Incidentally Diagnosed Osteopoikilosis: A Case Report

Osteopoikilosis is a rare bone dysplasia. Patients are generally asymptomatic and are frequently identified during radiological imaging by coincidence. The epiphysis and metaphysis of the long bones, the phalanges and pelvic bones are the localizations which frequently involved. It appears as oval or round, homogeneous, symmetrical, uniform, radio-opacities. Osteoblastic metastases, mastocytosis, tuberous sclerosis and Paget’s disease should be considered in differential diagnosis. Keywords: Osteopoikilosis, metastases, bone dysplasia

Introduction

Osteopoikilosis, also known as osteopathia condensans disseminata, is a rare bone dysplasia of uncertain etiology and pathogenesis (1). Hereditary or sporadic cases have been reported, and the genetic transmission has been shown to be autosomal dominant (2). It occurs as a result of abnormality in the bone maturation process. The condition is seen at similar levels in both sexes, with an estimated incidence of 1/50,000 (3). Patients are generally asymptomatic and are identified incidentally during radiological examinations performed for different causes (4). Numerous well-defined sclerotic lesions are observed at radiological examination. Sclerotic bone lesions, particularly bone metastasis must be thought in differential diagnosis (5). In this report we describe a case incidentally identified as osteopoikilosis at computerized tomography (CT) performed due to trauma.

Case Report

A 19-year-old male patient with a history of trauma four days previously presented to the emergency department with headache and tenderness in the lower abdomen and right thigh. He had no history of chronic disease. Apart from alanine aminotransferase elevation (52.9 U/L), no pathology were present except for swelling and tenderness in the soft tissue of the right thigh. Cranial and thoracoabdominal CT were performed due to post-traumatic headache and abdominal tenderness. Non-displaced fracture was detected on the left side of the occipital bone. Abdominal CT and pelvis radiography revealed numerous uniform, homogeneous, millimetric and symmetrically distributed hyperdense nodular lesions in the sacrum, acetabulum and bilateral femoral epiphysis and metaphysis in this patient with no identified emergency pathology (Figures 1, 2). Osteopoikilosis was diagnosed on the basis of typical radiological findings in the absence of any pathology in the physical examination and laboratory results. The joints were evaluated as normal at physical examination, and no
radiological pathology was observed in the shoulder or hip joints entering the imaging field. No pathological finding was determined at inspection and skin examination. The patient was followed-up for head trauma and discharged in a healthy condition.

**Discussion**

Osteopoikilosis is a bone dysplasia that can be seen at any age and with a similar incidence in the two sexes. Genetic transmission is thought to be autosomal dominant, and mutations in the *LEMD3* gene have been implicated (2,6). Publications in the literature have reported that osteopoikilosis is associated with dacryocystitis, keloid, syndactyly, scleroderma, endocrine and metabolic diseases and cardiac anomalies (6,7). Comorbid connective tissue lesions known as dermatofibrosis lenticularis have been reported on the skin in some cases. This combination is known as Buschke-Ollendorff syndrome (8). We determined no dermatological pathology during systemic examination in our case. While patients with osteopoikilosis are generally asymptomatic, cases with effusion in the joint spaces and joint pain have been reported. Joint symptoms affect approximately 20% of this patient group (9). Joint pain can be incidental in osteopoikilosis, or increased bone metabolism in this location or irritation of the joint capsule in association with venous stasis may be involved (9). There was no joint dysfunction or joint pain in our case, and those joints included the imaging field were radiologically normal. Typical radiological findings are round or oval, homogeneous, symmetrical, uniform radio-opacities 2-10 mm in diameter. Typically affected locations include the epiphysis and metaphysis of the long bones, the phalanges, the pelvic bones and the carpal and tarsal bones (5). Osteoblastic metastases in particular must be excluded at differential diagnosis. Metastasis must be considered in the presence of primary malignity, bone destruction, periosteal reaction, axial skeleton involvement or impaired uniformity. Rare involvement of the vertebra, costa and cranium in osteopoikilosis, and normal bone scintigraphy is important in differentiating from metastasis. Normal laboratory findings and typical morphological features of opacities can assist in differentiating from mastocytosis, tuberous sclerosis and Paget’s disease (4,7). In conclusion, osteopoikilosis is a frequently asymptomatic benign bone dysplasia. Its clinical significance is that it can be differentiated from other pathologies proceeding with sclerosis. While pain is rare, when symptomatic cases, analgesics such as acetaminophen and non-steroid anti-inflammatories can be used in treatment (10).

**Figure 1.** Millimetric, hyperdense sclerotic foci in the sacrum, acetabulum and neck of the femur on axial computerized tomography sections

**Figure 2.** Radiopaque foci compatible with osteopoikilosis on radiography image
Ethics

Informed Consent: It was taken.

Peer-review: Externally and internally peer-reviewed.

Authorship Contributions


Conflict of Interest: No conflict of interest was declared by the authors.

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References