D-Mannose

®

MedChemExpress

Cat. No.:	HY-N0379		
CAS No.:	3458-28-4		
Molecular Formula:	C ₆ H ₁₂ O ₆		
Molecular Weight:	180.16		
Target:	Endogenous Metabolite		
Pathway:	Metabolic Enzyme/Protease		
Storage:	4°C, stored under nitrogen * In solvent : -80°C. 6 months: -20°C. 1 month (stored under nitrogen)		

Product Data Sheet

HO

OH OH

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SOLVENT & SOLUBILITY

In Vitro	H ₂ O : ≥ 50 mg/mL (277.53 mM) DMSO : 50 mg/mL (277.53 mM; Need ultrasonic) * "≥" means soluble, but saturation unknown.						
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	5.5506 mL	27.7531 mL	55.5062 mL		
		5 mM	1.1101 mL	5.5506 mL	11.1012 mL		
		10 mM	0.5551 mL	2.7753 mL	5.5506 mL		
	Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: PBS Solubility: 150 mg/mL (832.59 mM); Clear solution; Need ultrasonic						
	2. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (13.88 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (13.88 mM); Clear solution						
	4. Add each solvent c Solubility: ≥ 2.5 mg	one by one: 10% DMSO >> 90% cor g/mL (13.88 mM); Clear solution	n oil				

BIOLOGICAL ACTIVITY					
Description	D-Mannose is a C-2 superpolymer of glucose that occurs naturally in many plants and fruits. D-Mannose has anti- inflammatory and antitumor activity. D-Mannose plays an important role in immune regulation ^{[1][2][3]} .				
IC ₅₀ & Target	Human Endogenous Metabolite				

In Vitro	D-Mannose (5.5, 11, 25 mM, 24 h) inhibits macrophage activation by inhibiting the production of IL-1β ^[1] . D-Mannose (25 mM, 24-72 h) stimulates Treg cell differentiation in human and mouse cells by promoting TGF-β activation and inhibiting immunopathology ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. RT-PCR ^[1]			
	Cell Line:	BMDMs		
	Concentration:	5.5, 11, 25 mM		
	Incubation Time:	24 h		
	Result:	Decreased LPS-induced Il1b gene expression dose-dependently		
In Vivo	D-Mannose (2g/kg, intraperitoneally injected 6 times per hour, killed 3 h later) can protect mice from DSS-induced ulcerative colitis ^[1] . D-Mannose (1.1 M D-mannose dissolved in water for 2 months) alleviates bone loss in mice through Treg cell proliferation and intestinal microbiome-dependent anti-inflammatory effects ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			
	Animal Model:	C57BL/6 mice ^[1]		
	Dosage:	20% (w/v)		
	Administration:	i.g. pre-treated for 2 weeks		
	Result:	Opposed loss of body weight induced by DSS and delayed colitis progression.		

REFERENCES

[1]. Torretta S, et al. D-mannose suppresses macrophage IL-1β production. Nat Commun. 2020 Dec 11;11(1):6343.

[2]. Zhang D, et al. D-mannose induces regulatory T cells and suppresses immunopathology. Nat Med. 2017 Sep;23(9):1036-1045.

[3]. Liu H, et al. D-mannose attenuates bone loss in mice via Treg cell proliferation and gut microbiota-dependent anti-inflammatory effects. Ther Adv Chronic Dis. 2020 Apr 17;11:2040622320912661.

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

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA