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Novel Immunotherapy for Malignant Melanoma with a Monoclonal Antibody That Blocks CEACAM1 Homophilic Interactions
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Glycolytic Inhibition Alters Anaplastic Thyroid Carcinoma Tumor Metabolism and Improves Response to Conventional Chemotherapy and Radiation
Vlad C. Sandulache, Heath D. Skinner, Yuan Wang, Yunyun Chen, Cristina T. Dodge, Thomas J. Ow, James A. Bankson, Jeffrey N. Myers, and Stephen Y. Lai

Correction: Microtubule Inhibitors: Differentiating Tubulin-Inhibiting Agents Based on Mechanisms of Action, Clinical Activity, and Resistance

ABOUT THE COVER

The CEACAM1 protein protects melanoma cells from cytotoxic lymphocytes in vitro via homophilic intercellular interactions. Immunohistochemistry of a human lymph node infiltrated with melanoma cells for CEACAM1 (brown pigmentation) and CD8 (pink pigmentation) showed that almost all CD8-positive lymphocytes in the tumor and its close vicinity were CEACAM1⁺, while most lymphocytes in other areas distant from tumor edge were mostly CEACAM1⁻. This strongly suggests that CEACAM1-mediated inhibition occurs in vivo and thus its blockade is a promising strategy for cancer immunotherapy. For details, see article by Ortenberg and colleagues on page 1300.