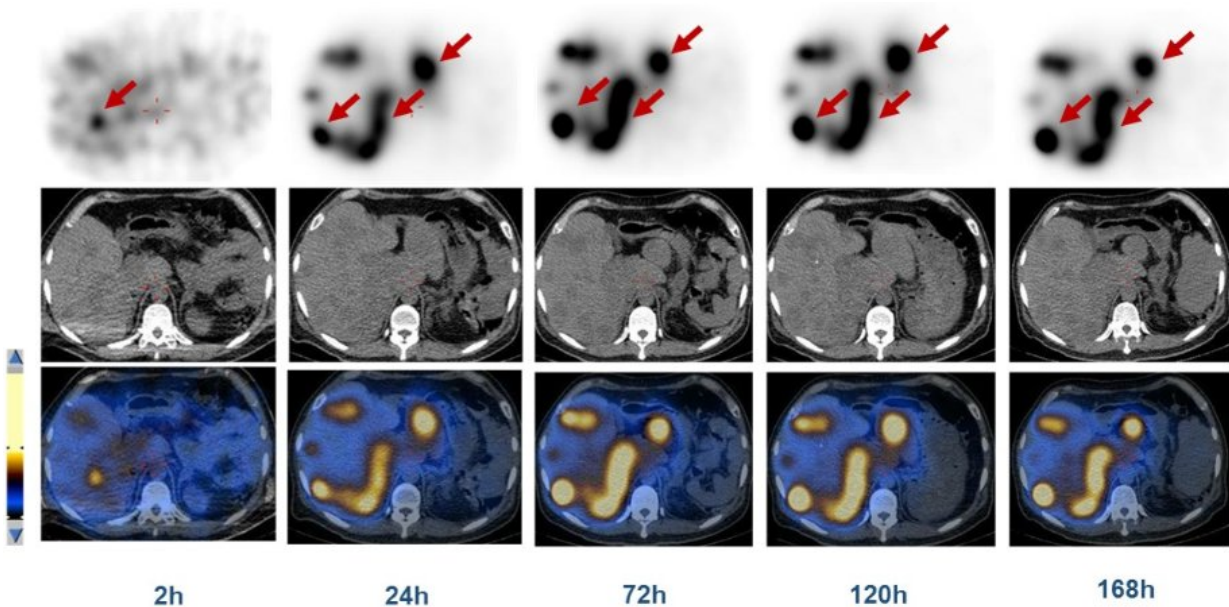


Long-lasting radionuclide therapy for advanced neuroendocrine tumors proves effective

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Scans were done at 2, 24, 72, 120 and 168 hours after the administration of ^{177}Lu -DOTA-EB-TATE. The radiopharmaceutical cleared from the blood pool over time and persistently retained in the tumors (arrows). Credit: J Zhang et al., Peking Union Medical College Hospital, Beijing, China; X Chen et al., Laboratory of Molecular Imaging and Nanomedicine, NIBIB/NIH, Bethesda, MD

A first-in-human study presented at the 2018 Annual Meeting of the

Society of Nuclear Medicine and Molecular Imaging (SNMMI) demonstrates the benefits and safety of a new, long-lasting type of radionuclide therapy for patients with advanced, metastatic neuroendocrine tumors (NETs).

Lutathera-177 (^{177}Lu)-DOTATATE (trade name Lutathera), a peptide receptor radionuclide therapy (PRRT) with radiolabeled somatostatin analogues (peptides), was recently approved by the U.S. Food and Drug Administration for the treatment of NETs. It is the therapeutic part of a [nuclear medicine](#) theranostic pairing. Gallium-68 (^{68}Ga)-DOTATATE is the diagnostic agent used in positron emission tomography/computed tomography (PET/CT) scans that first locates and marks the lesions for follow-up with targeted PRRT delivery directly to the [tumor cells](#) which express high levels of somatostatin receptors (SSTRs). Because the PRRT binds to receptors expressed by the [tumor](#) cells, healthy cells are unharmed.

However, the peptide quickly clears from the blood through the kidneys limiting the accumulation of radioactivity within tumors and making additional treatment cycles necessary to provide the therapeutic dose.

This first-in-human, first-in-class, Phase I trial (ID: NCT03308682) investigated the safety and dosimetry of a novel long-lasting radiolabeled [somatostatin analogue](#) that adds an albumin-binding Evans blue (EB, an azo dye) derivative to ^{177}Lu -DOTATATE. Albumin, the most abundant plasma protein in human blood, is a natural transport protein and has a long circulatory half-life.

" ^{177}Lu -DOTA-EB-TATE is a "three-in-one" therapeutic compound, with an octreotate peptide to find the tumor, an Evans blue motif, which uses endogenous albumin as a reversible carrier to effectively extend the half-life in the blood and substantially increase targeted accumulation and retention within the tumor, and a therapeutic radionuclide to kill the

tumor cells, to finally provide effective treatment of NETs," explains Shawn(Xiaoyuan) Chen, Ph.D., senior investigator, of National Institute of Biomedical Imaging and Bioengineering at the National Institutes of Health , Bethesda, Maryland.

For the study, conducted in collaboration with researchers at the U.S. National Institute of Biomedical Imaging and Bioengineering, 8 patients (6 men and 2 women ranging in age from 27 to 61 years old) with advanced metastatic neuroendocrine tumors were recruited from Peking Union Medical College Hospital and the Chinese Academy of Medical Sciences in Beijing, China.

Each patient underwent whole-body ^{68}Ga -DOTATATE PET/CT. Five of the patients then accepted intravenous injection with a single dose of 0.35-0.70 GBq of ^{177}Lu -DOTA-EB-TATE within one week, and were monitored at 2, 24, 72, 120 and 168 hours after ^{177}Lu -DOTA-EB-TATE administration with serial whole-body planar and single photon emission computed tomography (SPECT)/CT images acquired. The other 3 patients accepted a dose of 0.28-0.41 GBq of ^{177}Lu -DOTATATE and were monitored at 1, 3, 4, 24 and 72 hours with the same imaging procedures. Complete physical examinations, including vital signs, blood count, biochemistry, and immunology analyses were performed immediately before and 1, 3, and 7 days, as well as 3 months, after treatment.

Administration of ^{177}Lu -DOTA-EB-TATE was well tolerated, with no adverse symptoms reported throughout the procedure and follow-up. The total effective dose equivalent and effective dose were 0.2048 ± 0.1605 and 0.0804 ± 0.0500 mSv/MBq for ^{177}Lu -DOTA-EB-TATE and 0.1735 ± 0.0722 and 0.0693 ± 0.0317 mSv/MBq for ^{177}Lu -DOTATATE. The liver, kidneys, bone marrow and total body received slightly higher doses (mGy/MBq) with ^{177}Lu -DOTA-EB-TATE than with ^{177}Lu -DOTATATE, while the spleen received lower doses with

Lu-DOTA-EB-TATE. Blood clearance of ^{177}Lu -DOTA-EB-TATE was also slower. Most importantly, ^{177}Lu -DOTA-EB-TATE lasted in the tumors more than 4 times longer than ^{177}Lu -DOTATATE.

Jingjing Zhang and Zhaohui Zhu of Peking Union Medical College Hospital point out, "By introducing an albumin binding moiety, this long-lasting radiolabeled somatostatin analogue has remarkably enhanced uptake and retention in SSTR-positive tumors, which is important to increase the therapeutic efficacy in patients. With proper selection of patients with advanced metastatic neuroendocrine tumors, ^{177}Lu -DOTA-EB-TATE has great potential to be a highly effective treatment, while providing a safe dose with less frequency of administration than is possible with ^{177}Lu -DOTATATE."

More information: Abstract 118: "Safety, Pharmacokinetics and Dosimetry of a Long-lasting Radiolabeled Somatostatin Analogue ^{177}Lu -DOTA-EB-TATE in Patients with Advanced Metastatic Neuroendocrine Tumors: A Phase 1 First-in-human Study," Jingjing Zhang, MD,Ph.D., Yuejuan Cheng, MD,Hao Wang, MD, Jie Zang, Ph.D., Fang Li, MD, Chunmei Bai, MD, and Zhaohui Zhu, MD, Peking Union Medical College Hospital; Gang Niu, MD, Orit Jacobson, Ph.D.4, and Xiaoyuan Chen, Ph.D., U.S. National Institutes of Health, Bethesda, MD. SNMMI's 65th Annual Meeting, June 23-26, Philadelphia.
[jnm.snmjournals.org/content/59 ... a2-8c75-621d18d0d7fa](http://jnm.snmjournals.org/content/59...a2-8c75-621d18d0d7fa)

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