New Targeted Hormone-Radiation Treatment Slows Growth of Midgut Neuroendocrine Tumors

For immediate release
January 15, 2016

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ASCO Perspective

“It’s exciting to see a novel therapy have such a significant effect on tumor growth, particularly for these patients who have limited treatment options,” said Smitha Krishnamurthi, MD, ASCO Spokesperson and moderator of today’s presscast.

ALEXANDRIA, Va. – Early results from a phase III study of patients with previously treated, advanced midgut neuroendocrine tumors show that a novel therapy, 177Lutetium-DOTATATE (Lutathera), may substantially slow tumor growth. Patients treated with the experimental drug had a 79% lower risk of disease progression or death compared to those treated with octreotide LAR 60 mg every four weeks. The study will be presented at the upcoming 2016 Gastrointestinal Cancers Symposium in San Francisco.

“Our results indicate that Lutathera appears to be a safe and effective treatment option. On average, tumors responded to the drug for several years before they began growing again,” said lead study author Jonathan R. Strosberg, MD, a medical oncologist at the Moffitt Cancer Center in Tampa, FL. “The new therapy is also more convenient – it requires only four treatments, as opposed to medications that patients have to take daily over long periods of time.”

177Lutetium-DOTATATE belongs to a new class of drugs known as Peptide Receptor Radionuclide Therapy (PRRT), which combine hormone therapy and radiotherapy. In the case of 177Lutetium-DOTATATE, a somatostatin analog is attached to a radioactive molecule, enabling targeted delivery of cancer-killing radiation to tumors.

Most patients with metastatic midgut neuroendocrine tumors (i.e., those that begin in the small intestine and proximal colon) receive hormone therapy with a somatostatin analog, such as octreotide or lanreotide. There are currently no effective systemic second-line treatment options for
According to the authors, this is the first prospective, randomized trial to evaluate the efficacy of 
\textsuperscript{177}Lutetium-DOTATATE in patients with advanced midgut neuroendocrine tumors. All 230 trial 
participants had advanced tumors that worsened despite first-line somatostatin analog therapy. The patients were randomly assigned to treatment with \textsuperscript{177}Lutetium-DOTATATE or high-dose octreotide LAR.

At the time of data analysis, 23 patients in the \textsuperscript{177}Lutetium-DOTATATE group experienced 
disease progression, compared to 67 in the octreotide LAR group. The median progression-free 
survival was eight months in the octreotide LAR group and was not reached in the \textsuperscript{177}Lutetium-
DOTATATE group.

The preliminary findings of the study also suggest that \textsuperscript{177}Lutetium-DOTATATE may extend 
patient survival. There were substantially fewer deaths in the \textsuperscript{177}Lutetium-DOTATATE group vs. 
the standard care group (13 vs. 22). However, longer follow-up is needed to determine the 
definitive impact of \textsuperscript{177}Lutetium-DOTATATE on long-term survival.

Overall, the numbers of adverse events, including serious side effects, were similar between the 
two treatment groups. Octreotide often causes gas and bloating and can lead to development of 
gallstones with long-term use. \textsuperscript{177}Lutetium-DOTATATE can cause low blood cell counts, which 
are usually transient.

Neuroendocrine tumors are a group of cancers that begin in hormone-producing (neuroendocrine) 
cells of various organs in the body. Each year about 8,000 people in the United States are 
diagnosed with a neuroendocrine tumor of the gastrointestinal tract. \[1\] While they are rare overall, 
the incidence of neuroendocrine tumors is on the rise.

This study received funding from Advanced Accelerator Applications (AAA).

\[1\] \url{http://www.cancer.org/cancer/gastrointestinalcarcinoidtumor/detailedguide/gastrointestinal-

View the full abstract.

For your readers:

- Guide to Neuroendocrine Tumors
- Clinical Trials
- Interactive History of Cancer Advances

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Click here to view the disclosures for the News Planning Team.

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