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197 p73 and IGF1R Regulate Emergence of Aggressive Cancer Stem–like Features via miR-885-5p Control  
Claudia Meier, Philip Hardstock, Sophie Joost, Vijay Alla, and Brigitte M. Pützer  

Précis: These findings offer major new insight into how the p53 family member p73 promotes the pathogenesis of highly malignant cancers, with implications for how to eradicate tumor-initiating cells and overcome drug resistance.

## Clinical Studies

206 Germline BAP1 Mutational Landscape of Asbestos-Exposed Malignant Mesothelioma Patients with Family History of Cancer  
Jill A. Ohar, Mitchell Cheung, Jacqueline Talarchek, Suzanne E. Howard, Timothy D. Howard, Mary Hesdorffer, Hongzhuang Peng, Frank J. Rauscher, and Joseph R. Testa  

Précis: BAP1 genetic testing may help identify individuals from families with a history of mesothelioma who are at higher risk of developing this cancer, but also with greater chances at long-term survival, an unusual pattern.

## Integrated Systems and Technologies

216 Transcriptome Analysis of Recurrently Deregulated Genes across Multiple Cancers Identifies New Pan-Cancer Biomarkers  
Bogumil Czakowski, Yuji Tanaka, Hideya Kawaji, Albin Sandelin, Robin Andersson, Masayoshi Itoh, Timo Lassmann, the FANTOM5 consortium, Yoshihide Hayashizaki, Piero Carninci, and Alistair R.R. Forrest  

Précis: This genome-wide expression profiling approach identified new perspectives on DNA repetitive elements, often activated during cancer progression, as candidate biomarkers with pan-cancer potential.

## Microenvironment and Immunology

227 Control of PD-L1 Expression by Oncogenic Activation of the AKT–mTOR Pathway in Non–Small Cell Lung Cancer  
Kristin J. Lastwika, Willie Wilson III, Qing Kay Li, Jeffrey Norris, Haiying Xu, Sharon R. Ghazarian, Hiroshi Kitagawa, Shigeru Kawabata, Janis M. Taube, Sheng Yao, Linda N. Liu, Joel J. Gillis, and Phillip A. Dennis  

Précis: This study contributes to the rapidly accumulating evidence that oncogene signaling drives immune escape, implying that anti-oncogenic therapeutic strategies may be useful primarily for leveraging immunochemotherapy combinations.
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239 PD-1 Blunts the Function of Ovarian Tumor–Infiltrating Dendritic Cells by Inactivating NF-xB
Lavakumar Karyampudi, Purushottam Lamichhane, James Krempski, Kimberly R. Kalli, Marshall D. Behrens, Doris M. Vargas, Lynn C. Hartmann, Jo Marie T. Janco, Haidong Dong, Karen E. Hedlin, Allan B. Diets, Ellen L. Goode, and Keith L. Knutson
Précis: These findings reveal how the immunosuppressive molecule PD-1 blunts the activity of tumor-infiltrating dendritic cells, with important implications for the ongoing development of therapeutic strategies to correct immune escape, including by re-engaging innate immune cells.

251 A Preclinical Model of Malignant Peripheral Nerve Sheath Tumor-like Melanoma Is Characterized by Infiltrating Mast Cells
Michael Holzel, Jennifer Landsberg, Nicole Glodde, Tobias Rald, Meri Rogava, Stefanie Riesenberg, Albert Becker, Goran Jonsson, and Thomas Tuting
Précis: These findings highlight the ability to study human melanoma heterogeneity in a mouse model, revealing how melanocyte-immune cell interactions contribute to the development of distinct subsets of melanomas within a single individual.

264 Improved Treatment of Breast Cancer with Anti-HER2 Therapy Requires Interleukin-21 Signaling in CD8+ T Cells
Deepak Mittal, Franco Caramia, Stefan Michiels, Heikki Jorsn, Peikko-Lisa Kellokumpu-Lehtinen, Chistos Sotiriou, Sherene Loi, and Mark J. Smyth
Précis: These findings offer a preclinical rationale to boost IL21 signaling in HER2-positive breast cancer patients as a strategy to improve trastuzumab responses and limit the development of drug resistance.

275 PRC2 Epigenetically Silences Th1-Type Chemokines to Suppress Effector T-Cell Trafficking in Colon Cancer
Nisha Nagarsheth, Dongjun Peng, Ilona Kryczek, Ke Wu, Wei Li, Ende Zhao, Lili Zhao, Shuang Wei, Timothy Frankel, Linda Vatan, Wojciech Szeliga, Yuli Dou, Scott Owens, Victor Marquez, Kaixiong Tao, Emina Huang, Guobin Wang, and Weiping Zou
Précis: A repressive epigenetic program that operates in colon cancer limits the efficiency of effector T cell trafficking to the tumor microenvironment, with implications for improving the efficacy of cancer immunotherapy.

MOLECULAR AND CELLULAR PATHOBIOLOGY

283 Combined MYC Activation and Pten Loss Are Sufficient to Create Genomic Instability and Lethal Metastatic Prostate Cancer
Grethchen K. Hubbard, Laura N. Mutton, May Khalili, Ryan P. McMullin, Jessica L. Hicks, Daniella Bianchi-Frias, Lucas A. Horn, Ibrahim Kulac, Michael S. Moubarek, Peter S. Nelson, Srinivasan Veynusbramanian, Angelo M. De Marzo, and Charles J. Bieberich
Précis: The mouse model described can recapitulate key histopathologic and molecular features of human prostate cancer, including development of genomic instability and overt metastases to lymph nodes, liver, and lung.

293 HBXIP and LSD1 Scaffolded by IncRNA Hotair Mediate Transcriptional Activation by c-Myc
Yinghui Li, Zhen Wang, Hui Shi, Hang Li, Leilei Li, Rumping Fang, Xiaoli Cai, Bowen Liu, Xiaodong Zhang, and Li Hong Ye
Précis: This seminal study defines an oncogenic RNA/protein complex that serves as an effector for c-Myc in activating transcription of its target genes, illuminating long-standing questions concerning how c-Myc drives carcinogenesis.

305 EPHA2 Blockade Overcomes Acquired Resistance to EGFR Kinase Inhibitors in Lung Cancer
Katherine R. Amato, Shan Wang, Li Tan, Andrew K. Hastings, Wenqiang Song, Christine M. Lovly, Catherine B. Meador, Fei Ye, Pengcheng Liu, Justin M. Bal, Daniel C. Colvin, Justin M. Cates, William Pao, Nathanael S. Gray, and Jin Chen
Précis: Targeting a cell surface receptor kinase involved in cell-cell interactions appears to mitigate an important pathway of drug resistance in preclinical models of lung cancer, with immediate impact on clinical testing of the discovery.

319 Gender-Specific Molecular and Clinical Features Underlie Malignant Pleural Mesothelioma
329 Cyclin D1 Promotes Androgen-Dependent DNA Damage Repair in Prostate Cancer Cells
Mathew C. Casimiro, Gabriele Di Sante, Xiaoming Ju, Zhiping Li, Ke Chen, Marco Crosariol, Ismail Yaman, Michael Gormley, Hui Meng, Michael P. Lisanti, and Richard G. Pestell

Précis: These findings shed light on how cyclin D1 promotes DNA damage repair mediated by androgens in the prostate, with potential clinical implications for treating therapy-resistant prostate cancer.

339 Obesity-Induced Colorectal Cancer Is Driven by Caloric Silencing of the Guanylin–GUCY2C Paracrine Signaling Axis
Jieru E. Lin, Francheska Colon-Gonzalez, Erik Blomain, Gilbert W. Kim, Amanda Aing, Brian Stoecker, Justin Rock, Adam E. Snook, Tingting Zhao, Terry M. Hyslop, Michal Tomczak, Richard S. Blumberg, and Scott A. Waldman

Précis: These seminal findings offer the first mechanistic connection between obesity and negation of a universal tumor suppressor pathway in colon tumorigenesis, with immediate implications for a hormone replacement strategy to prevent colorectal cancer in high-risk obese patients.

347 Peritoneal Dissemination Requires an Sp1-Dependent CXCR4/CXCL12 Signaling Axis and Extracellular Matrix–Directed Spheroid Formation
Yuta Kasagi, Yui Harada, Yosuke Morodomi, Toshiki Iwai, Satoru Saito, Kumi Yoshida, Eiji Oki, Hiroshi Saeki, Kippei Ohgaki, Masahiko Sugiyama, Mitsuho Onimaru, Yoshihiko Maehara, and Yoshikazu Yonemitsu

Précis: These findings illuminate mechanisms of peritoneal cancer dissemination, highlighting the Sp1/CXCR4/CXCL12 signaling axis as a rational target for the development of therapeutics to manage this intractable form of malignancy.

358 SKAP2 Promotes Podosome Formation to Facilitate Tumor-Associated Macrophage Infiltration and Metastatic Progression
Masamitsu Tanaka, Shintaro Shimamura, Sei Kuriyama, Daichi Maeda, Akiteru Goto, and Namiko Aiba

Précis: These findings reveal how macrophages infiltrate tumors and enhance metastasis, suggesting a new way to attenuate the tumor-promoting effects of this innate immune cell population in the tumor microenvironment.

370 Skin Cancer Risk Is Modified by KIR/HLA Interactions That Influence the Activation of Natural Killer Immune Cells
Karin A. Vineretsky, Margaret R. Karagas, Brock C. Christensen, Jacqueline K. Kuriger-Laber, Ann E. Perry, Craig A. Storm, and Heather H. Nelson

Précis: These findings reveal associations between keratinocyte-based cancers and activating signals for natural killer immune cells, demonstrating a role for natural killer cell tumor immunity in controlling the most common types of skin cancer.

377 Characterization of a c-Rel Inhibitor That Mediates Anticancer Properties in Hematologic Malignancies by Blocking NF-kB–Controlled Oxidative Stress Responses
Yusuke Shono, Andrea Z. Tuckett, Hsiou-Chi Liu, Ekaterina Doubrovina, Enrico Derenzini, Samedy Ouk, Jennifer J. Tsai, Odette M. Smith, Emily R. Levy, Fabiana M. Kreines, Carly G.K. Ziegler, Mary I. Scallion, Mikhail Doubrovin, Glenn Heller, Anas Younes, Richard J. O’Reilly, Marcel R.M. van den Brink, and Johannes L. Zakrzewski

Précis: These results provide mechanistic insight and preclinical proof of concept for a novel small molecule to treat human lymphoid malignancies, with additional application in recipients of allogeneic bone marrow transplantations to ameliorate graft-versus-host disease, a major clinical challenge.

390 Maximizing the Efficacy of MAPK-Targeted Treatment in PTENLOF/BRAFMT Melanoma through PI3K and IGF1R Inhibition

Précis: These findings reveal that the efficacy of MAPK inhibitors for the treatment of PTEN-deficient and BRAF mutant melanoma can be significantly improved by incorporating pharmacological strategies that also inhibit PI3K and IGF1R signaling.
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**TUMOR AND STEM CELL BIOLOGY**

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Interactions between Adipocytes and Breast Cancer Cells Stimulate Cytokine Production and Drive Src/Sox2/miR-302b–Mediated Malignant Progression


Précis: This study identifies feed-forward signaling loops triggered by adipocyte-cancer cell interactions, driving inflammation and malignant growth and offering new therapeutic strategies to target breast cancers associated with obesity.

Correction: A Modeling Approach to Explain Mutually Exclusive and Co-Occurring Genetic Alterations in Bladder Tumorigenesis

Overall structure of a survivin dimer with deeply buried dimerization core residues shown by their molecular surface in gray and noncore residues shown by sticks in green. Qi and colleagues show that the deeply buried dimerization core residues in undruggable oncogenic dimeric proteins can be targeted using computational approaches for drug discovery to destroy these proteins. Specifically, a lead small-molecule inhibitor LQZ-7F targeting the dimerization core residues of survivin was discovered that induced proteasome-dependent survivin degradation, mitotic arrest, apoptosis, and blocked the growth of human xenograft tumors. For details, see article by Qi and colleagues on page 453.
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