Data Sheet (Cat.No.T12491)



Pitolisant oxalate

Chemical Properties

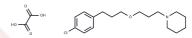
CAS No.: 362665-57-4

Formula: C19H28ClNO5

Molecular Weight: 385.88

Appearance: no data available

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year



Biological Description

Description	Pitolisant oxalate is a potent and selective nonimidazole inverse agonist of the recombinant human histamine H3 receptor, exhibiting a Ki of 0.16 nM.
Targets(IC50)	Others
In vitro	Pitolisant behaves as a competitive antagonist with a Ki value of 0.16 nM and as an inverse agonist with an EC50 value of 1.5 nM and an intrinsic activity ~50% higher than that of ciproxifan.
In vivo	Administering Pitolisant at a single dose of 10 mg/kg 30 minutes before a 2 mg/kg b.w. dose of Olanzapine significantly affects immobility time in the FST. This drug sequence in mice significantly increases immobility duration compared to the control group in the FST and also decreases locomotor activity [3].

Solubility Information

Solubility	DMSO: 50 mg/mL (129.57 mM), Sonication is recommended.	
	(< 1 mg/ml refers to the product slightly soluble or insoluble)	

Preparing Stock Solutions

-				
	1mg	5mg	10mg	
1 mM	2.5915 mL	12.9574 mL	25.9148 mL	
5 mM	0.5183 mL	2.5915 mL	5.183 mL	
10 mM	0.2591 mL	1.2957 mL	2.5915 mL	
50 mM	0.0518 mL	0.2591 mL	0.5183 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.

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Reference

Ligneau X, et al. BF2.649 [1-{3-[3-(4-Chlorophenyl)propoxy]propyl}piperidine, hydrochloride], a nonimidazole inverse agonist/antagonist at the human histamine H3 receptor: Preclinical pharmacology. J Pharmacol Exp Ther. 2007 Jan; 320(1):365-75.

Dudek M, et al. H3 histamine receptor antagonist pitolisant reverses some subchronic disturbances induced by olanzapine in mice. Metab Brain Dis. 2016 Oct;31(5):1023-9.

Uguen M, et al. Preclinical evaluation of the abuse potential of Pitolisant, a histamine H? receptor inverse agonist/antagonist compared with Modafinil. Br J Pharmacol. 2013 Jun;169(3):632-44.

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