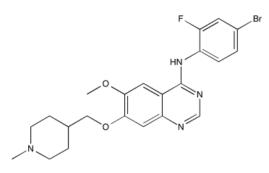


Catalog # 10-2185 Vandetanib

CAS# 443913-73-3 4-(4-Bromo-2-fluoroanilino)-6-methoxy-7-(1-methylpiperidin-4-yl)methoxy]quinazoline ZD6474 Lot # X105611



A potent and selective inhibitor of VEGFR2, (KDR $IC_{50} = 40 \text{ nM}$). Also inhibits fms-like tyrosine kinase 4 (VEGFR3, $IC_{50} = 110 \text{ nM}$) and epidermal growth factor receptor (EGFR/HER1, $IC_{50} = 500 \text{ nM}$) but shows selectivity against a panel of other tyrosine and serine-threonine kinases. Inhibits colony formation in seven cancer cell lines.² Displays anti-angiogenesis activity.³

- 1) Wedge et al. (2002), ZD6474 inhibits vascular endothelial growth factor signaling, angiogenesis and tumor growth following oral administration; Cancer Res., **62** 4645
- 2) Ciardiello et al. (2003), Antitumor effects of ZD6474, a small molecule vascular endothelial growth factor receptor tyrosine kinase inhibitor, with additional activity against epidermal growth factor receptor tyrosine kinase; Clin. Cancer Res., **9** 1546
- 3) Herbst et al. (2007), Vandetanib (ZD6474): an orally available receptor tyrosine kinase inhibitor that selectively targets pathways critical for tumor growth and angiogenesis; Expert Opin. Investig. Drugs, **16** 239

PHYSICAL DATA

Molecular Weight:	475.35
Molecular Formula:	$C_{22}H_{24}BrFN_4O_2$
Purity:	98% by TLC
	NMR: (Conforms)
Solubility:	DMSO (up to 30 mg/ml), or Ethanol (up to 10 mg/ml with warming)
Physical Description:	White or off-white solid
Storage and Stability:	Store as supplied at -20°C for up to 2 years from the date of purchase. Solutions in
	DMSO or ethanol may be stored at -20°C for up to 1 month.

Materials provided by Focus Biomolecules are for laboratory research use only and are not intended for human or veterinary applications.