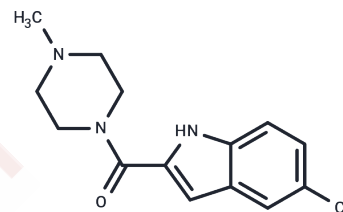


JNJ-777120

Chemical Properties

CAS No. : 459168-41-3
Formula: C₁₄H₁₆ClN₃O
Molecular Weight: 277.75
Appearance: no data available
Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year



Biological Description

Description	JNJ-777120 is the first potent and selective non-imidazole histamine H ₄ receptor antagonist with K _i of 4.5 nM, exhibits >1000-fold selectivity over the other histamine receptors.
Targets(IC ₅₀)	Histamine Receptor
In vitro	JNJ 777120 administration in mice inhibits the migration of tracheal mast cells from connective to epithelial tissues induced by histamine. Furthermore, JNJ 777120 treatment in mice-derived bone marrow mast cells suppresses histamine-induced chemotaxis and calcium influx. In a peritonitis model induced by yeast polysaccharides in mice, JNJ 777120 markedly inhibits neutrophil infiltration. The compound demonstrates oral bioavailability, with a 30% rate in rats and 100% in dogs, and a half-life of 3 hours for both species.
In vivo	JNJ 777120 is an effective, selective antagonist for the H ₄ receptor, with little to no affinity for over 50 other targets. It binds with high affinity to the H ₄ receptor, exhibiting over a thousand times greater selectivity compared to other histamine receptors.

Solubility Information

Solubility	DMSO: 40 mg/mL (144.01 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	3.6004 mL	18.0018 mL	36.0036 mL
5 mM	0.7201 mL	3.6004 mL	7.2007 mL
10 mM	0.360 mL	1.8002 mL	3.6004 mL
50 mM	0.072 mL	0.360 mL	0.7201 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

Jablonowski JA, et al. J Med Chem, 2003, 46(19), 3957-3960.

Thurmond RL, et al. J Pharmacol Exp Ther, 2004, 309(1), 404-413.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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