Preemptive Management of Dermatologic Toxicities Associated With Epidermal Growth Factor Receptor Inhibitors

Jean Boucher, PhD, RN, ANP, Linnea Olson, RN, BSN, and Bilal Piperdi, MD

Epidermal growth factor receptor inhibitors (EGFRIs) are a treatment option for patients diagnosed with advanced-stage gastrointestinal, lung, and head and neck cancers. The most prevalent complications associated with EGFRIs are dermatologic toxicities, which may result in either disruption or discontinuation of treatment and adversely affect patients' quality of life. Nurses play a vital role in educating patients about EGFRI-related dermatologic toxicities; therefore, nurses must continue to educate themselves on the various aspects of EGFRI treatment. An overview of the EGF signaling pathway is provided, and dermatologic toxicities associated with EGFRI treatment are described. A review of several studies evaluating reactive skin treatment regimens also are discussed. Nurses play a critical role in providing patient support. Informing patients about potential EGFRI-related symptoms and dermatologic toxicities will help prepare patients for their course of treatment. In addition, nurses should provide patients with a variety of coping strategies to help manage dermatologic toxicities that will assist in enhancing patients' adherence to EGFRI treatment.

ncology nurses treat an increasing number of patients diagnosed with cancer who receive epidermal growth factor receptor inhibitors (EGFRIs). EGFRIs, either alone or in combination with chemotherapy, are the standard of care for the treatment of advanced-stage gastrointestinal (colorectal and pancreatic) (Jonker et al., 2007; Moore et al., 2007; Van Cutsem et al., 2007), lung (Shepherd et al., 2005), and head and neck cancers (Bonner et al., 2006). Treatment with EGFRIs may result in dermatologic toxicities that lead to interruption or discontinuation of patient treatment (Lacouture, Cotliar, & Mitchell, 2007; Molinari, De Quatrebarbes, Andre, & Aractingi, 2005); therefore, management of dermatologic toxicities in patients treated with EGFRIs is crucial to help promote patient adherence with therapy. The purpose of this article is to inform nurses about preemptive management strategies for dermatologic toxicities associated with EGFRI treatment. In addition, the important role nurses play in preemptive care working with healthcare providers will be addressed; including educating,

At a Glance

- Dermatologic toxicities are the most common side effects associated with epidermal growth factor receptor inhibitor (EGFRI) treatment.
- Nurses should be educated about the benefits of administering a preemptive skin treatment regimen to patients to potentially prevent or minimize dermatologic toxicities associated with EGFRI treatment.
- Oncology nurses can collaborate with physicians to perform a thorough assessment of EGFRI-related dermatologic toxicities and provide support and education to patients.

monitoring, and providing supportive care that promotes better understanding of EGFRI-related dermatologic toxicities and coping strategies for patients. Results from several studies evaluating the use of preemptive treatments (e.g., topical steroids,

Jean Boucher, PhD, RN, ANP, is an oncology nurse practitioner and an assistant professor, Linnea Olson, RN, BSN, is a clinical research nurse, and Bilal Piperdi, MD, is an assistant professor of medicine, all at the University of Massachusetts Memorial Medical Center in Worcester. The authors take full responsibility for the content of the article. Editorial support was provided by Amgen, Inc. 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. (Submitted September 2010. Revision submitted February 2011. Accepted for publication February 25, 2011.)

Digital Object Identifier: 10.1188/11.CJON.501-508