GNE-0439

Cat. No.:	HY-123824	O
CAS No.:	1241902-40-8	N
Molecular Formula:	C ₂₁ H ₃₁ NO ₃	H
Molecular Weight:	345.48	O
Target:	Sodium Channel	O
Pathway:	Membrane Transporter/Ion Channel	H
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)	

SOLVENT & SOLUBILITY

In Vitro	DMSO : 130 mg/mL (376.29 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg	
		1 mM	2.8945 mL	14.4726 mL	28.9452 mL	
		5 mM	0.5789 mL	2.8945 mL	5.7890 mL	
		10 mM	0.2895 mL	1.4473 mL	2.8945 mL	
	Please refer to the sol	ubility information to select the app	propriate solvent.			
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 3.25 mg/mL (9.41 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 3.25 mg/mL (9.41 mM); Clear solution					
	3. Add each solvent o Solubility: ≥ 3.25 n	one by one: 10% DMSO >> 90% cor ng/mL (9.41 mM); Clear solution	n oil			

Description	GNE-0439 is a novel Nav1.7-selective inhibitor with IC ₅₀ of 0.34 uM and inhibits Nav1.5 with an IC ₅₀ of 38.3 μM. GNE-0439 inhibits mutant N1742K channels (IC ₅₀ =0.37 uM) in membrane potential assays. GNE-0439 possesses a carboxylic acid group, binds outside of the channel pore, and is unique compared with known selective VSD4 binders ^[1] .			
IC ₅₀ & Target	IC50: 0.34 uM (Nav1.7) ^[1]			

REFERENCES



[1]. Chernov-Rogan T, et al. Mechanism-specific assay design facilitates the discovery of Nav1.7-selective inhibitors. Proc Natl Acad Sci U S A. 2018 Jan 23;115(4):E792-E801.

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

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