Tumor-Specific Cytotoxic T Cells Are Crucial for Efficacy of Immunomodulatory Antibodies in Patients with Lung Cancer
Joachim G. Aerts and Joost P. Hegmans

Understanding Phenotypic Variation in Rodent Models with Germline Apc Mutations
Maged Zeineidin and Kristi L. Neufeld

Siah: A Promising Anticancer Target
Christina S.F. Wong and Andreas Müller

The Model Muddle: In Search of Tumor Growth Laws
Philip Gerlee

Challenges and Key Considerations of the Enhanced Permeability and Retention Effect for Nanomedicine Drug Delivery in Oncology
Uma Prabhakar, Hiroshi Maeda, Rakesh K. Jain, Eva M. Sevick-Muraca, William Zamboni, Omid C. Farokhzad, Simon T. Barry, Alberto Gabizon, Piotr Grodzinski, and David C. Blakey

Epithelial-to-Mesenchymal Transition and Autophagy Induction in Breast Carcinoma Promote Escape from T-cell–Mediated Lysis
Intissar Akalay, Bassam Janji, Meriem Hasnim, Muhammad Zaeeem Noman, Fabrice André, Patricia De Cremoux, Philippe Bertheau, Cécile Badoual, Philippe Vielh, Annette K. Larsen, Michele Sabbah, Tuan Zea Tan, Joan Herr Keira, Nicole Tsang Ying Hung, Jean Paul Thiery, Fathia Mami-Chouaib, and Salem Chouaib

Précis: This seminal study shows how the acquisition of EMT and autophagy in cancers promotes their ability to escape T-cell immunity, revealing new mechanistic perspectives on how immune escape and metastatic progression are linked.

EGFR-TKI Resistance Due to BIM Polymorphism Can Be Circumvented in Combination with HDAC Inhibition
Takayuki Nakagawa, Shinya Takeuchi, Tadakii Yamada, Hiromichi Ebi, Takako Sano, Shigeki Nanjo, Daisuke Ishikawa, Mitsuo Satoh, Yoshinori Hasegawa, Yoshitake Sekido, and Seiji Yano

Précis: This study unravels the mechanistic basis for EGFR inhibitor resistance in patients who harbor a common BIM polymorphism that prevents BIM induction needed for a robust drug response and shows how to restore sensitivity by epigenetic reprogramming with an HDAC inhibitor.

Higher Frequencies of GARP+CTLA-4 Foxp3+ T Regulatory Cells and Myeloid-Derived Suppressor Cells in Hepatocellular Carcinoma Patients Are Associated with Impaired T-Cell Functionality
Suresh Kalathil, Amit A. Lugade, Austin Miller, Renuka Iyer, and Yasmin Thanavala

Précis: This is one of the first studies to confirm the expectation that multiple immunosuppressive phenotypes are present in patients with advanced cancer, with results that also greatly encourage testing of combinatorial approaches to degrade these phenotypes in this most challenging population as a rational strategy of immunotherapy.

Automated Tracking of Nanoparticle-labeled Melanoma Cells Improves the Predictive Power of a Brain Metastasis Model
Terje Sundstrom, Inderjit Daphu, Ingvild Wendelbo, Erlend Hodneland, Arvid Lundervold, Heike Immervoll, Kai Ove Skaftnesmo, Michal Babic, Pavla Jendelova, Eva Sykova, Morten Lund-Johansen, Rolf Bjerkvig, and Frits Thorsen

Précis: This study describes a model that can improve upon the predictive elements derived from preclinical studies of brain metastases, perhaps ultimately helping reduce the number of clinical trials that fail to show patient benefit.
### Microenvironment and Immunology

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This study offers preclinical proof-of-concept for a novel element in the homologous recombination pathway of DNA repair widely elevated in pancreatic cancer as a candidate therapeutic target to treat this deadly disease.
2551  **Endocrine Fibroblast Growth Factor FGF19 Promotes Prostate Cancer Progression**
Shu Feng, Olga Dakhova, Chad J. Creighton, and Michael Ittmann
*Précis: Expression of an endocrine FGF in prostate cancers is found to promote tumor progression, suggesting it may offer a novel therapeutic target.*

2563  **CYP24A1 and CYP27B1 Polymorphisms Modulate Vitamin D Metabolism in Colon Cancer Cells**
Elizabeth T. Jacobs, Chad Van Pelt, Ryan E. Forster, Wasiq Zaidi, Elizabeth A. Hibler, Michael A. Galligan, Mark R. Haussler, and Peter W. Jurutka
*Précis: These results illustrate how naturally occurring genetic variations in vitamin D metabolic pathways may influence the risk of colon cancer.*

2574  **Characterization of Torin2, an ATP-Competitive Inhibitor of mTOR, ATM, and ATR**
*Précis: An mTOR inhibitor with superior pharmacologic properties in vivo is found to inhibit PIKK family kinases involved in DNA damage signaling and to cooperate strongly with MEK kinase inhibition in killing mouse and human cancer cells.*

2587  **MDM2 Small-Molecule Antagonist RG7112 Activates p53 Signaling and Regresses Human Tumors in Preclinical Cancer Models**
*Précis: The first p53 activator to reach clinical trials is shown in preclinical testing to shrink human tumors, offering a proof-of-concept for eradication of tumors with wild-type forms of this tumor suppressor.*

2598  **Ex Vivo Expansion of Highly Cytotoxic Human NK Cells by Cocultivation with Irradiated Tumor Cells for Adoptive Immunotherapy**
Seon Ah Lim, Tae-Jin Kim, Jung Eun Lee, Chung Hee Sonn, Kwanghee Kim, Jiyoung Kim, Jong Gwon Choi, Il-Kyu Choi, Chae-Ok Yun, Jae-Hong Kim, Cassian Yee, Vinay Kumar, and Kyung-Mi Lee
*Précis: This article shows how to activate and expand human NK cells, providing the basis for a straightforward strategy to expand a highly cytotoxic effector cell population that may help treat advanced cancers refractory to conventional therapy.*

2608  **Ex Vivo Activation of CD56+ Immune Cells That Eradicate Neuroblastoma**
Piya Rujkijyanant, Wing Keung Chan, Paul W. Eldridge, Timothy Lockey, Martha Holladay, Barbara Rooney, Andrew M. Davidoff, Wing Keung, and Queenie Yong
*Précis: These results show a clinically expedient strategy to generate activated NK cells that are highly cytotoxic to neuroblastoma with minimal risk of GvHD.*

2619  **scFv-Based "Grababody'' as a General Strategy to Improve Recruitment of Immune Effector Cells to Antibody-Targeted Tumors**
Zheng Cai, Ting Fu, Yasuhiro Nagai, Lian Lam, Marla Yee, Zhiqiang Zhu, and Hongtao Zhang
*Précis: A novel recombinant molecule combining the specificity of an antibody variable chain with the IgG binding domain facilitates the ability to enable tumor specific killing by endogenous immune effector cells.*

2628  **AMPK Activation by Oncogenesis Is Required to Maintain Cancer Cell Proliferation in Astrocytic Tumors**
Marcos Rios, Marc Foretz, Benoit Viollet, Angel Prieto, Maximo Fraga, Jose A. Costoya, and Rosa Señarís
*Précis: Whether AMPK is essential or not for cancer cell proliferation has been somewhat controversial, but this preclinical study offers a clear rationale for its further exploration as a therapeutic target in brain cancers.*
Rhythmic Control of the ARF-MDM2 Pathway by ATF4 Underlies Circadian Accumulation of p53 in Malignant Cells
Michiko Horiguchi, Satoru Koyanagi, Ahmed M. Hamdan, Keisuke Kakimoto, Naoya Matsunaga, Chikamasa Yamashita, and Shigehiro Ohdo
Précis: Circadian rhythms that determine the accumulation of p53 in malignant cells explain how temporal changes in their chemosensitivity can result, with potential implications for increasing therapeutic efficacy by optimizing the time of drug administration in patients.

Menin Epigenetically Represses Hedgehog Signaling in MEN1 Tumor Syndrome
Buddha Gurung, Zijie Feng, Daniel V. Iwamoto, Austin Thiel, Guanghui Jin, Chen-Min Fan, Jessica M.Y. Ng, Tom Curran, and Xianxin Hua
Précis: Mechanistic results suggest a way to treat parathyroid, pituitary, pancreatic, and other tumors that characterize the MEN1 syndrome, offering a unified treatment approach.

Hepatocyte Growth Factor Activator Inhibitor Type 1 Is a Suppressor of Intestinal Tumorigenesis
Shinri Hoshiko, Makiko Kawaguchi, Tsunoshu Fukushima, Yukihito Haruyama, Kenji Yorita, Hiroyuki Tanaka, Motoharu Seiki, Haruhiko Inatsu, Kazuo Kitamura, and Hiroaki Kataoka
Précis: A membrane-associated tumor suppressor in the intestinal tract is defined that may stimulate novel therapeutic approaches to prevent progression of benign colon tumors.

Inherited Variation in miR-290 Expression Suppresses Breast Cancer Progression by Targeting the Metastasis Susceptibility Gene Arid4b
Natalie Goldberger, Renard C. Walker, Chang Hee Kim, Scott Winter, and Kent W. Hunter
Précis: This is the first study to show that inherited differences in microRNA expression can modify susceptibility to metastatic progression in breast cancer.

CD44-Positive Cancer Stem Cells Expressing Cellular Prion Protein Contribute to Metastatic Capacity in Colorectal Cancer
Lei Du, Guanhua Rao, Hongyi Wang, Baowei Li, WeiLi Tian, Jiantao Cui, Leya He, Brian Laffin, Xiuyun Tian, Chunyi Hao, Hongmin Liu, Xin Sun, Yushan Zhu, Dean G. Tang, Maryam Mehrpour, Youyong Lu, and Quan Chen
Précis: Prions may be functional markers of a highly metastatic subpopulation of cancer stem cells in colorectal cancer, where they might be targeted to treat metastatic disease.

Regulation of Lung Cancer Metastasis by Klf4-Numb–like Signaling
Valentina Vaira, Alice Faversani, Nina M. Martin, David S. Garlick, Stefano Ferrero, Mario Nosotti, Joseph L. Kissil, Silvano Bosari, and Dario C. Altieri
Précis: These findings uncover a novel signaling network centered on the polarity protein Numb-like, which dually promotes abnormal cell motility needed for metastasis along with the persistence of cancer-initiating, stem-like cells that reduce overall survival in lung cancer.

ABOUT THE COVER
Cancer stem cells are implicated in tumor metastasis although the exact mechanisms remain poorly understood. The expression of cellular prion protein (PrPc), a highly conserved glycoprotein that has the same protein sequence as the scrapie prion protein, is positively correlated with an increased risk of metastasis in colorectal cancer. By double immunofluorescence staining of CD44 (green) and PrPc (red), CD44⁺PrPc⁺ cells were detected in the cryosections of colorectal cancers. PrPc⁺CD44⁺ colorectal cancer stem cells displayed high liver metastatic capability. For details, see article by Du and colleagues on page 2682.