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- 1135 | **Vasohibin-2 Expressed in Human Serous Ovarian Adenocarcinoma Accelerates Tumor Growth by Promoting Angiogenesis**
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- 1147 | **Inhibition of the Hedgehog Pathway Targets the Tumor-Associated Stroma in Pancreatic Cancer**
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- 1158 | **Cancer-Associated Fibroblasts Induce Matrix Metalloproteinase-Mediated Cetuximab Resistance in Head and Neck Squamous Cell Carcinoma Cells**
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CELL CYCLE, CELL DEATH, AND SENEESCENCE

- 1169 | **RBM38 Is a Direct Transcriptional Target of E2F1 that Limits E2F1-Induced Proliferation**
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- 1178 | **Group I p21-Activated Kinases (PAKs) Promote Tumor Cell Proliferation and Survival through the AKT1 and Raf-MAPK Pathways**
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- 1189 | **The p38 MAPK-MK2 Axis Regulates E2F1 and FOXM1 Expression after Epirubicin Treatment**
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- 1203 | **S-Nitrosylation of EGFR and Src Activates an Oncogenic Signaling Network in Human Basal-Like Breast Cancer**
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- 1216 | **Hepatocyte Growth Factor Enhances Alternative Splicing of the Krüppel-like Factor 6 (KLF6) Tumor Suppressor to Promote Growth through SRSF1**
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- 1228 | **KRAS^{G12D}- and BRAF^{V600E}-Induced Transformation of Murine Pancreatic Epithelial Cells Requires MEK/ERK-Stimulated IGF1R Signaling**
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ABOUT THE COVER

Group I p21-activated kinases (PAKs) regulate cell survival, proliferation and motility, all factors that contribute to tumorigenesis. The tumor suppressor NF2 negatively regulates group I PAKs, and mutation or loss of NF2 leads to subsequent PAK activation. Using immunohistochemistry, PAK was found to be phosphorylated/activated in asbestos-induced malignant mesotheliomas from *Nf2*-deficient mice. Inhibition of group I PAKs in patient-derived mesothelioma cell lines was sufficient to inhibit tumor cell proliferation and viability via inactivation of the AKT and Raf-MAPK pathways, suggesting that PAKs represent novel targets for therapeutic intervention in NF2-deficient malignancies. For details, see article by Menges and colleagues on page 1178.

