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Precise: Reinforcing the need to study cancer in an organismal context to gain deeper understanding, a core scaffolding protein in lipid rafts is found to regulate the growth of primary tumors and metastases quite differently.

The SUMO E3-ligase PIAS1 Regulates the Tumor Suppressor PML and Its Oncogenic Counterpart PML-RARA
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Precise: Findings offer mechanistic insights into how the sumoylation machinery modifies oncogenic signals regulated by the tumor suppressor PML, and also the therapeutic response to leukemias involving PML mutations.

Intragenic ATM Methylation in Peripheral Blood DNA as a Biomarker of Breast Cancer Risk
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Precise: As cancer risk studies move from genome to epigenome associations, the use of DNA isolated from peripheral blood cells offers an easily accessible sample type for epigenome-wide association studies.
Effects of a Caloric Restriction Weight Loss Diet and Exercise on Inflammatory Biomarkers in Overweight/Obese Postmenopausal Women: A Randomized Controlled Trial
Ikuyo Imayama, Cornelia M. Ulrich, Catherine M. Alfano, Chiachi Wang, Liren Xiao, Mark H. Wener, Kristin L. Campbell, Catherine Duggan, Karen E. Foster-Schubert, Angela Kong, Caitlin E. Mason, Ching-Yun Wang, George L. Blackburn, Carolyn E. Bain, Henry J. Thompson, and Anne McTiernan
Précis: Findings suggest that weight loss with or without exercise may reduce risk of breast cancer, possibly due to a reduction in systemic inflammation that may support tumor development or progression.

Impact of Intertumoral Heterogeneity on Predicting Chemotherapy Response of BRCA1-Deficient Mammary Tumors
Précis: Studies of BRCA1-deficient mammary cancers suggest that tumor heterogeneity makes it difficult to define gene expression signatures that could predict chemotherapy responses.

Immune Response Is an Important Aspect of the Antitumor Effect Produced by a CD40L-Encoding Oncolytic Adenovirus
Iulia Diaconu, Vincenzo Cerullo, Mari L.M. Hirvinen, Sophie Escutenaire, Matteo Ugolini, Salla K. Pesonen, Simona Bramante, Sari Parviainen, Anna Kanerva, Angelica S.J. Loskog, Aristides G. Elioopoulos, Sari Pesonen, and Akseli Hemminki
Précis: Findings detail the development of a new generation of oncolytic adenovirus that is armed with CD40L, which results in the induction of a Th1-type immune response that causes accumulation of cytotoxic T cells at the tumor site and increased antitumor efficacy.

Novel MT1-MMP Small-Molecule Inhibitors Based on Insights into Hemopexin Domain Function in Tumor Growth
Albert G. Remacle, Vladislav S. Golubkov, Sergey A. Shiryayev, Russell Dab, John L. Stebbins, Andrei V. Chernov, Anton V. Chelslov, Maurizio Pellecchia, and Alex Y. Strongin
Précis: Findings reveal that targeting a regulatory domain of increasing pharmacologic interest in matrix metalloproteases and other proteins can exert potent antitumor properties.

Expression of the p53 Target CDIP Correlates with Sensitivity to TNFα-Induced Apoptosis in Cancer Cells
Lauren Brown-Endres, David Schoenhfeld, Fang Tian, Hyung-Gu Kim, Takushi Namba, César Muñoz-Fontela, Anna Mandinova, Stuart A. Aaronson, and Sam W. Lee
Précis: This study suggests that the product of a p53 target gene may serve as a predictive biomarker for TNF-based cancer therapeutics.

S-Glutathionylated Serine Proteinase Inhibitors as Plasma Biomarkers in Assessing Response to Redox-Modulating Drugs
Précis: Novel blood-based biomarkers will assist in pharmacogenetic design of protocols that test new drugs.

Translation Initiation Factor eIF4E Is a Target for Tumor Cell Radiosensitization
Thomas J. Hayman, Eli S. Williams, Mohammad Jamal, Uma T. Shunkavaram, Kevin Camphausen, and Philip J. Tofilon
Précis: Findings suggest that an existing agent that targets a critical component of the translation machinery might be repositioned as a general neoadjuvant strategy to heighten radiotherapeutic responses in cancer, with the potential to exert a broad impact in radiation oncology.
Dithiolethiones Inhibit NF-κB Activity via Covalent Modification in Human Estrogen Receptor–Negative Breast Cancer
Christopher H. Switzer, Robert Y.-S. Cheng, Lisa A. Ridnour, Margaret C. Murray, Valerio Tazzari, Anna Sparatore, Piero Del Soldato, Harry B. Hines, Sharon A. Glynn, Stefan Ambs, and David A. Wink

Precis: A novel chemical mechanism to inhibit NF-κB activation in aggressive estrogen receptor-negative breast cancers may blunt their invasive capabilities.

TUMOR AND STEM CELL BIOLOGY

p120RasGAP-Mediated Activation of c-Src Is Critical for Oncogenic Ras to Induce Tumor Invasion
Po-Chao Chan and Hong-Chen Chen

Precis: The requirement for c-Src in tumor invasion evoked by oncogenic Ras has implications for the development of therapies to target the Ras pathway, long a goal of the field.

Estrogen Receptor Alpha Mediates Progestin-Induced Mammary Tumor Growth by Interacting with Progesterone Receptors at the Cyclin D1/MYC Promoters
Sebastián Giulianielli, José P. Vaqué, Rocío Soldati, Victoria Wargon, Silvia I. Vanzulli, Rubén Martins, Eduardo Zeitlin, Alfredo A. Molinolo, Luisa A. Helguero, Caroline A. Lamb, J. Silvio Gutkind, and Claudia Lanari

Precis: Antiestrogens block progesterone-induced tumor growth because they disrupt estrogen receptor-progesterone receptor interactions that are essential for target gene transcription.

Proteomic Portrait of Human Breast Cancer Progression Identifies Novel Prognostic Markers
Tamar Geiger, Stephen F. Madden, William M. Gallagher, Juergen Cox, and Matthias Mann

Precis: In performing the deepest proteomic analysis of breast cancer progression to date, this study identifies novel prognostic markers for overall survival that function in metabolic and secretory processes.

Suppression of the Epithelial–Mesenchymal Transition by Grainyhead-like-2
Benjamin Cieply, Philip Riley IV, Phillip M. Pifer, Joseph Widmeyer, Joseph B. Addison, Alexey V. Ivanov, James Denvir, and Steven M. Frisch

Precis: A gene involved in wound healing and neural tube closure is found to be a suppressor of oncogenic epithelial–mesenchymal transition, a pivotal process in cancer cells that is tightly associated with the capacity for metastatic progression.

ABOUT THE COVER

Galectin-3 binding protein, a glycoprotein produced by neuroblastoma cells, upregulates the expression of interleukin-6 in bone marrow mesenchymal cells by interacting with galectin-3. Using immunofluorescence, it was found that the galectin-3 binding protein colocalizes with galectin-3 at the surface and in the cytosol of mesenchymal cells. This interaction generates a Ras/MEK/ERK-dependent signal that transcriptionally upregulates the production of interleukin-6 in the bone marrow microenvironment. Activation of this pathway contributes to neuroblastoma bone metastasis. For details, see article by Silverman and colleagues on page 2228 of this issue.