Inhibitors

RIPK1-IN-4

Cat. No.: HY-18901 CAS No.: 1481641-08-0 Molecular Formula: $C_{23}H_{23}N_5O_2$ Molecular Weight: 401.46 RIP kinase Target:

Storage: Powder

3 years $4^{\circ}C$ 2 years

-80°C In solvent 2 years

-20°C

Apoptosis

-20°C 1 year

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

Pathway:

DMSO: 250 mg/mL (622.73 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.4909 mL	12.4545 mL	24.9091 mL
	5 mM	0.4982 mL	2.4909 mL	4.9818 mL
	10 mM	0.2491 mL	1.2455 mL	2.4909 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 (5.18 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (5.18 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 1.44 mg/mL (3.59 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	RIPK1-IN-4 (compound 8) is a potent and selective type II kinase inhibitor of receptor interacting protein 1 (RIP1) kinase and		
	binds to a DLG-out inactive form of RIP1 with an IC $_{50}$ s of 16 nM and 10 nM for RIP1 and ADP-Glo kinase $^{[1]}$.		

RIPK1 IC₅₀ & Target

REFERENCES

1]. Harris PA, et al. Discovery of Small Molecule RIP1 Kinase Inhibitors for the Treatment of Pathologies Associated with Necroptosis. ACS Med Chem Lett. 2013 Nov 4;4(12):1238-43.						
	Caution: Product has no	ot been fully validated for me	edical applications. For research use only.			
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Page 2 of 2 www.MedChemExpress.com