## Guanfacine hydrochloride

Cat. No.:	HY-17416		
CAS No.:	29110-48-3	∽ .Cl	
Molecular Formula:	C <sub>9</sub> H <sub>10</sub> Cl <sub>3</sub> N <sub>3</sub> O	O NH	
Molecular Weight:	282.55		
Target:	Adrenergic Receptor	H $H$ $H$	
Pathway:	GPCR/G Protein; Neuronal Signaling		
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 : ≥ 30 mg/mL (106.18 mM) H <sub>2</sub> O : 20 mg/mL (70.78 mM; Need ultrasonic) * "≥" means soluble, but saturation unknown.				
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
		1 mM	3.5392 mL	17.6960 mL	35.3920 mL
		5 mM	0.7078 mL	3.5392 mL	7.0784 mL
		10 mM	0.3539 mL	1.7696 mL	3.5392 mL
	Please refer to the sol	ubility information to select the ap	propriate solvent.		
In Vivo	<ol> <li>Add each solvent of Solubility: ≥ 2.5 mg</li> <li>Add each solvent of Solubility: ≥ 2.5 mg</li> <li>Add each solvent of Solubility: ≥ 2.5 mg</li> </ol>	one by one: 10% DMSO >> 40% PE( g/mL (8.85 mM); Clear solution one by one: 10% DMSO >> 90% (20 g/mL (8.85 mM); Clear solution one by one: 10% DMSO >> 90% cor g/mL (8.85 mM); Clear solution	G300 >> 5% Tween-8 % SBE-β-CD in saline) n oil	0 >> 45% saline	

Description	Guanfacine hydrochloride is an orally active noradrenergic α2A agonist and has high selective for the α2A receptor subtype. Guanfacine has effects in producing hypotension and sedation. Guanfacine can be used for the research of a variety of prefrontal cortex (PFC) cognitive disorders, including tourette's syndrome and attention deficit hyperactivity disorder (ADHD) <sup>[1][2][3]</sup> .			
IC <sub>50</sub> & Target	α adrenergic receptor			



In Vitro	Guanfacine (hydrochloride) increases the delay-related neuronal firing needed for working memory on dIPFC neurons at the cellular level <sup>[1][2]</sup> . Guanfacine improves PFC cognitive function by inhibiting the production of CAMP, closing HCN channels, and strengthening the PFC networks <sup>[1][2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	Guanfacine (hydrochloride) enhances PFC working memory function in aged monkeys and improves impulse control in monkeys performing a delayed discounting task <sup>[1][2]</sup> . Guanfacine improves cognitive performance when infused directly into the rat or monkey PFC <sup>[1][2]</sup> . Guanfacine blocks 2A receptors in the monkey dIPFC markedly impairs working memory, behavioral inhibition and greatly reduces persistent neuronal firing <sup>[1][2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## CUSTOMER VALIDATION

• Patent. US20230147129A1.

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## REFERENCES

[1]. Amy FT Arnsten, et al. Guanfacine for the treatment of cognitive disorders: a century of discoveries at Yale. Yale J Biol Med. 2012 Mar;85(1):45-58. Epub 2012 Mar 29.

[2]. P. A. Van Zwieten, et al. The pharmacology of centrally acting antihypertensive drugs. Br J Clin Pharmacol. 1983; 15(Suppl 4): 455S–462S.

[3]. Min Wang, et al. Alpha2A-adrenoceptors strengthen working memory networks by inhibiting cAMP-HCN channel signaling in prefrontal cortex. Cell. 2007 Apr 20;129(2):397-410.

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

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