Bone Scintigraphy: Differentiating Benign Cortical Irregularity of the Distal Femur from Malignancy

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Two cases of benign cortical irregularity of the distal femur (BCIDF), which radiologically simulate malignancy, are presented. The use of bone scintigraphy in differentiating this entity from malignancy is described.


Malignant bone tumors typically appear as abnormal foci of intense radionuclide uptake on bone image, whereas benign cortical irregularity of the distal femur (BCIDF), being more diffuse in nature, characteristically has normal or only minimally increased uptake of the tracer. This dramatic difference in scintigraphic appearance should obviate unnecessary amputations and biopsies in patients with BCIDF, which radiologically masquerades as malignancy. Avulsive cortical irregularity of the distal femur (1) was first described in 1951 by Kimmelstiel and Rapp (2); it is also referred to as cortical desmoid, peristomal desmoid (2), subperiosteal desmoid, subperiosteal abrasion, cortical abrasion, medial distal metaphysical femoral irregularity (3), and subperiosteal cortical defect. There is very little in the nuclear medicine literature concerning this topic. Conway et al. found normal bone images in several children with this entity and strongly recommended the use of bone scintigraphy in such cases (4).

CASE REPORTS

Case 1. A 101/2-yr-old white male had a history of left knee pain beginning approximately 2 mo before admission. Physical examination was remarkable for fullness in the popliteal space of the left knee and about the medial femoral condyle in the region of the adductor tubercle. The impression on admission was a mass in the region of the left knee, possibly from osteosarcoma. Plain radiographs revealed an area of cortical irregularity and periosteal reaction in the posterior medial aspect of the distal left femur in the region of the adductor tubercle, with a mild degree of associated soft-tissue fullness and no evidence of effusion (Fig. 1). A single-phase bone image, performed with 10 mCi of Tc-99m MDP and including pinhole views of both knees, was normal (Fig. 2). An arteriogram demonstrated normal vascularity. Since the plain radiographs were suspicious for a malignant tumor of the distal femur, a biopsy of the adductor tubercle was performed. Microscopy revealed normal-appearing bone, cartilage, vascular fibrous tissue, and giant cells suggestive of reactive changes without any evidence of malignancy. Plain radiographs obtained at the 4-mo follow-up demonstrated healing of the biopsy site without any evidence of bone destruction.

Case 2. A 13-yr-old black male presented with a 3- or 4-hr history of pain in the left knee associated with swelling and difficulty in weight-bearing after playing basketball. There was a history of an effusion in this knee 6 mo before admission. Physical examination of the left knee revealed swelling and...
tenderness of the distal thigh in its mid aspect, without evidence of an effusion.

Plain radiographs showed a mild degree of cortical irregularity in the posterior medial aspect of the distal left femur in the region of the adductor tubercle, with a slight degree of periosteal elevation (Fig. 3).

The patient was admitted to the hospital with a tentative diagnosis of a tumor involving the distal aspect of the left femur. A bone image was ordered for diagnostic purposes and in order to locate the lesion and mark its extent, should an amputation be necessary. The single-phase bone image demonstrated a minimal increase in activity diffusely throughout the distal left femur, consistent with the history of trauma and with no intense focus to suggest tumor (Fig. 3).

The patient underwent arthroscopy with stress testing under anesthesia. At surgery, all of his intraarticular structures were normal but he had a massive chylous hematoma. The diagnosis was a nondisplaced Salter I fracture of the distal femoral growth plate. The knee was flexed at 45° and placed in a cylinder cast for 3 wk.

Three weeks later, follow-up radiographs of the left knee continued to show a minimal degree of periosteal elevation in the posterior medial aspect of the distal left femur. The secondary diagnosis was a benign lesion of the posterior medial aspect of the distal left femur, most likely representing benign cortical irregularity.

One month later, the patient was progressing well with physical therapy, and follow-up radiographs showed continued resolution.

**DISCUSSION**

BCIDF occurs in the age range of 3–17 yr, most commonly between 10 and 15 yr (1–3). It is rarely seen in the preschool child, and not at all after epiphyseal closure (3). There is a male-to-female ratio of approximately 3:1 (3,5). It occurs about twice as often in the left femur as in the right (3), and bilaterally in 25–35% of cases (1,5). It is usually asymptomatic, discovered incidentally (1,3,5). In general there is no associated mass, soft-tissue swelling, pain, or loss of soft-tissue planes (3). However, occasionally pain may be present and may be the presenting complaint (5).

Radiographically, the BCIDF may simulate malignancy (1,3,5–9). Kimmelstiel et al. reported a case in which an amputation was performed because BCIDF mimicked a malignant neoplasm (2). Biopsy of the lesion is unnecessary and generally not helpful (1,5,8). Furthermore, it may result in the false-positive diagnosis of malignancy (most commonly osteosarcoma or fibrosarcoma) (1,5,7,8).

BCIDF characteristically occurs in the posterior medial aspect of the distal femur along the medial supracondylar ridge of the linea aspera, just above the adductor tubercle at the insertion of
the adductor magnus (1, 3, 5, 7, 9). It is best demonstrated by obtaining an oblique anteroposterior radiograph with 30° of external rotation (Fig. 1C). It measures approximately 1–3 cm in length, with its long axis parallelizing the long axis of the femur (3). It appears as a lytic area of cortical irregularity with associated periosteal reaction. Sometimes there is reactive bone formation and spiculation, with small cortical fragments being located in the adjacent soft tissue (5). These features may raise suspicion of malignancy. With healing, a sclerotic margin and cortical thickening may develop along with progressive migration of the lesions away from the epiphysis (1, 5).

Benign cortical irregularity has been observed rarely in other locations such as the humerus, fibula, radius, metatarsus, metacarpus, and distal phalanx (1). Fulton et al. (6) described its counterpart in the proximal humerus at the insertion of the pectoralis major muscle as "ringman's shoulder lesion", occurring primarily in gymnasts.

The knee region is a common area for both benign and malignant neoplasms in children. BCIDF is most often confused with an osteosarcoma. Radiologically, both may have cortical irregularity, periosteal reaction, spiculation, and cortical fragments or bony density within the adjacent soft tissues. However, these features are usually more pronounced in osteosarcoma. In addition, osteosarcomas usually have a significant associated soft-tissue component and loss of soft-tissue planes (3). Frequently, however, the two may be indistinguishable radiologically.

Malignant bone tumors characteristically appear scintigraphically as abnormal foci of intense radionuclide uptake. Of the six patients with BCIDF and scintigraphic correlation reported in the literature, five had normal bone images and one patient had a very slight increase in uptake, minimal in intensity (10).

It is readily apparent from a review of the literature that much confusion has existed about the pathogenesis and histologic nature of the lesion (1, 5). Macroscopically, there is periosteal and cortical thickening (1, 2, 3, 9). In addition, fragments of resorbing bone are often found in the adjacent soft tissue (5). Microscopically, there is evidence of fibrous-tissue proliferation and numerous osteoclasts (1, 2, 5). Thus, the pathologic features are those of reactive process rather than a neoplastic one (4). The periosteal reaction, cortical thickening, reactive bone formation, and bony fragments within the soft tissues may easily be confused with a malignant tumor such as an osteosarcoma or fibrosarcoma (1, 3, 5, 8, 9).

Currently the pathogenesis is thought to be related to excessive mechanical stress produced at the insertion of the adductor magnus, resulting in microavulsions, disparity between resorption and formation of reactive bone, hypervascularity, and a fibroblastic response, stimulating osteoclasts and leading in turn to bony resorption, erosion, and remodeling in an area of rapid bone growth (1, 2, 3, 7, 8) (Fig. 1). This has been described as a repetitive cycle of "micro-fracture-resorption-microfracture" (1, 5).

If untreated, BCIDF will follow a benign course and resolve spontaneously, as indicated by the absence of this finding after epiphyseal closure (3). Contrarily, the treatment of choice for osteosarcoma is amputation. This marked difference in therapy emphasizes the importance of making the correct diagnosis. In this regard, it is recommended that bone scintigraphy be performed, especially in those cases where a diagnosis cannot be made from the plain film with a high level of confidence. In addition, the bone image will act as a screening procedure for metastasis. Biopsy is unnecessary and may be misleading.

REFERENCES