

## Somatostatin receptor expressing tumors: Imaging with $^{68}\text{Ga}$ -DOTATOC-PET/CT versus $^{111}\text{In}$ -DTPAOC-SPECT/CT

P. Aschoff<sup>1</sup>, M. Oeksuez<sup>1</sup>, B. Kemke<sup>1</sup>, K. Zhernosekov<sup>2</sup>, M. Jennewein<sup>2</sup>, F. Roesch<sup>2</sup>, H. Bihl<sup>1</sup>

<sup>1</sup>Klinik für Nuklearmedizin und PET-Center, Klinikum Stuttgart, D-70022, Stuttgart

<sup>2</sup>Institut für Kernchemie, Johannes Gutenberg-Universität, D-55128 Mainz

**Introduction:** Imaging of somatostatin-receptor (SSTR) expressing tumors using  $^{111}\text{In}$ -DTPA-octreotide ( $^{111}\text{In}$ -DTPAOC) has proven to be helpful in the detection of SSTR-expressing tumors and their metastases. One of the drawbacks of this method - even if performed in SPECT-technique - is its limited sensitivity in detecting small lesions.  $^{68}\text{Ga}$ -DOTA-DPhe(1)-Tyr(3)-octreotide ( $^{68}\text{Ga}$ -DOTATOC) is a new PET-radiotracer with a higher specific binding to SSTR2 compared to  $^{111}\text{In}$ -DTPAOC [1].

This study compares the clinical performance of both methods in detecting SSTR2-positive tumor lesions.

**Experimental:** Thirty-six patients with metastatic neuroendocrine gastroenteropancreatic tumor were evaluated with both  $^{111}\text{In}$ -DTPAOC SPECT/CT (185 MBq) and  $^{68}\text{Ga}$ -DOTATOC-PET/CT (50-120 MBq). GE Millennium VG Hawkeye gamma camera and GE Discovery LS PET/CT-scanner were used to perform the respective scans.

400-200 MBq of  $^{68}\text{Ga}$  were obtained using a 20 mCi  $^{68}\text{Ge}/^{68}\text{Ge}$ -generator, based on a  $\text{TiO}_2$ -phase (Cyclotron Co., Obninsk, Russia). For syntheses of  $^{68}\text{Ga}$ -DOTATOC,  $^{68}\text{Ga}$  was pre-concentrated and purified on micro-chromatography column filled with Bio-Rad AG 50W-X8 [2].

**Results:** Whereas  $^{111}\text{In}$ -DTPAOC-scintigraphy was performed in a 2-d protocol,  $^{68}\text{Ga}$ -DOTATOC-PET/CT generated decisive images of high quality 20 min after injection. 95% of all lesions were already seen 10 min after injection (Fig.1). All " $^{111}\text{In}$ -DTPAOC-lesions" were detected with  $^{68}\text{Ga}$ -DOTATOC-PET/CT as well and additional lesions (bone, lymphnodes) could be identified in 24/36 patients (67%). In all patients, lesion localisation was significantly more precise with PET/CT than with SPECT/CT.

In this study,  $^{68}\text{Ga}$ -DOTATOC-PET/CT turned out to be superior to  $^{111}\text{In}$ -DTPAOC-SPECT/CT in the detection of SSTR2-positive tumors and metastases. For clinical practice, it is important that  $^{68}\text{Ga}$ -DOTATOC-PET/CT can be performed in a 1-d protocol, i.e. scans will start immediately after tracer injection (20 min) and will be achieved in about <30 min.

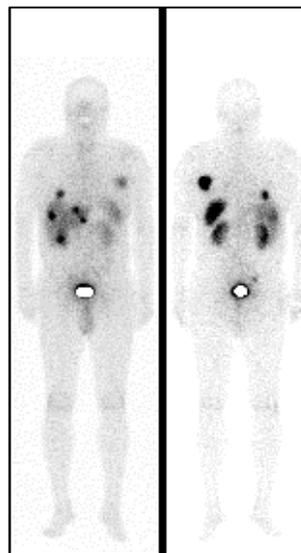
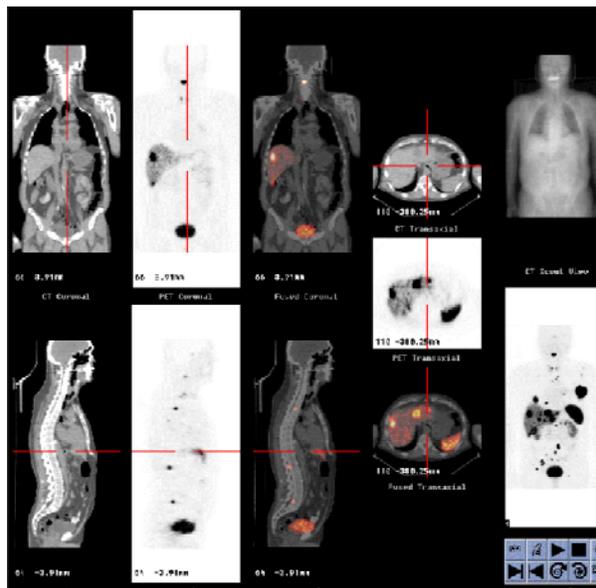


Figure 1

Figure 2



Patient with liver and bone metastasis:  
Figure 1)  $^{111}\text{In}$ -DTPAOC planar imaging  
Figure 2)  $^{68}\text{Ga}$ -DOATATOC PET/CT and image fusion of PET/CT

### References

- [1] Mäcke, H. R., Good, S.: "Radiometals (non-Tc, non-Re) and bifunctional labeling chemistry"; Handbook of Nuclear Chemistry, volume 4, Amsterdam, (2003)
- [2] Zhernosekov, K.P. et. al., this report