Highlights from Recent Cancer Literature

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 Zeineldin and Kristi L. Neufeld

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

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

Automated Tracking of Nanoparticle-labeled Melanoma Cells Improves the Predictive Power of a Brain Metastasis Model
Terje Sundstrøm, 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

Epithelial-to-Mesenchymal Transition and Autophagy Induction in Breast Carcinoma Promote Escape from T-cell–Mediated Lysis
Intissar Akalay, Bassam Janji, Meriem Hasmim, Muhammad Zaem Noman, Fabrice André, Patricia De Cremoux, Philippe Bertheau, Cécile Badoual, Philippe Vielh, Annette K. Larsen, Michele Sabbah, Tuan Z. 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, Shintaro Takeuchi, Tadakai Yamada, Hiromichi Ebi, Takako Sano, Shigeki Nanjo, Daisuke Ishikawa, Mitsuo Sató, Yoshinori Hasegawa, Yoshitaka 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.

Précis: This study describes a model that can improve upon the predictive elements derived from preclinical studies of brain metastasis, perhaps ultimately helping reduce the number of clinical trials that fail to show patient benefit.
Selective Blockade of Matrix Metalloprotease-14 with a Monoclonal Antibody Abrogates Invasion, Angiogenesis, and Tumor Growth in Ovarian Cancer

Rajani Kaimal, Raid Aljumaily, Sarah L. Tressel, Rutika V. Pradhan, Lidija Covic, Athan Kuliopulos, Corrine Zarwan, Young B. Kim, Sheida Sharifi, and Anika Agarwal

Precis: Targeting MMP-14 with a monoclonal antibody may be important for antiangiogenic therapy not only in ovarian cancer but also in other solid tumors.

Interleukin 21–Induced Granzyme B–Expressing B Cells Infiltrate Tumors and Regulate T Cells

Stefanie Lindner, Karen Dahlke, Kai Sontheimer, Magdalena Hagn, Christof Kaltenmeier, Thomas F. Barth, Thamara Beyer, Lisa T. C. Vogelpoel, Eveline M. Dijkgraaf, Moniek Heusinkveld, Hubert Schrezenmeier, and Bernd Jahrsdoerfer

Precis: This potentially seminal study establishes the existence and control of human B regulatory cells in the tumor microenvironment, which under the control of IL-21 may contribute significantly to Treg-dependent mechanisms of local immune escape.

Chemotherapy Alters Monocyte Differentiation to Favor Generation of Cancer-Supporting M2 Macrophages in the Tumor Microenvironment


Precis: Chemotherapy-induced activation of the NfkB pathway reinforces immune suppression in tumor microenvironments where prostanglandin E2 and IL-6 are being produced, suggesting combination strategies with inhibitors of COX-2 or IL-6 signaling to improve chemotherapeutic efficacy.

Chemotherapy Acts as an Adjuvant to Convert the Tumor Microenvironment into a Highly Permissive State for Vaccination-Induced Antitumor Immunity

Tae Heung Kang, Chih-Jen Wang, Crystal D. Lin, Yi-Hsin Tseng, Chung-Ying Tsai, Sheng-Yen Lin, Yu-Ting Hung, Chih-Jen Wang, Crystal D. Lin, and Kwang-Huei Lin

Precis: 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.

Thyroid Hormone Regulation of miR-21 Enhances Migration and Invasion of Hepatoma


Precis: Mechanistic findings reveal how the thyroid hormone can drive the growth and invasion of liver cancer cells by directly regulating an oncogenic microRNA that mediates the hormone’s procarcinogenic effects.

SRC Signaling Is Crucial in the Growth of Synovial Sarcoma Cells

Sebastian Michel, Marcel Trautmann, Elisabeth Sievers, Dagmar Kindler, Sebastian Huss, Marcus Remer, Nicolaus Friedrichs, Jutta Kirfel, Susanne Steiner, Elnar Endl, Peter Wurst, Lukas Heukamp, Roland Penzel, Olle Larsson, Akira Kawai, Shinya Tanaka, Hiroshi Sonobe, Peter Schirmacher, Gunhild Mecht舍r, Eva Wardemann, Reinhard Büttner, and Wolfgang Hartmann

Precis: SRC activation is a critical signaling component of synovial sarcomas and represents a promising therapeutic target in this disease setting.

PARI Overexpression Promotes Genomic Instability and Pancreatic Tumorigenesis

Kevin W. O’Connor, Donnphat Dejsuphong, Eunmi Park, Claudia M. Nicolai, Alec C. Kimmelman, Alan D. D’Andrea, and George-Lucian Moldovan

Precis: 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.

Inhibition of SRC Corrects GM-CSF Hypersensitivity That Underlies Juvenile Myelomonocytic Leukemia

Severa Bunda, Michelle W. Kang, Stephanie S. Sybingco, Julie Weng, Helene Favre, Danielle H. Shin, Meredith S. Irwin, Mignon L. Loh, and Michael Ohl

Precis: This study identifies Src as the critical oncogenic driver and potential druggable target of juvenile myelomonocytic leukemia, an aggressive myeloid malignancy without an effective cure.
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.

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.

Characterization of Torin2, an ATP-Competitive Inhibitor of mTOR, ATM, and ATR
Qingsong Liu, Chunxiao Xu, Swupriya Kirubakaran, Xin Zhang, Woonyoung Hur, Yan Liu, Nicholas P. Kwaśniewski, Jinhua Wang, Kenneth D. Westover, Peng Gao, Dalia Erkan, Mario Niepel, Carson C. Thoreen, Seong A. Kang, Matthew P. Patricelli, and Nathanael S. Gray

*Précis:* An mTOR inhibitor with superior pharmacologic properties in vivo is found to inhibit PI3K family kinases involved in DNA damage signaling and to cooperate strongly with MEK kinase inhibition in killing mouse and human cancer cells.

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.
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, Tsunoshit Fujishima, Yukihiro Haruyama, Kenji Yorita, Hiroyuki Tanka, Motobasu 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, Jiatao 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.