

Product Name : DCC-2618

Synonyms : Ripretinib; DCC2618

**Cat No.** : M11875

**CAS Number** : 1442472-39-0

Molecular Formula : C24H21BrFN5O2

Formula Weight : 510.37

Chemical Name : 1-(4-bromo-5-(1-ethyl-7-(methylamino)-2-oxo-1,2-dihydro-1,6-naphthyridin-3-yl)-2-fluorophenyl)-3-phenylurea

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; also inhibits PDGFR $\alpha/\beta$ , KDR and cFMS, robustly inhibits exon 17,

**Description** : exon 9/13, exon 9/14, and exon 9/17 KIT mutants, as well as exon 11/17 KIT mutants; inhibits wild type and mutant KIT

phosphorylation in cancer cells, demonstrates the potential to treat KIT mutant-driven cancers including GIST, systemic

mastocytosis, AML, or melanoma. Gastric Cancer Phase 3 Clinical

Pathway : Angiogenesis

Target : c-Kit

Receptor : c-Kit

Solubility : DMSO : 45 mg/mL 88.17 mM

**SMILES** : CCN1C2=CC(=NC=C2C=C(C1=0)C3=CC(=C(C=C3Br)F)NC(=0)NC4=CC=CC=C4)NC

**Storage** : (-20°C)

**Stability** :  $\geq 2$  years

Reference :

1. Mathias A Schneeweiss, et al. Blood 2016 128:1965. 2. Cancer Discov. 2017 Feb;7(2):121-122. doi: 10.1158/2159-8290.