# Mecamylamine hydrochloride

Cat. No.:	HY-B1395	•
CAS No.:	826-39-1	$\land$
Molecular Formula:	C <sub>11</sub> H <sub>22</sub> ClN	
Molecular Weight:	203.75	
Target:	nAChR; Histamine Receptor	
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling; GPCR/G Protein; Immunology/Inflammation	
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)	H-(

## SOLVENT & SOLUBILITY

In Vitro	H <sub>2</sub> O : 100 mg/mL (490.80 mM; Need ultrasonic) DMSO : 31.25 mg/mL (153.37 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg	
		1 mM	4.9080 mL	24.5399 mL	49.0798 mL	
		5 mM	0.9816 mL	4.9080 mL	9.8160 mL	
		10 mM	0.4908 mL	2.4540 mL	4.9080 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: ≥ 2.08 mg/mL (10.21 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (10.21 mM); Clear solution					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (10.21 mM); Clear solution					

Description	Mecamylamine hydrochloride is an orally active, nonselective, noncompetitive nAChR antagonist. Mecamylamine hydrochloride is also a ganglionic blocker. Mecamylamine hydrochloride can across the blood-brain barrier. Mecamylamine hydrochloride can be used in the research of neuropsychiatric disorders, hypertension, antidepressant area <sup>[1][2][5]</sup> .		
IC <sub>50</sub> & Target	nAChR <sup>[1]</sup> , histamine receptor <sup>[2]</sup>		
In Vitro	Mecamylamine hydrochloride (0.5-9 $\mu$ M, bath administered) increases the firing frequency of identified 5-HT DRN (dorsal		

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**Product** Data Sheet

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raphe nucleus) neurons<sup>[1]</sup>.

Mecamylamine hydrochloride (0.5-9  $\mu$ M, bath administered) increases the glutamatergic and decreases the GABAergic input of 5-HT DRN neurons<sup>[1]</sup>.

Mecamylamine hydrochloride (1 mM, 5 min) blocks the histamine receptor and the histamine-induced contractions in helically cut strips of rabbit aorta<sup>[2]</sup>.

Mecamylamine hydrochloride (1-100 nM, 30 min) dose-dependently attenuates endothelial tube formation in HDMVECs<sup>[4]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

#### Western Blot Analysis<sup>[3]</sup>

Cell Line:	SCG neurons
Concentration:	10 μΜ
Incubation Time:	48 h
Result:	Reduced the nicotine-facilitated increase in ERK1/2.

#### In Vivo

Mecamylamine hydrochloride (subcutaneous pumps, 50 mg/kg/day, 2 days) inhibits Choroidal neovascularization (CNV) in CNV mice model<sup>[4]</sup>.

Mecamylamine hydrochloride (intraperitoneal injection, 0.5-1 mg/kg) has antidepressant-like effects in both the TST (tail suspension test) and FST (forced swim test) in C57BL/6J mice, which are dependent on both  $\beta$ 2 and  $\alpha$ 7 subunits<sup>[5]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Choroidal neovascularization (CNV) mice model <sup>[1]</sup>
Dosage:	50 mg/kg/day, 2 days
Administration:	Subcutaneous pumps implanted beneath the skin of the back, 200 $\mu L$ and mean pumping rate of 0.5 $\mu L/h.$
Result:	Suppressed the development of CNV at Bruch's membrane rupture sites in the absence of nicotine.
Animal Model:	C57BL/6J mice <sup>[5]</sup>
Dosage:	0.5-1 mg/kg
Administration:	Intraperitoneal injection
Result:	Had no effect in $\beta 2$ knockout mice and $\alpha 7$ knockout mice, but decreased immobility time in wildtype littermates in the FST.

#### **CUSTOMER VALIDATION**

- Nat Commun. 2023 Apr 17;14(1):2182.
- Acta Pharmacol Sin. 2024 Mar 4.
- bioRxiv. 2023 Jul 6.

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### REFERENCES

[1]. Omar Hernández-González, et al. Mechanisms of stimulatory effects of mecamylamine on the dorsal raphe neurons. Brain Res Bull. 2020 Nov;164:289-298.

[2]. C P Robinson, et al. The influence of mecamylamine on contractions induced by different agonists and on the role of calcium ions in the isolated rabbit aorta. J Pharmacol Exp Ther. 1976 Apr;197(1):57-65.

[3]. Mahadevappa P Badanavalu, et al. Nicotine is neuroprotective to neonatal neurons of sympathetic ganglion in rat. Auton Neurosci. 2019 Jan;216:25-32.

[4]. Katsuji Kiuchi, et al. Mecamylamine suppresses Basal and nicotine-stimulated choroidal neovascularization. Invest Ophthalmol Vis Sci. 2008 Apr;49(4):1705-11.

[5]. Rabenstein RL, et al. The nicotinic antagonist mecamylamine has antidepressant-like effects in wild-type but not beta2- or alpha7-nicotinic acetylcholine receptor subunit knockout mice. Psychopharmacology (Berl). 2006 Dec;189(3):395-401.

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

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