Src Inhibitor 3

Cat. No.:	HY-130254				
CAS No.:	2380027-49-4				
Molecular Formula:	C ₃₄ H ₃₂ CIFN ₈ O ₄				
Molecular Weight:	671.12				
Target:	Src				
Pathway:	Protein Tyrosine Kinase/RTK				
Storage:	Powder	-20°C	3 years		
		4°C	2 years		
	In solvent	-80°C	6 months		
		-20°C	1 month		

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SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (149.00 mM; Need ultrasonic)					
	Preparing Stock Solutions	Mass Solvent Concentration	1 mg	5 mg	10 mg	
		1 mM	1.4900 mL	7.4502 mL	14.9005 mL	
		5 mM	0.2980 mL	1.4900 mL	2.9801 mL	
		10 mM	0.1490 mL	0.7450 mL	1.4900 mL	
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (3.73 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (3.10 mM); Clear solution					

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Description	Src Inhibitor 3 is a potent, orally active c-terminal Src kinase (CSK) with IC ₅₀ values below 3 nM and 4 nM in CSK HTRF and Caliper assay, respectively. Src Inhibitor 3 shows the ability to increase T cell proliferation induced by T cell receptor signaling ^[1] .			
IC ₅₀ & Target	IC50: <3 nM (c-terminal Src kinase in CSK HTRF assay), 4 nM (c-terminal Src kinase in Caliper assay) ^[1]			
In Vivo	Src Inhibitor 3 reduces inhibitory LCK phosphorylation in vivo upon oral dosing and shows the ability to enhance T cell activation in response to antigen stimulation ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

Product Data Sheet

ΞN

CUSTOMER VALIDATION

• Int J Mol Sci. 2021 Aug 27;22(17):9289.

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REFERENCES

[1]. O'Malley DP, et al. Discovery of Pyridazinone and Pyrazolo[1,5-a]pyridine Inhibitors of C-Terminal Src Kinase. ACS Med Chem Lett. 2019 Sep 25;10(10):1486-1491.

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

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