Chronic Diseases and Non-Melanoma Skin Cancer: Is There an Association?

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Non-melanoma skin cancer (NMSC) has been associated with several other conditions, including melanoma and non-Hodgkin's lymphoma (Hjalgri *et al.*, 2000; Hu *et al.*, 2005). In addition, the risk of NMSC has been found to be greater in various chronic disease groups (Frisch *et al.*, 1996; Frisch and Melbye, 1995). It is uncertain whether this association is related to the treatments for those chronic diseases (such as radiation therapy or immunosuppressant therapy) rather than the disease itself (Saladi and Persaud, 2005; Karagas *et al.*, 2001). Moreover, using a Danish nationwide cohort of patients with NMSC, Jensen *et al.* (2006) found a lower mortality in basal cell carcinoma (BCC) patients compared with the general population, reflecting what the authors suggested to be a selection bias to that data set.



In a follow-up study utilizing a nationwide Danish patient registry linked

to the nationwide cohort of patients with NMSC, the researchers attempted to analyze the development of NMSC in patients hospitalized for various chronic diseases (Jensen *et al.*, 2008). To control for potential surveillance bias, they also evaluated the severity of NMSC diagnosed. Several chronic diseases were associated with increased incidence of NMSC, including an association between BCC and transplantation as well as lymphoma. The risk of squamous cell carcinoma was found to be higher in patients previously hospitalized for leukemia as well as other skin diseases.

Through the following questions, we examine this paper in greater detail. For brief answers, please refer to http://network.nature.com/group/jidclub.

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QUESTIONS

- 1. Why would patients with other conditions be at risk for skin cancer?
- 2. What is the Charlson index, and how was it used in this study?
- 3. How did the investigators control for potential bias in this study?
- 4. What were the major findings of this study?
- 5. How did the investigators explain their findings?
- 6. What may be the clinical implications of this article?
- 7. What future studies would help confirm these hypothesis-generating results?

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