Current Preclinical Models for the Advancement of Translational Bladder Cancer Research
David J. DeGraff, Victoria L. Robinson, Jay B. Shah, William D. Brandt, Guru Sonpavde, Yibin Kang, Monica Liebert, Xue-Ru Wu, and John A. Taylor III for the Translational Science Working Group of the Bladder Advocacy Network Think Tank

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Ezia Bello, Giulia Taraboletti, Gennaro Colella, Massimo Zucchetto, Daniele Forestieri, Simonetta A. Licandro, Alexander Berndt, Petra Richter, Maurizio D’Incalci, Ennio Cavalletti, Raffaella Giavazzi, Gabriella Camboni, and Giovanna Damia

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Hiroyuki Sumi, Masato Yabuki, Kenichi Iwai, Megumi Morimoto, Ryosuke Hibino, Masakazu Inazuka, Kentaro Hashimoto, Yohei Kosugi, Kazunobu Aoyama, Shunsuke Yamamoto, Mie Yoshimatsu, Hideki Yamasaki, Ryuichi Tozawa, Tomoyasu Ishikawa, and Sei Yoshida

Correction: TPI-287, a New Taxane Family Member, Reduces the Brain Metastatic Colonization of Breast Cancer Cells

ABOUT THE COVER

_In silico_ molecular modeling of the EGFR exon 20 A763_Y764 insertion mutation. Mutations in this region are predicted to cause significant rearrangement of the C helix (yellow) but do not affect the erlotinib binding pocket directly (erlotinib shown in green). Insertions in _EGFR_ exon 20 that are more distal (3') are expected to result in a greater obstructive effect on erlotinib binding. These predictions suggest a basis for the observed variability of response to EGFR inhibition in patients with different types of _EGFR_ exon 20 insertions. For details, see article by Arcila and colleagues on page 220.
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