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The Vigorous Immune Microenvironment of Microsatellite Instable Colon Cancer Is Balanced by Multiple Counter-Inhibitory Checkpoints ................. 43
Précis: Mismatch repair-deficient colorectal cancers counteract Th1/CTL immune responses by upregulating immune checkpoint proteins, including PD-1 and PD-L1. See commentary, p. 16

Mutant KRAS-Induced Expression of ICAM-1 in Pancreatic Acinar Cells Causes Attraction of Macrophages to Expedite the Formation of Precancerous Lesions ............... 52
Précis: Crosstalk between pancreatic acinar cells and proinflammatory macrophages promotes initiation of acinar-to-ductal metaplasia via KRASG12D-induced expression of the macrophage chemoattractant ICAM1.
Prospective Blinded Study of BRAF<sup>V600E</sup> Mutation Detection in Cell-Free DNA of Patients with Systemic Histiocytic Disorders


Précis: Cell-free DNA testing using plasma and urine samples may be a reliable, noninvasive method to identify mutations and monitor treatment response in histiocytic disorders.

Measuring Residual Estrogen Receptor Availability during Fulvestrant Therapy in Patients with Metastatic Breast Cancer

M. van Kruchten, E.G. de Vries, A.W. Glaudemans, M.C. van Lanschot, M. van Faassen, I.P. Kema, M. Brown, C.P. Schröder, E.F. de Vries, and G.A. Hospers

Précis: Decreased [<sup>18</sup>F]fl uoroestradiol uptake visualized by PET/CT provides a measure of tumor ER availability and correlates with fulvestrant treatment outcome in patients with metastatic breast cancer.

Induction of Telomere Dysfunction Mediated by the Telomerase Substrate Precursor 6-Thio-2'-Deoxyguanosine

I. Mender, S. Gryaznov, Z.G. Dikmen, W.E. Wright, and J.W. Shay

Précis: 6-thio-2'-deoxyguanosine is a precursor of a telomerase substrate that is incorporated into newly synthesized telomeres, leading to telomere dysfunction and death in telomerase-expressing cells.

See commentary, p. 19

Hyman, Diamond, and colleagues carried out a prospective, blinded study to quantitatively detect the BRAF<sup>V600E</sup> mutation in circulating tumor cell-free DNA (cfDNA) from the urine and plasma of patients with Langerhans cell histiocytosis or Erdheim-Chester disease. Urinary cfDNA analysis defined the BRAF genotype of all 30 patients and was 100% concordant with tissue genotypes among treatment-naïve patients. Furthermore, serial urinary cfDNA analyses in patients treated with a BRAF inhibitor or immunomodulatory therapy showed a progressive decrease in BRAF<sup>V600E</sup> allele burden, consistent with radiographic evidence of disease improvement. Tissue and cfDNA genotyping also identified a previously unreported somatic KRAS<sup>G12S</sup> mutation in a BRAF wild-type patient. These data suggest cfDNA testing as a reliable, noninvasive method to detect BRAF<sup>V600E</sup> mutations and monitor response to therapy in histiocytic disorders. For details, please see the article by Hyman, Diamond, and colleagues on page 64.