

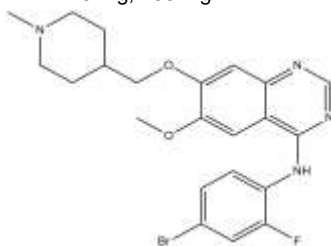
Vandetanib

ALTERNATE NAME: N-(4-Bromo-2-fluorophenyl)-6-methoxy-7-((1-methylpiperidin-4-yl)methoxy)quinazolin-4-amine; ZD6474, Zactima

CATALOG #: 1751-25, 100

AMOUNT: 25 mg, 100 mg

STRUCTURE:



MOLECULAR FORMULA: C₂₂H₂₄BrFN₄O₂

MOLECULAR WEIGHT: 475.35

CAS NUMBER: 443913-73-3

APPEARANCE: White solid

SOLUBILITY: DMSO (30 mg/ml) or EtOH (10 mg/ml)

PURITY: ≥98% by HPLC

STORAGE: Store at -20 °C

DESCRIPTION: A potent inhibitor of kinase insert domain-containing receptor [KDR/vascular endothelial growth factor receptor (VEGFR) 2] tyrosine kinase activity (IC₅₀ = 40 nM). This compound also inhibits fms-like tyrosine kinase 4 (VEGFR3, IC₅₀ = 110 nM) and epidermal growth factor receptor (EGFR/HER1, IC₅₀ = 500 nM) but shows selectivity relating to a range of other tyrosine and serine-threonine kinases.

REFERENCE: Wedge, S.R., *et al.* (2002). *Cancer Res.* 62, 4645-4655.

HANDLING: Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of exposure.

RELATED PRODUCTS:

- Bosutinib (Cat. No. 1584-5, 25)
- Dasatinib (Cat. No. 1586-25, 100)
- Erlotinib, Hydrochloride (Cat. No. 1588-100, 1000)
- Gefitinib (Cat. No. 1589-5, 25)
- Imatinib Mesylate (Cat. No. 1625-100, 1000)
- Lapatinib Ditosylate (Cat. No. 1624-25, 100)
- Nilotinib (Cat. No. 1750-25, 100)
- Sorafenib (Cat. No. 1594-25, 100)
- Sunitinib Malate (Cat. No. 1611-100, 1000)

FOR RESEARCH USE ONLY! Not to be used on humans.