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Integrative Survival-Based Molecular Profiling of Human Pancreatic Cancer

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Human Prostate Cancer in a Clinically Relevant Xenograft Mouse Model: Identification of β(1,6)- Branched Oligosaccharides as a Marker of Tumor Progression

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High-Risk Ovarian Cancer Based on 126-Gene Expression Signature Is Uniquely Characterized by Downregulation of Antigen Presentation Pathway


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Correction: Noninvasive Detection of Breast Cancer Lymph Node Metastasis Using Carbonic Anhydrases IX and XII Targeted Imaging Probes

Correction: Glutamatergic Pathway Targeting in Melanoma: Single-Agent and Combinatorial Therapies
ABOUT THE COVER

Following treatment with a G-quadruplex ligand, telomestatin, glioma stem cells rapidly developed punctate nuclear 53BP1 foci. Of note, some of these foci colocalized with nontelomeric DNA, thereby representing both telomeric and nontelomeric dysfunction-induced foci, a hallmark of deprotected DNA damage. The loss of tumor stemness is likely associated with a failure in the DNA damage response elicited by telomestatin in glioma stem cells. For details, see the article by Miyazaki and colleagues on page 1268 of this issue.
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