## Dibucaine

Cat. No.:	HY-B0552				
CAS No.:	85-79-0				
Molecular Formula:	C <sub>20</sub> H <sub>29</sub> N <sub>3</sub> O <sub>2</sub>				
Molecular Weight:	343.46				
Target:	Sodium Channel				
Pathway:	Membrane Transporter/Ion Channel				
Storage:	Powder	-20°C	3 years		
		4°C	2 years		
	In solvent	-80°C	2 years		
		-20°C	1 vear		

### SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (291.15 mM; Need ultrasonic)					
Preparing Stock Solutions		Solvent Mass Concentration	1 mg	5 mg	10 mg	
	1 mM	2.9115 mL	14.5577 mL	29.1155 mL		
		5 mM	0.5823 mL	2.9115 mL	5.8231 mL	
		10 mM	0.2912 mL	1.4558 mL	2.9115 mL	
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	<ol> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 40% PEG300 &gt;&gt; 5% Tween-80 &gt;&gt; 45% saline Solubility: ≥ 2.5 mg/mL (7.28 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2.5 mg/mL (7.28 mM); Clear solution</li> </ol>					

BIOLOGICAL ACTIVITY					
Description	Dibucaine (Cinchocaine) is a sodium channel inhibitor. Dibucaine is a potent SChE inhibitor <sup>[1][2]</sup> .				
In Vitro	Dibucaine (Cinchocaine) reduces the degradation of BSA-gold complex in the reservosomes, which was not caused either by an inhibition of the whole proteolytic activity of the parasite or by a reduction on the expression levels of cruzipain <sup>[1]</sup> . Dibucaine, a quaternary ammonium compound, inhibited SChE to a minimum within 2 min in a reversible manner. The inhibition was very potent. It had an IC(50) of 5.3 microM with BuTch or 3.8 microM with AcTch. The inhibition was competitive with respect to BuTch with a K(i) of 1.3 microM and a linear-mixed type (competitive/noncompetitive) with respect to AcTch with inhibition constants, K(i) and K(I) of 0.66 and 2.5 microM, respectively. Dibucaine possesses a butoxy side chain that is similar to the butryl group of BuTch and longer by an ethylene group from AcTch <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.				

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Product Data Sheet

### **CUSTOMER VALIDATION**

- PLoS Biol. 2024 Mar 25;22(3):e3002565.
- Biomed Pharmacother. 2023 Aug, 164, 114903.
- Stem Cell Res Ther. 2021 Feb 4;12(1):107.
- Platelets. 2021 May 12;1-9.

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#### REFERENCES

[1]. Souto-Padron, T., A.P. Lima, and O. Ribeiro Rde, Effects of dibucaine on the endocytic/exocytic pathways in Trypanosoma cruzi. Parasitol Res, 2006. 99(4): p. 317-20.

[2]. Elamin, B., Dibucaine inhibition of serum cholinesterase. J Biochem Mol Biol, 2003. 36(2): p. 149-53.

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

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