Pyomyositis is rarely seen in temperate climates. Typically, it presents with the formation of an abscess requiring surgical drainage and it has been reported as a differential diagnosis for septic arthritis of the hip.

We describe the occurrence of pyomyositis of the iliacus muscle in a ten-year-old girl which was diagnosed by MRI and blood culture. Formation of an abscess did not occur despite marked focal inflammation and swelling of the muscle. Conservative treatment with antibiotics alone led to complete clinical and radiological resolution of the infection.

We could find no previous description of pyomyositis in a child in the British orthopaedic literature. Orthopaedic surgeons, particularly those with a paediatric interest, should be aware of this condition and its presentation, diagnosis and treatment.

**Case report**

A ten-year-old Caucasian girl was referred by her general practitioner with a two-week history of increasing pain in the left side of her lower back radiating to the left buttock, the posterior area of the thigh and the knee. The pain had begun after a vigorous physical exercise (PE) lesson at school and had gradually increased until she was unable to bear weight. She was admitted to hospital.

There was no history of recent foreign travel, prodromal illness or any other significant medical or family history.

On examination she was distressed, in pain and afebrile. The left lower back and the sacroiliac and buttock areas were tender. The range of movement of the left hip was markedly reduced, particularly flexion, although rotation was free in extension. On the right the straight-leg raise was to 90° at which point pain was felt in the left lower back. That on the left was limited to 10° because of severe pain in the left lower back and buttock. Neurological examination was normal.

Initially, haematological investigations showed a leucocytosis of $16.6 \pm 10^3$ per mm$^3$, a neutrophil count of $13.50 \pm 10^3$ per mm$^3$ (normal 2.0 to $7.5 \times 10^3$ per mm$^3$), an ESR of 51 mm/hr and a C-reactive protein level of 90 mg/l (normal <10 mg/l). Urine microscopy and culture proved negative.

Overnight, she had a pyrexia of 38.5°C and the following day her condition remained unchanged. We carried out blood cultures and screening tests for anti-streptolysin O and rheumatoid factor. A provisional diagnosis of acute infective lumbar discitis or acute sacroiliitis was made.

An MR scan was carried out. This showed an enlarged left iliaceus muscle with a low signal on the T1-weighted and a high signal on the T2-weighted images (Fig. 1). The findings were thought to be consistent with localised inflammation of the muscle, in the absence of an abscess or any other abnormality in the surrounding bones and joints including the left hip and sacroiliac joint. The diagnosis of pyomyositis was confirmed when *Staphylococcus aureus* was isolated from the blood cultures.

She was initially treated by bed rest and with intravenous Magnapen, 500 mg every six hours, which was changed to flucloxacillin 500 mg intravenously when *Staphylococcus aureus* was isolated. By the fifth day after admission her
pain had markedly improved; she was able to walk independently partially weight-bearing on crutches, and had been afebrile for 48 hours. At this stage she was discharged home on oral flucloxacillin. One week later her pain had almost completely resolved and at six weeks she was asymptomatic. Her full blood count, ESR, CRP and MRI at six weeks were all normal (Fig. 2).

Discussion

Pyomyositis is a condition usually seen in hot climates and commonly affects the muscles around the hip. It has rarely been reported in adults in the UK. A review of the literature revealed only one case previously described in the UK, in a nine-year-old boy who presented with an abscess in the psoas muscle which was diagnosed by CT and required surgical drainage. MRI was crucial in making an early diagnosis. The high signal on the T2-weighted and the low signal on the T1-weighted images confirmed the florid inflammation within the iliacus muscle, without formation of an abscess. It has been recognised as the radiological investigation of choice in the diagnosis of pyomyositis. In addition to demonstrating the abnormal iliacus muscle, it was possible to rule out associated infection in the pelvis and hips. Our findings contrast with those of other reports in which presentation has often been with an abscess requiring surgical drainage. Most of these cases did not have the benefit of MRI. Predisposing factors were often present such as skin conditions, pyarthrosis, recent surgery and wound dehiscence, diabetes, childbirth and epidural cannulation, and HIV disease. There were no such features in our case, which appeared to be a primary infection of muscle with no secondary spread from adjacent structures.

Pyomyositis affecting the iliacus muscle and the adductor muscles of the hip has been described as a differential diagnosis for septic arthritis of the hip. Our patient presented with left-sided low back, buttock and thigh pain and the free movement of the left hip in extension made septic arthritis unlikely. The iliacus muscle is closely associated with the psoas muscle, the femoral and lateral femoral cutaneous nerves, the hip and pelvic and intra-abdominal structures. Pathological abnormality in the iliacus muscle may present, as in our patient, with lower abdominal pain, pain in the hip or femoral neuropathy.
Staphylococcus aureus was the infecting organism and is responsible for pyomyositis in approximately 90% of cases. It is not clear why there was no progression to formation of an abscess since the condition had been developing over two weeks. This is comparable to the range of three days to three weeks which has been described elsewhere and in which the primary presentation was usually with an abscess.

It may be that primary pyomyositis occurring in a deeply-sited muscle is a different clinical entity from the formation of an abscess in more superficial muscles, which are more susceptible to the spread of infection from adjacent skin lesions or puncture wounds.

The history of previous physical exercise may or may not be relevant. It is possible that a ‘minor’ strain of the iliacus in some way made the muscle more susceptible to transient bacteraemia. This association has been recognised previously. The initial MR scan showing a low signal on the T1-weighted and a high signal on the T2-weighted images in the iliacus is consistent with inflammation and not haemorrhage.

Pyomyositis has been reported in children in temperate climates, and in a British child in the paediatric literature. It should be considered in the differential diagnosis of any child with pyrexia complaining of joint pain or muscle aches, particularly as a differential diagnosis in septic arthritis of the hip. Orthopaedic surgeons should be aware of this condition. MRI is invaluable in establishing the diagnosis and its anatomical site and in helping to exclude other possible diagnoses.

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References