An Antibody Targeted to VEGFR-2 Ig Domains 4-7 Inhibits VEGF-2 Activation and VEGFR-2-Dependent Angiogenesis without Affecting Ligand Binding

Determinants of Mitotic Catastrophe on Abrogation of the G2 DNA Damage Checkpoint by UCN-01

(−)-Gossypol Suppresses the Growth of Human Prostate Cancer Xenografts via Modulating VEGF Signaling–Mediated Angiogenesis

Dependence on the MUC1-C Oncoprotein in Non–Small Cell Lung Cancer Cells

Antitumor Activity of the Hsp90 Inhibitor IPI-504 in HER2-Positive Trastuzumab-Resistant Breast Cancer

High-Throughput Screen Identifies Novel Inhibitors of Cancer Biomarker α-Methylacyl Coenzyme A Racemase (AMACR/P504S)
PRECLINICAL DEVELOPMENT

839  Caspase-3–Dependent Mitotic Checkpoint Inactivation by the Small-Molecule Inducers of Mitotic Slippage SU6656 and Geraldol
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850  A Human Model of Epithelial to Mesenchymal Transition to Monitor Drug Efficacy in Hepatocellular Carcinoma Progression
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861  Assessing the Activity of Cediranib, a VEGFR-2/3 Tyrosine Kinase Inhibitor, against VEGFR-1 and Members of the Structurally Related PDGFR Family

874  2-Methoxyestradiol Analogue ENMD-1198 Reduces Breast Cancer-Induced Osteolysis and Tumor Burden Both In Vitro and In Vivo
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883  Curcumin Inhibition of the Functional Interaction between Integrin α6β4 and the Epidermal Growth Factor Receptor
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883  Bortezomib Sensitizes HCC Cells to CS-1008, an Antihuman Death Receptor 5 Antibody, through the Inhibition of CIP2A

892  Therapeutic Potential and Molecular Mechanism of a Novel, Potent, Nonpeptide, Smac Mimetic SM-164 in Combination with TRAIL for Cancer Treatment
Jianfeng Lu, Donna McEachern, Haiying Sun, Longchuan Bai, Yuefeng Peng, Su Qiu, Rebecca Miller, Jinhui Liao, Han Yi, Meiian Liu, Anita Bellail, Chunhai Hao, Shi-Yong Sun, Adrian T. Ting, and Shaomeng Wang

ABOUT THE COVER

Migration of hepatocellular carcinoma (HCC) cells that have undergone epithelial to mesenchymal transition (EMT). The 3sp cells transdifferentiated from malignant hepatocytes in the HCC patient via EMT show a migratory potential as determined by Platypus technology that can be modulated by pharmacological interference. Migrating cells are visualized by staining with CellTracker. For details, see article by van Zijl and colleagues on page 850.