

## **Data Sheet**

WWW.PROBECHEM.COM

Global Supplier of Chemical Probes, Inhibitors & Agonists.

 Product Name
 :
 PF-114

 Cat. No.
 :
 PC-20215

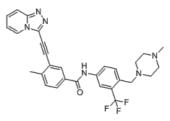
 CAS No.
 :
 1416241-23-0

 Molecular Formula
 :
 C<sub>29</sub>H<sub>27</sub>F<sub>3</sub>N<sub>6</sub>O

 Molecular Weight
 :
 532.57

**Solubility** : 10 mM in DMSO

: Bcr-Abl



## **Biological Activity**

**Target** 

Vamotinib (PF-114) is a potent, selective and orally available inhibitor of native (IC50=0.49 nM) and mutated **BCR/ABL** (IC50=0.7-4 nM, ABL (T315I) IC50=0.78 nM).

PF-114 potently inhibits ABL2, DDR1, DDR2, FMS, FRK, LCK, LYN and PDGFR kinases, but did not inhibit c-SRC, CSK, or c-KIT.

PF-114 is a potent TKI with a more restricted selectivity profile as compared to ponatinib or dasatinib.

PF-114 inhibits the autophosphorylation of BCR/ABL and BCR/ABL-T315I and abolishes factor-independent growth of Ba/F3 cells mediated by BCR/ABL and its resistance mutants.

PF-114 potently inhibits Ph+ patient derived cell lines, abolishes tumor growth in a K562 nude-mouse xenograft model.

PF-114 suppresses growth of Ph+ PD-LTC with non mutational resistance as well as T315I mutation.

PF-114 prolongs the survival of mice with both BCR/ABL- and BCR/ABL-T315I-driven CMLlike disease.

## References

A A Mian, et al. *Leukemia*. 2015 May;29(5):1104-14.

Ivanova ES, et al. Int J Oncol. 2019 Jul;55(1):289-297.

Caution: Product has not been fully validated for medical applications. Lab Use Only!

E-mail: tech@probechem.com