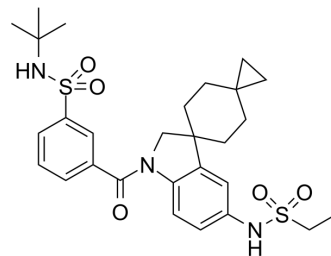


KIF18A-IN-6

Cat. No.:	HY-153065		
CAS No.:	2914879-10-8		
Molecular Formula:	C ₂₈ H ₃₇ N ₃ O ₅ S ₂		
Molecular Weight:	559.74		
Target:	Microtubule/Tubulin		
Pathway:	Cell Cycle/DNA Damage; Cytoskeleton		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 125 mg/mL (223.32 mM; Need ultrasonic)						
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg	
				1 mM	1.7865 mL	8.9327 mL	17.8654 mL
				5 mM	0.3573 mL	1.7865 mL	3.5731 mL
				10 mM	0.1787 mL	0.8933 mL	1.7865 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: ≥ 2.08 mg/mL (3.72 mM); Clear solution						

BIOLOGICAL ACTIVITY

Description	KIF18A-IN-6 (Compound 134) is an orally active KIF18A inhibitor with an IC ₅₀ of 0.016 μM against KIF18A microtubule-dependent ATPase activity ^[1] .				
IC ₅₀ & Target	IC ₅₀ : 0.016 μM (KIF18A microtubule-dependent ATPase activity) ^[1]				
In Vitro	KIF18A-IN-6 (Compound 134; 7 days) inhibits JIMT-1, HCC-15 and NIH-OVCAR3 cells viability with IC ₅₀ s of 0.0040, 0.0051 and 0.0051 μM, respectively ^[1] .				
	MCE has not independently confirmed the accuracy of these methods. They are for reference only.				
	Cell Viability Assay ^[1]				
	Cell Line:	HCC-15, JIMT-1 and NIH-OVCAR3			
	Concentration:				

	Incubation Time:	7 days
	Result:	Inhibited cell viability with IC ₅₀ s of 0.0040, 0.0051 and 0.0051 μM against JIMT-1, HCC-15 and NIH-OVCAR3 cells, respectively.
In Vivo	KIF18A-IN-6 (Compound 134; 10-60 mg/kg; p.o.; twice or once a day for 1 month) inhibits HCC15 and OVCAR3 tumor growth in mice ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	SCID Beige mice, HCC15 tumor model ^[1]
	Dosage:	10, 30 and 60 mg/kg
	Administration:	PO, twice a day for 1 month
	Result:	Inhibited tumor growth by 61±10%, 89±7% and 94±5% at 10, 30 and 60 mg/kg, respectively.
	Animal Model:	Balb/C nude mice, OVCAR3 tumor model ^[1]
	Dosage:	10, 30 and 60 mg/kg
	Administration:	PO, twice or once a day for 1 month
	Result:	Completely inhibited tumor growth (over 100%) over 30 mg/kg.

REFERENCES

[1]. COGAN, et al. SPIRO INDOLINE INHIBITORS OF KIF18A. Patent WO2023028564.

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

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