Golgicide A

Cat. No.:	HY-100540			
CAS No.:	1139889-93-2			
Molecular Formula:	C ₁₇ H ₁₄ F ₂ N ₂			
Molecular Weight:	284.3			
Target:	Enterovirus			
Pathway:	Anti-infection			
Storage:	Powder	-20°C	3 years	
		4°C	2 years	
	In solvent	-80°C	2 years	
		-20°C	1 year	

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In Vitro	DMSO : 100 mg/mL (351.74 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg	
		1 mM	3.5174 mL	17.5871 mL	35.1741 mL	
		5 mM	0.7035 mL	3.5174 mL	7.0348 mL	
		10 mM	0.3517 mL	1.7587 mL	3.5174 mL	
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	 Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (8.79 mM); Clear solution Add each solvent each base and 10% DMSO are 20% each if 					
	Solubility: ≥ 2.5 mg/mL (8.79 mM); Clear solution					

Description	Golgicide A (GCA) is a potent, highly specific, and reversible inhibitor of the cis-Golgi ADP-ribosylation factor guanine nucleotide exchange factors (ArfGEF) GBF1 ^[1] . Golgicide A drastically reduced replication of coxsackievirus B3 (CVB3) and other human enterovirus species ^[2] .			
IC ₅₀ & Target	GBF1 ^[1]			
In Vitro	Golgicide A (GCA) is a potent and highly effective inhibitor of shiga toxin activity. Golgicide A (GCA) inhibits the effect of shiga toxin on protein synthesis with an IC ₅₀ of 3.3 μM ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

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CUSTOMER VALIDATION

- Mol Metab. 2021 Dec;54:101329.
- bioRxiv. 2024 Jan 2.

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REFERENCES

[1]. Sáenz JB, et al. Golgicide A reveals essential roles for GBF1 in Golgi assembly and function. Nat Chem Biol. 2009 Mar;5(3):157-65.

[2]. van der Linden L, et al. Differential effects of the putative GBF1 inhibitors Golgicide A and AG1478 on enterovirus replication. J Virol. 2010 Aug;84(15):7535-42.

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

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