Patient with Dactylitis and Enthensopathy- Forestier Disease - RADISH Variant

Luis Arturo Gutiérrez-Gonzalez¹* and Salvador Malave²

¹Rheumatology and Internal Medicine Unit, El-Avila Clinic, Altamira, Caracas 1060, Venezuela
²Department of Radiology, Doximity, The Professional Network for Clinicians, Miami, Florida, USA

*Corresponding author: Luis Arturo Gutiérrez-Gonzalez, Rheumatologist, Rheumatology and Internal Medicine Unit, El-Avila Clinic, Altamira, Caracas 1060, Venezuela, Tel: +58212276.1582, Fax: +58212276.12583; E-mail: Umircaracas@yahoo.es

Received: January 23, 2018; Accepted: January 29, 2017; Published: January 31, 2017


Abstract

Diffuse idiopathic skeletal hyperostosis (DISH) is a bony hardening (calcification) of ligaments in areas where they attach to your spine. Also known as Forestier's disease, diffuse idiopathic skeletal hyperostosis may cause no symptoms and require no treatment. The most common symptoms are mild to moderate pain and stiffness in your upper back. DISH can also affect your neck and lower back. Some people have DISH in other areas, such as shoulders, elbows, knees and heels. A non-inflammatory condition, DISH can be progressive. As it worsens, it can cause serious complications.

This is a case report of a patient with dactylitis and enthensopathy - Forestier disease - RADISH variant

Keywords: Diffuse Idiopathic Skeletal Hyperostosis (DISH); Spondyloarthritis; Rheumatoid arthritis; Spinal symptomatology

Introduction

Diffuse Idiopathic Skeletal Hyperostosis (DISH) or Forestier’s disease is a condition whose primary manifestation is the ossification of the anterior longitudinal ligament (ALL) and the development of intervertebral bony bridges. Diffuse Idiopathic Skeletal Hyperostosis (DISH), also called Forestier’s disease is a rheumatological condition mostly affecting the axial skeleton. However, a subset of these patients presents with peripheral arthritides and enthensopathies that mimic spondyloarthritides and even rheumatoid arthritis. This group of patients is called RADISH (Rheumatism articular in Diffuse Idiopathic Skeletal Hyperostosis), a systemic disease characterized by bone neoformation in the ligament, tendon and capsular attachments.

Case Presentation

This is a 75-year old male patient presenting with a 10-year history of axial pain and mixed characteristics. The pain improves through the morning (<2 hours), and then gets better after resting at the end of the afternoon – evening [1,2]. The patient self-medicates with standard painkillers such as acetaminophen 1 gram, alternating with various NSAIDs (sodium diclofenac, ibuprofen and naproxen) using on demand dosing. Since the pain fails to improve and the patient experienced limited neck rotation, the patient decides to visit the rheumatologist. A complete Spine X-ray is required which upon studying the cervical spine evidences diffuse hyperostosis along the anterior aspect of the cervical spine, with bony bridges at various levels. The hyperostotic bone is present at the edges of the cortex of the original vertebral body at >45° angles; the intervertebral disc spaces are intact and there are no erosions or romanus sign (Figure 1).

Figure 1 Cervical bony bridges.

A routine lab test in addition to an immune-rheumatological profile (ANA, Ra Test, Anti CCP and HLA B27) were requested, which were reported negative or within normal, and the patient was prescribed meloxicam 15 mg per day for 6 weeks, together with physical therapy resulting in satisfactory improvement [3,4].
14 months later the patient comes back presenting with boxing glove arthritis in the right hand only, in addition to arthralgia in both shoulders and ankles, with dactylitis of the 3rd left foot toe. A routine lab test was prescribed (CBC, blood chemistry, PPD (Tuberculin), feet X-rays, X-rays of both hands, Chest X-ray [5,6]. A recommendation was given to repeat the Ra Test and Anti CCP, just reporting elevation of acute phase reactants (VSG 78 mm; PCR 4.32 mg/dl) [7]. An ultrasound of the joints in both hands showed inflammation of all the tendon sheaths of the extensors of the fingers, with no erosions but just osteodegenerative changes consistent with age, but with a power/Doppler (+) sign. The patient started with deflazacort at a 30mg dose tapering by 6mg/week until a minimum dose of 6mg was reached, which was maintained for 3 months, with good response and tolerance [8].

Discussion

In the classical DISH the basic lesion is the ossification of the anterior longitudinal ligament (ALL) developing intervertebral bridges. Forestier and Rotès-Quérol in 1950 described the disease and called it ankylosing hyperostosis of the spine, as opposed to disc arthrosis or ankylosing spondyloarthritis. Resnick in 1975 recommended the term diffuse idiopathic skeletal hyperostosis, insisting on its systemic nature with extra-vertebral involvement and defined the specific radiological criteria currently used [9,10]. This pathology has been described in routine autopsies and presents more often in males between sixty and seventy years old. The spinal symptomatology is the most frequently described clinic, with pain, cervical and dorsal stiffness, but most of the patients are asymptomatic (Figure 2).

A DISH peripheral variant has been described called RA-DISH (Rheumatoid arthritis in diffuse idiopathic skeletal hyperostosis), where the patient presents with arthralgias and in some cases arthritis of the small and large joints that mimic rheumatoid arthritis, except for the absence of Ra Test and other biomarkers. From the clinical point of view, the age of presentation is mostly between 60 and 70 years old and is usually asymmetrical [11,12].

Among other radiological findings, RADISH differs from rheumatoid arthritis (RA) because usually osteoporosis is absent, but there is often bone sclerosis and proliferation of erosions, in addition to osteophytes and bone ankyloses. Some of the atypical clinical characteristics include a high incidence of flexion contractures of the elbows, wrists, ankles and knees. It is no surprise that the production of bone takes place on the joints involved in patients with RA-DISH, since the latter disorder is characterized by bone proliferation at the bone attachment sites of ligaments and tendons in the axial and extra-axial skeleton, probably stress related. One of the extra-axial manifestations of RA-DISH is dactylitis (Figure 3).

Criteria according to Resnick:

1. Presence of ossification along the anterolateral aspect of four contiguous vertebral bodies leading to the formation of bone bridges.
2. The relative preservation of the intervertebral disc height in the affected region and the absence of evolving radiological changes, hence confirming degenerative disc pathology.
3. The lack of erosion, sclerosis or bone ankyloses of the sacroiliac joints or the posterior interapophyseal joints.

Dactylitis is a typical manifestation of spondyloarthopathies and in particular of psoriatic arthritis, though there are other causes that may cause this peculiar inflammation. There are inflammatory dactylitis (spondiloarthopathies, gout or sarcoidosis), infectious dactylitis (tuberculosis, syphilis or blistering distal dactylitis) or non-inflammatory dactylitis (sickle cell anemia) [13,14].

Conclusion

Using conventional radiology, no vascular infarction of the bone marrow was identified in our patient, which is the typical lesion in sickle cell anemia dactylitis; even the blood test only reported VSG and PCR elevation [15]. In case of infection...
secondary to syphilis or brucellosis, no radiological signs of osteomyelitis were observed which the result of soft tissue damage is. There was no disruption either of the fat pad of the volar surface of the fingers (typical radiological lesion secondary to beta-hemolytic Streptococcus Group A infection) [16,17]. The only finding in our patient was a Tenosynovitis of the flexor sheath, typical of non-infectious inflammatory dactyilitis and not due to crystal deposits or granulomatous tenosynovitis.

References