# Data Sheet (Cat.No.T31488)



## Dimethindene

ties	
5636-83-9	H <sub>3</sub> C
C20H24N2	
292.42	
no data available	
store at low temperature Powder: -20°C for 3 years   In solvent: -80°C for 1 year	СН3
	5636-83-9 C20H24N2 292.42 no data available store at low temperature

#### **Biological Description**

Description	Dimethindene (Dimetindeno) is a selective antagonist of H1 receptor and blocks K+ current. Dimethindene exhibits antihistamine and anticholinergic effects.
Targets(IC50)	Potassium Channel,Endogenous Metabolite,AChR,Histamine Receptor
In vitro	In follicle-enclosed Xenopus oocytes, Dimethindene (5-500 $\mu$ M) decreases Cromakalim cromakalim-induced K+ currents (IC50 = 29.5 $\mu$ M)[1].
In vivo ©	In C57BL/6 mice with skin wound healing, Dimethindene (0.25 mg; i.p.) impairs cutaneous wound healing and delays skin wound closure[2].

### Solubility Information

Solubility	DMSO: 55 mg/mL (188.09 mM)
	(< 1 mg/ml refers to the product slightly soluble or insoluble)

### Preparing Stock Solutions

	1mg	5mg	10mg	
1 mM	3.4197 mL	17.0987 mL	34.1974 mL	
5 mM	0.6839 mL	3.4197 mL	6.8395 mL	
10 mM	0.342 mL	1.7099 mL	3.4197 mL	
50 mM	0.0684 mL	0.342 mL	0.6839 mL	

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

#### Reference

Sakuta H. Inhibition by histamine H1 receptor antagonists of endogenous glibenclamide-sensitive K+ channels in follicle-enclosed Xenopus oocytes. Eur J Pharmacol. 1994 Jan 1;266(1):99-102.

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