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PREDICTIVE BIOMARKERS AND PERSONALIZED MEDICINE

Inactivation of the CDKN2A Tumor-Suppressor Gene by Deletion or Methylation Is Common at Diagnosis in Follicular Lymphoma and Associated with Poor Clinical Outcome
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A Functional Germline Variant in GLI1 Implicates Hedgehog Signaling in Clinical Outcome of Stage II and III Colon Carcinoma Patients

Noninvasive Detection of Response and Resistance in EGFR-Mutant Lung Cancer Using Quantitative Next-Generation Genotyping of Cell-Free Plasma DNA

Correction: Carbonic Anhydrase IX Promotes Tumor Growth and Necrosis In Vivo and Inhibition Enhances Anti-VEGF Therapy

ABOUT THE COVER
The cover shows a section of an intracranial glioblastoma (GBM). GBM cells express the fluorescent protein citrine (green) and the high-mobility group protein B1 (HMGB1) fused to the red fluorescent protein cherry. In living cells HMGB1 is located in the nucleus; upon cell death, HMGB1 is translocated to the cytoplasm and is eventually secreted. Circulating levels of HMGB1 may constitute a noninvasive surrogate biomarker of therapeutic efficacy. For details, see the article by Candolfi and colleagues on page 1555 of this issue.