

Product Name : Vatalanib
Synonyms : PTK787;PTK/ZK;CGP-79787D;CGP-79787;ZK-222584

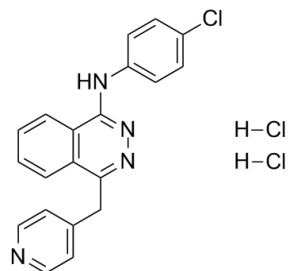
Cat No. : M13374

CAS Number : 212141-51-0

Molecular Formula : C₂₀H₁₇Cl₃N₄

Formula Weight : 419.73

Chemical Name : 1-Phthalazinamine, N-(4-chlorophenyl)-4-(4-pyridinylmethyl)-, hydrochloride (1:2)



Description : A potent, orally available class III receptor tyrosine kinases inhibitor with IC₅₀ of <1 μM for VEGFR, Flt-1, KDR and PDGFRβ; shows no activity against EGFR, FGFR-1, c-Met, and Tie-2, or c-Src, c-Abl, and PKC-α; blocks the VEGF-induced receptor autophosphorylation in CHO cells ectopically expressing the KDR receptor with IC₅₀ of 34 nM; inhibits EGF and PDGF-induced angiogenesis in a growth factor implant model (25-100 mg/kg). Ovarian Cancer Phase 1 Discontinued

Pathway : Angiogenesis

Target : VEGFR

Receptor : PDGFRβ; VEGFR1/FLT1; VEGFR2/Flk1; VEGFR2/KDR; VEGFR3/FLT4

Solubility : 10 mM in DMSO

SMILES : ClC1=CC=C(NC2=NN=C(CC3=CC=NC=C3)C4=C2C=CC=C4)C=C1.[H]Cl.[H]Cl

Storage : (-20°C)

Stability : ≥ 2 years

Reference :

1. Bold G, et al. J Med Chem. 2000 Jun 15;43(12):2310-23. 2. Wood JM, et al. Cancer Res. 2000 Apr 15;60(8):2178-89. 3. Dreves J, et al. Cancer Res. 2000 Sep 1;60(17):4819-24. 4. Hess C, et al. Br J Cancer. 2001 Dec 14;85(12):2010-6.