

# NVP-ACC789

# Catalog No: tcsc0015997

Available Sizes

Size: 5mg

Size: 10mg

Size: 25mg

Size: 50mg

**Specifications** 

### CAS No:

300842-64-2

### Formula:

 $\mathsf{C}_{21}^{}\mathsf{H}_{17}^{}\mathsf{BrN}_{4}^{}$ 

## Pathway:

Protein Tyrosine Kinase/RTK;Protein Tyrosine Kinase/RTK

# Target:

VEGFR;PDGFR

## Purity / Grade:

>98%

## Solubility:

DMSO : 60 mg/mL (148.04 mM; Need ultrasonic and warming)

## Alternative Names:

ACC-789;ZK202650

# **Observed Molecular Weight:**

405.29

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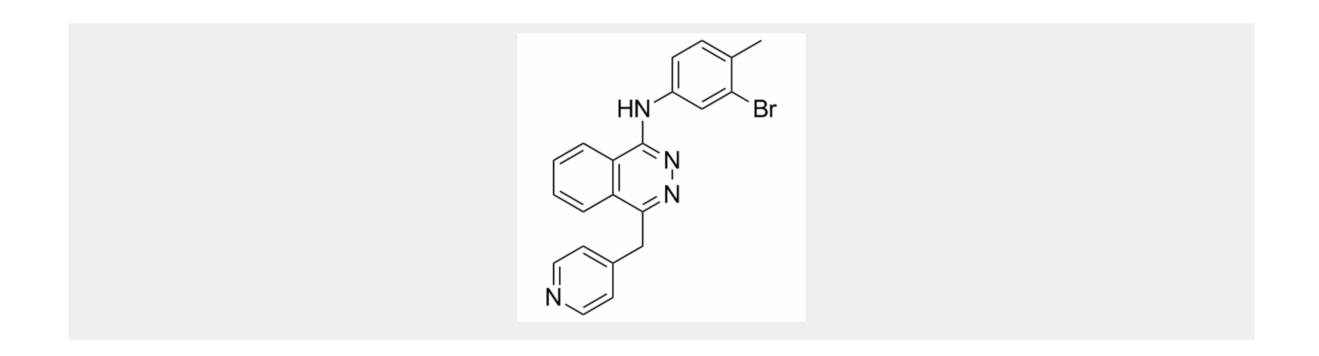
## **Product Description**

NVP-ACC789 is an inhibitor of human VEGFR-1, VEGFR-2 (mouse VEGFR-2), VEGFR-3 and PDGFR- $\beta$  with IC<sub>50</sub>s of 0.38, 0.02 (0.23), 0.18, 1.4  $\mu$ M, respectively.

IC50 & Target: IC50: 0.38  $\mu$ M (human VEGFR-1), 0.02  $\mu$ M (human VEGFR-2), 0.23  $\mu$ M (mouse VEGFR-2), 0.18  $\mu$ M (human VEGFR-3), 1.4  $\mu$ M (human PDGFR- $\beta$ )<sup>[1]</sup>

*In Vitro:* The enzymatic kinase assays demonstrate that NVP-ACC789 is an inhibitor of human VEGFR-1, VEGFR-2 (mouse VEGFR-2), VEGFR-3 and PDGFR- $\beta$  with IC<sub>50</sub>s of 0.38, 0.02 (0.23), 0.18, 1.4  $\mu$ M, respectively. In VEGF-treated cultures, addition of the VEGFR-2 inhibitor NVP-ACC789 reduces BME cell number to baseline levels from 1  $\mu$ M. Likewise, bFGF-induced BME cell proliferation is reduced markedly by NVP-ACC789 from 1 to 10  $\mu$ M, without however reaching basal levels. NVP-ACC789 is found to be a potent inhibitor of VEGF-induced HUVE cell proliferation with an IC<sub>50</sub> of 1.6 nM. NVP-ACC789 also completely inhibits VEGF-induced BME and BAE cell invasion and VEGF-C-induced BAE cell invasion. The inhibition is dose-dependent in both cell types with a maximal effect from 1  $\mu$ M<sup>[1]</sup>.

*In Vivo:* NVP-ACC789 which is given in daily oral doses for 6 days blocks VEGF-induced angiogenesis in a dose-dependent manner. NVP-ACC789 also inhibits the response to bFGF to some extent, but the dose-response curve is not linear for NVP-ACC789<sup>[1]</sup>.



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