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Bilateral rapidly progressive osteoarthritis of the hip: Case report and review

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Abstract

Currently rapidly progressive osteoarthritis of the hip (RPOH) is an infrequent syndrome that compromises the femoral head and acetabulum described by Forestier in 1957.

There is no standardized definition of the characteristics of this disease. Other diseases such as osteonecrosis or inflammatory and infectious arthropaties process may appear similar or present concurrently with osteoarthritis.

Clinically it is characterized by hip pain characterized by chondrolysis with massive destruction of the femoral head, with or without acetabular involvement. The associated radiographic changes normally occur within 12 months after the onset of symptoms.

The reported incidence is 7.2 to 15.7%. The patient is frequently a woman and the involvement in the vast majority is unilateral. The etiology remains uncertain, but increasing attention in recent years suggesting subchondral fractures as a contributing factor to the development this disease. We described a case review of a male patient with the development of bilateral RPOH.

Keywords: Dysplasia, osteoarthritis, osteonecrosis, rapidly progressive arthritis

Introduction

Currently rapidly progressive osteoarthritis of the hip (RPOH) is an infrequent syndrome that compromises the femoral head and acetabulum, first reported by Forestier in 1957^[1]. There is no standardized definition of the characteristics of this disease, accepting in many cases the definition proposed by Lequesne as the loss of joint space of 2 mm or more per year or the loss of 50% or more of joint space in one year in the absence of another cause of destructive arthropathy ^[2]. Other disease process such as osteonecrosis (painful condition that can be especially difficult to distinguish), inflammatory and infectious arthropaties, may appear similar or present concurrently with osteoarthritis.

Clinically it is characterized by hip pain characterized by chondrolysis associated with massive destruction of the femoral head, with or without acetabular involvement, which is evidenced in a radiographic study in the absence of signs of sepsis, neurological, metabolic or inflammatory disease ^[3]. The associated radiographic changes normally occur within 12 months after the onset of symptoms ^[4].

The reported incidence is 7.2 to 15.7% ^[5]. The patient is frequently a woman and the involvement, in the vast majority of reported cases, is unilateral⁶. The etiology remains uncertain, but subchondral fractures have gained increasing attention in recent years, suggesting subchondral fractures as a contributing factor to the development of rapidly destructive osteoarthritis⁷. With the aim of providing more clinical data, and given the low incidence with which bilaterality is described in this disease, the clinical case of a male patient with the development of bilateral RPOH is reported below.

Case report

We present the clinical case of a 73-year-old man; due to an exacerbation of hip pain of 2 months of evolution, where he reported impossibility to walk without mechanical aids.

Personal history of AHT, Heart Failure with an ejection fraction of 54% and Sleep Apnea-Hypopnea Syndrome under treatment with nocturnal CPAP, as well as a history of prostate cancer with treatment He was referred to an outpatient trauma clinic for right hip pain. He has a body mass index (BMI) of 36.44 and smoking as a toxic habit.

Complementary examinations were performed that included routine laboratory tests with an inflammatory profile and lumbar and pelvic x-rays (Fig. 1), which showed degenerative changes at the level of both hips with decreased joint space, without osteophytosis, with

maintenance of joint morphology, osteoarthritis of the hip, the process to which the symptoms were attributed; no other relevant findings.

Twelve months later, the patient was admitted to the Orthopaedic Service for scheduled total hip arthroplasty (THA) surgery, having suffered an even greater exacerbation of hip pain during the time on the waiting list, limiting mobility to a wheelchair. He was admitted for surgery arriving in a wheelchair.

The right hip surgery was performed throught posterior approach, a remarkable destruction of the right femoral head as well as the acetabulum was evidenced, take samples for strip leucocyte esterase assay and alpha-defensin test both negative. Take samples for microbiology and histology studies. Performing a surgery to cementless total hip arthroplasty replacement implantation acetabular Trident with double mobility and stem cementless Accolade II (Strycker^R).

The anatomopathological and microbiological studies of bone and synovial tissue extracted were negative. Histological sections revel severe degenerative joint disease, bone avascular necrosis, and chronic synovial inflammation. The postoperative radiographic control is reported below (Figure 2), evidencing a destruction of the contralateral head with significant acetabular involvement.

The postoperative period was uneventful, with adequate pain control and evolution of the surgical wound, recovering the ability to walk with canes, but with persistent left hip pain. Months later, he underwent a second surgical intervention, in this time in the left hip, performing THA with cementless total hip arthroplasty replacement, acetabular revisión with Delta Revision TT cup (Lima^R) and Corail cementless collar stem (Johnson^R) (figure 3). Again, samples were sent for a microbiological study that were negative and the anatomopathological study did not differ from the previous one.

The patient evolved without complications, with adequate evolution of the surgical wound and adequate pain control, being referred to a rehabilitation treatment program.

Surgical and rehabilitative treatment were effective, and the discomfort of the patient practically disappeared. He maintained an adequate joint balance in the prosthetic joints and walked with the help of two canes with few restrictions the first three months post-surgery, carrying out progressive withdrawal of aids for ambulation.

Discussion

Although RPOH was described in 1957, there are few references in the literature of it being considered a variation of primary hip osteoarthritis. RPOH remains a poorly understood disease of unknown etiology, where systematic reviews define a more frequent involvement of women, a mean age of 70 years and unilateral involvement ^[8]. The national registry of England suggests an increase in the incidence of RPOH that may be related to longer surgery waiting lists, since the time from symptoms to destruction can be as short as 12 months ^[9-11], as exemplified by the clinical case reported.

Patogenesis of the disease remains uncertain though immunologic effects cytokine mediated, subcondral insuficiency fractures or toxicity NSAIDs consumption are suspec-ted pathologic mechanism, suggesting they impair bone turnover. At first, x-ray studies show normal anatomy or mild osteoarthritic changes, secondly, months after the appearance of the symptoms appear femoral head and acetabulum destruction, with sclerosis and subchondral cysts with minimal or absence ostophytes. The rapid progression of this disease makes it difficult to obtain sequential radiographs in early stages ^[3].

RPHO was classified into three types by Lequesne and Amoroux in 1970, it depends on the amount of bone loss and time period over which this takes place ^[12]. Posteriorly Zagyva *et al.* ^[7] describe a new gradding system and classification for patients with RPHO.

The disease developes, first time the joint space was narrowing (grade I), and progresses to a complete disappearance of the articular joint space (grade II), in this moment the femoral head and acetabulum could be deformed; partial osteolysis appears in grade III, with superolateral ascension of the femoral head in more of them. Histology sections mostly reveal extensive fibrosis of the joint capsule, disappearance of the cartilage in all cases, acute or chronic inflammation of soft tissues, in most cases the synovial membrane revealed imflammation and hyperplasia; and bone with absence of specific inflammatory cells, bone resorption and focal loci of osteonecrosis in the subchondral bone and also of a distinction between healthy and necrotic tissue.

The review carried out by Charrois *et al.* ^[13] found that THAs carried out in patients with RPOH have a higher degree of difficulty and frequently require additional procedures due to bone loss, longer duration of surgery and need for further procedures, particularly acetabular reconstruction, increased transfusion requirements and longer surgical time. This is consistent with the clinical case reported in which he required dual mobility THA, as well as a revision cup on the left side due to the significant bone loss described. In this line, Kawai *et al.* ^[10] used kerboull-type acetabular reinforcement devices and graft in all their reported cases, Yuasa *et al.* ^[11] and Peters *et al.* ^[14] report the use of a revision acetabulum in all their cases.



Fig 1: Initial pelvic radiograph shows narrowing of the joint space and subchondral sclerosis, without osteophytosis



Fig 2: The pelvic radiograph shows postoperative control of the right CTA and shows acetabular and femoral destruction of the left hip



Fig 3: The pelvic radiograph shows postoperative radiographic control

Conclusion

The RPOH is a disease that causes a rapid unfavourable evolution of the symptoms with a significant affectation in mobility and severe disability. Bilaterality is a rare but possible feature in this disease.

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