Hedgehog Fights Back: Mechanisms of Acquired Resistance against Smoothened Antagonists
Ciara Metcalfe and Frederic J. de Sauvage

Chemotherapeutic Resistance: Surviving Stressful Situations
Luke A. Gilbert and Michael T. Hemann

Combinatorial Treatments That Overcome PDGFRβ-Driven Resistance of Melanoma Cells to V600E-RAF Inhibition
Hubing Shi, Xiangju Kong, Antoni Ribas, and Roger S. Lo

Mechanical Stiffness Grades Metastatic Potential in Patient Tumor Cells and in Cancer Cell Lines

Keap1 Mutations and Nrf2 Pathway Activation in Epithelial Ovarian Cancer
Panagiotis A. Konstantinopoulos, Dimitrios Spentzos, Elena Fountzilas, Nancy Francoeur, Srisowmya Sanisetty, Alexandros P. Grammatikos, Jonathan L. Hecht, and Stephen A. Cannistra

Tumor Microenvironment–Derived Proteins Dominate the Plasma Proteome Response during Breast Cancer Induction and Progression
Sharon J. Pitteri, Karen S. Kelly-Spratt, Kay E. Gurley, Jacob Kennedy, Tina Busald Buson, Alice Chin, Hong Wang, Qing Zhang, Chee-Hong Wong, Lewis A. Chodosh, Peter S. Nelson, Samir M. Hanash, and Christopher J. Kemp

Myeloid-Derived Suppressor Cell Inhibition of the IFN Response in Tumor-Bearing Mice
PD-1/PD-L1 Interactions Contribute to Functional T-Cell Impairment in Patients Who Relapse with Cancer After Allogeneic Stem Cell Transplantation

Précis: Findings show how leukemia cells can escape from graft-versus-tumor immune responses, and how interfering with this escape mechanism can restore antitumor immunity and possibly prevent relapses.

Systemic Cancer Therapy with a Small Molecule Agonist of Toll-like Receptor 7 Can Be Improved by Circumventing TLR Tolerance
Carole Bourquin, Christian Hotz, Daniel Noerenberg, Andreas Voelkl, Simon Heidegger, Bettina Storch, Nadja Sandholzer, Cornelia Wurzenberger, David Anz, and Stefan Endres

Précis: Findings offer insight into a cellular mechanism of immune tolerance that occurs during cancer immunotherapy with a Toll-like receptor agonist and provide a strategy to bypass this tolerance.

Antibody-Dependent Cell Cytotoxicity Synapses Form in Mice during Tumor-Specific Antibody Immunotherapy
Pascale Hubert, Adèle Heitzmann, Sophie Viel, André Nicolas, Xavier Sastré-Garau, Pablo Oppezzo, Otto Pritsch, Eduardo Osinaga, and Sebastien Amigorena

Précis: Findings demonstrate that a therapeutic monoclonal antibody specific for a ubiquitous tumor-associated antigen inhibits tumor growth via antibody-dependent cell toxicity.

IGFBP-3 Is a Metastasis Suppression Gene in Prostate Cancer

Précis: This study reports the first transgenic mouse model of spontaneous metastatic prostate cancer, a milestone that may help advance studies of progression and treatment at this deadly late stage of disease.

MOLECULAR AND CELLULAR PATHOBIOLOGY

Combining Betulinic Acid and Mithramycin A Effectively Suppresses Pancreatic Cancer by Inhibiting Proliferation, Invasion, and Angiogenesis
Yong Gao, Zhiliang Jia, Xiangyu Kong, Qiang Li, David Z. Chang, Daoyan Wei, Xiangdong Le, Shengdong Huang, Liwei Wang, Suyun Huang, and Keping Xie

Précis: Findings demonstrate that targeting the transcription factor Sp1 can inhibit pancreatic tumor growth with greater efficacy and fewer side effects than traditional treatment.

IGFBP-3 Is a Metastasis Suppression Gene in Prostate Cancer

Précis: This study reports the first transgenic mouse model of spontaneous metastatic prostate cancer, a milestone that may help advance studies of progression and treatment at this deadly late stage of disease.

THERAPEUTICS, TARGETS, AND CHEMICAL BIOLOGY

Inhibition of miR-193a Expression by Max and RXRa Activates K-Ras and PLAU to Mediate Distinct Aspects of Cellular Transformation
Dimitrios Ilipoulos, Asaf Rotem, and Kevin Struhl

Précis: Findings describe a molecular pathway involved in cellular transformation that focuses on miR-193a as a regulator of both tumorigenicity and invasiveness.
BMP4 Promotes Prostate Tumor Growth in Bone through Osteogenesis
Yu-Chen Lee, Chien-Jui Cheng, Mehmet A. Bilen, Jing-Fang Lu, Robert L. Satcher, Li-Yuan Yu-Lee, Gary E. Gallick, Sankar N. Maity, and Sue-Hwa Lin

Précis: Osteogenesis is shown to be necessary for prostate cancer progression in bone and inhibition of this process may offer a treatment strategy for bone metastases.

Akt-Dependent Glucose Metabolism Promotes Mcl-1 Synthesis to Maintain Cell Survival and Resistance to Bcl-2 Inhibition
Jonathan L. Coloff, Andrew N. Macintyre, Amanda G. Nichols, Tingyu Liu, Catherine A. Gallo, David R. Plas, and Jeffrey C. Rathmell

Précis: Inhibition of glucose metabolism alters expression of a key Bcl-2 family member, sensitizing cancer cells to apoptosis and illustrating how strategies to disrupt glycolysis disruptions may disable tumors.

MicroRNA Replacement Therapy for miR-145 and miR-33a Is Efficacious in a Model of Colon Carcinoma
Ahmed Fawzy Ibrahim, Ulrike Weirauch, Maren Thomas, Arnold Grünweller, Roland K. Hartmann, and Achim Aigner

Précis: This study establishes nonviral miRNA replacement therapy for cancer treatment.

Combined Gene Expression Profiling and RNAi Screening in Clear Cell Renal Cell Carcinoma Identify PLK1 and Other Therapeutic Kinase Targets
Yan Ding, Dan Huang, Zhongla Zhang, Josh Smith, David Petillo, Brendan D. Looyenga, Kristin Feenstra, Jeffrey P. MacKeigan, Kyle A. Furge, and Bin T. Teh

Précis: RNAi screening and gene expression profiling reveal new potential therapeutic targets to treat highly aggressive kidney cancers.

Exosome Targeting of Tumor Antigens Expressed by Cancer Vaccines Can Improve Antigen Immunogenicity and Therapeutic Efficacy
Ryan B. Bountree, Stefanie J. Mandl, James M. Nachtwey, Katie Dalpozzo, Lisa Do, John R. Lombardo, Peter L. Schoonmaker, Kay Brinkmann, Ulrike Dirmeier, Reiner Laut, and Alain Delcayre

Précis: This study shows how engineering the MVA-BN-PRO vaccine to express its antigens in secreted exosomes could greatly improve its immunotherapeutic activity.

Retinoic Acid Enhances TRAIL-Induced Apoptosis in Cancer Cells by Upregulating TRAIL Receptor 1 Expression
Latha Dhandapani, Ping Yue, Suresh S. Ramalingam, Fadlo R. Khuri, and Shi-Yong Sun

Précis: By establishing that retinoic acid can stimulate the proapoptotic activity of TRAIL by upregulating the TRAIL receptor DR4, this study suggests a strategy to augment the anticancer activity of TRAIL being studied in clinical trials.

A Drug Resistance Screen Using a Selective MET Inhibitor Reveals a Spectrum of Mutations That Partially Overlap with Activating Mutations Found in Cancer Patients
Ralph Tiedt, Elisa Degenkolbe, Pascal Furet, Brent A. Appleton, Sabrina Wagner, Joseph Schoepfer, Emily Buck, David A. Buddy, John E. Monahan, Michael D. Jones, Jutta Blank, Dorothea Haasen, Peter Drueckes, Markus Wartmann, Clive McCarthy, William R. Sellers, and Francesco Hofmann

Précis: Drug resistance mutation profiles reported in this manuscript are likely to predict clinical responses to MET kinase inhibitors, a number of which are currently in development.

Overcoming Hypoxia-Induced Apoptotic Resistance through Combinatorial Inhibition of GSK-3β and CDK1

Précis: Findings elucidate a novel therapeutic strategy to overcome the inherent resistance of hypoxic cancer cells to cell death, by targeting two kinases not previously linked to cell death protection in hypoxic tumor microenvironments.
TUMOR AND STEM CELL BIOLOGY

5276 Interleukin-1α Mediates the Antiproliferative Effects of 1,25-Dihydroxyvitamin D₃ in Prostate Progenitor/Stem Cells
Sophia L. Maund, Wendy W. Barclay, Laura D. Hover, Linara S. Axanova, Guangchao Sui, Jason D. Hipp, James C. Fleet, Andrew Thorburn, and Scott D. Cramer

Précis: Supporting applications of vitamin D as a chemopreventative agent for prostate cancer, this study shows that the metabolically active form of vitamin D can induce differentiation and senescence of prostate progenitor/stem cells and that its antiproliferative effects rely upon interleukin-1 alpha.

5287 PTEN Positively Regulates UVB-Induced DNA Damage Repair
Mei Ming, Li Feng, Christopher R. Shea, Keyoumars Soltani, Baozhong Zhao, Weinong Han, Robert C. Smart, Carol S. Trembus, and Yu-Ying He

Précis: Findings explain how failure to repair DNA damage caused by UVB sunlight radiation can cause skin carcinogenesis, due to inactivation of the tumor suppressor PTEN that destroys its key gatekeeper function in supporting DNA repair in the skin.

5296 IL-8 Signaling Plays a Critical Role in the Epithelial–Mesenchymal Transition of Human Carcinoma Cells
Romaine I. Fernando, Marianne D. Castillo, Mary Litzinger, Duane H. Hamilton, and Claudia Palena

Précis: Findings elucidate the role of epithelial-to-mesenchymal transition in the modulation of the tumor microenvironment, suggesting that IL-8 signaling blockades might be very effective at targeting invasive tumor cells.

5307 ABCB5 Identifies a Therapy-Refractory Tumor Cell Population in Colorectal Cancer Patients
Brian J. Wilson, Tobias Schatton, Qian Zhan, Martin Gasser, Jie Ma, Karim R. Saab, Robin Schanche, Ana-Maria Waaga-Gasser, Jason S. Gold, Qin Huang, George F. Murphy, Markus H. Frank, and Natasha Y. Frank

Précis: Findings point to the need to eradicate a particular tumor cell population to improve outcomes in colorectal cancer therapy.

5317 Cancer Stem Cells in Squamous Cell Carcinoma Switch between Two Distinct Phenotypes That Are Preferentially Migratory or Proliferative
Adrian Biddle, Xiao Liang, Luke Gammon, Bilal Fazil, Lisa J. Harper, Helena Emich, Daniela Elena Costea, and Ian C. Mackenzie

Précis: Findings suggest that cancer stem cells can switch their phenotype between two states that can either drive tumor cell proliferation or metastatic dissemination, implying a need for therapeutic approaches that are able to eradicate cancer stem cells in both states.

5327 ADP-Ribosylarginine Hydrolase Regulates Cell Proliferation and Tumorigenesis
Jiro Kato, Jianfeng Zhu, Chengyu Liu, Mario Stylianou, Victoria Hoffmann, Martin J. Lizak, Connie G. Glasgow, and Joel Moss

Précis: Findings point to an important role for posttranslational protein modification by ADP-ribosylation in supporting cell proliferation and tumorigenesis.

5336 Sequential Activation of Snail1 and N-Myc Modulates Sonic Hedgehog–Induced Transformation of Neural Cells
Leah E. Colvin Wanshura, Katherine E. Galvin, Hong Ye, Martin E. Fernandez-Zapico, and Cynthia Wetmore

Précis: N-Myc activation by a key target of the Sonic Hedgehog signaling pathway may be an essential step information of an aggressive class of pediatric brain tumors, with implications for therapeutic targeting strategies.

5346 Microvesicles Released from Human Renal Cancer Stem Cells Stimulate Angiogenesis and Formation of Lung Premetastatic Niche
Cristina Grange, Marta Tapparo, Federica Collino, Lorian Vitillo, Christian Damasco, Maria Chiara Deregibus, Ciro Tetta, Benedetta Bussolati, and Giovanni Camussi

Précis: Cancer stem cells may promote metastatic progression by secreting a class of microvesicles known as exosomes that can transfer proangiogenic RNAs to endothelial cells and directly stimulate angiogenesis.
LETTERS TO THE EDITOR

5357 MicroRNA Expression and Outcome in Resected NSCLC—Letter
Shun-ichiro Kageyama, Yusuke Takagi, Takeshi Sawada, Natsuko Kageyama-Yahara, and Masahiko Shibuya

5358 MicroRNA Expression and Outcome in Resected NSCLC—Response
Johannes Voortman, Aaron J. Schetter, Curtis C. Harris, and Giuseppe Giaccone

CORRECTIONS

5360 Correction: Effects of Carbon Ion Beam on Putative Colon Cancer Stem Cells and Its Comparison with X-rays

5361 Correction: A Requirement of STAT3 DNA Binding Precludes Th-1 Immunostimulatory Gene Expression by NF-κB in Tumors

ABOUT THE COVER

The search for biomarkers of cancer has focused on the tumor cells themselves. Pitteri and colleagues, using an unbiased and in depth proteomic analysis of plasma from a model of Her2/neu driven breast cancer, have identified the signaling between the tumor cells and microenvironment as a primary source of biomarkers. Shown are tumor cells embedded within extensive extracellular collagen matrix stained with trichrome blue. For details, see the article by Pitteri and colleagues on page 5090 of this issue.

Updated version
Access the most recent version of this article at:
http://cancerres.aacrjournals.org/content/71/15

E-mail alerts
Sign up to receive free email-alerts related to this article or journal.

Reprints and Subscriptions
To order reprints of this article or to subscribe to the journal, contact the AACR Publications Department at pubs@aacr.org.

Permissions
To request permission to re-use all or part of this article, contact the AACR Publications Department at permissions@aacr.org.