RedChemExpress

Product Data Sheet

GsMTx4

Cat. No.:	HY-P1410			
CAS No.:	1209500-46-8			
Molecular Formula:	C ₁₈₅ H ₂₇₃ N ₄₉ O ₄₅ S ₆			
Molecular Weight:	4095.84 GCLEFWWKCNPNDDKCCRPKLKCSKLFKLCNFSF-NH2 (Disulfide bridge:Cys2-Cys17, Cys9-Cys23, Cys16-Cys30)			
Sequence:	Gly-Cys-Leu-Glu-Phe-Trp-Trp-Lys-Cys-Asn-Pro-Asn-Asp-Asp-Lys-Cys-Cys-Arg-Pro-Lys- Leu-Lys-Cys-Ser-Lys-Leu-Phe-Lys-Leu-Cys-Asn-Phe-Ser-Phe-NH2 (Disulfide bridge:Cy s2-Cys17, Cys9-Cys23, Cys16-Cys30)			
Sequence Shortening:	GCLEFWWKCNPNDDKCCRPKLKCSKLFKLCNFSF-NH2 (Disulfide bridge:Cys2-Cys17, Cys 9-Cys23, Cys16-Cys30)			
Target:	TRP Channel; Piezo Channel			
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling			
Storage:	Sealed storage, away from moisture and light Powder -80°C 2 years -20°C 1 year * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)			

In Vitro		H ₂ O : 50 mg/mL (12.21 mM; Need ultrasonic) DMSO : 50 mg/mL (12.21 mM; Need ultrasonic)					
		Mass Solvent Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	0.2442 mL	1.2208 mL	2.4415 mL		
		5 mM	0.0488 mL	0.2442 mL	0.4883 mL		
		10 mM	0.0244 mL	0.1221 mL	0.2442 mL		
	Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 1.25 mg/mL (0.31 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 1.25 mg/mL (0.31 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1.25 mg/mL (0.31 mM); Clear solution						

BIOLOGICAL ACTIVITY

Description	GsMTx4 is a spider venom peptide that selectively inhibits cationic-permeable mechanosensitive channels (MSCs) belonging to the Piezo and TRP channel families. GsMTx4 also blocks cation-selective stretch-activated channels (SACs), attenuates lysophosphatidylcholine (LPC)-induced astrocyte toxicity and microglial reactivity. GsMTx4 is an important pharmacological tool for identifying the role of these excitatory MSCs in normal physiology and pathology ^{[1][2][4]} .				
IC ₅₀ & Target	MSCs ^[1]				
In Vitro	 GsMTx4 (5 μM) reduces Piezo1-mediated charge transfer to 38% of its initial levels in HEK293 cells transfected with Piezo1 cDNA^[1]. GsMTx4 (5 μM) blocks cation-selective stretch-activated channels in astrocytes, cardiac cells, and smooth and skeletal muscle cells^[2]. GsMTx4 (2.5 μM, 16 h) significantly diminishes both the leptin-induced AMPK and MLC-2 phosphorylation in breast epithelial cells (MCF10A)^[3]. GsMTx4 (500 nM, 48 h) attenuates demyelination induced by the cytotoxic lipid and psychosine (organotypic cerebellar slices)^[4]. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis^[3] 				
	Cell Line:	MCF10A cells			
	Concentration:	2.5 μΜ			
	Incubation Time:	16 h			
	Result:	Diminished both the leptin-induced AMPK and MLC-2 phosphorylation.			
In Vivo	GsMTx4 (stereotactic injection, 3 μM of 1 μL, a single dose) is neuroprotective and inhibits lysophosphatidylcholine- induced astrocyte toxicity and demyelination in the cerebral cortex ^[4] . GsMTx-4 (intraperitoneal injection, 270 μg/kg for a single dose) reduces mechanical allodynia induced by inflammation and by sciatic nerve injury in Von Frey test ^[6] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.				
	Animal Model:	Male C57BL/6 mice (toxin-induced focal demyelination of cortical brain tissue) ^[4]			
	Dosage:	3 μM for 1 μL, a single dose.			
	Administration:	Stereotactic injection in the left and right cerebral hemispheres (sacrificed 4 days post- injection)			
	Result:	Prevented the enhanced increase in microglial reactivity and microglial cell numbers induced by lysophosphatidylcholine (LPC). Prevented LPC-mediated astrocyte toxicity by attenuating the decrease in GFAP+ cells and GFAP fluorescence intensity.			
	Animal Model:	Sciatic nerve injury model of male Sprague-Dawley rats ^[6]			
	Dosage:	270 μg/kg, a single dose			
	Administration:	Intraperitoneal injection			
	Result:	Reduced inflammation-evoked mechanical allodynia.			

CUSTOMER VALIDATION

- Cell Discov. 2022 Sep 6;8(1):84.
- Neuron. 2022 Nov 8;S0896-6273(22)00954-0.
- Research (Wash D C). 2023 Jan 20.
- Materials Today Advances. 2023, 17: 100325.
- Hypertension. 2021 Sep;78(3):647-660.

See more customer validations on <u>www.MedChemExpress.com</u>

REFERENCES

[1]. Gnanasambandam R, et al. GsMTx4: Mechanism of Inhibiting Mechanosensitive Ion Channels. Biophys J. 2017 Jan 10;112(1):31-45.

[2]. T M Suchyna, et al. Identification of a peptide toxin from Grammostola spatulata spider venom that blocks cation-selective stretch-activated channels. J Gen Physiol. 2000 May;115(5):583-98.

[3]. Anna Acheva, et al. Adipokine Leptin Co-operates With Mechanosensitive Ca 2 +-Channels and Triggers Actomyosin-Mediated Motility of Breast Epithelial Cells. Front Cell Dev Biol. 2021 Jan 6;8:607038.

[4]. María Velasco-Estevez, et al. Inhibition of Piezo1 attenuates demyelination in the central nervous system. Glia. 2020 Feb;68(2):356-375.

[5]. Medha M Pathak, et al. Stretch-activated ion channel Piezo1 directs lineage choice in human neural stem cells. Proc Natl Acad Sci U S A. 2014 Nov 11;111(45):16148-53.

[6]. Seung Pyo Park, et al. A tarantula spider toxin, GsMTx4, reduces mechanical and neuropathic pain. Pain. 2008 Jul;137(1):208-217.

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

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA