BREAKING ADVANCES

5363 Highlights from Recent Cancer Literature

REVIEWS

5365 Long Noncoding RNA, Polycomb, and the Ghosts Haunting INK4b-ARF-INK4a Expression
Francesca Aguilo, Ming-Ming Zhou, and Martin J. Walsh

5370 B-Myb, Cancer, Senescence, and MicroRNAs
Ivan Martinez and Daniel DiMaio

PRIORITY REPORTS

5374 Indirubins Decrease Glioma Invasion by Blocking Migratory Phenotypes in Both the Tumor and Stromal Endothelial Cell Compartments
Shanto P. Williams, Michal O. Nowicki, Fang Liu, Rachael Press, Jakub Godlewski, Mahmoud Abdel-Rasoul, Balveen Kaur, Soledad A. Fernandez, E. Antonio Chiocca, and Sean E. Lawler

Précis: Preclinical studies validate a novel class of small molecule inhibitors of the enzyme GSK-3, which exert potent antitumor properties by blocking both tumor invasion and angiogenesis.

5381 Imatinib Sensitivity in BCR-ABL1–Positive Chronic Myeloid Leukemia Cells Is Regulated by the Remaining Normal ABL1 Allele
Anna Virgili, Mateusz Kopytya, Yashodhara Dasgupta, Eliza Gadowska-Mrowka, Tomasz Stoklosa, Elisabeth P. Nacheva, and Tomasz Skorski

Précis: The lack of a complete cytogenetic response in chronic phase CML patients treated with the ABL kinase inhibitor imatinib can be explained by loss of the remaining normal ABL1 allele in CML cells.

INTEGRATED SYSTEMS AND TECHNOLOGIES

5400 Comparing Signaling Networks between Normal and Transformed Hepatocytes Using Discrete Logical Models

Précis: Findings illustrate a novel approach to creating cell-specific computational models of signaling networks based on biochemical data, applying it in this study to define deregulated pathways in hepatocellular carcinoma.
Profound Coordinated Alterations of Intratumoral NK Cell Phenotype and Function in Lung Carcinoma
Sophia Platonova, Julien Cherfils-Vicini, Diane Damotte, Lucile Crozet, Vincent Vieillard, Pierre Validire, Pascale André, Marie-Caroline Dieu-Nosjean, Marco Alifano, Jean-François Régnard, Wolf-Herman Fridman, Catherine Saulès-Fridman, and Isabelle Cremer

Precis: The tissue microenvironment in human lung carcinomas suppresses the tumoricidal activity of natural killer cells, thereby contributing to immune escape and progression.

Quantitative and Functional Alterations of Plasmacytoid Dendritic Cells Contribute to Immune Tolerance in Ovarian Cancer
Sana Intidhar Labidi-Galy, Vanja Sisirak, Pierre Meeus, Michael Gobert, Isabelle Treilleux, Agathe Bajard, Jean-Damien Combes, Alexandre Cassignol, Olivier Tredan, Isabelle Durand, Christine Ménétrier-Caux, Christophe Caux, Jean-Yves Blay, Isabelle Ray-Coquard, and Nathalie Bendriss-Vermare

Precis: Findings define a critical cellular mechanism of immune escape in human ovarian cancers, offering new insights into the pathophysiology of this often fatal cancer.

A Novel Tumor Antigen Derived from Enhanced Degradation of Bax Protein in Human Cancers
Claudia Trindade Nunes, Kelly L. Miners, Garry Dolton, Chris Pepper, Chris Fegan, Malcolm D. Mason, and Stephen Man

Precis: Study findings suggest that useful peptide antigens for cancer vaccination might be derived from proteins commonly expressed in normal cells but abnormally proteolyzed in cancer.

Comparison of Increased Aromatase versus ERα in the Generation of Mammary Hyperplasia and Cancer
Edgar S. Díaz-Cruz, Yasuro Sugimoto, G. Ian Gallicano, Robert W. Brueggemeier, and Priscilla A. Furth

Precis: Findings show that increased aromatase levels may play an even larger role in breast cancer progression than overexpression of estrogen receptor α.

Maximal T Cell–Mediated Antitumor Responses Rely upon CCR5 Expression in Both CD4+ and CD8+ T Cells
Alicia González-Martín, Lucio Gómez, Joseph Lastgarten, Emilia Mira, and Santos Mañes

Precis: A chemokine receptor that is critical to organize leukocyte trafficking responses to infection is also found to be critical for T cell-mediated responses against tumors, suggesting mechanistic similarities in the way that the immune system interprets cancer cells and infectious pathogens.

TLR4 Engagement during TLR3-Induced Proinflammatory Signaling in Dendritic Cells Promotes IL-10–Mediated Suppression of Antitumor Immunity
Dusan Bogunovic, Olivier Manches, Emmanuelle Godfroy, Alice Yewdall, Anne Gallois, Andres M. Salazar, Isabelle Marie, David E. Levy, and Nina Bhardwaj

Precis: Findings suggest that the antitumor properties of a potent immune stimulatory Toll receptor ligand with therapeutic potential may be reduced or negated if the Toll receptor TLR4 is also activated, with implications for the design of immunotherapy trials.
**THERAPEUTICS, TARGETS, AND CHEMICAL BIOLOGY**

**CXCL12/CXCR4 Blockade Induces Multimodal Antitumor Effects That Prolong Survival in an Immunocompetent Mouse Model of Ovarian Cancer**

Elda Righi, Satoshi Kashiwagi, Jianping Yuan, Michael Santosuosso, Pierre Leblanc, Rachel Ingraham, Benjamin Forbes, Beth Edelblute, Brian Collette, Deyin Xing, Magdalena Kowalski, Maria Cristina Mingari, Fabrizio Vianello, Michael Birrer, Sandra Orsulic, Glenn Tranoff, and Mark C. Poznansky

**Précis:** A druggable target implicated in promoting metastasis is found to coordinate immune escape, illustrating the important linkage between these two processes in ovarian cancer progression.

**Identification of the MEK1(F129L) Activating Mutation as a Potential Mechanism of Acquired Resistance to MEK Inhibition in Human Cancers Carrying the B-RafV600E Mutation**

Huisheng Wang, Sherif Daouti, Wen-hui Li, Yang Wen, Christine Rizzo, Brian Higgins, Kathryn Packman, Neal Rosen, John F. Boylan, David Heimbrook, and Huifeng Niu

**Précis:** Combined inhibition of Raf and MEK may offer a clinical strategy to bypass or overcome acquired resistance to MEK inhibitors that can arise as the result of a powerful activating mutation in MEK1.

**GPR56 Regulates VEGF Production and Angiogenesis during Melanoma Progression**

Liquan Yang, Guangchun Chen, Sonali Mohanty, Glynnis Scott, Fabeha Fa zal, Arshad Rahman, Shahinooor Begum, Richard O. Hynes, and Lei Xu

**Précis:** Findings identify a novel G protein-coupled receptor that regulates VEGF production, offering a new therapeutic target for angiogenesis inhibition.

**YB-1 Bridges Neural Stem Cells and Brain Tumor–Initiating Cells via Its Roles in Differentiation and Cell Growth**


**Précis:** A transcription factor required for embryonic brain development also contributes in later life to brain tumor development, due to its roles in normal and malignant neural stem cells.
Protein Arginine Methyltransferase 5 Accelerates Tumor Growth by Arginine Methylation of the Tumor Suppressor Programmed Cell Death 4
Matthew A. Powers, Marta M. Fay, Rachel E. Factor, Alana L. Welm, and Katharine S. Ullman

Précis: This article reports a new regulatory node in cancer in which a protein methyltransferase works in conjunction with the tumor suppressor PDCD4 to cause accelerated tumor growth.

p53-Dependent Regulation of Mitochondrial Energy Production by the RelA Subunit of NF-κB
Renée F. Johnson, Ini-Ishabelle Witzel, and Neil D. Perkins

Précis: This study defines an important new link in the control of mitochondrial function by oncogenes that influence cellular metabolism.

ABOUT THE COVER

The Y-box binding protein (YB-1) is an oncogenic transcription factor known for its ability to cause drug resistance and cancer recurrence. Fotovati and colleagues report that YB-1 supports brain tumor–initiating cells by inhibiting differentiation through the maintenance of proteins associated with stem cells. In cancer-derived neurospheres grown from pediatric glioblastoma multiforme cells, YB-1 was highly expressed along with the stem cell markers nestin and Bmi-1. For details, see the article by Fotovati and colleagues on page 5569 of this issue.
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