Bone tumor mimickers: A pictorial essay

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Abstract

Focal lesions in bone are very common and many of these lesions are not bone tumors. These bone tumor mimickers can include numerous normal anatomic variants and non-neoplastic processes. Many of these tumor mimickers can be left alone, while others can be due to a significant disease process. It is important for the radiologist and clinician to be aware of these bone tumor mimickers and understand the characteristic features which allow discrimination between them and true neoplasms in order to avoid unnecessary additional workup. Knowing which lesions to leave alone or which ones require workup can prevent misdiagnosis and reduce patient anxiety.

Key words: Bone tumors; MRI artifacts; osteomyelitis; traumatic lesions; tumor mimickers

Introduction

Focal lesions in bone are very common and are frequently encountered in routine imaging studies. While many lesions are true neoplasms, a number of these abnormalities in bone are not tumors. These lesions can include normal variants, congenital abnormalities, traumatic/iatrogenic lesions, metabolic and arthritic changes, infection, and artifacts related to imaging technique. It is important for the radiologist and clinician to be aware of this possibility and to identify the characteristic features which allow discrimination between bone tumors and bone tumor mimickers. Subjecting the patient to an inappropriate workup can lead to misdiagnosis, poor management, and anxiety for both the patient and physician. In many instances, these tumor mimickers can be left alone and no treatment is necessary; however, in other cases, they can indicate a significant disease process. Although there are innumerable processes that can lead to focal lesions in bone, we present here a review of commonly encountered bone lesions [Table 1] that can mimic bone tumors and discuss the key imaging and clinical features that can help distinguish these entities from neoplasms. For the purpose of this pictorial essay, we performed a systematic search of the electronic database PubMed to identify relevant studies published in the literature from 1991 to 2014 using the terms “bone tumor mimickers,” “bone tumor mimics,” and “pseudolesions of bone.” Additional targeted searches were performed for the specific disease conditions.

Normal Variants

Red marrow

Erythropoietic or red marrow can be a common cause for concern on magnetic resonance imaging (MRI). This can be particularly problematic if the area of red marrow is mass-like in appearance. Red marrow should be hyperintense to fatty marrow on fat-suppressed T2-weighted (T2W) MRI sequences and hypointense on T1-weighted (T1W) MRI sequences.¹ The key feature is that the low signal intensity on T1W MRI sequences should be higher than that of skeletal muscle or the intervertebral discs.² In-phase and out-of-phase T1W MRI images can be helpful in equivocal cases as red marrow should have some intermixed fatty marrow and, consequently, should lose signal (become darker) on out-of-phase compared to in-phase MRI.³ On the other hand, marrow-replacing tumors, such as many metastases, should replace all the fatty marrow and should not lose signal on out-of-phase T1W imaging [Figure 1]. Thus, when approaching marrow abnormalities on MRI, it is important to have T1W images that include skeletal muscle for comparison and in-phase and out-of-phase T1W images.
to show the presence or absence of fat. Yellow marrow can reconvert to red marrow with physiologic stressors such as anemia.\(^4\) Moreover, red marrow should not extend past the physeal scar into the epiphysis and should not distort normal trabecular pattern.\(^5\)

**Humeral pseudocyst**

A radiolucent area in the humeral head may be seen due to a normal decrease in the trabeculae often associated with an increase in the amount of fat.\(^6\) This radiolucency is seen in the superolateral humeral head and may be misdiagnosed as a chondroblastoma, giant cell tumor, Langerhans cell histiocytosis, or even an osteolytic metastasis on radiographs.\(^7\) The increased fat in this region can be readily seen on MRI and helps make the diagnosis [Figure 2]. On radiographs, this pseudolesion will be seen on an external rotation view of the shoulder and there is usually a sharp line of demarcation inferiorly between the pseudolesion and adjacent marrow, which is due to the line of fusion between the epiphysis in the greater tuberosity and the shaft of the humerus. The remainder of the margin is usually ill-defined.\(^8\)

**Ward's triangle**

A focal area of increased lucency is often seen in the femoral neck at the junction of the compressive and tensile trabeculae [Figure 3]. As with the humeral pseudocyst, this radiolucency can become less apparent if the patient is osteoporotic due to attenuation of the adjacent trabeculae.\(^7,8\)

**Table 1: Common lesions mimicking bone tumors**

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MRI: Magnetic resonance imaging

**Figure 1 (A and B):** Island of red marrow in the sacrum. A 49 year old man with recurrent bloating underwent a MR enterography, which demonstrated an incidental lesion in the sacrum. He was recalled for in-phase and out-of-phase T1W MRI imaging. (A) In-phase T1W MRI image demonstrates the lesion (arrow) is slightly hyperintense to skeletal muscle (B) On the out-of-phase T1W MRI image, there is loss of signal due to the presence of intermixed fatty marrow (arrow)

**Figure 2 (A-C):** Humeral pseudocyst. A 47 year old female with left shoulder pain. A round radiolucency in the greater tuberosity (arrow) on the external rotation shoulder radiograph (A) corresponds to normal fatty marrow (arrow) which is hyperintense on the (B) T1W and hypointense on the (C) T2W fat-saturated coronal MRI images

**Figure 3 (A and B):** Ward's triangle. A 67 year old female with left hip pain. (A) Anteroposterior (AP) radiograph of the hip demonstrates a triangular radiolucency (arrows) in the femoral neck (B) The coronal CT image shows a paucity of trabecular lines in the femoral neck (arrows)
Calcaneal pseudocyst

In a similar pattern to Ward’s triangle, a radiolucency in the anterior aspect of the calcaneus can be outlined by the major trabecular groups [Figure 4]. Although this is a normal appearance, a number of pathologic lesions can occur in this location and form a radiolucent region on radiographs. These tumors include simple bone cyst, giant cell tumor, chondroblastoma, and intraosseous lipoma. Intraosseous lipomas often develop central necrosis which can cause a central dystrophic calcification and tends to have well-defined sclerotic margins.

Congenital and Developmental Abnormalities

Dorsal defect of the patella

A subarticular abnormality in the superolateral aspect of the patella is known as the dorsal defect of the patella. It is seen in approximately 1% of the population and can be bilateral. The dorsal patellar defect can appear as a 1-2 cm rounded area of lucency in the same location as a bipartite patella and is believed to be due to incomplete fusion of the patellar ossification centers [Figure 5]. Another potential etiology is that it is due to traction at the insertion of vastus lateralis. Occasionally, this lesion may be symptomatic.

Synovial herniation pit in the proximal femur

A well-defined round or oval radiolucency in the proximal superior femoral neck is known as a synovial herniation pit or Pitt’s pit. It is thought to represent herniation of the synovium into cortical defects created by abrasion of the hip joint capsule against the femoral neck, although it may represent a normal variant. Typically these lesions are less than 1 cm in size, but can grow up to 2-3 cm and may be lobulated. Although these lesions have been considered asymptomatic, an association with femoracetabular impingement has been described.

Avulsive cortical irregularity of the posterior femur

An avulsive cortical irregularity of the posterior femur, known as a cortical desmoid, appears as an irregular focal...
radiolucent lesion along the posteromedial aspect of the distal femur in children [Figure 7]. Differential diagnosis for this appearance includes osteomyelitis and surface osteosarcoma, especially if the lesion has an aggressive appearance. It has been proposed that this lesion may be caused by traction due to the medial head of gastrocnemius or adductor magnus. This lesion should not be seen in skeletally mature individuals.

**Supracondylar process of the humerus**
A supracondylar process in the humerus is a bony spur that arises from the anteromedial aspect of the humerus in about 1-3% of the population. It is usually an incidental finding and should not be mistaken for an osteochondroma or surface osteosarcoma. Osteochondromas point away from the joint, whereas the supracondylar process points toward the elbow joint [Figure 8]. Occasionally, a ligament extends from the supracondylar process to the medial epicondyte (the ligament of Struthers), forming a tunnel that can entrap the median nerve and even the brachial artery, leading to symptoms.

**Soleal line**
The soleal line is a bony “tug lesion” that can form on the tibia at the attachment of the soleus and mimics periostitis from a tumor, infection, or stress fracture [Figure 9]. The soleal line begins 1-2 cm below the fibular facet and may present as a line or a ridge. This can arise from the tibial head of the soleus, with cortical thickening extending lateral to medial along the posterior upper one-third of the tibia. Similar bony changes can be seen at the fibular attachment of the soleus.

**Trauma and Iatrogenic Lesions**

**Subperiosteal hematoma**
The periosteum is a highly vascular thick fibrous membrane that is closely adherent to the bone. Injury to the periosteum can result in a subperiosteal hematoma, which lifts the periosteum off the bone and can resemble a focal mass such as a parosteal osteosarcoma or osteochondroma [Figure 10]. Most often, they resolve without treatment; however, they may ossify and persist. On imaging, these lesions have a non-aggressive appearance and are centered in the subperiosteum. If they ossify, they can contain fatty marrow.

**Stress fracture**
Stress fractures may be related to fatigue, when excessive repetitive force is applied to a normal bone, or insufficiency, when normal stress is applied to abnormal bone such as in osteoporosis or Paget’s disease. Common sites for stress fractures include the metatarsals, tarsals, and tibia. Initially, stress fractures may not be visible on radiographs and are better detected on technetium-99 m pyrophosphate bone scintigraphy (bone scan) or MRI [Figure 11]. With time, periosteal reaction and cortical resorption may be seen. A fracture line may be visible on radiographs, but could be better seen on computed tomography (CT). The fracture line is usually perpendicular to the cortex, and vertically oriented fractures can be difficult to detect. Radiographic features of stress fracture in the tibia can resemble a soleal line or osteoid osteoma, but can be differentiated from one another on CT [Figure 12]. Moreover, if the periosteal reaction appears aggressive, it can mimic infection or an aggressive tumor. The presence of a fracture line, lack of a soft tissue mass, and evidence of healing on follow-up studies should help distinguish stress fractures from other entities.

**Myositis ossificans**
Myositis ossificans is heterotopic ossification that occurs in muscle usually following trauma, although the patient may be unable to recall the precipitating trauma. This commonly occurs in the upper and lower extremities, usually in the lateral muscles. Patients may be asymptomatic or present with pain, swelling, or an
elevated erythrocyte sedimentation rate (ESR). Ossification develops 3-8 weeks after onset, beginning peripherally and progressing centrally. Initially, myositis ossificans forms faint irregular densities; but with time, a rim of mature lamellar bone and central osteoid matrix can develop [Figure 13]. The MRI appearance is variable depending on the stage of development, and earlier on, can mimic a sarcoma as there may be enhancement following contrast administration. Differentiation from an osteochondroma or osteosarcoma may also be difficult if the area of ossification is adherent to the adjacent bone. CT can be helpful in demonstrating a plane of soft tissue between the mass and the bony cortex. Myositis ossificans may be difficult to distinguish from an osteosarcoma even on biopsy specimens.

**Biceps tenodesis**

In biceps tenodesis, the intra-articular portion of the long head of the biceps tendon is cut and the proximal portion of the tendon is reattached to the proximal humeral diaphysis. The site of attachment can mimic a radiolucent lesion with a sclerotic border [Figure 14]. This classic location along the proximal humerus should raise suspicion for this tumor mimicker, which can be confirmed by reviewing patients’ surgical notes.

**Bone marrow biopsy and bone graft donor sites**

Bone marrow aspiration and biopsy for hematological diagnosis is most often obtained from the iliac bone via a posterior approach. If the biopsy has been recently performed, there may be marrow edema or cystic changes in the region, which can be mistaken for a focal lesion [Figure 15]. Similarly, bone graft donor sites can demonstrate edema in the early post-procedure period. In both cases, review of the patient’s clinical history is essential to confirm that a procedure has been performed.

**Particle disease**

Particle disease can present as areas of radiolucency surrounding the hardware components, usually following arthroplasty. However, unlike mechanical loosening, the lucent areas seen with particle disease typically do not follow the outline of the prosthesis [Figure 16]. The arthroplasty components can incite a macrophage-mediated granulomatous response, which then stimulates osteoclast activity. Particle disease can mimic osteolytic tumors or infection; however, particle disease can be distinguished by the presence of hardware and the fact that abnormal lucencies are seen on both sides of a joint.
Radiation changes
Initially, radiotherapy causes vascular congestion, edema, and decreased cellularity in the bone marrow. This will cause decreased signal on T1W sequences and increased signal on T2W sequences [Figure 17]. With time, the bone marrow will be replaced with fat and occasionally with fibrosis, with high signal on T1W and intermediate signal on T2W sequences. There can be a clear line of demarcation along the borders of the radiation field. Irradiated bone can be at increased risk for insufficiency fractures, osteonecrosis, and radiation-induced sarcomas.
Metabolic Disease and Arthritic Changes

Brown tumor of hyperparathyroidism
Longstanding untreated hyperparathyroidism can result in osteolytic lesions known as brown tumors (osteoclastomas) [Figure 18]. They can be seen in either primary or secondary hyperparathyroidism and are seen in 5% of patients with hyperparathyroidism. However, the incidence has decreased with improved early diagnosis of the disease. The typical appearance of a brown tumor is a well-defined osteolytic lesion, which may have septations, be expansile, and can sometimes have aggressive features. Common sites include the long bones, ribs, pelvis, and facial bones. The lesions improve with treatment, often becoming sclerotic. If lesions fail to improve in appearance with treatment, an alternative diagnosis should be considered.

Melorheostosis
Melorheostosis is a benign bone dysplasia characterized by sclerotic bone lesions, often described as “dripping candle wax.” Melorheostosis is not a hereditary disorder and is often asymptomatic; however, when symptoms do...
Osteonecrosis
Ischemic necrosis of the bone and marrow can be due to a variety of causes including trauma, steroids, hemoglobinopathies, alcoholism, radiotherapy, and chemotherapy. When osteonecrosis involves the epiphysis (avascular necrosis), it can lead to subchondral bony collapse and osteoarthritis. Initially, osteonecrosis may be occult on radiography; but over time, it can manifest as a central radiolucency with a sclerotic margin. It may mimic enchondromas, but lacks central calcifications. MRI is sensitive for detection of bone infarcts. Initially, these areas appear as non-specific regions of marrow edema; but with time, the characteristic features of an outer band of low signal associated with an inner band of high signal on non-fat-saturated T2W images (double line sign) can develop.

Calcific tendinitis (resorptive phase)
Calcific tendinitis is a common cause of joint pain and stiffness, and is caused by the deposition of calcium hydroxyapatite crystals in the tendons. The tendons of the rotator cuff and around the hip are most commonly involved; however, it can involve any tendon. During the resorptive phase, calcific tendinitis can mimic an aggressive process such as infection or neoplasm. Calcific tendinitis can be associated with erosions of the adjoining bone, mimicking a destructive bone lesion. This aggressive pattern is common along the posterior proximal femoral diaphysis. The process is typically self-limiting, but needle barbotage and steroid injection can provide symptomatic relief.

Subchondral cyst (geode)
In osteoarthritis, defects in the overlying cartilage can allow synovium and joint fluid to enter the subchondral bone causing subchondral cysts (geodes). They are typically small, about the articular surface, and have a sclerotic margin. However, they can be large, but may extend down the shaft of a tubular bone mimicking a neoplasm. CT can be helpful in demonstrating the sclerotic margin. On MRI, the lesion behaves like a cyst and is typically isointense to muscle on T1W images and hyperintense on T2W images. High T1 signal may occur in lesions that contain proteinaceous material, and internal enhancement may be seen if the lesions contain fibrous material.

Infection
Osteomyelitis/Brodie’s abscess
In acute osteomyelitis, the radiographic findings include areas of aggressive periostitis, cortical destruction, endosteal scalloping, and intracortical tunneling. There may be soft tissue swelling or gas formation. However, the radiographic findings may not be present for 1-2 weeks. MRI and technetium-99 m pyrophosphate bone scintigraphy (bone scintigraphy) are more sensitive in the detection of early osteomyelitis. Subacute or chronic osteomyelitis can cause an intraosseous abscess (Brodie’s abscess), commonly in the metaphysis of tubular bones. On radiographs, these lesions appear as single or multiple round or oval areas of medullary destruction with sclerotic margins.
multilobulated radiolucent lesions with surrounding sclerosis that fades toward the periphery. These lesions can mimic an osteoid osteoma or osteosarcoma.\footnote{Lesions without significant sclerosis can mimic Langerhans cell histiocytosis, chondroblastoma, giant cell tumor, and Ewing’s sarcoma.} CT can be helpful to delineate a sinus tract extending away from the central abscess, excluding other lesions.\footnote{Systemic signs of infection can be helpful; however, several of the lesions listed in the differential can also present with fever, pain, and other clinical signs of infection. Bone biopsy is often necessary for diagnosis and to identify an organism to guide appropriate antibiotic therapy.} Tuberculosis infection of bone deserves special mention and has been called “the great mimicker.”\footnote{Tuberculosis infection of bone deserves special mention and has been called “the great mimicker.”} Most prevalent in underdeveloped countries, tuberculous osteomyelitis differs from pyogenic osteomyelitis as fever and pain can be absent and the symptoms are more insidious in onset.\footnote{Nearly any bone can be affected [Figure 23] and it is primarily caused by hematogenous spread from other sites, most commonly lung.} Bony destruction, loss of normal T1 marrow signal, marrow enhancement, and adjacent...
abscess or septic arthritis can occur. Spinal involvement by tuberculosis is not uncommon and can differ from bacterial spinal infection in that the disc spaces are preserved until late in the disease due to the lack of proteolytic destructive enzymes by Mycobacterium tuberculosis.\cite{47,48} Finally, due to the hematogenous nature of spread, multifocal lesions can occur in the spine and appendicular skeleton, mimicking malignancy.\cite{47,48}

**Technical Artifacts**

**Humeral head - internal rotation view**
On internal rotation radiographs of the shoulder, a pseudolesion with a sclerotic border and radiolucent center can appear in the humeral head [Figure 24]. A sharp sclerotic border is seen at the humeral neck as the diameter of the bone changes abruptly. The pseudolesion should not be seen on the external rotation or other views and should not be mistaken for an osteolytic lesion.

**Radial tuberosity - lateral view**
The radial tuberosity is a normal anatomic structure in the proximal radius; however, on lateral projections, it is imaged en face and can appear as an ovoid radiolucent lesion [Figure 25]. On other projections, the tuberosity becomes clear and the artifactual radioluency disappears. The bony protuberance of an osteochondroma can mimic a radiolucent lesion when seen en face as well. To avoid this pitfall, it is important to review additional projections.

**Wrap-around/aliasing in MRI**
The field of view (FOV) in MRI refers to the anatomic region being imaged. Deciding on an appropriate FOV depends on the size of the structure being imaged and taking into account the trade-offs between spatial resolution and the signal-to-noise ratio. If a FOV is chosen which is smaller than the anatomy being imaged, wrap-around or aliasing artifacts can occur.\cite{49} This can lead to image data that are outside the FOV being “wrapped around” and artifactually included within the image [Figure 26]. This can be corrected by using a large enough FOV in the phase-encoding direction to include the entire body part or by using phase oversampling techniques during imaging.

**Pulsation artifact on MRI**
Pulsation of vascular structures can cause “ghosting” on MRI.\cite{49} This can mimic bone lesions as artificial image data from the vessels are superimposed onto bone [Figure 27]. Repeating the imaging sequence after swapping the phase- and frequency-encoding directions can help to determine whether or not the lesion is real. To reduce pulsation artifact, one can place a saturation band over the vessels or not align the vessel and target lesion in the same phase-encoding direction.\cite{50}

**External objects**
External objects lying on a patient’s skin can mimic bone...
lesions [Figure 28]. This commonly occurs in the acute trauma setting when urgent imaging is required and the technique may be suboptimal.

Conclusion
Numerous normal anatomic variants and non-neoplastic lesions can have an imaging appearance, which raises concern for a bone tumor. Awareness of these lesions and an understanding of their discriminating features are essential to avoid unnecessary additional imaging and procedures. Knowing which lesions to leave alone or which ones require workup can prevent misdiagnosis and reduce patient anxiety.

References

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