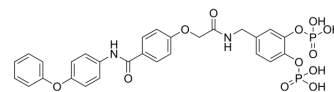


Stafib-2

Cat. No.:	HY-112648		
CAS No.:	2097938-74-2		
Molecular Formula:	C ₂₈ H ₂₆ N ₂ O ₁₂ P ₂		
Molecular Weight:	644.46		
Target:	STAT		
Pathway:	JAK/STAT Signaling; Stem Cell/Wnt		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 50 mg/mL (77.58 mM; Need ultrasonic)					
		Solvent Concentration	Mass	1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM		1.5517 mL	7.7584 mL	15.5169 mL
		5 mM		0.3103 mL	1.5517 mL	3.1034 mL
10 mM			0.1552 mL	0.7758 mL	1.5517 mL	
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (3.88 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (3.88 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (3.88 mM); Clear solution 					

BIOLOGICAL ACTIVITY

Description	Stafib-2 is a potent and selective inhibitor of the transcription factor STAT5b, with an IC ₅₀ of 82 nM and 1.7 μM for STAT5b and STAT5a, respectively. Stafib-2 exhibits poor cell permeability ^[1] .
IC ₅₀ & Target	IC ₅₀ : 82 nM (STAT5b) ^[1]
In Vitro	Stafib-2 has an extremely high affinity for STAT5b (K _i =8.8 nM) ^[1] . Stafib-2 (3-10 μM; 4-48 h) does not show significant activity in K562 and MDA-MB-231 cells ^[1] .

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Elumalai N, et, al. Rational development of Stafib-2: a selective, nanomolar inhibitor of the transcription factor STAT5b. Sci Rep. 2017 Apr 11;7(1):819.

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

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