

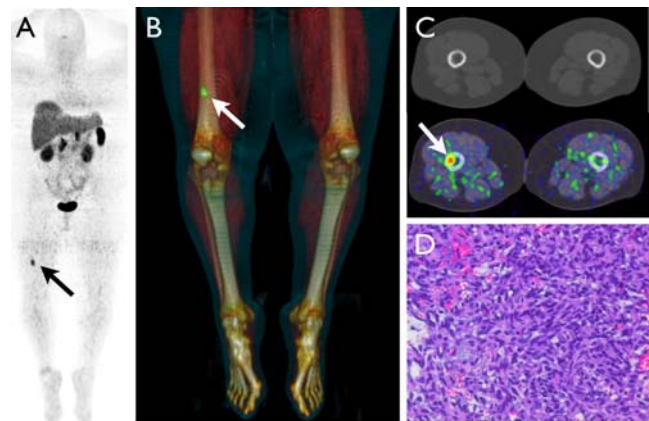
## <sup>68</sup>Ga-DOTANOC PET/CT for the detection of a mesenchymal tumor causing oncogenic osteomalacia

Christian von Falck · Thomas Rodt ·  
Herbert Rosenthal · Florian Länger ·  
Thomas Goesling · Wolfram H. Knapp ·  
Michael Galanski

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A 54-year-old female patient had presented with clinical features of hyperphosphaturia, hypophosphatemia and osteomalacia. These findings were suggestive of oncogenic osteomalacia, a rare paraneoplastic disorder that is usually associated with a phosphaturic mesenchymal tumor [1]. Conventional morphologic imaging including whole-body computed tomography (CT) failed to localise the primary tumor. The patient underwent additional positron emission tomography (PET)/CT using <sup>68</sup>Ga-DOTANOC, a highly sensitive and specific tracer for imaging of somatostatin receptor overexpression, which has recently proven potential in oncogenic osteomalacia [2, 3].

Abnormal focal tracer uptake was seen in the right distal femur (A). Using image fusion and three-dimensional volume-rendering techniques, the localisation of the suspected primary tumor was clearly visualised (B). Notably, no morphologic correlative was observed in the corresponding low-dose CT (C). Based on the PET/CT findings, the patient underwent segmental resection and



compound osteosynthesis of the distal femur. The hematoxylin and eosin-stained section (D) demonstrated randomly organised spindle cells with slight cellular and nuclear atypia and a sparse intercellular matrix. Immunohistochemistry was negative for myogenic, neural, vascular and epithelial markers. These histopathologic findings were consistent with the diagnosis of a benign phosphaturic, mesenchymal tumor.

C. von Falck (✉) · T. Rodt · H. Rosenthal · M. Galanski  
Department of Diagnostic Radiology, Hannover Medical School,  
Carl-Neuberg-Strasse 1,  
30625 Hannover, Germany  
e-mail: c.v.falck@gmx.de

F. Länger  
Institute of Pathology, Hannover Medical School,  
Hannover, Germany

T. Goesling  
Department of Trauma Surgery, Hannover Medical School,  
Hannover, Germany

W. H. Knapp  
Department of Nuclear Medicine, Hannover Medical School,  
Hannover, Germany

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