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519 Systems Analysis of BCL2 Protein Family Interactions Establishes a Model to Predict Responses to Chemotherapy
Andreas U. Lindner, Caoimhin G. Concannon, Gerhardt J. Boukes, Mary D. Cannon, Fabien Llambi, Deborah Ryan, Karen Boland, Joan Kehoe, Deborah A. McNamara, Frank Murray, Elaine W. Kay, Suzanne Hector, Douglas R. Green, Heinrich J. Huber, and Jochen H.M. Prehn

Précis: Findings provide insights into how an important oncometabolite is produced by a particularly deadly subset of gliomas, with implications for prognosis and therapy.

Précis: This study offers preclinical proof of concept for a safe and effective strategy to target the many types of tumors driven by a deregulated Wnt pathway.

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Précis: This study offers preclinical proof of concept for a safe and effective strategy to target the many types of tumors driven by a deregulated Wnt pathway.

Précis: An intestinal homeodomain protein widely overexpressed in liver cancer may provide a key connection between long-term chronic inflammation and malignant development.

Précis: This study reports the development of a computational tool that can reliably predict colon cancer patient responses to chemotherapy and might also be useful for prognosis and patient stratification in the future.
Hyperpolarized 13C-Pyruvate Magnetic Resonance Reveals Rapid Lactate Export in Metastatic Renal Cell Carcinomas

Kayvan R. Keshari, Renuka Sriram, Bertram L. Koelsch, Mark Van Cwikking, David M. Wilson, John Kurhanewicz, and Zhen J. Wang

Precis: This preclinical study offers proof of concept for a noninvasive characterization of dynamic cellular metabolism and transporter expression in cancer, with the potential to determine tumor aggressiveness and treatment responses.

VEGFA-VEGFR Pathway Blockade Inhibits Tumor-Induced Regulatory T-cell Proliferation in Colorectal Cancer

Magali Terme, Simon Pernot, Elie Marcheteau, Federico Sandoval, Nadine Benhamouda, Orianne Colussi, Olivier Dubreuil, Antoine F. Carpentier, Eric Tartour, and Julien Taieb

Precis: A central pathway of tumor angiogenesis also has a pivotal role in driving immune escape, illustrating a tight integration of pathways driving malignant progression.

Cell Surface Receptor FPR2 Promotes Antitumor Host Defense by Limiting M2 Polarization of Macrophages

Ying Liu, Keqiang Chen, Chunyan Wang, Wanghua Gong, Teizo Yoshimura, Mingyong Liu, and Ji Ming Wang

Precis: A G protein-coupled receptor implicated in airway inflammation responses supports immune surveillance by restraining macrophages to the tumor cell-killing M1 phenotype.

CD40-Mediated Activation of Chronic Lymphocytic Leukemia Cells Promotes Their CD44-Dependent Adhesion to Hyaluronan and Restricts CCL21-Induced Motility

Tamara Girbl, Elisabeth Hintserer, Eva Melanie Grossinger, Daniela Asdlaber, Karin Oberrascher, Lukas Weiss, Cornelia Hauser-Kronberger, Daniel Neureiter, Hubert Kerschbaum, David Naor, Ronen Alon, Richard Greil, and Tanja Nicole Hartmann

Precis: The evolution of a specific adhesion event in the lymph node microenvironment where a particular leukemia arises is found to be a critical event in determining the outgrowth of these cancers.

CXCR2 Expression in Tumor Cells is a Poor Prognostic Factor and Promotes Invasion and Metastasis in Lung Adenocarcinoma


Precis: A logical strategy to treat smoking-associated lung cancers is suggested by the definition of a proinflammatory signaling axis that cooperates with Box mutations commonly found in this disease.

Lysyl Oxidase Plays a Critical Role in Endothelial Cell Stimulation to Drive Tumor Angiogenesis

Ann-Marie Baker, Demelza Bird, Jonathan C. Welti, Morgane Gourlaouen, Georgina Lang, Graeme I. Murray, Andrew R. Reynolds, Thomas R. Cox, and Janine T. Erler

Precis: Findings point to a new general mechanism for restricting VEGF-driven tumor angiogenesis, with important clinical and therapeutic implications in a wide variety of solid tumor types.

High-Avidity T Cells Are Preferentially Tolerized in the Tumor Microenvironment


Precis: T cells that are designed to be more potent for antitumor activity appear to lose their ability to control tumor growth when they persist in the tumor, offering new insights into why T-cell immunotherapies may fail.

Durable Adoptive Immunotherapy for Leukemia Produced by Manipulation of Multiple Regulatory Pathways of CD8+ T-Cell Tolerance


Precis: Simultaneous inhibition of the CTLA4, PD1, and LAG3 pathways of T-cell tolerance can relieve immune escape in leukemia and empower durable responses to adoptively transferred tumor-reactive T cells.
CD103 or LFA-1 Engagement at the Immune Synapse between Cytotoxic T Cells and Tumor Cells Promotes Maturation and Regulates T-cell Effector Functions
Katarzyna Franciszkiewicz, Audrey Le Floc’h, Marie Boutet, Isabelle Vergnon, Alain Schmitt, and Fathia Mami-Chouaib
Précis: This study shows how integrins that regulate the strength of the interaction between tumor cells and tumor-reactive T cells can alter their killing efficiency.

Tissue Damage–Associated "Danger Signals'' Influence T-cell Responses That Promote the Progression of Preneoplasia to Cancer
Ying He, Jikun Zha, Yamin Wang, Wenhua Liu, Xuanming Yang, and Ping Yu
Précis: Release of the tissue damage-associated molecule HMGB1 by budding prostate tumors exerts an early impact on immune escape and malignant progression.

Homing of Human B Cells to Lymphoid Organs and B-Cell Lymphoma Engraftment Are Controlled by Cell Adhesion Molecule JAM-C
Carmen Doñate, Christiane Ody, Thomas McKee, Sylvie Ruault-Jungblut, Nicolas Fischer, Patricia Ropraz, Beat A. Imhof, and Thomas Matthes
Précis: Findings suggest an important new therapeutic target that may have general applications in the treatment of a variety of types of human lymphoma.

Expression of CD137 on Hodgkin and Reed–Sternberg Cells Inhibits T-cell Activation by Eliminating CD137 Ligand Expression
Weng Tong Ho, Wan Lu Pang, Siew Meng Chong, Antonio Castella, Subail Al-Salam, Teng Ee Tan, Mei Chung Moh, Liang Kai Koh, Shu Uin Gan, Cheong Kin Cheng, and Herbert Schwarz
Précis: Findings show how Hodgkin lymphomas escape immune control, with implications for leveraging more effective immunotherapy to treat this disease.

Twist1 Induces CCL2 and Recruits Macrophages to Promote Angiogenesis
Janine M. Low-Marchelli, Veronica C. Ardi, Edward A. Vizcarra, Nico van Rooijen, James P. Quigley, and Jing Yang
Précis: Findings show how a gene implicated in EMT and metastasis also promotes angiogenesis by recruiting macrophages into the tumor microenvironment.
Identification of an Aurora Kinase Inhibitor Specific for the Aurora B Isoform
Hua Xie, Mee-Hyun Lee, Feng Zhu, Kanamata Reddy, Cong Peng, Yan Li, Do Young Lim, Dong Joon Kim, Xiang Li, Soouk Kang, Haitao Li, Weiya Ma, Ronald A. Lubet, Jian Ding, Ann M. Bode, and Zigang Dong

Précis: Although a variety of Aurora kinase inhibitors have been described, with some now in clinical trials, this is the first study to identify an inhibitor specific for the Aurora B isoform that appears to exert special features in cancer.

SMAD2, SMAD3 and SMAD4 Mutations in Colorectal Cancer

Précis: Through a combined analysis of SMAD gene mutations in colorectal cancer, this article shows a common pathogenic mechanism and a new way to understand how SMAD mutations compromise the TGF-β pathway.

Epigenetic Regulation by Z-DNA Silencer Function Controls Cancer-Associated ADAM-12 Expression in Breast Cancer: Cross-talk between MeCP2 and NF1 Transcription Factor Family

Précis: Findings highlight a novel epigenetic regulatory process involving Z-DNA and several epigenetic factors that contribute to overexpression of breast cancer–associated genes.

Bcl3 Selectively Promotes Metastasis of ERBB2-Driven Mammary Tumors
Alison Wakefield, Jitka Soukupova, Amelie Montagne, Jill Ranger, Rhiannon French, William J. Muller, and Richard W. E. Clarkson

Précis: The study reveals that the oncogene Bcl3 is critical for breast cancer metastasis by controlling tumor cell motility.

miR-186 Downregulation Correlates with Poor Survival in Lung Adenocarcinoma Where It Interferes with Cell-Cycle Regulation
Junchao Cai, Jueheng Wu, Huizhong Zhang, Lishan Fang, Yongbo Huang, Yi Yang, Xun Zhu, Rong Li, and Mengfeng Li

Précis: Results define a tumor suppressor function for a microRNA that blocks cell-cycle progression by directly targeting the pivotal cell-cycle regulators cyclin D1, CDK2, and CDK6.

Common Single-Nucleotide Polymorphisms in the Estrogen Receptor β Promoter Are Associated with Colorectal Cancer Survival in Postmenopausal Women

Précis: Estrogen receptor β is suspected to play a role in colorectal cancer progression by modifying pathways that influence malignant invasion and metastasis.

Combination Therapy Targeting the Chk1 and Wee1 Kinases Shows Therapeutic Efficacy in Neuroblastoma

Précis: Findings offer preclinical proof of concept that inhibitors of a key mitotic regulator can be highly efficacious for treating neuroblastoma, one of the most aggressive pediatric tumors.

DNA Damage–Specific Control of Cell Death by Cryptochrome in p53-Mutant Ras–Transformed Cells
Jin Hyup Lee, Shobhan Gaddameedhi, Nuri Ozturk, Rui Ye, and Aziz Sancar

Précis: Findings show how different regulators of the circadian clock affect distinct pathways of intrinsic apoptosis in malignant cells.
Histone Deacetylase Inhibitor AR-42 Differentially Affects Cell-cycle Transit in Meningeal and Meningioma Cells, Potently Inhibiting NF2-Deficient Meningioma Growth

Sarah S. Burns, Elena M. Akhmametyeva, Janet L. Oblinger, Matthew L. Bush, Jie Huang, Volker Senner, Ching-Shih Chen, Abraham Jacob, D. Bradley Welling, and Long-Sheng Chang

Précis: Preclinical investigations establish the potential utility of a pan-HDAC inhibitor in treatment of meningioma, a type of brain tumor.

804

De Novo Design of a Tumor-Penetrating Peptide

Luca Alberici, Lise Roth, Kazuki N. Sugahara, Lilach Agemy, Venkata R. Kotamraju, Tambet Teesalu, Claudio Bordignon, Catia Traversari, Gian-Paolo Rizzardi, and Eriki Ruoslahi

Précis: Nanotherapies incorporating effective tumor-penetrating peptides may improve drug delivery to tumor vessels and into the extravascular tumor tissue.

813

Polo-like Kinase 1: A Potential Therapeutic Option in Combination with Conventional Chemotherapy for the Management of Patients with Triple-Negative Breast Cancer

Virginie Maire, Fariba Nemati, Marion Richardison, Anne Vincent-Salomon, Bruno Tesson, Guillaume Rigail, Éléonore Gravier, Bérénice Marty-Prouvost, Leanne De Koning, Guillaume Lang, David Gentien, Aurélie Dumont, Emmanuel Barillot, Elisabetta Marangoni, Didier Decaudin, Guerrin C. Tucker, and Thierry Dubois

Précis: This study suggests a promising therapeutic target for chemotherapy combination treatment of patients with triple-negative breast cancers, which tend to be more aggressive and less treatable than other types of breast cancer.

824

Dual Targeting of EGFR and HER3 with MEHD7945A Overcomes Acquired Resistance to EGFR Inhibitors and Radiation

Shyhmin Huang, Chunrong Li, Eric A. Armstrong, Chimera R. Peet, Jarob Saker, Lukas C. Amler, Mark X. Sliwkowski, and Paul M. Harari

Précis: By anticipating a common resistance mechanism to EGFR inhibitors, a new agent that can overcome HER3 signaling as well as target the EGFR has great potential to address a current therapeutic barrier in the field.

834

Resistance to Irreversible EGF Receptor Tyrosine Kinase Inhibitors through a Multistep Mechanism Involving the IGF1R Pathway

Alexis B. Cortot, Claire E. Repellin, Takeshi Shimamura, Marzia Capellelli, Kreshnik Zejnullahu, Dalia Ercan, James G. Christensen, Kwock-Kin Wong, Nathanael S. Gray, and Pasi A. Jänne

Précis: Prophylactic blockade of known mechanisms of drug resistance before they can emerge will be critical for generating more effective strategies of targeted combination therapy.

844

MDA-9/Syntenin and IGFBP-2 Promote Angiogenesis in Human Melanoma


Précis: This work highlights novel targets to reverse the production of new blood vessels by tumors.

855

Real-time In Vivo Molecular Detection of Primary Tumors and Metastases with Ratiometric Activatable Cell-Penetrating Peptides

Elamprakash N. Savarir, Csilla N. Felsen, Nadia Nashi, Tao Jiang, Lesley G. Ellies, Paul Steinbach, Roger Y. Tsien, and Quyen T. Nguyen

Précis: Improved methods to detect tumors and metastases in preclinical models are needed for longitudinal animal studies and are vital to move the field beyond increasingly irrelevant cell culture models of cancer.

865

Specific Elimination of CD133+ Tumor Cells with Targeted Oncolytic Measles Virus


Précis: Preclinical studies show how to improve the efficacy of oncolytic viruses by targeting them to CD133+ stem cells in multiple orthotopic and xenograft mouse models of cancer.
Therapeutic Effects of Deleting Cancer-Associated Fibroblasts in Cholangiocarcinoma


Précis: The prototypical demethylating drug 5-aza-2C exhibits strong single-agent activity in an aggressive mouse model of pancreatic cancer characterized by an abundant stroma that resembles human tumors, with the drug efficacy reflecting effects on both the stromal and the malignant epithelial compartments of the tumor.

Therapeutic Effects of Deleting Cancer-Associated Fibroblasts in Cholangiocarcinoma


Précis: Preclinical observations suggest that the Bax mimetic drug navitoclax may be useful for triggering apoptosis of cancer-associated fibroblasts in the tumor microenvironment as a generalized strategy to attack solid tumors that rely on these cells.

Convergence of the ZMIZ1 and NOTCH1 Pathways at C-MYC in Acute T Lymphoblastic Leukemias

Lesley A. Rakowski, Derek D. Garagiola, Choi M. Li, Margaret Decker, Sarah Caruso, Morgan Jones, Rork Kuick, Tomasz Cierpicki, Ivan Maillard, and Mark Y. Chiang

Précis: A novel oncogene that cooperates with NOTCH1 to drive C-MYC expression is critical to license the emergence of diverse solid and liquid tumors by NOTCH1.

Definition of Molecular Determinants of Prostate Cancer Cell Bone Extravasation


Précis: This study offers the first comprehensive mechanism of how metastatic cancer cells traverse the vascular endothelium.

Chromosome Instability Modulated by BMI1–AURKA Signaling Drives Progression in Head and Neck Cancer

Chun-Hung Chou, Neng-Kai Yang, Ting-Yun Liu, Shyh-Kuan Tai, Dennis Shin-Shian Hsu, Chun-Hung Chou, Neng-Kai Yang, Ting-Yun Liu, Shyh-Kuan Tai, Dennis Shin-Shian Hsu, Muh-Hwa Yang

Précis: This important study reveals a mechanistic link between cancer progression and chromosomal instability, and it provides a rationale for clinical evaluation of Aurora kinase inhibitors to treat a type of cancer with a rapidly rising incidence.

DKK2 Mediates Osteolysis, Invasiveness, and Metastatic Spread in Ewing Sarcoma


Précis: A family of antagonists of the Wnt pathway drives bone invasion and osteolysis in Ewing sarcoma, an aggressive pediatric malignancy, with implications for understanding bone invasion more generally.
**Novel Oncogene–Induced Metastatic Prostate Cancer Cell Lines Define Human Prostate Cancer Progression Signatures**
Xiaoming Ju, Adam Ertel, Mathew C. Casimiro, Zuoren Yu, Hui Meng, Peter A. McCue, Rhonda Walters, Paolo Fortina, Michael P. Lisanti, and Richard G. Pestell

**Précis:** Murine prostate cancer cell lines that can metastasize in immunocompetent hosts accurately reflect the genomic and transcriptomic changes that occur in human prostate cancer and provide an improved set of tools for the field to test new therapies.

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**Loss of miR-204 Expression Enhances Glioma Migration and Stem Cell-like Phenotype**
Zhe Ying, Yun Li, Juengeng Wu, Xun Zhu, Yi Yang, Han Tian, Wei Li, Bo Hu, Shi-Yuan Cheng, and Mengfeng Li

**Précis:** Findings define a microRNA that controls the development of stem cell-like phenotypes and cell motility in malignant brain cancer cells, with possible implications in cancer stem cell-like functions generally.

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**Dual Functions of the Homeoprotein DLX4 in Modulating Responsiveness of Tumor Cells to Topoisomerase II-Targeting Drugs**
Bon Q. Trinh, Song Yi Ko, Nicolas Barengo, Shiaw-Yih Lin, and Honami Naora

**Précis:** This provocative study may explain why some tumors respond poorly to drugs that target topoisomerase II despite expressing high levels of this chromatin maintenance factor.

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**Novel Mechanism of Apoptosis Resistance in Cancer Mediated by Extracellular PAR-4**
Ravshan Burikhanov, Tripti Shrestha-Bhattarai, Shirley Qiu, Nithil Shukla, Nikhil Hebbar, Subodh M. Lele, Craig Horbinski, and Vivek M. Rangnekar

**Précis:** Provocative findings offer new insights into how NFκB regulates ER stress and extrinsic apoptosis signaling in cancer cells, with implications for cross-talk with pro-inflammatory signaling in cancer cells.
ABOUT THE COVER

Twist1 is a key inducer of epithelial-mesenchymal transition during tumor metastasis and is upregulated in aggressive human cancers. A new role for Twist1 in tumor progression was uncovered, showing that Twist1 is capable of stimulating angiogenesis via CCL2 production and subsequent recruitment of macrophages from the tumor stroma. Using a Matrigel plug assay, the images show that mouse mammary tumor cells endogenously expressing high levels of Twist1 stimulated a strong angiogenic response, seen as red blood vessels in the isolated plugs. When Twist1 expression was suppressed in these cells, the angiogenic response was diminished. For details, see the article by Low-Marchelli and colleagues on page 662.
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