ST91

Cat. No.:	HY-103203		
CAS No.:	4749-61-5	<	
Molecular Formula:	C ₁₃ H ₂₀ ClN ₃) H	
Molecular Weight:	253.77		-0
Target:	Adrenergic Receptor	HN_/ "	01
Pathway:	GPCR/G Protein; Neuronal Signaling)	
Storage:	4°C, sealed storage, away from moisture and light * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)		

SOLVENT & SOLUBILITY

In Vitro	DMSO : 125 mg/mL (492.57 mM; Need ultrasonic)						
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	3.9406 mL	19.7029 mL	39.4058 mL		
		5 mM	0.7881 mL	3.9406 mL	7.8812 mL		
		10 mM	0.3941 mL	1.9703 mL	3.9406 mL		
	Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (8.20 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (8.20 mM); Clear solution						

Description	ST91 is a α_2 -adrenoceptor (α_2 AR) agonist. ST91 activates both α_{2A} AR and non- α_{2A} AR subtypes to produce spinal antinociception ^{[1][2][3]} .			
IC ₅₀ & Target	α2-adrenergic receptor			
In Vitro	ST91 decreases the viability, proliferation and mitochondrial function of B16F10 cells ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			
In Vivo	ST91 (intrathecal administration) produces antinociception in rats ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

Product Data Sheet



REFERENCES

[1]. Graham BA, et, al. Synergistic interactions between two alpha(2)-adrenoceptor agonists, dexmedetomidine and ST-91, in two substrains of Sprague-Dawley rats. Pain. 2000 Mar;85(1-2):135-43.

[2]. Maccari S, et, al. α-Adrenoceptor stimulation attenuates melanoma growth in mice. Br J Pharmacol. 2022 Apr;179(7):1371-1383.

[3]. Stone LS, et, al. ST91 [2-(2,6-diethylphenylamino)-2-imidazoline hydrochloride]-mediated spinal antinociception and synergy with opioids persists in the absence of functional alpha-2A- or alpha-2C-adrenergic receptors. J Pharmacol Exp Ther. 2007 Dec;323(3):899-906.

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