MARK4 inhibitor 4

Cat. No.:	HY-154988		
Molecular Formula:	C ₃₃ H ₂₇ ClN ₄ O)3	
Molecular Weight:	563.05		
Target:	Others		
Pathway:	Others		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month

SOLVENT & SOLUBILITY

	Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	1.7760 mL	8.8802 mL	17.7604 mL
	5 mM	0.3552 mL	1.7760 mL	3.5521 mL
	10 mM	0.1776 mL	0.8880 mL	1.7760 mL
Please refer to the sol	ubility information to select the app	propriate solvent.		i
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	Stock Solutions Please refer to the sol 1. Add each solvent c	Stock Solutions 5 mM 10 mM Please refer to the solubility information to select the app	Stock Solutions 5 mM 0.3552 mL 10 mM 0.1776 mL Please refer to the solubility information to select the appropriate solvent. 1. Add each solvent one by one: 10% DMSO >> 90% corn oil	Stock Solutions 5 mM 0.3552 mL 1.7760 mL 10 mM 0.1776 mL 0.8880 mL Please refer to the solubility information to select the appropriate solvent. 1. Add each solvent one by one: 10% DMSO >> 90% corn oil

BIOLOGICAL ACTIV	ІТҮ		
Description	MARK4 inhibitor 4 is a MARK4 inhibitor with an IC ₅₀ of 1.49 μM. MARK4 inhibitor 4 can be used in cancer and tau lesion- related research ^[1] .		
IC ₅₀ & Target	IC ₅₀ : 1.49 μM (MARK4)		
In Vitro	EC ₅₀ s value of 2.16 and 3.	ound 23c) (1-20 μM; 24 h) has obvious inhibitory effect on the growth of Hela and U87MG cells, with 51 μM, respectively ^[1] . tly confirmed the accuracy of these methods. They are for reference only.	
	Cell Line:	Hela, U87MG, U251, MDA-MB-435, HGFs cells	
	Concentration:	1-20 μΜ	

Product Data Sheet

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Incubation Time:	24 h
Result:	Showed obvious cytotoxicity in Hela and U87MG cell lines.

REFERENCES

[1]. Voura M, et al. Synthesis, Structural Modification, and Bioactivity Evaluation of Substituted Acridones as Potent Microtubule Affinity-Regulating Kinase 4 Inhibitors[J]. ACS Pharmacology & Translational Science, 2023.

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

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