

Catalog # 10-1027 SU-4312

CAS# 5812-07-7
3-(4-Dimethylaminobenzylidene)-1,3-dihydroindol-2-one
Lot # X101207

Potent selective inhibitor of VEGFR kinase (Flk-1 and KDR). Inhibition of the activated kinase (IC₅₀ = 0.4 μ M) is 100-fold less than inhibition of the unactivated kinase (IC₅₀ = 0.04 μ M). Also inhibits PDGFR kinase (IC₅₀ = 19.4 μ M).

- 1) Kendall et al. (1999), Vascular endothelial growth factor receptor KDR tyrosine kinase activity is increased by autophosphorylation of two activation loop tyrosine residues; J. Biol. Chem., **274** 6453
- 2) Sun et al. (1998), Synthesis and biological evaluations of 3-substituted indolin-2-ones: a novel class of tyrosine kinase inhibitors that exhibit selectivity toward particular receptor tyrosine kinases; J. Med. Chem., **41** 2588

PHYSICAL DATA

 $\begin{array}{lll} \mbox{Molecular Weight:} & 264.33 \\ \mbox{Molecular Formula:} & C_{17}\mbox{H}_{16}\mbox{N}_2\mbox{O} \\ \mbox{Purity:} & 98\% \ \mbox{by TLC} \end{array}$

NMR: (Conforms)

Solubility: DMSO (up to 25 mg/ml)

Physical Description: Orange solid

Storage and Stability: Store as supplied at -20°C for up to 1 year from the date of purchase. Solutions

in DMSO may be stored at -20°C for up to 3 months.

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