

PICTORIAL ESSAY

Sacral Tumours on MRI: A Pictorial Essay

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Tumours of the sacrum can be primary or secondary. Since the sacrum is rich in haematopoietic bone marrow, bone metastases are the most frequent aetiologies. However, tumours can arise from all components of the sacrum and primary bone tumours should be considered in case of a solitary lesion and absence of oncologic history. As the clinical signs are usually non-specific, magnetic resonance imaging has become an indispensable tool in narrowing the differential diagnosis and determining the therapeutic approach. This pictorial essay illustrates specific features of the most common sacral tumours on magnetic resonance (MR) imaging.

Keywords: Sacrum; Bone tumour; MRI; musculoskeletal imaging

Introduction

Metastases are the most common tumours in the sacrum. However, all components of the sacrum can give rise to benign or malignant tumours and some primary bone tumours present a particular predilection for the sacrum, especially chordoma and giant cell tumour of bone [1, 2]. Magnetic resonance (MR) imaging helps narrowing the differential diagnosis and plays an important role in further management of the patient [2, 3]. This pictorial review highlights the specific imaging features of sacral tumours on MRI.

Primary Tumours

Approximately 5% of the sacral tumours are primary tumours [4]. A detailed overview of the distribution of these tumours is given in **Figure 1** [5].

Chordoma

Chordomas are the most common primary sacral tumours, accounting for almost 40% [2, 3]. The highest incidence is noted in the fifth and sixth decades, with a male-to-female ratio of 2:1. Derived from notochord remnants, the malignant tumour has a predilection for sacral locations and often develops midline or paramedian, with posterior extension and soft tissue invasion [6]. On MRI, the lesions appear as a lobulated mass with heterogeneous hypo- to isointense signal on T1-weighted images, hyperintense signal on T2-weighted sequences (**Figure 2**). Moderate enhancement is observed after gadolinium injection. Hyperintense foci on T1-weighted images indicates either haemorrhage or proteinaceous content, while hypointense signal on T2-weighted images should be interpreted as old haemorrhage or septations [1, 6, 7].



Figure 1: Prevalence of primary sacral bone tumours.

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Giant cell tumour

Giant cell tumour of bone, also known as osteoclastoma, is the second most common primary tumour in the sacrum [5]. Patients aged 15–50 are most frequently affected, and women are twice as likely to develop the tumour compared to men [3, 8]. Giant cell tumours are usually located on both sides of the midline and extension across the sacroiliac joint is frequent (**Figure 3**). The solid components show a low to intermediate signal on T1-weighted images and usually enhance intensely after administration of gadolinium. The heterogeneous signal on T2-weighted sequences is variable and non-specific. Cystic changes and fluid-fluid levels are occasionally observed [8].

Haemangioma

Although vertebral haemangiomas occur in more than 11% of the population, sacral involvement is rare. The tumour is most prevalent during the fifth decade, with a male to female ratio of 1:1.5. Typical haemangiomas are

fat predominant, creating a hyperintense well-defined signal on T1- and T2-weighted images. Due to the presence of vascular elements the signal remains high on fluid-sensitive sequences. Linear areas of lower signal correspond to thickened vertical trabeculae and are referred to as the polka dot sign (**Figure 4**). Atypical haemangiomas are composed of less fatty and more vascular components creating a hypointense to isointense signal on T1-weighted images with variable enhancement and heterogeneous hyperintense signal on both T2-weighted images and STIR-sequences. As a result the polka dot sign is more difficult to distinguish. Maintained vertebral height, a sharp margin, intact bone cortex and enlarged paraspinal vessels are additional findings that favour the diagnosis of haemangioma [9].

Enchondroma/Chondrosarcoma

Enchondromas originate from the medullary cavity and are particularly common in long bones. Sacral location is rare. They are most often diagnosed during the fourth



Figure 2: A 79-year-old male with sacral chordoma. **(a)** Sagittal T1-weighted MR image shows a hypointense mass arising from the lower sacrum with a large soft tissue component. **(b)** Sagittal T2-weighted MR image shows a lobulated mass with a heterogeneous hyperintense signal, invading the bone and soft tissues.



Figure 3: A 40-year-old female with giant cell tumour of the sacrum. **(a)** Axial T1-weighted MR image shows a homogeneous hypointense mass located on both sides of the midline with extension into the presacral soft tissues. **(b)** Axial fat saturated T2-weighted MR image shows heterogeneous high signal intensity. **(c)** Axial post-contrast T1-weighted MR image shows moderate enhancement.

and fifth decade of life with an equal gender distribution [10]. On MRI, the lobules of firm matured cartilage create a hypointense to isointense signal on T1-weighted sequences and a hyperintense signal on T2-weighted sequences. Regions of calcification and osseous content translate as a low signal on T2-weighted images with either high, intermediate or low signal on T1-weighted

images. Enhancement along the fibrovascular septa surrounding the cartilaginous lobules may be observed after administration of gadolinium [11] (**Figure 5**). Bone or soft tissue oedema are rare, as well as soft tissue masses, cortical destruction or solid enhancement. When present, these findings are highly suggestive of chondrosarcoma and a biopsy should be performed [12].



Figure 4: A 65-year-old female with sacral hemangioma. **(a)** Sagittal T1-weighted MR image and **(b)** sagittal fat saturated T2-weighted MR image shows a S1 sacral hemangioma with an atypical extra-osseous component extending into the spinal canal (asterix). Note the maintained vertebral height and cortical integrity (arrow). **(c)** Axial T2-weighted MR image shows hyperintense signal with multiple punctuate hypointense areas (polka-dot sign) (arrow). **(d)** Axial post-contrast T1-weighted MR image shows heterogeneous enhancement of the tumour.



Figure 5: A 51-year-old male with small sacral enchondroma (arrow). **(a)** Coronal T1-weighted MR image shows a hypointense mass in the body of the sacrum. **(b)** Coronal fat-saturated T2-weighted MR image shows a lobulated hyperintense mass. Note absence of bone marrow oedema. **(c)** Coronal fat-saturated post-contrast T1-weighted MR image shows septal enhancement.

Aneurysmal bone cyst

Aneurysmal bone cysts are primarily seen in children and adolescents with a slight female predominance (56%) [2]. The multiseptated lesion has a heterogeneous appearance on T1- and T2-weighted sequences with a low-intensity rim in the periphery, indicating a thin shell of bone. The typical fluid-fluid levels are diagnostic [13] (**Figure 6**). As there are usually no solid structures within the cysts, only the septations enhance on T1Gd-sequences [2]. When solid



Figure 6: A 43-year-old female with aneurysmal bone cyst. Axial fat saturated T2-weighted MR image shows a lobulated cyst with multiple fluid-fluid levels (arrow). Remark the sedimentation of blood cells caused by slow flow of the blood in the cystic spaces of the tumour.

components are present, the MRI appearance is very similar to that of a giant cell tumour, albeit with a heterogeneous signal intensity on both T1- and T2-weighted images.

Ewing sarcoma

Ewing sarcoma is a high-grade malignancy composed of uniform small round blue cells. It occurs most frequently during the teenage years, with a male predominance of 62% [1]. The tumour often presents as a destructive osteolytic lesion with a soft tissue component. The lesion has a homogeneous hypo- to isointense signal on T1-weighted images and isointense signal on T2-weighted images. The extraosseous component is typically larger than the intraosseous lesions and invasion into the paraspinal area and spinal cord is common (**Figure 7**). After the administration of gadolinium mild to moderate enhancement is observed [1, 14].

Metastases

Lung, breast, prostate, kidney, head and neck, skin (melanoma) and gastro-intestinal cancers are the most common tumours that can produce sacral metastases. Most metastases are osteolytic except for prostate cancer, where metastases are mainly osteoblastic [1]. Osteolytic lesions have a hypointense signal on T1-weighted sequences and an iso- to hyperintense signal on T2-weighted sequences compared to normal bone marrow (**Figure 8**). On STIR sequences the signal is usually hyperintense. Administration of gadolinium most often



Figure 7: A 14-year-old male with sacral Ewing sarcoma. (a) Axial T1-weighted MR image shows a hypointense mass arising from left side of the sacral body. (b) Axial T2-weighted MR image shows a large soft-tissue mass, invading into the spinal canal (arrow) and presacral area. Note the absence of hyperintense cerebrospinal fluid signal. (c) Axial fat saturated post-contrast T1-weighted MR image shows enhancement of the tumour and the intraspinal component.



Figure 8: A 13-year-old female with hemangiosarcoma and sacral metastasis. **(a)** Axial T1-weighted MR-image shows multiple hypointense lesions in sacral and iliac bone. **(b)** Axial fat-saturated post-contrast T1-weighted MR image shows strong enhancement of the lesions with central necrosis in the largest lesions (arrow).

leads to intense uptake although moderate, heterogeneous and low uptake are not uncommon. A peripheral hyperintense rim on T2-weighted images is a rather rare but highly specific finding. Cortical disruption is seen in 57% of all osteolytic metastases. In osteosclerotic metastases, the sclerotic areas appear hypointense on all sequences [15, 16].

Conclusions

The differential diagnosis of sacral tumours is extensive, and although metastases are the most common lesions, a broad spectrum of primary bone tumours can arise from sacral components. MR imaging allows us to detect bone marrow invasion, and changes in morphology and intensity will help to determine the most probable diagnosis. In addition, imaging has an important role in the staging of the tumour and further follow-up.

Competing Interests

The authors have no competing interests to declare.

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