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913  “(Not) All (Dead) Things Share the Same Breath”: Identification of Cell Death Mechanisms in Anticancer Therapy
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930  Emergence of Drug Tolerance in Cancer Cell Populations: An Evolutionary Outcome of Selection, Nongenetic Instability, and Stress-Induced Adaptation
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940  Cell Division Patterns in Acute Myeloid Leukemia Stem-like Cells Determine Clinical Course: A Model to Predict Patient Survival
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950  CSF1 Receptor Targeting in Prostate Cancer Reverses Macrophage-Mediated Resistance to Androgen Blockade Therapy
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Tumor-Derived Osteopontin Reprograms Normal Mammary Fibroblasts to Promote Inflammation and Tumor Growth in Breast Cancer
Yoray Sharon, Yael Raz, Noam Cohen, Amir Ben-Shmuel, Hila Schwartz, Tamar Geiger, and Neta Erez

Précis: These findings deepen the causative influence of a proinflammatory protein secreted by breast carcinoma cells in programming a supportive role for fibroblasts in the tumor microenvironment, with implications for understanding the etiology of advanced cancer and its therapeutic management by renormalization of the tissue microenvironment.

RAGE Mediates S100A7-Induced Breast Cancer Growth and Metastasis by Modulating the Tumor Microenvironment
Mohd W. Nasser, Nissar Ahmad Wani, Dinesh K. Ahirwar, Catherine A. Powell, Janani Ravi, Mohamad Elbaz, Helong Zhao, Laura Padilla, Xiaoli Zhang, Konstantin Shilo, Michael Ostrowski, Charles Shapiro, William E. Carson III, and Ramesh K. Ganju

Précis: A signaling axis that conditions the inflammatory microenvironment in breast cancer helps drive the aggressive growth of these tumors, including deadly triple-negative tumors that occur more commonly in premenopausal women.

TLR2 Limits Development of Hepatocellular Carcinoma by Reducing IL18-Mediated Immunosuppression
Shinan Li, Rui Sun, Yongyan Chen, Haiming Wei, and Zhigang Tian

Précis: These findings illuminate a mechanism of immunosuppression in liver carcinogenesis that may assist the design of effective immunotherapies to treat hepatocellular carcinoma.

Carbonic Anhydrase IX Promotes Myeloid-Derived Suppressor Cell Mobilization and Establishment of a Metastatic Niche by Stimulating G-CSF Production
Shawn C. Chafe, Yuanmei Lou, Jaclyn Seneay, Marylou Vallejo, Melissa J. Hamilton, Paul C. McDonald, Kevin L. Bennewith, Andreas Möller, and Shoukat Dedhar

Précis: Targeting the function of CAIX, an enzyme that is upregulated by hypoxia in the primary tumor, affects the mobilization of immunosuppressive myeloid cells that promote metastasis.
1056 Starvation Promotes REV1 SUMOylation and p53-Dependent Sensitization of Melanoma and Breast Cancer Cells
Hong Seok Shim, Min Wei, Sebastian Brandhorst, and Valter D. Longo
Précis: These findings suggest how dietary fasting may offer a nontoxic strategy to increase the efficacy of cytotoxic therapies that act in part by activating p53.

1068 Tumorigenic Activity of Merkel Cell Polyomavirus T Antigens Expressed in the Stratified Epithelium of Mice
Megan E. Spurgeon, Jingwei Cheng, Roderick T. Bronson, Paul F. Lambert, and James A. DeCaprio
Précis: Use of a new mouse model of Merkel cell polyomavirus-associated tumorigenesis provides deeper insights into how viral tumor antigens alter the cellular microenvironment in vivo, with potential relevance to various human cancers that involve virus infection.

THERAPEUTICS, TARGETS, AND CHEMICAL BIOLOGY

1080 Targeting LINX Inhibits Non–Small Cell Lung Cancer Growth and Metastasis
Xiaohu Zheng, Min Cheng, Binqing Fu, Xiaolei Fan, Qing Wang, Xiaoqing Yu, Rui Sun, Zhigang Tian, and Haiming Wei
Précis: These results offer preclinical proof of concept for a candidate immunotherapy target in non–small cell lung cancers.

1091 Genetic Events That Limit the Efficacy of MEK and RTK Inhibitor Therapies in a Mouse Model of KRAS-Driven Pancreatic Cancer
Précis: Understanding the activation status of various oncogenic drivers as they exist in specific treatment contexts in vivo may be important to achieve beneficial outcomes, increasing the complexity in how to use targeted drugs that were designed only with cancer cells in mind.

1102 Genetic and Pharmacologic Inhibition of eIF4E Reduces Breast Cancer Cell Migration, Invasion, and Metastasis
Filippa Pettersson, Sonata V. del Rincon, Audrey Emond, Bonnie Huo, Elaine Ngan, Jonathan Ng, Monica C. Dobocan, Peter M. Siegel, and Wilson H. Miller Jr
Précis: These findings offer a powerful rationale to broaden the clinical evaluation of ribavirin, a small molecule inhibitor of the translation initiation factor eIF4E currently being tested in leukemia patients, as a strategy to treat advanced solid tumors such as metastatic breast cancer in which eIF4E is commonly overexpressed.

TUMOR AND STEM CELL BIOLOGY

1113 Constitutive Activation of Myosin-Dependent Contractility Sensitizes Glioma Tumor-Initiating Cells to Mechanical Inputs and Reduces Tissue Invasion
Sophie Y. Wong, Theresa A. Ulrich, Loic P. Deleyrolle, Joanna L. MacKay, Jung-Ming G. Lin, Regina T. Martuscello, Musa A. Jundi, Brent A. Reynolds, and Sanjay Kumar
Précis: Because recurrences of brain cancer are tied to local invasion of tumor cells, strategies to restrict the motility of stem-like cells by increasing their cellular contractility may help limit relapses and prolong survival.

1123 miR340 Suppresses the Stem-like Cell Function of Glioma-Initiating Cells by Targeting Tissue Plasminogen Activator
Daisuke Yamashita, Toru Kondo, Shiro Ohue, Hisaaki Takahashi, Madoka Ishikawa, Ryo Matoba, Satoshi Suehiro, Shuhei Kohno, Hironobu Harada, Junya Tanaka, and Takamori Ohnishi
Précis: A tumor suppressor gene that functions in glioma stem-like cells acts to inhibit their expression of tissue plasminogen activator, with the provocative implication that targeting this central coagulation factor may ablate cancer stem-like functions in the brain.

1134 Host Age Is a Systemic Regulator of Gene Expression Impacting Cancer Progression
Afshin Beheshti, Sébastien Benzekry, J. Tyson McDonald, Lili Ma, Michael Peluso, Philip Hahnfeldt, and Lynn Hlatky
Précis: This study offers direct support for age dependence in determining the host tumor control dynamic and provides initial mechanism-based insights into how aging modulates tumor progression in ways that may be actionable for therapy or prevention.

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Haksier Ehedego, Mark V. Boekschoten, Wei Hu, Carina Doler, Johannes Haybaeck, Nikolaus Gaßler, Michael Müller, Christian Liedke, and Christian Trautwein

Précis: These findings illuminate the ways in which loss of the NF-κB pathway regulator NEMO (also known as IKKγ) promotes liver inflammation and carcinogenesis by elevating p21, which acts in this setting to protect against the generation of DNA damage that contributes to chronic hepatitis and hepatocarcinoma formation in patients.

LETTERS TO THE EDITOR

1156 Bufalin Is a Steroid Receptor Coactivator Inhibitor—Letter
José Manuel Calderón-Montaño, Estefanía Burgos-Morón, Manuel Luis Orta, Irene García-Domínguez, Dolores Maldonado-Navas, and Miguel López-Lázaro

1157 Bufalin Is a Steroid Receptor Coactivator Inhibitor—Response
David M. Lonard, Jianming Xu, and Bert W. O’Malley

ABOUT THE COVER

miR340 is a tumor suppressor whose overexpression in human glioma-initiating cells (GIC) inhibits their proliferation, invasive, and migratory properties. Transplantation of miR340-overexpressed GICs in NOD/SCID mouse brains completely suppressed the tumor formation of malignant gliomas. Among factors related to antitumorigenesis, the transplanted GICs with miR340 overexpression showed a positive immunostain for an active form of caspase-3 in the early stage of transplantation, suggesting that antitumorigenic activity of miR340 in GICs may be due to tumor cell apoptosis. For details, see the article by Yamashita and colleagues on page 1123.
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