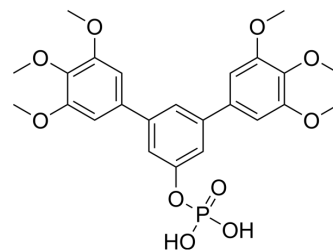


Stafia-1

Cat. No.:	HY-136546		
CAS No.:	2582757-90-0		
Molecular Formula:	$C_{24}H_{27}O_{10}P$		
Molecular Weight:	506.44		
Target:	STAT		
Pathway:	JAK/STAT Signaling; Stem Cell/Wnt		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 250 mg/mL (493.64 mM; Need ultrasonic)					
	Preparing Stock Solutions	<div><div>Solvent</div><div>Concentration</div></div>	Mass	1 mg	5 mg	10 mg
		1 mM		1.9746 mL	9.8728 mL	19.7457 mL
		5 mM		0.3949 mL	1.9746 mL	3.9491 mL
		10 mM		0.1975 mL	0.9873 mL	1.9746 mL
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (4.11 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (4.11 mM); Clear solution					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.11 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	Stafia-1 is a potent STAT5a inhibitor ($K_i=10.9 \mu\text{M}$, $\text{IC}_{50}=22.2 \mu\text{M}$). Stafia-1 displays high selectivity over STAT5b and other STAT family members ^[1] .	
IC ₅₀ & Target	STAT5a	STAT5a
	10.9 μM (K_i)	22.2 μM (IC_{50})
In Vitro	Stafia-1 shows at least 9-fold selectivity over STAT5b and higher selectivity against other STAT family members ^[1] .	

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

REFERENCES

[1]. Natarajan K, et al. Stafia-1: a STAT5a-Selective Inhibitor Developed via Docking-Based Screening of in Silico O-Phosphorylated Fragments. Chemistry. 2020;26(1):148-154.

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

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