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- 1820 **Continuous Docetaxel Chemotherapy Improves Therapeutic Efficacy in Murine Models of Ovarian Cancer**
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- 1864 **Zinc Protoporphyrin IX Stimulates Tumor Immunity by Disrupting the Immunosuppressive Enzyme Indoleamine 2,3-Dioxygenase**
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1884 **Preclinical Evaluation of Differentially Targeting Dual Virotherapy for Human Solid Cancer**

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1894 **Oligodeoxyribozymes That Cleave β -Catenin Messenger RNA Inhibit Growth of Colon Cancer Cells via Reduction of β -Catenin Response Transcription**

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1903 **Endocytosis of PEGylated Agents Enhances Cancer Imaging and Anticancer Efficacy**

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1913 **High-Content Phenotypic Profiling of Drug Response Signatures across Distinct Cancer Cells**

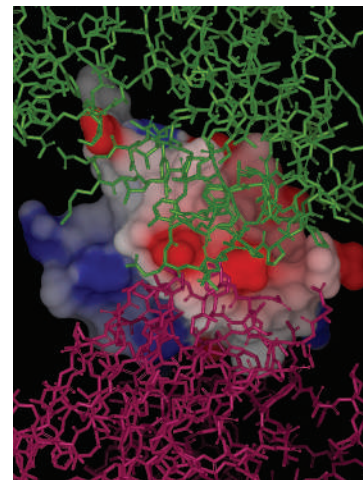
Peter D. Caie, Rebecca E. Walls, Alexandra Ingleston-Orme, Sandeep Daya, Tom Houslay, Rob Eagle, Mark E. Roberts, and Neil O. Carragher

CORRECTION

1927 **Correction: Analysis of Food and Drug Administration-Approved Anticancer Agents in the NCI60 Panel of Human Tumor Cell Lines**

ABOUT THE COVER

A recombinant human phage display-derived antibody, DX-2647, targeting IGF-II blocks the growth of human hepatocellular tumor cells *in vitro* and *in vivo*. DX-2647 blocks the binding of IGF-II to a panel of IGF-BPs and the mannose-6-phosphate receptor (IGF-IIR). A comparison of the crystal complexes of IGF-II with the Fab from DX-2647 (burgundy) and IGF-II with the IGF-IIR (green) reveals a small area of overlap in the two binding domains (Fab versus IGF-IIR) contributing mechanistically to the observed neutralizing biologic activity of this antibody. For details, see article by Dransfield and colleagues on page 1809.



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