

Neuroendocrine Tumors and Lanreotide Depot: Clinical Considerations and Nurse and Patient Preferences

Pamela Ryan, RN, BSN, Alexandria T. Phan, MD, Daphne T. Adelman, RN, BSN, MBA, and Michiko Iwasaki, RN, CCRC



© andrewsafonov/iStock/Thinkstock

Background: Somatostatin analogs (SSAs) are a mainstay therapy for the treatment of carcinoid syndrome associated with neuroendocrine tumors (NETs). They are effective for a range of gastroenteropancreatic NETs (GEP-NETs). Lanreotide depot (Somatuline®) is an SSA that is approved for the treatment of GEP-NETs to improve progression-free survival (PFS).

Objectives: The article reviews the efficacy, safety, and administration of lanreotide depot and relates those attributes to considerations and preferences of oncology nurses and their patients.

Methods: A review of the literature on the use of lanreotide for the treatment of NETs and carcinoid syndrome was conducted. In addition, the literature on drug delivery and routes of administration was surveyed to provide context for comparative studies related to clinical and patient preferences.

Findings: Lanreotide depot prolongs PFS and is well tolerated by patients who expressed satisfaction in the ability to control symptoms related to carcinoid syndrome. Nurses cited several benefits to using lanreotide depot in the clinical setting, including more time saved to address other patient care issues. Attributes of lanreotide depot—including its efficacy, safety and tolerability, dosing and administration, and cost—may contribute to healthcare decisions regarding the treatment and management of NETs.

Pamela Ryan, RN, BSN, is a nurse manager in the Neuroendocrine Tumor Program and Infusion Center at the Ochsner Medical Center in Kenner, LA; Alexandria T. Phan, MD, is the director of Gastrointestinal Medical Oncology at Houston Methodist Hospital in Texas; Daphne T. Adelman, RN, BSN, MBA, is a clinical nurse specialist in the Feinberg School of Medicine at Northwestern University in Chicago, IL; and Michiko Iwasaki, RN, CCRC, is a research nurse supervisor at the University of Texas MD Anderson Cancer Center in Houston. The authors take full responsibility for the content of the article. Ryan has previously consulted for and received honorarium from Ipsen and Novartis, including payment for lectures or services on speakers bureaus, and payment for writing or reviewing this article. Phan has previously consulted for and received honorarium from Ipsen. Adelman has previously consulted for and received honorarium from Novartis, Ipsen, Pfizer, and Concept and Charisma. Writing and editorial support was provided by ClinicalThinking through support from Ipsen Biopharmaceuticals. The content of this article has been reviewed by independent peer reviewers to ensure that it is balanced, objective, and free from commercial bias. No financial relationships relevant to the content of this article have been disclosed by the independent peer reviewers or editorial staff. Mention of specific products and opinions related to those products do not indicate or imply endorsement by the *Clinical Journal of Oncology Nursing* or the Oncology Nursing Society. Ryan can be reached at pryan@ochsner.org, with copy to editor at CJONEditor@ons.org. (Submitted July 2015. Revisions submitted February 2016. Accepted for publication February 15, 2016.)

Key words: lanreotide; neuroendocrine tumors; carcinoid syndrome; nurse preferences; patient preferences; subcutaneous injection

Digital Object Identifier: 10.1188/16.CJON.E139-E146

Neuroendocrine tumors (NETs) arise from secretory cells of the neuroendocrine system and are predominantly found in the gastrointestinal tract and pancreas, although they can occur in virtually every organ (Cives & Strosberg, 2014). The incidence of NETs may be as high as 5.86 in 100,000 (95% confidence interval [CI] [5.4, 6.35]), based on 2009 data (Hallet et al., 2014). Advanced metastatic NETs are associated with a relatively poor prognosis and may be unresectable. Treatment has traditionally focused on management of the symptoms of this chronic condition (Caplin, Pavel, et al., 2014; Oberg, 2012; Wolin, 2012).

Patients who present with well-differentiated (low and intermediate grade) NETs of the stomach, intestine, and pancreas are usually first treated with surgical resection

whenever possible and/or ablation. Medical management for gastroenteropancreatic (GEP)-NETs may also involve chemotherapeutic agents, monoclonal antibodies, and/or biotherapy with somatostatin analogs (SSAs) (Falconi et al., 2012; Kulke et al., 2010; National Comprehensive Cancer Network [NCCN], 2015; Oberg, 2012). Long-acting SSAs have been used successfully to treat GEP-NETs (Caplin, Pavel, et al., 2014; NCCN, 2015). Generally well tolerated, SSAs are used for the relief of symptoms associated with carcinoid syndrome, a condition that typically presents in advanced neuroendocrine cancer that has metastasized to the liver or with secretory pancreatic or midgut NETs (Pavel et al., 2012).

This article reviews a long-acting SSA, lanreotide depot (Somatuline®), and examines factors such as efficacy, tolerability, dosing and administration, patient and nurse preferences,