Targeting Oncogenic ALK: A Promising Strategy for Cancer Treatment
Enrique Grande, Maria-Victoria Bolós, and Edurne Arriola

GPR54 Is a Target for Suppression of Metastasis in Endometrial Cancer
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Targeting the Replication Checkpoint Using SCH 900776, a Potent and Functionally Selective CHK1 Inhibitor Identified via High Content Screening
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Targeted Delivery of an Antibody–Mutant Human Endostatin Fusion Protein Results in Enhanced Antitumor Efficacy
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Rapamycin Reverses Splenomegaly and Inhibits Tumor Development in a Transgenic Model of Epstein-Barr Virus–Related Burkitt’s Lymphoma
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Gene Expression Profiling Provides Insights into Pathways of Oxaliplatin-Related Sinusoidal Obstruction Syndrome in Humans
Laura Rubbia-Brandt, Sébastien Tauzin, Catherine Brezault, Céline Delucinge-Vivier, Patrick Descombes, Bertrand Dousset, Pietro E. Majno, Gilles Mentha, and Benoit Terris

Evasion Mechanisms to Igf1r Inhibition in Rhabdomyosarcoma

Correction: Interstitial Infusion of Glioma-Targeted Recombinant Immunotoxin 8H9scFv-PE38

ABOUT THE COVER
Genetically-engineered mouse models often represent some of the most physiologically accurate models of cancer from which to understand the tumor microenvironment and with which to perform preclinical trials. Abraham and colleagues present studies of a prototypic insulin-like growth factor receptor inhibitor using both genetically-engineered mouse models and the shell-free quail chorioallantois membrane (CAM) assay. Remarkably, the inexpensive short term (2 week) CAM assay offers xenografted tumors a scaffold of lymphatics, arteries, and veins that mimic short-term in vivo growth with all the advantages of intravital imaging. Photograph credits, Elaine Huang and Audra Lee. For details, see article by Abraham and colleagues on page 697.
Molecular Cancer Therapeutics

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