65

With Melanoma, Risk of Other Cancers Is High

BY DENISE NAPOLI

FROM THE ANNUAL MEETING OF THE AMERICAN COLLEGE OF MOHS SURGERY

NEW YORK — The risk of developing a primary malignancy of the salivary gland, bone, prostate, and other areas is significantly—and in some cases dramatically—increased following a cutaneous melanoma diagnosis.

That's according to research by Dr. Joshua Spanogle, a resident in the department of dermatology at the Mayo Clinic in Rochester, Minn., presented during an abstract session at the meeting.

Dr. Spanogle performed an analysis of data from the Surveillance, Epidemiology, and End Results database (SEER) from 1973 to 2003, which included "over 1.3 million person-years of observation," he said.

A total of 151,996 patients were found to have a cutaneous melanoma diagnosis during that period; 16,591 (11%) of these patients had a second documented primary malignancy sometime in the next 120 months. That observed rate was 32% higher than what would be expected among a healthy population in that time period, he said.

Further analysis of the results revealed that particular cancers carried a significantly higher risk than others.

The most striking result of the analysis, perhaps not surprisingly, was for a second primary cutaneous melanoma: There were 3,923 through 120 months of follow-up, for a standardized incidence ratio of 8.99.

"This is the 600-pound elephant in the room," said Dr. Spanogle.

But other cancers had high standardized incidence ratios (SIRs) as well. Salivary gland malignancies following carcinoma had a SIR of 2.18 overall. Prostate cancer had a SIR of 1.13 following melanoma. Breast cancer showed a SIR of 1.07. And soft tissue cancers, including malignancies of the heart, had a SIR of 2.80.

On the other hand, "quite a few cancers had a decreased incidence following melanoma," said Dr. Spanogle, including cancer of the liver (SIR 0.77), lungs (0.83), cervix (0.57), and pharynx (0.61).

The lower incidence of those kinds of cancer following melanoma could be because risk factors for melanoma are associated with higher socioeconomic sta-



tus, like fair skin and intermittent high-intensity UV exposure (tanning), said Dr. Spanogle.

In contrast, risk factors for lung cancer and liver cancer are associated with comorbidities commonly found in lower socioeconomic patients, like smoking and hepatitis. Moreover, according to Dr. Spanogle, the risks of secondary cancers of the prostate gland, bone, soft tissue, and salivary gland remained elevated throughout the study period, "implying no surveillance bias."

Instead, he speculated that the link could be genetic, and said that future research into the possibility is warranted. ■

Major Finding: Of 151,996 patients with a cutaneous melanoma diagnosis, 16,591 went on to have a second documented primary malignancy.

Data Source: A retrospective analysis of data from the SEER database for 1973-2003, encompassing over 1.3 million person-years of observation.

Disclosures: Dr. Spanogle reported having no relevant disclosures. His study has been accepted for publication in the Journal of the American Academy of Dermatology.



And help your patients improve glycemic control with NovoLog[®], the #1 selling rapid-acting mealtime insulin.¹

NovoLog[®] is an insulin analog indicated to improve glycemic control in adults and children with diabetes mellitus.

Important safety information

NovoLog[®] is contraindicated during episodes of hypoglycemia and in patients hypersensitive to NovoLog[®] or one of its excipients. NovoLog[®] has a more rapid onset and shorter duration of action than regular human insulin. An injection of NovoLog[®] should be immediately followed by a meal within 5 to 10 minutes. Because of the short duration of action of NovoLog[®], a longer-acting insulin also should be used in patients with type 1 diabetes and may be needed in patients with type 2 diabetes. When used in an external subcutaneous insulin infusion pump, NovoLog[®] should not be mixed with any other insulin or diluent. Hypoglycemia is the most common adverse effect of all insulin therapies, including NovoLog[®]. The timing of hypoglycemia usually reflects the time-action profile of the administered insulins.

Any change of insulin dose should be made cautiously and only under medical supervision. Glucose monitoring is recommended for all patients with diabetes and is particularly important for patients using external pump infusion therapy. As with all insulin preparations, the time course of action of NovoLog[®] may vary in different individuals or at different times in the same individual and is dependent on many conditions, including injection site, local blood supply, temperature, and level of physical activity.

Severe, life-threatening generalized allergy, including anaphylactic reaction, may occur with any insulin product, including NovoLog[®]. Adverse reactions observed with NovoLog[®] include hypoglycemia, allergic reactions, local injection site reactions, lipodystrophy, rash, and pruritus. Insulin, particularly when given intravenously or in settings of poor glycemic control, may cause hypokalemia. Like all insulins, NovoLog[®] requirements may be reduced in patients with renal impairment or hepatic impairment.

To access complimentary e-learning programs, visit novomedlink.com/NovoLog.



Please see brief summary of Prescribing Information on adjacent page. FlexPen® and NovoLog® are registered trademarks of Novo Nordisk A/S. © 2009 Novo Nordisk Inc. 137970 June 2009

