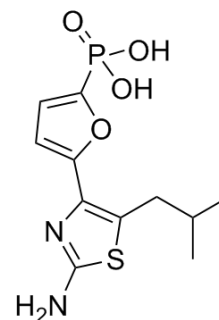


MB05032

Cat. No.:	HY-16307		
CAS No.:	261365-11-1		
Molecular Formula:	C ₁₁ H ₁₅ N ₂ O ₄ PS		
Molecular Weight:	302.29		
Target:	Phosphatase		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 50 mg/mL (165.40 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	3.3081 mL	16.5404 mL	33.0808 mL
		5 mM	0.6616 mL	3.3081 mL	6.6162 mL
10 mM		0.3308 mL	1.6540 mL	3.3081 mL	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (8.27 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (8.27 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (8.27 mM); Clear solution 				

BIOLOGICAL ACTIVITY

Description	MB05032 is a special and efficacious GNG inhibitor targeted the AMP binding site of fructose 1,6-bisphosphatase (FBPase) with an IC ₅₀ value of 16 nM.
IC₅₀ & Target	IC ₅₀ : 16 nM (Human Liver FBPase) ^[1]
In Vivo	MB06322 (3/6-300 mg/kg (young/aged Zucker diabetic fatty (ZDF) rats); p.o.; once) results in dose-dependent glucose lowering in young ZDF rats with mild diabetes and aged ZDF rats with overt diabetes ^[1] .

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	ZDF rats (8- to 9-week-old (mild diabetes) and 12- to 13-week-old (overt diabetes)) ^[1]
Dosage:	3, 6, 10, 30, 300 mg/kg (young ZDF rats); 6, 10, 30, 300 mg/kg (aged ZDF rats)
Administration:	Oral administration; once
Result:	Results in dose-dependent glucose lowering.

CUSTOMER VALIDATION

- Nat Med. 2018 Sep;24(9):1395-1406.
- Cell Metab. 2018 Aug 7;28(2):243-255.e5.

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REFERENCES

- [1]. Erion MD, et al. MB06322 (CS-917): A potent and selective inhibitor of fructose 1,6-bisphosphatase for controlling gluconeogenesis in type 2 diabetes. Proc Natl Acad Sci U S A. 2005 May 31;102(22):7970-5.
- [2]. Zhang Y, et al. Fructose-1,6-bisphosphatase regulates glucose-stimulated insulin secretion of mouse pancreatic beta-cells. Endocrinology. 2010 Oct;151(10):4688-95.

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

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