

A better way to diagnose and manage neuroendocrine tumors

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In-111-DTPA-octreotide scan (A) demonstrates recurrent mesenteric nodule (arrow) on the anterior view in a patient with a mid-gut carcinoid tumor resected 5 years previously, with no other tumor seen with planar or SPECT/CT (B) images. Ga-68 DOTATATE PET/CT anterior 3D MIP (C) and axial fused images (D) demonstrate other metastases (arrows), changing the surgical plan for resection. Findings were verified at surgery. Credit: Ronald C. Walker, MD

A recent study reported in the May issue of the Journal of Nuclear



Medicine demonstrates that Ga-68 DOTATATE PET/CT scans are superior to In-111 pentetreotide scans, the current imaging standard in the United States for detecting neuroendocrine tumors (NETS), and could significantly impact treatment management.

NETS occur mostly in the respiratory and digestive tracts and are usually slow-growing. They can be difficult to diagnose, and many treatment options exist. It's therefore critical to delineate the extent of disease accurately for proper management. While the incidence of NETS is relatively low, with 2.5-5 cases per 100,000 in the United States, data from the National Cancer Institute show a five-fold increase worldwide from 1973 to 2004. NETS can be malignant and, although they comprise less than two percent of gastrointestinal cancers, they are more prevalent than stomach and pancreatic cancers combined.

Ronald C. Walker, MD, corresponding author for the study and professor of clinical radiology and radiological sciences at Vanderbilt University School of Medicine, explains, "Our purpose was to evaluate the safety and efficacy of Ga-68 DOTATATE PET/CT compared to In-111 pentetreotide imaging for diagnosis, staging and re-staging of pulmonary and gastroenteropancreatic neuroendocrine tumors." With concerns for patient safety, detailed toxicity data were also collected.

The two imaging methods were performed in 78 of 97 consecutively enrolled patients with known or suspected pulmonary or gastroenteropancreatic (GEP) NETs. The study found that Ga-68 DOTATATE PET/CT combined with CT and/or liver MRI changed care in 28 of 78 (36 percent) patients. In addition, Ga-68 DOTATATE PET/CT correctly identified three patients for peptide receptor radiotherapy who had been incorrectly classified by In-111 pentetreotide. Demonstrating no significant toxicity, lower radiation exposure and improved accuracy, the study makes a strong case for the use of Ga-68 DOTATATE imaging over the current standard, where



available.

While Ga-68 DOTATATE PET/CT is in widespread use outside of the United States, the U.S. Food and Drug Administration (FDA) has not yet approved its use for the diagnosis, staging and treatment management of NETS. As a result, insurance companies do not cover its use.

Walker says, "Hopefully, our investigation will provide sufficient evidence on the safety and efficacy of Ga-68 DOTATATE to the U.S. FDA to allow approval. If so, then patients throughout the United States could soon have access to a higher-quality scan, allowing better patient management decisions while also lowering radiation exposure and shortening examination time." Such approval would also open up the possibility of reimbursement for the scans by third-party payers.

These findings could have an impact on the future of <u>nuclear medicine</u>, as well, as Walker explains: "If our evidence results in FDA approval of Ga-68 DOTATATE imaging for routine use in the treatment management of patients with <u>neuroendocrine tumors</u>, it would represent the first approval of a Ga-68-labeled PET imaging radiopharmaceutical in the U.S. This could help pave the way for similar studies to allow approval in the U.S. and elsewhere of other Ga-68-labeled radiopharmaceuticals, such as Ga-68 DOTATOC and Ga-68 PSMA."

More information: S. A. Deppen et al, Safety and Efficacy of 68Ga-DOTATATE PET/CT for Diagnosis, Staging, and Treatment Management of Neuroendocrine Tumors, *Journal of Nuclear Medicine* (2016). <u>DOI: 10.2967/jnumed.115.163865</u>

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