



The PSMA-11-derived hybrid molecule PSMA-914 specifically identifies prostate cancer by preoperative PET/CT and intraoperative fluorescence imaging

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Received: 6 November 2020 / Accepted: 27 December 2020 / Published online: 23 January 2021
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Resection of tumor tissue represents one of the standard curative treatment options for the clinical management of prostate cancer [1]. However, intraoperative localization and precise delineation of malignant tissue from surrounding healthy structures still remain challenging [2]. The development of PSMA-targeting hybrid molecules enabling the pre- and intraoperative detection of tumor tissue supported by both radioactivity (e.g., using DROP-IN technology) and fluorescence might help to overcome these limitations [3–5].

Here, we report for the first time preoperative PET/CT imaging and subsequent fluorescence-guided surgery aided by a PSMA-11-derived peptidomimetic PSMA-targeting hybrid molecule [6, 7].

A 71-year-old patient with high-risk prostate carcinoma (Gleason score 9 (4 + 5), initial PSA level 7 ng/ml) underwent preoperative PET/CT imaging with ⁶⁸Ga-Glu-urea-Lys-(HE)₃-HBED-CC-IRDye800CW (⁶⁸Ga-PSMA-914) 1 h

after intravenous tracer injection, revealing strong tracer uptake of the primary tumor located in the left prostate lobe (A, arrow).

DaVinci-assisted radical prostatectomy was subsequently performed the day after PET imaging under fluorescence guidance (PSMA-914 administration: intravenously 1 h prior to surgery). The primary tumor was clearly visualized by the specific fluorescence signal (green) and delineated from surrounding tissue (B). Consecutive ex situ fluorescence detection after radical prostatectomy (C) verified the intraoperative findings of tumor-specific hybrid molecule enrichment (D) resulting in high contrast to surrounding healthy structures (E).

Our initial experience with the novel PSMA-11-derived hybrid molecule PSMA-914 demonstrates its potential to precisely detect PSMA-expressing lesions pre- and intraoperatively.

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

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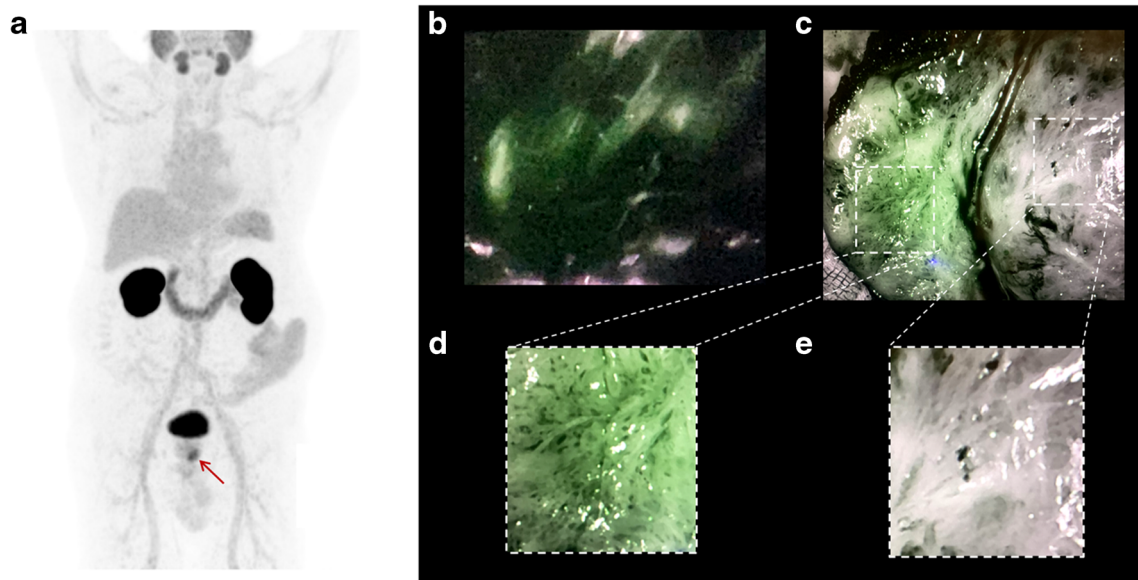
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Funding Open Access funding enabled and organized by Projekt DEAL. We gratefully acknowledge support by the VIP+ grant VP00130, Federal Ministry of Education & Research (BMBF), Germany.

Compliance with ethical standards

Conflict of interest ACE, KK, and ME hold patent rights on dual-labeled PSMA inhibitors. All other authors declare no competing interests.

Ethics approval All procedures performed in studies involving patients were in accordance with the ethical standards of the institutional research and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. The local ethics committee approved the corresponding data analysis (N°562/15).

Consent to participate/for publication Written informed consent for participation and publication was obtained from the patient.

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