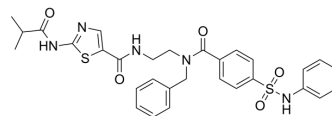


TH470

Cat. No.:	HY-151539		
CAS No.:	2834739-51-2		
Molecular Formula:	C ₃₀ H ₃₁ N ₅ O ₅ S ₂		
Molecular Weight:	605.73		
Target:	LIM Kinase (LIMK)		
Pathway:	Cell Cycle/DNA Damage		
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 : 100 mg/mL (165.09 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
			10 mg	
	Preparing Stock Solutions	1 mM	5 mM	10 mM
		1.6509 mL	8.2545 mL	16.5090 mL
		0.3302 mL	1.6509 mL	3.3018 mL
		0.1651 mL	0.8255 mL	1.6509 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: 2.5 mg/mL (4.13 mM); Clear solution; Need ultrasonic 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (4.13 mM); Clear solution; Need ultrasonic			

BIOLOGICAL ACTIVITY

Description	TH470 is a highly selective LIMK1/2 (LIM kinase1/2) inhibitor (LIMK1 IC ₅₀ =9.8 nM; LIMK2 IC ₅₀ =13 nM), and can be used in orphan disease research ^[1] .	
IC₅₀ & Target	LIMK1 9.8 nM (IC ₅₀)	LIMK2 13 nM (IC ₅₀)
In Vitro	TH470 (0.05-5 μM; 12 h) inhibits the growth of neurite ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay ^[1]	

Cell Line:	N1E-115 cells transfected with siRNAs against FMR1 (siFMR1) or control siRNAs (siCtr)
Concentration:	0.05, 0.5, or 5 μ M
Incubation Time:	12 hours
Result:	Inhibited neurite outgrowth in a dose-dependent manner.

REFERENCES

[1]. Hanke T, et al. Development and Characterization of Type I, Type II, and Type III LIM-Kinase Chemical Probes. J Med Chem. 2022 Oct 13;65(19):13264-13287.

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

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