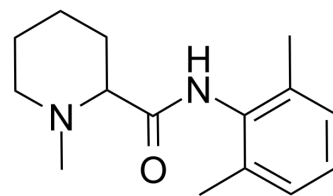


## Mepivacaine hydrochloride

<b>Cat. No.:</b>	HY-B0517A
<b>CAS No.:</b>	1722-62-9
<b>Molecular Formula:</b>	C <sub>15</sub> H <sub>23</sub> ClN <sub>2</sub> O
<b>Molecular Weight:</b>	282.81
<b>Target:</b>	Sodium Channel
<b>Pathway:</b>	Membrane Transporter/Ion Channel
<b>Storage:</b>	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)



HCl

### SOLVENT & SOLUBILITY

<b>In Vitro</b>	H <sub>2</sub> O : 100 mg/mL (353.59 mM; Need ultrasonic)				
		Solvent Concentration	Mass		
	<b>Preparing Stock Solutions</b>		1 mg	5 mg	10 mg
		1 mM	3.5359 mL	17.6797 mL	35.3594 mL
		5 mM	0.7072 mL	3.5359 mL	7.0719 mL
	10 mM	0.3536 mL	1.7680 mL	3.5359 mL	
Please refer to the solubility information to select the appropriate solvent.					
<b>In Vivo</b>	1. Add each solvent one by one: PBS Solubility: 100 mg/mL (353.59 mM); Clear solution; Need ultrasonic				

### BIOLOGICAL ACTIVITY

<b>Description</b>	Mepivacaine hydrochloride binds to specific voltage-gated sodium ion channels in neuronal cell membranes, which inhibits both sodium influx and membrane depolarization <sup>[1][2]</sup> .
<b>In Vitro</b>	<p>Mepivacaine hydrochloride binds to specific voltage-gated sodium ion channels in neuronal cell membranes, which inhibits both sodium influx and membrane depolarization. This leads to a blockage of nerve impulse initiation and conduction and results in a reversible loss of sensation. Compared to other local anesthetics, this agent has a more rapid onset and moderate duration of action<sup>[2]</sup>.</p> <p>Mepivacaine hydrochloride has a reasonably rapid onset (more rapid than that of procaine) and medium duration of action (shorter than that of procaine)<sup>[3]</sup>.</p> <p>Mepivacaine hydrochloride displays a preferential use-dependent block of Na(v)1.8, S(-)-bupivacaine displays a preference for TTXs Na(+) channels<sup>[4]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>

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## REFERENCES

- [1]. Burm, A.G., et al., Pharmacokinetics of the enantiomers of mepivacaine after intravenous administration of the racemate in volunteers. *Anesth Analg*, 1997. 84(1): p. 85-9.
- [2]. Leffler, A., J. Reckzeh, and C. Nau, Block of sensory neuronal Na<sup>+</sup> channels by the secretolytic ambroxol is associated with an interaction with local anesthetic binding sites. *Eur J Pharmacol*, 2010. 630(1-3): p. 19-28.
- [3]. Froehle M, et al. ECMO for Cardiac Rescue after Accidental Intravenous Mepivacaine Application. *Case Rep Pediatr*. 2012;2012:491692.
- [4]. mepivacaine hydrochloride.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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