 Highlights from Recent Cancer Literature

MAX and MYC: A Heritable Breakup
Alberto Cascón and Mercedes Robledo

Cellular Constituents of Immune Escape within the Tumor Microenvironment
Sid P. Kerkar and Nicholas P. Restifo

The Continuum Model of Selection in Human Tumors: General Paradigm or Niche Product?
Simon Leedham and Ian Tomlinson

Stat3 Activation in Urothelial Stem Cells Leads to Direct Progression to Invasive Bladder Cancer
Philip Levy Ho, Erica Julienne Lay, Weiguo Jian, Diana Parra, and Keith Syson Chan

Antigen Shedding May Improve Efficiencies for Delivery of Antibody-Based Anticancer Agents in Solid Tumors
Youngshang Pak, Yujian Zhang, Ira Pastan, and Byungkook Lee

Breast Cancer Cell Uptake of the Inflammatory Mediator Neutrophil Elastase Triggers an Anticancer Adaptive Immune Response
Elizabeth A. Mittendorf, Gheath Alatrash, Na Qiao, Yun Wu, Pariya Sukhumalchandra, Lisa S. St. John, Anne V. Philips, Haile Xiao, Mao Zhang, Kathryn Ruisaard, Karen Clise-Dwyer, Sijie Lu, and Jeffrey J. Moldrem
Précis: Findings reveal an intriguing mechanism through which tumor-associated neutrophils present in the tumor microenvironment can trigger an antigen-specific cytolytic T-cell response against the malignant cells present there.

Radiotherapy Increases the Permissiveness of Established Mammary Tumors to Rejection by Immunomodulatory Antibodies
Inge Verbrugge, Jim Hagekyriakou, Leslie L. Sharp, Mara Galli, Alison West, Nicole M. McLaughlin, Hélène Duret, Hideo Yagita, Ricky W. Johnstone, Mark J. Smyth, and Nicole M. Haynes
Précis: Findings show how systemically delivered monoclonal antibodies that can concomitantly stimulate antitumor immunity and degrade tumoral immunosuppression can act as powerful adjuvants to improve the efficacy of cancer radiotherapy, with immediate translational potential as immunoradiotherapy.

CXCL10 Promotes Osteolytic Bone Metastasis by Enhancing Cancer Outgrowth and Osteoclastogenesis
Jong-Ho Lee, Ha-Neui Kim, Kyung-Os Kim, Won Jong Jin, Seungbok Lee, Hong-Hee Kim, Hyunil Ha, and Zang Hee Lee
Précis: Findings advance understanding of the microenvironmental contributions in the bone that permit the development of osteolytic bone metastasis, with implications for treatment of this common aspect of malignant progression in some common cancers.

Précis: This study challenges our expectations by presenting a mathematical model suggesting that antigen shedding will enhance rather than inhibit the delivery of recombinant immunotoxins.
MOLECULAR AND CELLULAR PATHOLOGY

3207 Induction of the RNA Regulator LIN28A Is Required for the Growth and Pathogenesis of RESTless Breast Tumors
Kearney T.W. Gunsalus, Matthew P. Wagoner, Kassandra Meyer, Wyatt B. Potter, Barry Schoenike, Soyoung Kim, Caroline M. Alexander, Andreas Friedl, and Avtar Roopra

3208 Precis: This study offers pivotal mechanistic insights into how the loss of a transcriptional repressor that occurs frequently in breast cancer leads to an increase in tumor growth.

3217 Histone Lysine Methyltransferase SETD8 Promotes Carcinogenesis by Deregulating PCNA Expression

3218 Loss of Fibroblast HIF-1α Accelerates Tumorigenesis
Jung-whan Kim, Colin Evans, Alexander Weidemann, Norihiko Takeda, Yun Sok Lee, Christian Stockmann, Cristina Branco-Price, Filip Brandberg, Gustavo Leone, Michael C. Ostrowski, and Randall S. Johnson

3228 MEK Inhibition Leads to PI3K/AKT Activation by Relieving a Negative Feedback on ERBB Receptors
Alexa B. Turke, Youngchul Song, Carlotta Costa, Rebecca Cook, Carlos L. Arteaga, John M. Asara, and Jeffrey A. Engelman

3238 Combination Therapy with HSP90 Inhibitor 17-DMAG Reconditions the Tumor Microenvironment to Improve Recruitment of Therapeutic T cells
Aparna Rao, Jennifer L. Taylor, Nina Chi-Sabins, Mayumi Kawabe, William E. Gooding, and Walter J. Storkus

3239 Precis: Used in combinational studies for immunochemotherapy, a chaperone small-molecule inhibitor might generally heighten antitumor responses mediated by specific CD8-positive T cells.

3251 Chk2 Phosphorylation of Survivin-ΔEx3 Contributes to a DNA Damage-Sensing Checkpoint in Cancer
Qi Lv, Wei Wang, Jianfei Xue, Fang Hua, Rong Mu, Heng Lin, Jun Yan, Xiaoxi Lv, Xiaoguang Chen, and Zhuo-Wei Hu

3260 Oncogenic PI3K Mutations Lead to NF-κB–Dependent Cytokine Expression following Growth Factor Deprivation
Jessica E. Showe, and Dario C. Altieri

3267 Deregulating PCNA Expression

3270 Precis: This important study furthers a deepening trend in the field by enhancing our understanding of how the most common mutations in cancer not only exert autonomous effects in cancer cells but also powerfully influence the local tumor microenvironment to support the outgrowth of these cells.

3272 TMEM16A Induces MAPK and Contributes Directly to Tumorigenesis and Cancer Progression
Umamaheswar Duruvuri, Daniel J. ShiWARDS, Dong Xiao, Carol Bertrand, Xin Huang, Robert S. Edinger, Jason R. Rock, Brian D. Harfe, Brian J. Henson, Karl Kunzelmann, Rainer Schreiber, Raja S. Seethala, Ann Marie Egloff, Xing Chen, Vivian W. Lui, Jennifer R. Grandis, and Susanne M. Gollin

Precis: A calcium-activated chloride channel is shown to promote tumorigenesis in head and neck cancers via signaling through the Rac/Rho/MEK pathway and therefore may represent a novel therapeutic target.
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<td>3282</td>
<td>Increased Caffeine Intake Is Associated with Reduced Risk of Basal Cell Carcinoma of the Skin</td>
<td>Fengju Song, Abrar A. Qureshi, and Jiali Han</td>
<td>Précis: This prospective analysis shows an inverse association between caffeine intake and the risk of basal cell carcinoma in 2 large national cohorts.</td>
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<td>3290</td>
<td>Cancer Cell Secretion of the DAMP Protein HMGB1 Supports Progression in Malignant Mesothelioma</td>
<td>Sandro Jube, Zeyana S. Rivera, Marco E. Bianchi, Amy Powers, Ena Wang, Ian Pagano, Harvey I. Pass, Giovanni Gaudino, Michele Carbone, and Haining Yang</td>
<td>Précis: This study offers a preclinical proof-of-principle that therapies aimed at blocking HMGB1-driven inflammation may be beneficial to patients with mesothelioma, especially at early stages of tumor growth.</td>
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<td>3312</td>
<td>Identification of Anaplastic Lymphoma Kinase as a Potential Therapeutic Target in Ovarian Cancer</td>
<td>Hong Ren, Zhi-Ping Tan, Xin Zhu, Katherine Crosby, Herbert Haack, Jian-Min Ren, Sean Beausoleil, Albrecht Moritz, Gregory Innocenti, John Rush, Yi Zhang, Xin-Min Zhou, Ting-Lei Gu, Yi-Feng Yang, and Michael J. Comb</td>
<td>Précis: Findings suggest the use of ALK kinase inhibitors now in clinical development as potential agents for treatment of some patients with serous ovarian carcinoma or stromal sarcoma.</td>
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<td>3324</td>
<td>Antibody Targeting of Cell-Bound MUC1 SEA Domain Kills Tumor Cells</td>
<td>Edward Pichinuk, Itai Benhar, Oded Jacob, Michael Chalik, Lotem Weiss, Ravit Ziv, Carolyn Sympon, Amolkumar Karwa, Nechama I. Smorodinsky, Daniel B. Rubinstein, and Daniel H. Wreschner</td>
<td>Précis: Findings explore the development of a universal class of anticancer antibodies, showing high-affinity binding and potent killing of cancer and cancer stem cells that highly express a common cancer antigen.</td>
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<td>3337</td>
<td>Glucose-Regulated Protein 78 Controls Cross-talk between Apoptosis and Autophagy to Determine Antiestrogen Responsiveness</td>
<td>Katherine L. Cook, Ayesha N. Shajahan, Anni Wärri, Lu Jin, Leena A. Hilakivi-Clarke, and Robert Clarke</td>
<td>Précis: Expression of an endoplasmic reticulum-located chaperone protein is shown to play a role in the development of acquired ant{&quot;\text{&quot;\text{e}}}str{&quot;\text{&quot;\text{g}}}}en resistance, suggesting that the unfolded protein response may be an adaptation to stress in estrogen receptor-positive breast cancer.</td>
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<td>3350</td>
<td>Sensitivity of Glioblastomas to Clinically Available MEK Inhibitors Is Defined by Neurofibromin 1 Deficiency</td>
<td>Wendy L. See, I-Li Tan, Joydeep Mukherjee, Theodore Nicolaides, and Russell O. Pieper</td>
<td>Précis: This study offers immediate implications for stratification of brain cancer patients who might be treated with MEK inhibitors.</td>
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<td>3360</td>
<td>The Nedd8-Activating Enzyme Inhibitor MLN4924 Induces Autophagy and Apoptosis to Suppress Liver Cancer Cell Growth</td>
<td>Zhongguang Luo, Guangyang Yu, Hyuk Woo Lee, Libui Li, Lingyan Wang, Dongjin Yang, Yongfu Pan, Chan Ding, Jing Qian, Lijun Wu, Yiwei Chu, Jing Yi, Xiangdong Wang, Yi Sun, Lak Shin Jeong, Jie Liu, and Lijun Jia</td>
<td>Précis: Findings offer preclinical evidence for therapeutic efficacy of a first-in-class neddylation inhibitor to treat liver cancer and provide a mechanistic rationale for its combination with an autophagy inhibitor.</td>
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### Acquired Resistance to Tamoxifen Is Associated with Loss of the Type I Insulin-like Growth Factor Receptor: Implications for Breast Cancer Treatment

Dedra H. Fagan, Ryan R. Uselman, Deepali Sachdev, and Douglas Yee

 précis: Findings in a preclinical breast cancer model of tamoxifen resistance may explain the reason for the lack of clinical benefit observed in trials that combine anti-IGF1 receptor and antiestrogen therapies.

### Definition of Genetic Events Directing the Development of Distinct Types of Brain Tumors from Postnatal Neural Stem/Progenitor Cells

Falk Hertwig, Katharina Meyer, Sebastian Braun, Sara Ek, Rainer Spang, Cosima V. Pfenninger, Isabella Artner, Gaëlle Prost, Xinbin Chen, Jaclyn A. Biegel, Alexander R. Judkins, Elisabet Englund, and Ulrike A. Nuber

 précis: This study reveals that, within malignant brain tumors derived from a common precursor cell pool, the temporal order of genetic events that accumulate is a critical determinant of the histopathologic phenotype that emerges.

### Distinct microRNA Expression Profiles in Prostate Cancer Stem/Progenitor Cells and Tumor-Suppressive Functions of let-7

Can Liu, Kevin Kelnner, Alexander V. Vlassov, David Brown, Junchen Wang, and Dean G. Tang

 précis: microRNA expression profiling performed in prostate cancer stem/progenitor cells identifies a relatively small signature of functionally significant tumor-suppressive genes in the control of prostate cancer stem cell activity.

### PTEN and NF1 Inactivation in Schwann Cells Produces a Severe Phenotype in the Peripheral Nervous System That Promotes the Development and Malignant Progression of Peripheral Nerve Sheath Tumors

Vincent W. Keng, Eric P. Rahmann, Adrienne L. Watson, Barbara R. Tschiida, Christopher L. Moertel, Walter J. Jessen, Tilat A. Rizvi, Margaret H. Collins, Nancy Ratner, and David A. Largaespada

 précis: This study establishes a mouse model that can rapidly recapitulate the histologies associated with onset of human neurofibromas and their malignant progression to high-grade malignant nerve tumors, based on genetic mutations that are relevant to the human tumor setting.

### CD44s Regulates the TGF-β–Mediated Mesenchymal Phenotype and Is Associated with Poor Prognosis in Patients with Hepatocellular Carcinoma

Kosuke Mima, Hirohisa Okabe, Takatsugu Ishimoto, Hiromitsu Hayashi, Shigeki Nakagawa, Hideyuki Kuroki, Masayuki Watanabe, Toru Beppu, Mayumi Tamada, Osamu Nagatomo, Hideyuki Saya, and Hideo Baba

 précis: The so-called standard isoform of a stem cell marker plays a crucial role in mesenchymal phenotypes associated with progression of hepatocellular carcinoma, with implications for therapeutic targeting of this deadly disease.

### FRMD4A Upregulation in Human Squamous Cell Carcinoma Promotes Tumor Growth and Metastasis and Is Associated with Poor Prognosis

Stephen J. Goldie, Klaas W. Mulder, David Wei-Min Tan, Scott K. Lyons, Andrew H. Sims, and Fiona M. Watt

 précis: A candidate stem cell marker implicated in epithelial polarization is found to be a key regulator of metastatic growth in head and neck cancers, with implications for therapeutic targeting of this aggressive and increasingly common disease.
ABOUT THE COVER

Malignant mesothelioma cells secrete the damage-associated molecular pattern protein HMGB1 that promotes tumor progression and sustains the malignant phenotype. HMGB1-secreting mesothelioma cells are addicted to HMGB1, and their anchorage independent growth is impaired by the antagonist BoxA or by a HMGB1 specific monoclonal antibody (mAb), or by a mAb against HMGB1 receptor RAGE, as indicated by the reduced number and size of colonies, compared to controls. For details, see article by Jube et al. on page 3290 of this issue.