Rare Bone Disorder

Zeljko Stepanovic
Assistant Professor Medical Sciences, University of Kragujevac, Serbia

Extended Abstract

Area of Interest: Musculoskeletal bone Musculoskeletal system

Procedure: Diagnostic procedure

Imaging Technique: CT-High Resolution

Imaging Technique: MR

Imaging Technique: Nuclear medicine conventional

Special Focus: Tissue characterisation Metastases

Case Type: Clinical Cases

Authors: Zeljko L Stepanovic1, Ana Rankovic2

Patient: 67 years, male

Clinical History:
A 67-year-old man was referred to the orthopaedic department for further evaluation of osteoblastic bone lesions incidentally found on pelvis radiograph. Recent examinations revealed benign prostate hyperplasia. 99mTc bone scan (Fig. 6) showed positive radionuclide uptake, indicating possible osteoblastic metastases. His only complaint was moderate pain in the right knee.

Imaging Findings:
Planeradiography of the pelvis (Fig. 1) revealed the 3rd grade osteoarthritis of hip joints according to Kellgren-Lowrence (KL) criteria associated with multiple, scattered, juxta-articularand metaphyseal, ovoid and longitudinal radiopacities.

Knee radiographs (Fig. 2) revealed KL grade 2 and KL grade 3 osteoarthritis of the left and the right knee. Ovoid sclerotic bone islands adjacent to inner cortices were detected in the metaphyseal regions of the femoral bones.

Shoulder joint radiographs (Fig. 3) detected the same sclerotic bone lesions in peri-articular and metaphyseal regions of the proximal humeral bones, clavicles and scapulae.

Control MDCT (Fig. 4) and MRI (Fig. 5) of the pelvis were performed to differentiate these lesions between osteoblastic metastases and benign bone dysplasias and to rule out prostate proliferation. The same ovoid and longitudinal symmetric sclerotic bone formations were detected in the pelvis and the proximal femurs.

Class imaging findings of osteopoikilosis were associated with normal clinical, laboratory and histology result of benign prostate hyperplasia, excluding osteoblastic bonemeta metastases.

Discussion: Osteopoikilosis represents an autosomal dominant bone dysplasia that mimics osteoblastic bone metastases, and may lead to misdiagnosis and unnecessary treatment. Its incidence is 1 in 50,000; it is seen in up to 6 per 100,000 radiographs. [1] A mutation in gene LEMD3 at position 12q13 is responsible for this benign bone condition. [2] Patients are usually asymptomatic or may complain of nonspecific articular pain in 15-20% of cases. Mostly, these lesions are discovered incidentally on radiographic imaging. [3] Radiographic findings of osteopoikilosis include the
appearance of numerous symmetric, ovoid, sclerotic foci, ranging from 2 to 10 mm, clustered in periarticular regions of tubular bones. The metaphyseal lesions may be longitudinal and located eccentrically, abutting the endosteal cortical surface. The main distribution of lesions in the pelvis is found about the acetabulum and mostly related to the orientation of the trabecular bone. The skull, mandible, clavicles and thoracic cage are spared. [4] Macroscopically, foci of compact bone within the cancellous bone are found. Histologically, the sclerotic nodular or star-like foci correspond to old and inactive condensation of cancellous bone with peripheral scant osteocytes. Osteoblasts and osteoclasts are in the central core of irregular trabeculae. [5] Osteopoikilosis could be associated with skeletal anomalies, such as fibrous changes in Buschke-Ollendorf syndrome. It has been associated with scleroderma, syringometastases, dwarfism, mental retardation, and tuberous sclerosis. [1, 2] Mixed sclerosing bone dysplasia represents another condition, whereas triads of benign bone disorders: osteopoikilosis, osteopathiastriata, and melorheostosis occur concomitantly.

The differential diagnosis includes conditions with radiographic appearance similar to those of osteopoikilosis, such as osteoblastic metastases, osteopathiastriata, mastocytosis, tuberous sclerosis and Paget disease. [4, 5] However, these bone conditions have additional distinctive physical findings. Bone scans are mostly normal in osteopoikilosis, which has been used to differentiate it from metastatic bone disease. [6] There are a few reports of abnormal scintigraphy findings in osteopoikilosis, mimicking osteoblastic bone metastases. [7] Bone scan of a patient with bone metastases usually reveals an increased uptake, with asymmetrical distribution when multiple sites throughout the skeleton are involved. The main problem is its low specificity for osteoblastic activity. Abnormal bone scans do not exclude the diagnosis of osteopoikilosis if the radiographic findings are characteristic of that entity.

Take Home Message:

Benign bone dysplasia that mimics osteoblastic bone metastases.

Usually asymptomatic. Radiographic findings are characteristic and diagnostic. Bone strength is normal.

Abnormal bone scan does not exclude the diagnosis of osteopoikilosis.

Differential Diagnosis List: Osteopoikilosis (osteopathy condensans disseminata), Osteoblastic bone metastases, Melorheostosis, Osteopathiastriata, Tuberous sclerosis, Mastocytosis

Final Diagnosis: Osteopoikilosis (osteopathy condensans disseminata)

References:


Baasanjav et al (2010) Osteopoikilosis and multiple exostoses caused by novel mutations in LEMD3 and EXT1 genes respectively—coincidence within one family. BMC Medical Genetics 11:110 (PMID: 20618940)

