Introduction

SAPHO syndrome (synovitis, acne, pustulosis, hyperostosis, and osteitis) includes a variety of rheumatic alterations associated with skin lesions. It generally presents during infancy and adolescence, mainly in females.

The distribution of the illness depends on the age of presentation. In adolescents and during middle adulthood, it mainly is localized to the sternoclavicular region, followed by the lumbar spine, the pelvis, and long bones. During infancy it affects long bones (tibia, femur, and ulna), the clavicle and the lumbar spine. The diagnosis of SAPHO syndrome is not difficult when the osteoarticular lesions are localized in the characteristic target areas. The gammagraphic bone scan image simulating a "cow horn" of the sternoclavicular joint is very specific to this syndrome.

Clinical Case

We describe a case of SAPHO (synovitis, acne, pustulosis, hyperostosis, and osteitis) with an unusual presentation in a 14 year-old girl with low grade fever which lasts 8 months, left low back pain and elevated erythrocyte sedimentation rates and protein C-reactive with chronic anaemia. A radiography of the lower limbs showed a lytic image with osteitis and hyperostose in the right fibula, as a casual finding. This information, in addition to the acne, pustulosis, sternoclavicular arthritis, and the studies got from the magnetic resonance image (MR) of spine, pointed out the diagnosis of SAPHO.

Key words: SAPHO syndrome. Spondyloarthritis. Chronic recurrent multifocal osteomyelitis (CRMO).
nerve root compression. She also presented pain on the left Achilleal enthesis. She did not have pain on the fibromyalgia trigger points and had no ulcers on the vulvae.

**Complementary testing**

Hemogram: hemoglobin, 11.7 g/dL; hematocrite, 33.9%; MCV, 81 fl; normal leucocytes; CRP, 20 mg/L; ESR, 42 mm/1st h.

Blood chemistry: uric acid, 6.5 mg/dL; ferritin, 65 ng/mL; immunoglobulin, C3, C4, and TSH, normal; rheumatoid factor, ANA, HLA-B27, urine culture, and a pharyngeal exudate were all negative; serology for mononucleosis and yersinia were negative; Schirmer’s test, saliva secretion, and pathergy testing were negative.

The following imaging techniques were carried out: peroneal x-ray, showing a lytic image with osteitis, and hyperostosis in the right peroneal bone (Figure 2); normal chest x-ray; lumbar MRI showing a protrusion of L5–S1, with a diminished signal and a nodular lesion <1 cm, isointense on T1 and hyperintense on T2, adjacent to the right vertebral margin of D8, non-specific, and a thoracic bone scan that presented an increased uptake of the sternoclavicular joint that simulated a “cows horn” as well as an increased uptake in the right sternoclavicular joint (Figure 3).

SAPHO syndrome was diagnosed and treatment was initiated with naproxen and cloxacyllin, showing a discreet improvement, after which the use of disease modifying anti-rheumatic drugs (DMARD) was proposed to the family (methotrexate and sulfasalazine) and pamidronate, which they rejected. The patient is currently receiving no treatment and symptoms, pain and elevated acute phase reactants persist.
Discussion

SAPHO syndrome is a controversial entity. It was described by Kahn et al to group a series of conditions with common findings, such as bone affection with aseptic osteitis that affects determined zones and the skin lesions in the form of palmar and plantar pustulosis as well as acne conglobata. In many of these cases, sacroileitis is found additionally, evolving as a spondyloarthropathy. On the other hand, chronic, recurrent, multifocal osteomyelitis (CRMO) is a chronic, aseptic, non supplicative bone inflammation that affects multiple localizations, generally long bones and the clavicle, and less frequently the spine and pelvis, frequently affecting children and adolescents. It is also frequently associated with palmar and plantar pustulosis and has a favorable evolution. This similarity between entities makes it impossible to reach a consensus on whether they are separate diseases or a spectrum of the same disease, though it seems that SAPHO syndrome carries a worse prognosis in the long term.

The fundamental component of SAPHO is an inflammatory osteitis that can or cannot be associated to skin lesions, presenting negative bacterial cultures. These skin lesions typically are palmar and plantar pustulosis and acne (55.7 and 19.3%, respectively). They can either precede, occur simultaneously or after the start of the skin lesions. The osteoarticular lesions include synovitis, hyperostosis, and osteitis. The distribution depends on the age of presentation, predominantly on the sternoclavicular region during adolescence, while during infancy it affects long bones (tibia, femur, and ulna), the clavicle and lumbar spine. Systemic manifestations are rare, but fever can sometimes be present.

In the SAPHO syndrome with skin lesions, a high percentage of cases have detectable Propionibacterium acnes in the synovial fluid, but its role in the pathogenesis of the disease is not clear. Imaging in this patient helped us orient the diagnosis, in spite of a negative joint fluid culture.

Conventional treatment of the SAPHO syndrome (with non steroidal antiinflammatories, steroids and methotrexate) have not proven effective. There are currently several studies that support the superiority of pamidronate for pain reduction and an improvement in functionality with few adverse events. This is a second-generation biphosphonate that suppresses bone resorption and has anti-inflammatory properties. We think that knowledge on this syndrome is important because its diagnosis is unique and requires a high index of suspicion. We hope to stimulate pediatric rheumatologists to the careful inspection of the skin and the anterior thorax that, in our opinion, are key points in elucidating this condition.

References