Mevalonic acid lithium salt

Cat. No.: CAS No.: Molecular Formula: Molecular Weight: Target: Pathway:	HY-113071A 2618458-93-6 C ₆ H ₁₁ LiO ₄ 154.09 Endogenous Metabolite Metabolic Enzyme/Protease	HO O HO OLi
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)	

SOLVENT & SOLUBILITY

	DMSO : 50 mg/mL (32	DMSO : 50 mg/mL (324.49 mM; Need ultrasonic)					
		Solvent Mass Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	6.4897 mL	32.4486 mL	64.8971 mL		
		5 mM	1.2979 mL	6.4897 mL	12.9794 mL		
	10 mM	0.6490 mL	3.2449 mL	6.4897 mL			
	Please refer to the so	lubility information to select the app	propriate solvent.				
In Vivo		1. Add each solvent one by one: PBS Solubility: 25 mg/mL (162.24 mM); Clear solution; Need ultrasonic					
		 Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (16.22 mM); Clear solution 					
		3. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (16.22 mM); Clear solution					
		4. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (16.22 mM); Clear solution					

BIOLOGICAL ACTIVITY				
Description	Mevalonic acid (MVA) lithium salt is a precursor substance of the mevalonate pathway, which is essential for cell growth and proliferation. Mevalonic acid lithium salt is effective in inhibiting <u>Simvastatin</u> (HY-17502)-induced decrease in C2C12 cell viability in vitro. Mevalonic acid lithium salt can be used in studies of myopathy and heart failure ^{[1][2]} .			
In Vitro	Mevalonic acid lithium salt (80, 90, 100, 110 μM; 72 h) shows prevention of simvastatin-induced loss of viability of C2C12myotube cells in vitro ^[1] .			

Product Data Sheet

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MCE has not independe Cell Viability Assay ^[1]	MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay ^[1]		
Cell Line:	C2C12 cells (simvastatin-induced)		
Concentration:	80, 90, 100, 110 μM		
Incubation Time:	72 h		
Result:	Showed no decline in cell viability.		

CUSTOMER VALIDATION

- Acta Pharm Sin B. 2023 Apr 17.
- Acta Pharmacol Sin. 2021 Feb 19.
- Cell Biosci. 2021 Oct 9;11(1):179.
- Biochim Biophys Acta Mol Cell Biol Lipids. 2022 Aug 16;159217.
- Biomedicines. 2022, 10(10), 2489.

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REFERENCES

[1]. Moschetti A, et al. Coenzyme Q nanodisks counteract the effect of statins on C2C12 myotubes. Nanomedicine. 2021 Oct;37:102439.

[2]. Soma MR, et al. Cholesterol and mevalonic acid modulation in cell metabolism and multiplication. Toxicol Lett. 1992 Dec;64-65 Spec No:1-15.

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

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