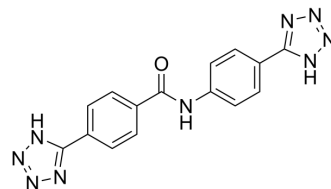


Andolast free base

Cat. No.:	HY-106358		
CAS No.:	132640-22-3		
Molecular Formula:	C ₁₅ H ₁₁ N ₉ O		
Molecular Weight:	333.31		
Target:	Phosphodiesterase (PDE)		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 50 mg/mL (150.01 mM; ultrasonic and warming and heat to 60°C)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	3.0002 mL	15.0010 mL	30.0021 mL
		5 mM	0.6000 mL	3.0002 mL	6.0004 mL
10 mM		0.3000 mL	1.5001 mL	3.0002 mL	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (7.50 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	Andolast (CR 2039) (free base) is an anti-allergic agent. Andolast can inhibit cAMP-phosphodiesterase with an IC ₅₀ value of 50 μM. Andolast can be used for the research of asthma ^[1] .	
In Vivo	Andolast (CR 2039) (i.v. or i.m.) inhibits rat passive cutaneous anaphylaxis (PCA) with an ED ₅₀ of 0.1 mg/kg ^[1] . CR 2039 (10-100 mg/kg; i.v. or i.m.) inhibits the microvascular permeability changes in a model of allergic conjunctivitis in sensitized guinea-pigs ^[1] . CR 2039 (0-1000 μM; i.v.) inhibits dose dependently guinea-pig lung cAMP-phosphodiesterase with an IC ₅₀ value of 50 μM ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	Male Hartley guinea-pigs (300-500 g) ^[1]

Dosage:	10-100 mg/kg
Administration:	I.M; I.V.
Result:	Showed dose-related significant protection against IgE-dependent bronchial anaphylaxis induced by aerosolized antigen in anesthetized guinea-pigs. Delayed dose dependently the onset of bronchoconstriction induced by aerosolized antigen.

REFERENCES

- [1]. Revel L, Colombo S, Ferrari F, Folco G, Rovati LC, Makovec F. CR 2039, a new bis-(1H-tetrazol-5-yl)phenylbenzamide derivative with potential for the topical treatment of asthma. *Eur J Pharmacol.* 1992;229(1):45-53.
- [2]. Czuczwar SJ, Gasior M, Kozicka M, Pietrasiewicz T, Turski WA, Kleinrok Z. A potential anti-asthmatic drug, CR 2039, enhances the anticonvulsive activity of some antiepileptic drugs against pentetrazol in mice. *Eur Neuropsychopharmacol.* 1998;8(3):233-238.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA