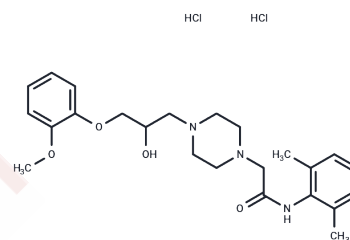


## Ranolazine dihydrochloride

## Chemical Properties

CAS No. : 95635-56-6  
Formula: C<sub>24</sub>H<sub>35</sub>Cl<sub>2</sub>N<sub>3</sub>O<sub>4</sub>  
Molecular Weight: 500.46  
Appearance: no data available  
Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year



## Biological Description

Description	Ranolazine dihydrochloride (Ranolazine 2HCl) , an antianginal agent, can treat arrhythmia via a novel mechanism of action (inhibition of the late phase of the inward sodium current), and do not affect blood pressure or heart rate.
Targets(IC50)	Calcium Channel,Autophagy,Sodium Channel
In vitro	Ranolazine (5 mM and 10 mM) reversibly shortened the duration of twitch contractions and abolished postcontractions.Ranolazine bound more to sodium channels in the inactivated state. In cardiomyocytes, selective inhibition of late I (sodium) by Ranolazine reduced sodium-dependent calcium overload and attenuated ventricular repolarization and contraction, which correlated with abnormalities in heart failure and ischemia/reperfusion injury. In dog left ventricular myocytes, in a concentration-dependent manner Ranolazine was able to reversibly shorten myocyte action potential duration in response to 0.25/0.5 Hz stimulation.
In vivo	Ranolazine (5 mM and 10 mM) reversibly shortened the duration of twitch contractions and abolished postcontractions.Ranolazine bound more to sodium channels in the inactivated state. In cardiomyocytes, selective inhibition of late I (sodium) by Ranolazine reduced sodium-dependent calcium overload and attenuated ventricular repolarization and contraction, which correlated with abnormalities in heart failure and ischemia/reperfusion injury. In dog left ventricular myocytes, in a concentration-dependent manner Ranolazine was able to reversibly shorten myocyte action potential duration in response to 0.25/0.5 Hz stimulation.

## Solubility Information

Solubility	DMSO: 45 mg/mL (89.92 mM),Sonication is recommended. H <sub>2</sub> O: 50.1 mg/mL (100.11 mM),Sonication is recommended. ( < 1 mg/ml refers to the product slightly soluble or insoluble)
------------	---

### Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	1.9982 mL	9.9908 mL	19.9816 mL
5 mM	0.3996 mL	1.9982 mL	3.9963 mL
10 mM	0.1998 mL	0.9991 mL	1.9982 mL
50 mM	0.040 mL	0.1998 mL	0.3996 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

Belardinelli L, et al. Heart, 2006, 92 Suppl 4, iv6-iv14.

Undrovinas AI, et al. J Cardiovasc Electrophysiol, 2006, 17 Suppl 1, S169-S177.

McCormack JG, et al. Circulation, 1996, 93(1), 135-142.

**Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins**

**This product is for Research Use Only · Not for Human or Veterinary or Therapeutic Use**

Tel: 781-999-4286    E\_mail: info@targetmol.com    Address: 36 Washington Street, Wellesley Hills, MA 02481