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Cancer Research
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Joseph W. Franses and Elazer R. Edelman

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7356 Targeting PI3K/mTOR Signaling in Cancer
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7362 5-Hydroxymethylcytosine Is Strongly Depleted in Human Cancers but Its Levels Do Not Correlate with IDH1 Mutations
Seong-Gi Jin, Yong Jiang, Runxiang Qiu, Tibor A. Rauch, Yinheng Wang, Gabriele Schackert, Dietmar Krex, Qiang Lu, and Gerd P. Pfeifer

Précis: This study identifies an intermediate step in the process of DNA demethylation as a new contributor to aberrant cancer epigenomes.

INTEGRATED SYSTEMS AND TECHNOLOGIES

7366 Quantifying the Role of Angiogenesis in Malignant Progression of Gliomas: In Silico Modeling Integrates Imaging and Histology
Kristin R. Swanson, Russell C. Rockne, Jonathan Claridge, Mark A. Chaplain, Ellsworth C. Alvord Jr, and Alexander R.A. Anderson

Précis: This modeling study shows that malignant progression of a tumor does not necessarily rely on changes in either the rate of proliferation or invasion.

7372 Metabolomic Profiling Reveals Potential Markers and Bioprocesses Altered in Bladder Cancer Progression

Précis: Metabolic profiling in bladder cancer reveals candidate biomarkers that may be effective in urine sample analysis, with the potential to offer a simple, cost-effective means of diagnosis and prognosis in this disease.

7383 In Vivo Magnetic Resonance Imaging of the Estrogen Receptor in an Orthotopic Model of Human Breast Cancer
Adi Pais, Chidambaram Gunanathan, Baanan Margalit, Inbal Etiti Biton, Ady Yosepovich, David Milstein, and Hadassa Degani

Précis: Findings offer preclinical proof-of-concept for a novel ER-targeting contrast agent that can detect ER-positive tumors in a noninvasive manner in vivo.
A Framework to Select Clinically Relevant Cancer Cell Lines for Investigation by Establishing Their Molecular Similarity with Primary Human Cancers
Garrett M. Dancik, Yuanbin Ru, Charles R. Owens, and Dan Theodorescu

Précis: Unfortunately, experimental findings from established human cancer cell lines often do not translate to clinical settings, but by selecting cell lines that better reflect human pathobiology it may be possible to improve this situation.

Functional Characterization of an scFv-Fc Antibody that Immunotherapeutically Targets the Common Cancer Cell Surface Proteoglycan CSPG4
Xinhui Wang, Akihiro Katayama, Yangyang Wang, Ling Yu, Elvira Favoino, Koichi Sakakura, Alessandra Favole, Takahiro Tsukichikawa, Susan Silver, Simon C. Watkins, Toshiro Kageshita, and Soldano Ferrone

Précis: Antibody targeting of cell surface proteoglycans that are widely overexpressed in human tumors offers an appealing general approach for passive immunotherapy in cancer patients.

Smad7 Expression in T cells Prevents Colitis-Associated Cancer
Angelamaria Rizzo, Maximilian J. Waldner, Carmine Stolfi, Massimiliano Sarra, Daniele Fina, Christoph Becker, Markus F. Neurath, Thomas T. Macdonald, Francesco Pallone, Giovanni Monte Leone, and Massimo C. Fantini

Précis: This study illustrates that chronic inflammation in colitis does not necessarily heighten risks of colorectal tumorigenesis, based on epigenetic differences in the immune microenvironment that dictate whether colitis-associated inflammation promotes or retards tumor development.

Hypoxia Induces Escape from Innate Immunity in Cancer Cells via Increased Expression of ADAM10: Role of Nitric Oxide
Ivraym B. Barsoum, Thomas K. Hamilton, Xin Li, Tiziana Cotechini, Ellen A. Miles, D. Robert Siemens, and Charles H. Graham

Précis: Nitric oxide can block hypoxia-mediated evasion from innate immunity, suggesting the immediate clinical exploration of nitric oxide mimetics like nitroglycerin for cancer treatment.

Dendritic Cell Internalization of α-Galactosyler ceramide from CD8 T Cells Induces Potent Antitumor CD8 T-cell Responses
Dong Hoon Choi, Kwang Soon Kim, Se Hwan Yang, Doo Hyun Chung, Boyeong Song, Jonathan Sprent, Jae Ho Cho, and Young Chul Sung

Précis: This study reports a strategy that might generally improve any adoptive T-cell immunotherapy, based on treatment of the T cells with an immune stimulatory lipid prior to transfer into cancer patients.

Peptide-Conjugated PAMAM Dendrimer as a Universal DNA Vaccine Platform to Target Antigen-Presenting Cells
Pirouz Daftarian, Angel E. Kaifer, Wei Li, Bonnie B. Blomberg, Daniela Frasca, Felix Roth, Raquibul Chowdhury, Eric A. Berg, Jordan B. Fishman, Husain A. Al Sayegh, Pat Blackwelder, Luca Inverardi, Victor L. Perez, Vance Lemmon, and Paolo Serafini

Précis: This article describes a new DNA vaccination technology that more efficiently targets dendritic cells in vivo, resulting in the production of high-affinity T cells that can more efficiently reject tumors.

PGE2-Induced CXCL12 Production and CXCR4 Expression Controls the Accumulation of Human MDSCs in Ovarian Cancer Environment
Nataša Obermajer, Ravikumar Muthuswamy, Kunle Odunsi, Robert P. Edwards, and Pawel Kalinski

Précis: This study provides a strong rationale for clinical testing of prostaglandin E2 antagonists, such as COX-2 inhibitors or EP2/EP4 blockers, in treatment of tumor ascites, a deadly feature of certain advanced cancers, including ovarian cancer.

PAX3-FoxO1 Induces Cannabinoid Receptor 1 to Enhance Cell Invasion and Metastasis
Amy D. Marshall, Irina Lagutina, and Gerard C. Grosveld

Précis: A targetable cell surface receptor is found to be essential for invasion and metastasis of a deadly type of muscle-derived pediatric tumor, with immediate implications to improve its treatment.
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PREVENTION AND EPIDEMIOLOGY

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Genome-Wide High-Density SNP Linkage Search for Glioma Susceptibility Loci: Results from the Gliogene Consortium

Precis: Linkage analysis of families with 2 or more cases of glioma identifies loci linked to glioma susceptibility, suggesting an inherited predisposition to development of this deadly cancer.

7576

Variations in HSPA1B at 6p21.3 Are Associated with Lung Cancer Risk and Prognosis in Chinese Populations
Huan Guo, Qifei Deng, Chen Wu, Lingmin Hu, Sheng Wei, Ping Xu, Dan Kuang, Li Liu, Zhibin Hu, Xiaoping Miao, Hongbing Shen, Dongxin Lin, and Tangchun Wu

Precis: Functional variants in a human member of the Hsp70 gene family are found to be significantly associated with lung cancer risk and patient survival, possibly offering new clinical biomarkers for risk and prognosis.

THERAPEUTICS, TARGETS, AND CHEMICAL BIOLOGY

7587

Glioblastoma-Derived Epidermal Growth Factor Receptor Carboxyl-Terminal Deletion Mutants Are Transforming and Are Sensitive to EGFR-Directed Therapies
Jeonghee Cho, Sandra Pastorino, Qing Zeng, Xiaoqin Xu, William Johnson, Scott Vandenberg, Roel Verhaak, Andrew D. Cherniack, Hideo Watanabe, Amit Dutt, Jihyun Kwon, Ying S. Chao, Robert C. Onofrio, Derek Chiang, Yuki Yuza, Santosh Kesari, and Matthew Meyerson

Precis: Genomic and functional analyses of primary gliomas reveal intragenic genomic deletions within the EGFR C-terminal domain that render the receptor oncogenic in vitro and in vivo.

7597

Dual IGF-1R/InsR Inhibitor BMS-754807 Synergizes with Hormonal Agents in Treatment of Estrogen-Dependent Breast Cancer
Xiaoman Hou, Fei Huang, Luciana F. Macedo, Sean C. Harrington, Karen A. Reeves, Ann Greer, Friedrich Graf Finckenstein, Angela Brodie, Marco M. Gottardis, Joan M. Carboni, and Paul Halaska

Precis: Findings offer preclinical proof-of-concept for a powerful new combinatorial approach to improve the eradication of estrogen-dependent breast cancers.

7608

miR-221 Silencing Blocks Hepatocellular Carcinoma and Promotes Survival
Jong-Kosk Park, Takayuki Kogure, Gerard J. Nuvoo, Jinmai Jiang, Lei He, Ji Hye Kim, Mitch A. Phelps, Tracey L. Papenfuss, Carlo M. Croce, Tushar Patel, and Thomas D. Schmittgen

Precis: In hepatocellular carcinomas with typically poor prognosis, antisense miR-221 treatments may be highly effective in blocking tumor proliferation.

7617

miR-221 Silencing Blocks Hepatocellular Carcinoma and Promotes Survival
Jong-Kosk Park, Takayuki Kogure, Gerard J. Nuvoo, Jinmai Jiang, Lei He, Ji Hye Kim, Mitch A. Phelps, Tracey L. Papenfuss, Carlo M. Croce, Tushar Patel, and Thomas D. Schmittgen

Precis: In hepatocellular carcinomas with typically poor prognosis, antisense miR-221 treatments may be highly effective in blocking tumor proliferation.

7628

Chemotherapeutic Properties of Phospho-Nonsteroidal Anti-Inflammatory Drugs, a New Class of Anticancer Compounds
Liqun Huang, Gerardo G. Mackenzie, Yu Sun, Nengtai Ouyang, Gang Xie, Kvetoslava Vrankova, Despina Komninou, and Basil Rigas

Precis: This important study addresses the quandary posed by NSAIDs, which exert useful anti-inflammatory effects but present safety and efficacy concerns that raise questions about their use in cancer prevention and treatment.

Wnt Inhibitor Screen Reveals Iron Dependence of β-Catenin Signaling in Cancers

Precis: In identifying a new class of compounds suitable for development as inhibitors of Wnt-driven cancers, this study led to the important discovery of a fundamental requirement for iron that might be immediately clinically exploitable.
SMAC Mimetic (JP1201) Sensitizes Non–Small Cell Lung Cancers to Multiple Chemotherapy Agents in an IAP-Dependent but TNF-α–Independent Manner
Rachel M. Greer, Michael Peyton, Jill E. Larsen, Luc Girard, Yang Xie, Adi F. Gazdar, Patrick Harran, Lai Wang, Rolf A. Brekken, Xiaodong Wang, and John D. Minna

Précis: Chemical mimetics of the proapoptotic molecule SMAC may be useful to broadly sensitize chemoresistant cancers to clinically relevant concentrations of various cytotoxic drugs, addressing an urgent need in oncology.

Thrombospondin-1 Triggers Cell Migration and Development of Advanced Prostate Tumors
Virginie Firlej, Jacques R.R. Mathieu, Christelle Gilbert, Loïc Lemonnier, Jessica Nakle, Catherine Gallou-Kabani, Basma Guarmit, Aurélie Morin, Natalia Prevarskaya, Nicolas Barry Delongchamps, and Florence Cabon

Précis: An extracellular protein known to be anti-angiogenic was found to stimulate invasion in prostate cancers, suggesting that disruption of this protein might actually confer therapeutic benefits.

Sorcin Induces a Drug-Resistant Phenotype in Human Colorectal Cancer by Modulating Ca2⁺ Homeostasis
Francesca Maddalena, Gabriella Laudiero, Annamaria Piscazzi, Agnese Secondo, Antonella Scorziello, Valentina Lombardi, Danilo Swann Matassa, Alberto Fersini, Vincenzo Neri, Franca Esposito, and Matteo Landriscina

Précis: Findings elucidate what appears to be a broadly active mechanism of multidrug resistance in human colorectal cancers, potentially responsible for much of the mortality in this common cancer in developed countries.

TUMOR AND STEM CELL BIOLOGY

Targets of the Tumor Suppressor miR-200 in Regulation of the Epithelial–Mesenchymal Transition in Cancer
Mark J. Schliekelman, Don L. Gibbons, Vitor M. Faca, Chad J. Creighton, Zain H. Rizvi, Qing Zhang, Chee-Hong Wong, Hong Wang, Christin Ungewiss, Young-Ho Ahn, Dong-Hoon Shin, Jonathan M. Kurie, and Samir M. Hanash

Précis: Findings offer insights into how a family of tumor suppressive microRNAs restricts metastatic progression, identifying key roles for a TGF-β protein network and extracellular cell adhesion molecules and proteases in the tumor microenvironment.

Letters to the Editor

STK33 Kinase Is Not Essential in KRAS-Dependent Cells—Letter
Stefan Frohling and Claudia Scholl

STK33 Kinase Is Not Essential in KRAS-Dependent Cells—Response
Isabelle Dussault, Josette Carnahan, Carol Babij, Yihong Zhang, Vivienne J. Watson, Kim Quon, and Paul D. Kassner

Correction

Correction: Definition of a FoxA1 Cistrome That Is Crucial for G1 to S-Phase Cell-Cycle Transit in Castration-Resistant Prostate Cancer
ABOUT THE COVER

Chronic inflammation plays important roles at different stages of cancer development, including carcinogenesis, tumor invasion, and metastasis. In this study, expression of angiopoietin-like protein 2 (Angptl2), recently identified as a chronic inflammation mediator in skin tissues, is highly correlated with the frequency of carcinogenesis in a chemically induced skin squamous cell carcinoma (SCC) mouse model. Furthermore, expression of Angptl2 in tumor cells promotes not only tumor angiogenesis and lymphangiogenesis, but also the acquisition of mesenchymal invasive characteristics in tumor cells through activation of TGF-β/Smad signaling, resulting in worse outcomes. This image represents a significantly increased tumor angiogenesis in SCC of Angptl2 transgenic (K14-Angptl2) mice. For details, see the article by Aoi and colleagues on page 7502 of this issue.
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