IMAGE OF THE MONTH



A rare case of polyostotic fibrous dysplasia detected on ¹⁸F-rhPSMA-7 PET/CT

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The prostate-specific membrane antigen (PSMA) has been proven to show high expression in prostate cancer cells [1]. The high binding affinity and internalization of PSMA radioligands makes it an excellent molecular target for theranostics of prostate cancer [1]. Positron emission tomography/computed tomography (PET/CT) using novel PSMA-targeting probes is increasingly used in recurrent and metastastic prostate cancer [2]. In addition, it is increasingly being used as an imaging modality for initial staging [3]. ¹⁸FrhPSMA-7 is a new theranostic PSMA-targeting agent which allows radiolabeling with ¹⁸F and radiometals and is associated with minimal renal excretion [4]. However, physiologic and other pathologic forms of tracer uptake have to be considered carefully as potential pitfalls to image interpretation, such as ganglia, healing fracture, adrenal adenoma, primary lung cancer, and metastatic renal cell carcinoma [5].

Fibrous dysplasia is an uncommon skeletal disorder, accounting for 5 to 7% of all benign bone tumors [6]. Around 75% of cases have a monostotic form with only one bone involved, commonly the craniofacial bones, but the ribs, femur, and tibia may also be involved [7, 8]. PSMA-ligand uptake has been reported in monostotic (rib) fibrous dysplasia using ⁶⁸Ga-PSMA-PET/CT, and this has to be differentiated from prostate cancer bone metastases [9]. Here, we present a

case of uncommon polyostotic fibrous dysplasia identified on PSMA PET/CT imaging.

An 80-year-old man with histologically confirmed prostate cancer (Gleason score 7a, iPSA 4.26 ng/mL) by transurethral resection of the prostate (TUR-P) performed in February 2018 underwent ¹⁸F-rhPSMA-7 PET/CT imaging for staging in October 2018. No focal uptake in the prostate bed and no evidence of pelvic lymph node metastases was seen. However, intense tracer uptake was observed in the right parietal skull, in the left 3rd, 8th, and 9th ribs, and in the left ilium (red arrows in a MIP and d + e PET). The corresponding bone scan performed 2 weeks earlier had shown hypermetabolic lesions in the same areas (b bone scan). Finally, typical CT morphologic findings, including expansive changes with ground-glass appearance, cortical erosion, diffuse sclerosis, and well-circumscribed margins could be observed and, therefore, clearly confirmed radiologically polyostotic fibrous dysplasia (c CT) [10].

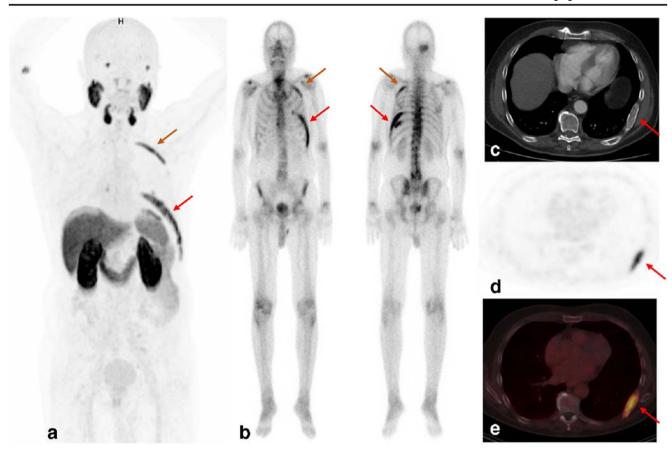
This case indicates that a thorough review of the CT dataset of a hybrid PSMA-ligand PET/CT is not only useful to correctly identify solitary or isolated lesions but also mandatory to avoid a potential pitfall and correctly differentiate extensive polyostotic fibrous dysplasia from metastatic bone lesions [11] when PSMA-ligand uptake in the bone is seen.

This article is part of the Topical Collection on Image of the month

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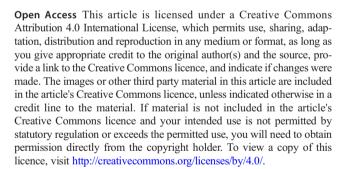
Compliance with ethical standards

Conflict of interest Matthias Eiber is named as an inventor on a patent application for rhPSMA. Matthias Eiber is a consultant for Blue Earth Diagnostics. No other potential conflict of interest relevant to this article was reported.

Ethics approval All procedures performed in studies involving patients were in accordance with the ethical standards of the institutional research committee and with the Helsinki declaration.

Consent to participate During our research, informed consent for each patient was obtained as patient privacy rights and they were always observed. Written informed consent was obtained from the patient.

Consent for publication Written informed consent for publication of their clinical details and/or clinical images was obtained from the patient.



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