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IEWS In The Spotlight

More T Cells versus Better T Cells in Patients with Breast Cancer 1062

D.E. Speiser and G. Verdeil

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Targeting the Noncoding Genome: Superenhancers Meet Their Kryptonite 1065

E. Wang and I. Aifantis

See article, p. 1136

Soils and Seeds That Initiate Pancreatic Cancer Metastasis .. 1067

C.R. Vakoc and D.A. Tuveson

See article, p. 1184

REVIEW **How Ribosomes Translate Cancer** 1069

S.O. Sulima, I.J.F. Hofman, K. De Keersmaecker, and J.D. Dinman

RESEARCH BRIEF **TCR Repertoire Intratumor Heterogeneity in Localized Lung Adenocarcinomas: An Association with Predicted Neoantigen Heterogeneity and Postsurgical Recurrence** 1088

A. Reuben, R. Gittelman, J. Gao, J. Zhang, E.C. Yusko, C.-J. Wu, R. Emerson, J. Zhang,

C. Tipton, J. Li, K. Quek, V. Gopalakrishnan, R. Chen, L.M. Vence, T. Cascone, M. Vignali, J. Fujimoto, J. Rodríguez-Canales, E.R. Parra, L.D. Little, C. Gumbs, M.-A. Forget, L. Federico, C. Haymaker, C. Behrens, S. Benzeno, C. Bernatchez, B. Sepesi, D.L. Gibbons, J.A. Wargo, W.N. William Jr, S. Swisher, J.V. Heymach, H. Robins, J.J. Lee, P. Sharma, J.P. Allison, P.A. Futreal, I.I. Wistuba, and J. Zhang

Précis: T-cell receptor sequencing of 45 tumor regions from 11 patients with NSCLC found T-cell repertoire intratumor heterogeneity that was associated with disease relapse and reduced disease-free survival.

RESEARCH ARTICLES **Immune Escape in Breast Cancer During *In Situ* to Invasive Carcinoma Transition** 1098



C.R. Gil Del Alcazar, S.J. Huh, M.B. Ekram, A. Trinh, L.L. Liu, F. Beca, X. Zi, M. Kwak, H. Bergholtz, Y. Su, L. Ding, H.G. Russnes, A.L. Richardson, K. Babski, E.M.H. Kim, C.H. McDonnell III, J. Wagner, R. Rowberry, G.J. Freeman, D. Dillon, T. Sorlie, L.M. Coussens, J.E. Garber, R. Fan, K. Bobolis, D.C. Allred, J. Jeong, S.Y. Park, F. Michor, and K. Polyak

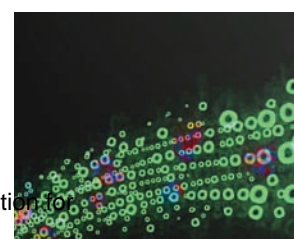
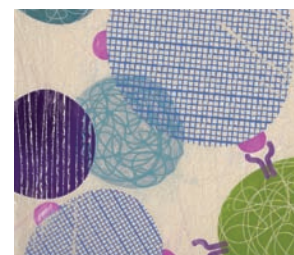
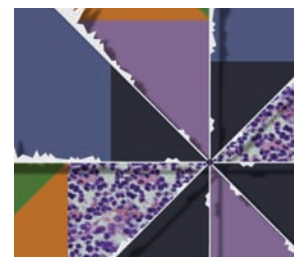
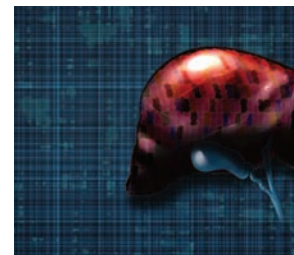
Précis: Progression from ductal carcinoma *in situ* to invasive ductal carcinoma is characterized by a switch to a more suppressive immune microenvironment.

See commentary, p. 1062

Whole-Genome and Epigenomic Landscapes of Etiologically Distinct Subtypes of Cholangiocarcinoma 1116



A. Jusakul, I. Cutcutache, C.H. Yong, J.Q. Lim, M.N. Huang, N. Padmanabhan, V. Nellore, S. Kongpetch, A.W.T. Ng, L.M. Ng, S.P. Choo, S.S. Myint, R. Thanan, S. Nagarajan, W.K. Lim, C.C.Y. Ng, A. Boot, M. Liu, C.K. Ong, V. Rajasegaran, S. Lie, A.S.T. Lim, T.H. Lim, J. Tan, J.L. Loh, J.R. McPherson, N. Khuntikeo, V. Bhudhisawasdi, P. Yongvanit, S. Wongkham, Y. Totoki, H. Nakamura, Y. Arai, S. Yamasaki, P.K.-H. Chow, A.Y.F. Chung, L.L.P.J. Ooi, K.H. Lim, S. Dima, D.G. Duda, I. Popescu, P. Broet, S.-Y. Hsieh, M.-C. Yu, A. Scarpa, J. Lai, D.-X. Luo, A.L. Carvalho, A.L. Vettore,



H. Rhee, Y.N. Park, L.B. Alexandrov, R. Gordán, S.G. Rozen, T. Shibata, C. Pairajkul, B.T. Teh, and P. Tan

Précis: In-depth genetic characterization defines cholangiocarcinoma subtypes and identifies previously undescribed drivers, noncoding promoter mutations, and structural variants.

Superenhancer Analysis Defines Novel Epigenomic Subtypes of Non-APL AML, Including an RAR α Dependency Targetable by SY-1425, a Potent and Selective RAR α Agonist 1136



M.R. McKeown, M.R. Corces, M.L. Eaton, C. Fiore, E. Lee, J.T. Lopez, M.W. Chen, D. Smith, S.M. Chan, J.L. Koenig, K. Austgen, M.G. Guenther, D.A. Orlando, J. Lovén, C.C. Fritz, and R. Majeti

Précis: Characterization of enhancer landscapes in patients with AML identified a subset of non-APL AML with an RARA superenhancer that confers sensitivity to treatment with the selective RAR α agonist SY-1425.

See commentary, p. 1065

Overcoming the Immunosuppressive Tumor Microenvironment of Hodgkin Lymphoma Using Chimeric Antigen Receptor T Cells 1154

M. Ruella, M. Klichinsky, S.S. Kenderian, O. Shestova, A. Ziober, D.O. Kraft, M. Feldman, M.A. Wasik, C.H. June, and S. Gill

Précis: Anti-CD123 chimeric antigen receptor T cells overcome the immunosuppressive

tumor microenvironment in Hodgkin lymphoma by targeting both malignant cells and tumor-associated macrophages.

Loss of MutL Disrupts CHK2-Dependent Cell-Cycle Control through CDK4/6 to Promote Intrinsic Endocrine Therapy Resistance in Primary Breast Cancer 1168



S. Haricharan, N. Punturi, P. Singh, K.R. Holloway, M. Anurag, J. Schmelz, C. Schmidt, J.T. Lei, V. Suman, K. Hunt, J.A. Olson Jr, J. Hoog, S. Li, S. Huang, D.P. Edwards, S.M. Kavuri, M.N. Bainbridge, C.X. Ma, and M.J. Ellis

Précis: Dysregulation of the mismatch repair complex MutL promotes intrinsic resistance to endocrine therapy in ER⁺ breast cancer model systems and patients and may confer sensitivity to CDK4/6 inhibitors.

BLIMP1 Induces Transient Metastatic Heterogeneity in Pancreatic Cancer 1184

S.-H. Chiou, V.I. Risca, G.X. Wang, D. Yang, B.M. Grüner, A.S. Kathiria, R.K. Ma, D. Vaka, P. Chu, M. Kozak, L. Castellini, E.E. Graves, G.E. Kim, P. Mourrain, A.C. Koong, A.J. Giaccia, and M.M. Winslow

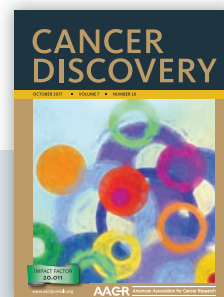
Précis: Pancreatic ductal adenocarcinoma metastasis is promoted by hypoxia and HIF-driven upregulation of the prometastatic transcription factor BLIMP1.

See commentary, p. 1067

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ON THE COVER To determine how intratumor heterogeneity in the T-cell landscape correlates with the genomic landscape and with patient outcome in non-small cell lung cancer (NSCLC), Reuben and colleagues characterized the T-cell repertoire in a cohort of 11 patients with NSCLC who had previously been subject to whole-exome sequencing. T-cell receptor (TCR) sequencing profiled 45 tumor regions across the 11 tumors and revealed a high level of intratumor heterogeneity, with differences in T-cell density, clonality, and repertoire. TCR intratumor heterogeneity was linked to neoantigen heterogeneity and was correlated with disease relapse and reduced disease-free survival in patients with NSCLC. These findings link T-cell repertoire heterogeneity to genomic intratumor heterogeneity and relapse in NSCLC. For details, please see the article by Reuben and colleagues on page 1088.



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