

## **Data Sheet**

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 Product Name
 :
 BFH-772

 Cat. No.
 :
 PC-42146

 CAS No.
 :
 890128-81-1

 Molecular Formula
 :
 C<sub>23</sub>H<sub>16</sub>F<sub>3</sub>N<sub>3</sub>O<sub>3</sub>

 Molecular Weight
 :
 439.3867

 Target
 :
 VEGFR

**Solubility** : DMSO: 7.75 mg/mL

## **Biological Activity**

A potent, selective and orally active VEGFR2 tyrosine kinase inhibitor with IC50 of 3 nM; shows 40-fold less potent activity B-RAF, RET and TIE-2, and >500-fold for FLK-1, FLT-1 and FLT-4; inhibits ligand-induced autophosphorylation of RET, PDGFR and KIT with IC50 of 30-160 nM, potently inhibits VEGF-driven angiogenesis in a chamber model and rodent tumor models at daily doses of less than 3 mg/kg.

**Psoriasis** 

Phase 2 Discontinued

## References

Bold G, et al. J Med Chem. 2016 Jan 14;59(1):132-46.

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

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