In The Spotlight

Making Sense of MEK1 Mutations in Intrinsic and Acquired RAF Inhibitor Resistance.............. 390
K.H.T. Paraíso and K.S.M. Smalley
Commentary on Shi et al., p. 414

Beta-Testing of PI3-Kinase Inhibitors: Is Beta Better? ... 393
P.R. Shepherd and W.A. Denny
Commentary on Ni et al., p. 425

Circulating Endothelial Progenitors and Tumor Resistance to Vascular-Targeting Therapies ........... 395
M. De Palma and S. Nucera
Commentary on Taylor et al., p. 434

Occupy EGFR ............... 398
J.H. Park and M.A. Lemmon
Commentary on Borkovich et al., p. 450 and Vivanco et al., p. 458

In Focus


REVIEW
Emerging Epigenetic Targets and Therapies in Cancer Medicine ............... 405
R. Popovic and J.D. Licht

RESEARCH BRIEFS
Preexisting MEK1 Exon 3 Mutations in V600E/KBRAF Melanomas Do Not Confer Resistance to RAF Inhibitors ............... 414
Précis: Mutation of the downstream RAF effector MEK1 is not a mechanism of innate resistance to targeted RAF inhibitors.

Functional Characterization of an Isoform-Selective Inhibitor of PI3K-p110β as a Potential Anticancer Agent ............... 425
Précis: A selective small-molecule inhibitor of the p110β isoform of PI3K is effective in a subset of PTEN-deficient tumor cell lines and xenografts.
Reversing Resistance to Vascular-Disrupting Agents by Blocking Late Mobilization of Circulating Endothelial Progenitor Cells


Précis: Vascular-disrupting agents induce a late surge in circulating endothelial progenitor cells that can be blocked by antiangiogenic agents.

Kinetics of Inhibitor Cycling Underlie Therapeutic Disparities between EGFR-Driven Lung and Brain Cancers


Précis: The glioma-derived EGFRvIII mutant releases erlotinib more quickly than non-small cell lung cancer–derived EGFR-mutant alleles.

Differential Sensitivity of Glioma-versus Lung Cancer–Specific EGFR Mutations to EGFR Kinase Inhibitors


Précis: Glioma cells with extracellular domain EGFR mutations are selectively sensitive to type II EGFR inhibitors that stabilize the inactive kinase conformation.

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ON THE COVER

Vivanco and colleagues demonstrated that glioma-specific EGFR extracellular domain mutants were more sensitive to type II EGFR inhibitors (e.g., lapatinib) that stabilize an inactive kinase conformation than type I EGFR inhibitors (e.g., erlotinib) that target the active kinase conformation more commonly found in EGFR-mutant lung cancers. In a related article, Barkovich and colleagues found that the rapid release of erlotinib by glioma-specific EGFR mutants rendered them less sensitive to erlotinib than lung cancer–derived EGFR mutants. Together, these studies provide explanations for the limited success of first-generation EGFR inhibitors in treatment of EGFR-mutant gliomas and suggest alternative EGFR inhibition strategies may work best in these tumors. For details, please see the article by Vivanco and colleagues on page 458 and the article by Barkovich and colleagues on page 450.