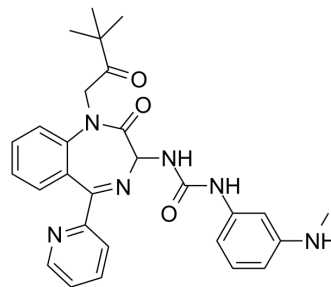


(Rac)-Sograzepide

Cat. No.:	HY-U00360		
CAS No.:	168161-71-5		
Molecular Formula:	C ₂₈ H ₃₀ N ₆ O ₃		
Molecular Weight:	498.58		
Target:	Cholecystokinin Receptor		
Pathway:	GPCR/G Protein; Neuronal Signaling		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 125 mg/mL (250.71 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.0057 mL	10.0285 mL	20.0570 mL
		5 mM	0.4011 mL	2.0057 mL	4.0114 mL
10 mM		0.2006 mL	1.0028 mL	2.0057 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: ≥ 2.08 mg/mL (4.17 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (4.17 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.17 mM); Clear solution 				

BIOLOGICAL ACTIVITY

Description	(Rac)-Sograzepide is an antagonist of cholecystokinin B (CCK-B) receptor, and has the potential of reducing the secretion of gastric acid.
IC₅₀ & Target	CCK-B receptor ^[1]
In Vitro	(Rac)-Sograzepide is an antagonist of CCK-B receptor, and has the potential of reducing the secretion of gastric acid ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Hamish Ryder, et al. Benzodiazepin derivatives useful as cck-receptor antagonists. WO 1993016999 A1.

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

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