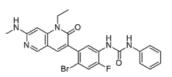


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| DCC-2618 |
|-------------------------|
| PC-62101 |
| 1442472-39-0 |
| $C_{24}H_{21}BrFN_5O_2$ |
| 510.367 |
| c-Kit |
| 10 mM in DMSO |
| |

Data Sheet

Global Supplier of Chemical Probes, Inhibitors & Agonists.



Biological Activity

DCC-2618 (Ripretinib, DCC2618) is a potent, oral inhibitor of singly and doubly mutated **KIT** with IC50 of WT (IC50=4 nM), V654A (8 nM), T670I (18 nM), D816H (5 nM), D816V.

DCC-2618 also inhibits PDGFR α/β , KDR and cFMS, robustly inhibits exon 17, exon 9/13, exon 9/14, and exon 9/17 KIT mutants, as well as exon 11/17 KIT mutants.

DCC-2618 inhibits wild type and mutant KIT phosphorylation in cancer cells, demonstrates the potential to treat KIT mutantdriven cancers including GIST, systemic mastocytosis, AML, or melanoma.

References

Mathias A Schneeweiss, et al. *Blood* 2016 128:1965.

Cancer Discov. 2017 Feb;7(2):121-122. doi: 10.1158/2159-8290.

Caution: Product has not been fully validated for medical applications. Lab Use Only! E-mail: tech@probechem.com