



Image-based dosimetry for ^{225}Ac -PSMA-I&T therapy using quantitative SPECT

A. Gosewisch¹ · M. Schleske¹ · F. J. Gildehaus¹ · I. Berg¹ · L. Kaiser¹ · J. Brosch¹ · P. Bartenstein¹ · A. Todica¹ · H. Ilhan¹ · G. Böning¹

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Targeted alpha therapy (TAT) using ^{225}Ac -PSMA ligands is a promising therapy option for advanced metastatic castration-resistant prostate cancer (mCRPC) [1]. The ^{225}Ac decay chain shows a noticeable gamma emission (440 keV, 25.9%; 218 keV, 11.4%). However, recommended low therapeutic activities (4–8 MBq) limit the clinical applicability of SPECT [2], although initial attempts for ^{225}Ac imaging exist [3, 4]. Particularly quantitative SPECT is a vital tool to assess dosimetry and therapy response. While the 218-keV-peak is characterized by a lower branching ratio and a higher scatter fraction, SPECT imaging of high-energy gammas such as 440 keV causes a complex detector point spread function (PSF) [5].

In this study, we would like to demonstrate the general feasibility of image-based dosimetry for ^{225}Ac radionuclide therapy using quantitative ^{225}Ac SPECT. For a mCRPC patient (65 years), imaging of the abdomen was performed 24 h p. i. of 8.1 MBq ^{225}Ac -PSMA-I&T on a Siemens Symbia Intevo T16 SPECT/CT (440 keV (width, 20%), lower adjacent window (width, 10%),

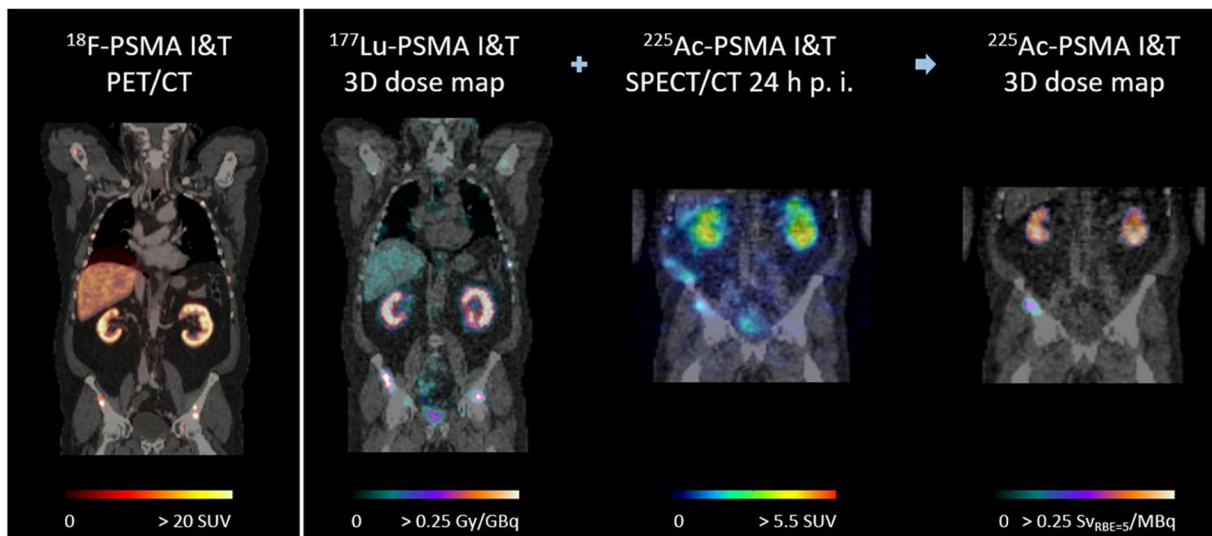
HEGP collimator, 16 projections/head, 128×128 pixel, 210 s/projection). Reconstruction was carried out via a MAP algorithm (30i1s) [6], including CT-based attenuation and dual-energy-window scatter correction and a simulated distance-dependent 2D PSF model (SIMIND). Final absorbed dose assessment was performed by combining the single ^{225}Ac image with the effective half-life information determined from a previous ^{177}Lu -PSMA-I&T imaging sequence [7]. This resulted in an absorbed dose of 0.18 and 0.17 $\text{Sv}_{\text{RBE}=5}/\text{MBq}$ for the left and right kidney, respectively, compared with 0.27 and 0.24 Gy/GBq for the preceding ^{177}Lu cycle (6.2 GBq). A comparison with the pre-therapy ^{18}F -PSMA-I&T PET/CT demonstrates that ^{225}Ac SPECT imaging for this patient was able to locate a small lesion in the right hip. The ^{225}Ac -absorbed dose was determined as 0.26 $\text{Sv}_{\text{RBE}=5}/\text{MBq}$, compared with 0.35 Gy/GBq for ^{177}Lu -PSMA-I&T.

Our analysis demonstrates the feasibility of dosimetry for ^{225}Ac -PSMA-I&T, which provides further insights into theranostic approaches using TAT in mCRPC patients.

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✉ H. Ilhan
harun.ilhan@med.uni-muenchen.de

¹ Department of Nuclear Medicine, University Hospital, LMU Munich, Munich, Germany



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Compliance with ethical standards

All procedures performed in this study involving human participants were in accordance with ethical standards of the institutional and/or national research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards. The retrospective evaluation was approved by the local ethic committee (20-178). Written informed consent was obtained prior to the exam.

Conflict of interest The authors declare that they have no conflict of interest.

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References

1. Kratochwil C, et al. Targeted alpha therapy of mCRPC with ^{225}Ac -PSMA-617: dosimetry estimate and empirical dose finding. *J Nucl Med*. 2017;8(10):1624–31 p. jnumed. 117.191395.
2. Robertson A, et al. Multi-isotope SPECT imaging of the ^{225}Ac decay chain: feasibility studies. *Phys Med Biol*. 2017;62(11):4406.
3. Usmani S, et al. ^{225}Ac prostate-specific membrane antigen posttherapy α imaging: comparing 2 and 3 photopeaks. *Clin Nucl Med*. 2019;44(5):401–3.
4. Vatsa R, et al. ^{225}Ac -PSMA-617 radioligand posttherapy imaging in metastatic castrate-resistant prostate cancer patient using 3 Photopeaks. *Clin Nucl Med*. 2020;45(6):437–8.
5. Chun SY, Fessler JA, Dewaraja YK. Correction for collimator-detector response in SPECT using point spread function template. *IEEE Trans Med Imaging*. 2012;32(2):295–305.
6. Delker A, et al. Dosimetry for ^{177}Lu -DKFZ-PSMA-617: a new radiopharmaceutical for the treatment of metastatic prostate cancer. *Eur J Nucl Med Mol Imaging*. 2016;43(1):42–51.
7. Gosewisch A, et al. Patient-specific image-based bone marrow dosimetry in ^{177}Lu -[DOTA 0, Tyr 3]-Octreotate and ^{177}Lu -DKFZ-PSMA-617 therapy: investigation of a new hybrid image approach. *EJNMMI Res*. 2018;8(1):76.

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