Voglibose

Cat. No.:	HY-B0025		
CAS No.:	83480-29-9		
Molecular Formula:	C ₁₀ H ₂₁ NO ₇		
Molecular Weight:	267.28		
Target:	Glucosidase		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 vear

SOLVENT & SOLUBILITY

In Vitro	H ₂ O : 250 mg/mL (935.35 mM; Need ultrasonic) DMSO : 100 mg/mL (374.14 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg	
		1 mM	3.7414 mL	18.7070 mL	37.4139 mL	
		5 mM	0.7483 mL	3.7414 mL	7.4828 mL	
	10 mM	0.3741 mL	1.8707 mL	3.7414 mL		
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: PBS Solubility: 100 mg/mL (374.14 mM); Clear solution; Need ultrasonic					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (9.35 mM); Clear solution					
	3. Add each solvent o Solubility: ≥ 2.5 m	one by one: 10% DMSO >> 90% cor g/mL (9.35 mM); Clear solution	n oil			

Description	Voglibose is an orally active alpha-glucosidase inhibitor that prevents the development of colorectal precancerous lesions induced by obesity and diabetes. Voglibose reduces oxidative stress in an inflammatory environment and inhibits the insulin-like growth factor/insulin-like growth factor-1 receptor (IGF/IGF-1R) functional axis.		
In Vitro	Voglibose (50 μM; 72 h) has no effect on SW480 and SW837 cells proliferation in both situation with 10 nM IGF-1 or not ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

Product Data Sheet

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In Vivo	Voglibose (10 mg/kg; po for 15 weeks) has an inhibitory effect on the development of colorectal pre-neoplastic lesions in Azoxymethane (HY-111375)-induced diabetic and obese db/db mice model ^[1] .
	MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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

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