

## Erratum to: A Phase 1 dose-escalation study of the safety and pharmacokinetics of once-daily oral foretinib, a multi-kinase inhibitor, in patients with solid tumors

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In the original version of this article, three authors are missing. The complete list of authors and their current and former (at the time the work was performed) affiliations are provided. The additional authors declare no conflicts of interest.

In addition, pharmacodynamics assays performed to obtain absolute plasma concentrations of HGF, VEGF, soluble KDR and soluble Met as described in the original article were incorrectly attributed in the Methods section.

The following text...

“Absolute plasma concentrations of HGF, VEGF, soluble KDR and soluble Met were determined by two-site electrochemiluminescent immunoassay as described previously [1].”

The online version of the original article can be found at <http://dx.doi.org/10.1007/s10637-012-9881-z>.

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...replaces the original text below:

“Plasma levels of soluble MET (sMET), HGF, soluble VEGFR2 (sVEGFR2), and VEGF-A were measured using enzyme-linked immunosorbent assay at Pathway Diagnostics, Malibu, California, USA (now Quest Diagnostics Biomarker Lab, Valencia, California, USA) and at Exelixis (for sMET).” The following citation refers to the text added, above.

[1] Athauda G, Giubellino A, Coleman JA, Horak C, Steeg PS, Lee MJ, Trepel J, Wimberly J, Sun J, Coxon A, Burgess TL, Bottaro DP. c-Met ectodomain shedding rate correlates with malignant potential. *Clin Cancer Res* 2006;12:4154-62.

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