Bone scintigraphic findings in a patient with β thalassemia major and knee pain

Haematologica 2004; 89:(5)e73

Skeletal abnormalities occours in human hemolytic disorders associated with thrombosis, such as beta-thalassemia and sickle cell disease.¹ The osteoarticular complications include avascular necrosis, common at all ages; acute septic arthritis and hematogenous osteomyelitis, that usually affect infant and children.² Tc-99m methylenediphosphonate (MDP) bone scintigraphy, eventually in combination with colloid marrow scan, could help clinician in the differential diagnosis between aseptic bone infarcts and secondary osteomyelitis.^{2,3} Magnetic resonance imaging (MRI) either in the rat model, imitating acute hemolysis and thrombosis or in humans, demonstrated to be useful in revealing avascular necrosis of bone.^{1,4}

We report a 29-year old female with beta-thalassemia major undergoing to transfusion therapy monthly. Due to the onset of left knee pain a bone scintigraphy was performed. Scan findings were consistent with postischemia recovery hyperactivity of left condylus medialis femoris following avascular necrosis. In addition, bone scan showed hyperactive marrow and nephromegaly; a renal abnormality described in patients with hemophilia related to multiple blood transfusions.⁵

In conclusion, bone scintighraphy should be considered in beta-thalassemic patients with recurrent pain attacks, in order to identify osteoarticular complications and other clinical manifestations of disease.

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Figure 1. A. Blood pool phase shows augmented vascularization at left condylus medialis femoris, consistent with proliferation of connective tissue and bone following postischemia recovery. B. Bone scan at 3 hours p.i. shows increased osteoblastic regenerative metabolism at left condylus medialis femoris. In addition, demonstrates a simmetrical-increased uptake of 99mTc-MDP at the epiphysis of humerus and diaphysis of femur. Similarly, an intense uptake of radiotracer was seen at the skull. These scan findings are typical of hyperactive marrow. Moreover, a significant nephromegaly was documented.

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