Web: www.taiclone.com
Tel: +886-2-2735-9682
Email: order@taiclone.com

# Vandetanib (hydrochloride)

## Catalog No: tcsc1592

### Available Sizes

**Size:** 25mg

**Size:** 100mg

**Size:** 200mg

### Specifications

**CAS No:**
524722-52-9

**Formula:**
$C_{22}H_{25}BrClFN_4O_2$

**Pathway:**
Protein Tyrosine Kinase/RTK;Autophagy

**Target:**
VEGFR;Autophagy

**Purity / Grade:**
>98%

**Solubility:**
10 mM in DMSO

**Alternative Names:**
ZD6474 hydrochloride

**Observed Molecular Weight:**
511.81

## Product Description

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Vandetanib hydrochloride is a potent inhibitor of **VEGFR2** with **$IC_{50}$** of 40 nM.

IC50 & Target: IC50: 40 nM (VEGFR2)

***In Vitro:*** Vandetanib inhibits VEGFR3 and EGFR with $IC_{50}$ of 110 nM and 500 nM, respectively. Vandetanib is not sensitive to PDGFRβ, Flt1, Tie-2 and FGFR1 with $IC_{50}$ of 1.1-3.6 μM, while almost has no activity against MEK, CDK2, c-Kit, erbB2, FAK, PDK1, Akt and IGF-1R with $IC_{50}$ above 10 μM. Vandetanib inhibits VEGF-, EGF- and bFGF-stimulated HUVEC proliferation with $IC_{50}$ of 60 nM, 170 nM and 800 nM, with no effect on basal endothelial cell growth. Vandetanib inhibits tumor cell growth with $IC_{50}$ of 2.7 μM (A549) to 13.5 μM (Calu-6)[1]. Odanacatib is a weak inhibitor of antigen presentation, measured in a mouse B cell line ($IC_{50}$=1.5±0.4 μM), compared to the Cat S inhibitor LHVS ($IC_{50}$=0.001 μM) in the same assay. Odanacatib also shows weak inhibition of the processing of the MHC II invariant chain protein Iip10 in mouse splenocytes compared to LHVS (minimum inhibitory concentration 1-10 μM versus 0.01 μM, respectively)[2]. Vandetanib suppresses phosphorylation of VEGFR-2 in HUVECs and EGFR in hepatoma cells and inhibits cell proliferation[4].

***In Vivo:*** Vandetanib (15 mg/kg, p.o.) has a superior anti-tumor effect than gefitinib in the H1650 xenograft model, and suppresses tumor growth with $IC_{50}$ of 3.5±1.2 μM[3]. In tumor-bearing mice, vandetanib (50 or 75 mg/kg) suppresses phosphorylation of VEGFR-2 and EGFR in tumor tissues, significantly reduces tumor vessel density, enhances tumor cell apoptosis, suppresses tumor growth, improves survival, reduces number of intrahepatic metastases, and upregulates VEGF, TGF-α, and EGF in tumor tissues[4].