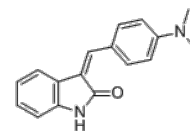


**Product Name** : SU4312  
**Cat. No.** : PC-61860  
**CAS No.** : 5812-07-7  
**Molecular Formula** : C<sub>17</sub>H<sub>16</sub>N<sub>2</sub>O  
**Molecular Weight** : 264.3  
**Target** : VEGFR  
**Solubility** : 10 mM in DMSO



## Biological Activity

SU 4312 (NSC 86429) is a selective, cell-permeable, brain-penetrant inhibitor of **VEGFR2** and PDGFR tyrosine kinase with IC<sub>50</sub> of 0.8 and 19.4 uM, respectively.

SU 4312 displays no significant inhibitory activity against EGFR, HER2, and IGF-1R (>100 uM).

SU 4312 exhibits neuroprotection against MPP(+) at least partly via selective and direct inhibition of nNOS.

SU 4312 inhibits VEGF-dependent angiogenesis in a zebrafish assay (IC<sub>50</sub>=1.8 uM) without affecting normal cells.

## References

Sun L, et al. J Med Chem. 1998 Jul 2;41(14):2588-603.

Tran TC, et al. Cancer Res. 2007 Dec 1;67(23):11386-92.

Cui W, et al. Br J Pharmacol. 2013 Mar;168(5):1201-14.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

E-mail: tech@probechem.com