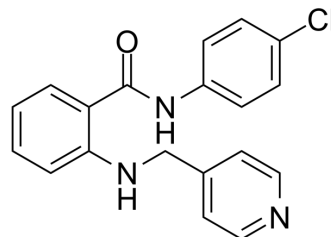


VEGFR-IN-1

Cat. No.:	HY-101219
CAS No.:	269390-69-4
Molecular Formula:	C ₁₉ H ₁₆ ClN ₃ O
Molecular Weight:	337.8
Target:	VEGFR; c-Kit; EGFR
Pathway:	Protein Tyrosine Kinase/RTK; JAK/STAT Signaling
Storage:	-20°C, protect from light, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light, stored under nitrogen)



SOLVENT & SOLUBILITY

In Vitro	DMSO : 125 mg/mL (370.04 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg
		Concentration				
		1 mM		2.9603 mL	14.8017 mL	29.6033 mL
		5 mM		0.5921 mL	2.9603 mL	5.9207 mL
	10 mM		0.2960 mL	1.4802 mL	2.9603 mL	
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.08 mg/mL (6.16 mM); Suspended solution; Need ultrasonic 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (6.16 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	VEGFR-IN-1 (compound 3) is a potent angiogenesis inhibitor with IC ₅₀ s of 0.02, 0.18, 0.24 7.3, and 7 μM for KDR, Flt-1, c-Kit, EGF-R, and c-Src, respectively ^[1] .		
IC₅₀ & Target	KDR 0.02 μM (IC ₅₀)	Flt-1 0.18 μM (IC ₅₀)	EGFR 0.24 μM (IC ₅₀)

REFERENCES

[1]. Furet P, et al. Identification of a new chemical class of potent angiogenesis inhibitors based on conformational considerations and database searching. Bioorg Med

Caution: Product has not been fully validated for medical applications. For research use only.

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