secondary to CMF. 6

To the best of our knowledge, there is not any case of aneurysmal bone cyst arising from an iliopubic chondromyxoid fibroma.

# Case Report

A 20-year-old man with no relevant medical history initially resort to General Surgery outpatient clinic appointment and presents pain in the left inguinal region with 6 months of evolution, precipitated and aggravated by efforts and relieved by rest. Physical examination demonstrated a palpable mass in the iliopubic ramus.

The conventional radiography (figure 1) revealed thinning of the left ischio-pubic ramus and a fracture line in the central region. MRI presents a weel defined lesion with lobulated contour and internal septation (figure 2) changes suggestive of aneurysmal cyst or osteosarcoma. The surgical biopsy suggested the diagnosis of chondromyxoid fibroma.

The surgical treatment was performed through an anterior approach and a Pfannenstiel incision was extended to the proximal region of the left thigh. Careful progression through fascia-muscular planes was performed, with isolation and protection of the vasculo-nervous bundle and spermatic cord, followed by en-bloc resection of iliopubic ramus and the soft tissues that surrounded it (figure 3). Abdominal wall was reinforced with Gore-Tex® mesh to prevent intra-pelvic herniation.

The patient was discharged home 2 days after surgery, with partial-weight bearing and no evidence of complications.

Microscopic examination showed complete excision of the lesion, with free margins.

Histological analysis revealed a benign mesenchymal neoplasm, well delimitated and organized in chondromyxoid lobes of variable cellularity, with stellate cells in myxoid areas and lacunar cells in chondroid areas, features consistent with chondromyxoid fibroma (fig. 4.1 and 4.2). There were also abundant hemorrhagic cavitated areas developing within the tumor, with septa rich in osteoclastic-type multinucleated giant cells and with deposition of linear immature osteoid, typical of aneurysmal bone cyst transformation (fig. 4.3 and 4.4), explaining the images seen in MRI. The neoplasm occupied and expanded the ischio-pubic ramus, reaching the subcartilaginous area of the pubic symphysis, but it didn’t invade the adjacent soft tissue, as it was surrounded by a thin layer of woven bone, consequence of cortical bone remodeling.

After 1 year of follow-up, the patient is pain-free, without functional impairment or complications. There is no recurrence until present.

# Discussion

This case represents an unusual localization for this type of tumor. A literature review of 278 cases of CMF of bone, reported that 30.3% of cases occurred in flat bones. Of these only 1 case (1.2%) was located on pubis and 2 on ischium (2.4%). 4

As the name indicates, chondromyxoid fibromas show a variety of histological features. The classic histological feature is stellate or spindle-shaped cells arranged in lobules in a myxoid background. 4 However, a lobulated pattern was seen in only 86.7% of cases, and can present a macro or microlobular pattern isolated or a mixture. 4

In our case, the tumor was accompanied by area of ABC. The origin of term ABC comes from an article by Jaffe and Lichtenstein in 1942.7 They postulated that ABC might be a secondary phenomenon due to hemorrhagic “blow-out” in a preexisting lesion, which may be destroyed in the process. 7 A preexisting lesion was identified in approximately one third. 6

ABCs conventional treatment has been the surgical removal of the entire lesion, or of as much as possible, with recurrence seen in 10-70% of cases.2,6 Spontaneous regression following incomplete removal is very unusual.2 More recently, non-surgical polidocanol sclerotherapy demonstrates to be a safe and effective option even for aggressive ABCs, depending on location and surgical risks.8 Appropriate treatment requires realizing that it has a specific pathophysiologic origin, and identifying the preexisting lesion. If no coexisting lesion is identified, lesion are usually treated with curettage and bone grafting. 6

The typical treatment of CMF by curettage has a 20% to 25% recurrence rate, lowering to 7% with addition of bone grafting.9,10 Resection provides lower recurrence rates but is not always feasible.

Based on the system proposed by Enneking WF and Dunham WK, this case of CMF involves zone III (involvement of pubis and ischium bones). 11 A review of 8 cases describing surgical management of CMF of pelvis, reported two cases of involvement of superior ramus of pubis. One was treated with resection without grafting and complicated with pelvic internal organs herniation. On second case reconstruction was done using fibular strut allograft, with none complication on follow-up. 12 Another case of CMF of iliopubic ramus presented good results with aggressive curettage and chip bone grafting. Author reclaims this procedure as safe, easy, and less morbid for this anatomic site. 13

In our case, we resected iliopubic ramus and used a Gore-tex® surgical mesh to prevent intra-pelvic herniation. Resection should be performed for large lesions or in anatomic sites such as the complex pelvis where such procedure does not increase the risk of fracture and reduce the likelihood of recurrence or a second surgery.

The secondary ABC arising from CMF was verified histologically. It is associated to a significant recurrence rate and the surgical strategy adopted was then confirmed the most appropriate.

This case represents a rare tumor, in an uncommon location, with an atypical histological transformation. This case shows the rare association between ABC and CMF. Although secondary ABCs are more frequently associated with highly vascularized tumors, it is important to consider the diagnosis of CMF in the presence of an ABC, as it may alter the clinical approach to minimize the risk of recurrence.

# Authors' Contributions

João Vale: Wrote the manuscript, Performed the literature review.

Sara Diniz: Edited the paper

Filipe Rodrigues: Edited the manuscript

Ana Ribau: Edited the paper

André Coelho: Edited the manuscript

Vânia Oliveira: Edited the manuscript

Pedro Cardoso: Edited the manuscript

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None.

# Conflicts of Interest

None.

# Consent for Publication

Written informed consent was obtained from the patient for publication of this case report and any accompanying images.

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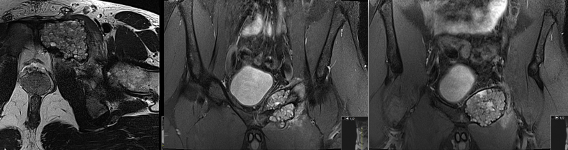
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# Figures



**Figure 1** – Preoperative radiographic image shows thinning of the iliopubic ramus and a fracture line in the central region.

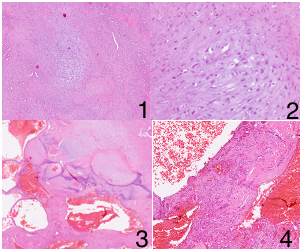


**Figure 2** - Preoperative MRI: weel defined lesion with lobulated contour and internal septation

Uma imagem com mesa, sentado, alimentação

Descrição gerada automaticamente

**Figure 3** – En-bloc resection of iliopubic ramus and the soft tissues that surrounded it



**Figure 4** – 1) The tumor has a zonal, vaguely lobulated architecture, with hypocellular chondromyxoid matrix at the center and a more cellular spindled-cell component at the periphery; 2) The chondroid central areas have oval to stellate cells, some of them residing within lacunae surrounded by cartilage-like matrix; 3) Secondary aneurysmal bone cyst-like areas are seen throughout the neoplasm, with hemorrhagic spaces; 4) In these areas, the septa contain numerous osteoclastic-type multinucleated giant cells, abundant hemosiderin and linear deposition of osteoid matrix.

Uma imagem com tatuagem, fotografia, branco, preto

Descrição gerada automaticamente

**Figure 5** - Postoperative x-ray, after the ressection of iliopubic rami