BNC375

Cat. No.:	HY-128575				
CAS No.:	1557240-80-8				
Molecular Formula:	C ₁₉ H ₂₃ ClN ₂ O ₃ S				
Molecular Weight:	394.92				
Target:	nAChR				
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling				
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 : 83.33 mg/mL (211.00 mM; Need ultrasonic)							
Preparing Stock Solutions	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg			
		1 mM	2.5322 mL	12.6608 mL	25.3216 mL			
	5 mM	0.5064 mL	2.5322 mL	5.0643 mL				
	10 mM	0.2532 mL	1.2661 mL	2.5322 mL				
	Please refer to the solubility information to select the appropriate solvent.							
In Vivo	1. Add each solvent o Solubility: ≥ 2.08 n	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (5.27 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (5.27 mM); Clear solution							
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (5.27 mM); Clear solution							

Description	BNC375 is a potent, selective, and orally available type I positive allosteric modulator of α7 nAChRs with an EC ₅₀ of 1.9 μM. BNC375 exhibits good CNS-agent like properties and clinical candidate potential. ^[1] .				
In Vitro	BNC375 significantly potentiates the acetylcholine signal without changing the rapid receptor desensitization ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.				
In Vivo	BNC375 (0.003-10.0 mg/kg, administered orally) exhibits the MED of 0.03 mg/kg, and achieves full reversal of the				



H₂N⁻O CI

Product Data Sheet

Scopolamine-induced impairment at 1.0 mg/kg in mouse T-maze model. BNC375 exhibits the plasm half-life $(t_{1/2})$ of 1.2 h^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Andrew J Harvey, et al. Discovery of BNC375, a Potent, Selective, and Orally Available Type I Positive Allosteric Modulator of α7 nAChRs. ACS Med Chem Lett. 2019 Mar 25;10(5):754-760.

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

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