

Product Name: Loxapine Revision Date: 05/14/2024

Product Data Sheet

Loxapine

Cat. No.: B1001

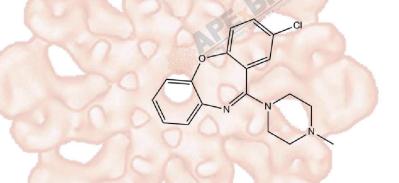
CAS No.: 1977-10-2

Formula: C18H18CIN3O

M.Wt: 327.81

Synonyms:

Target: Neuroscience
Pathway: 5-HT Receptor
Storage: Store at -20°C



Solvent & Solubility

≥55.6 mg/mL in DMSO; ≥55.5 mg/mL in EtOH; <2.78 mg/mL in H2O

In Vitro

Preparing Stock Solutions	Solvent Concentration	1mg	5mg	10mg
	1 mM	3.0505 mL	15.2527 mL	30.5055 mL
	5 mM	0.6101 mL	3.0505 mL	6.1011 mL
40	10 mM	0.3051 mL	1.5253 mL	3.0505 mL

Please refer to the solubility information to select the appropriate solvent.

Biological Activity

Shortsummary	5-HT receptor antagonist	
IC ₅₀ & Target		
In Vitro	Cell Viability Assay	
	Cell Line:	Rat glial and microglial cells
	Preparation method:	This compound is soluble in DMSO. General tips for obtaining a higher concentration: Please warm the tube at 37 °C for 10 minutes and/or shake it in the ultrasonic bath for a while. Stock solution can be stored below - 20 °C for several months.
	Reacting conditions:	0.2, 2 and 20 μM; 1 and 3 days
	Applications:	In LPS-activated rat mixed glial and microglial cell cultures, Loxapine at all

		indicated concentrations reduced IL-1β secretion. Besides, Loxapine also	
		reduced IL-2 secretion in mixed glia cultures. In addition, in LPS-induced	
		microglia cultures, Loxapine decreased both IL-1β and IL-2 secretion at the	
		concentrations of 0.2, 2 and 20 μM after the exposure of 1 and 3 days.	
	Animal experiment		
In Vivo	Animal models:	Wistar rats	
	Dosage form:	5 mg/kg; i.p.	
	Applications:	In Wistar rats, chronic administration of Loxapine for 4 weeks or 10 weeks did	
		not increase striatal dopamine receptor density. However, Loxapine	
		significantly reduced cortical serotonin receptor density by 50 ~ 60%. In	
		addition, a single dose of Loxapine exhibited the same potent effect. The	
		results suggested that Loxapine exerted its effect via the serotonin system.	
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may	
		slightly differ with the theoretical value. This is caused by an experimental	
	.10.	system error and it is normal.	

Product Citations

1. Perez-Gomez A, Carretero M, et al. "A phenotypic Caenorhabditis elegans screen identifies a selective suppressor of antipsychotic-induced hyperphagia." Nat Commun. 2018 Dec10;9(1):5272.PMID:30532051

See more customer validations on www.apexbt.com.

References

- [1]. Labuzek K1,Kowalski J,Gabryel B,Herman ZS. Chlorpromazine and loxapine reduce interleukin-1beta and interleukin-2 release by rat mixed glial and microglial cell cultures. Eur Neuropsychopharmacol. 2005 Jan; 15(1):23-30.
- [2]. Lee T,Tang SW. Loxapine and clozapine decreaseserotonin(S2) but do not elevatedopamine(D2)receptornumbers in the rat brain. Psychiatry Res. 1984 Aug; 12(4):277-85.

Caution

FOR RESEARCH PURPOSES ONLY.

NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

Specific storage and handling information for each product is indicated on the product datasheet. Most APExBIO products are stable under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Shortterm storage of many products are stable in the short-term at temperatures that differ from that required for long-term storage. We ensure that the product is shipped under conditions that will maintain the quality of the reagents. Upon receipt of the product, follow the storage recommendations on the product data sheet.

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