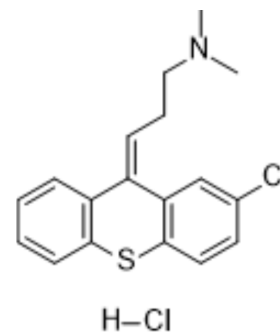


Chlorprothixene hydrochloride

Cat. No.:	HY-B0274A
CAS No.:	6469-93-8
Molecular Formula:	C ₁₈ H ₁₉ Cl ₂ NS
Molecular Weight:	352.32
Target:	Dopamine Receptor; Histamine Receptor; Bacterial
Pathway:	GPCR/G Protein; Neuronal Signaling; Immunology/Inflammation; Anti-infection
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



SOLVENT & SOLUBILITY

In Vitro

H₂O : 25 mg/mL (70.96 mM; Need ultrasonic)
Ethanol : 25 mg/mL (70.96 mM; Need ultrasonic)
DMSO : 2.5 mg/mL (7.10 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
	1 mM	2.8383 mL	14.1916 mL	28.3833 mL
	5 mM	0.5677 mL	2.8383 mL	5.6767 mL
	10 mM	0.2838 mL	1.4192 mL	2.8383 mL

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

In Vivo

- Add each solvent one by one: 10% EtOH >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: ≥ 2.5 mg/mL (7.10 mM); Clear solution
- Add each solvent one by one: 10% EtOH >> 90% (20% SBE-β-CD in saline)
Solubility: ≥ 2.5 mg/mL (7.10 mM); Clear solution
- Add each solvent one by one: 10% EtOH >> 90% corn oil
Solubility: ≥ 2.5 mg/mL (7.10 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Chlorprothixene hydrochloride is a dopamine and histamine receptors antagonist with K_is of 18 nM, 2.96 nM, 4.56 nM, 9 nM and 3.75 nM for hD₁, hD₂, hD₃, hD₅ and hH₁ receptors, respectively. Antipsychotic activity^[1].

IC₅₀ & Target

Human D ₁ Receptor 18 nM (K _i)	Human D ₂ Receptor 2.96 nM (K _i)	Human D ₃ Receptor 4.56 nM (K _i)	Human D ₅ Receptor 9 nM (K _i)
Human H ₁ Receptor			

	3.75 nM (Ki)
In Vitro	Chlorprothixene binds to 5-HT receptors with pK _i s of 8.3, 8.5, and 9.4 for 5-HT ₇ , 5-HT ₆ and 5-HT ₂ , respectively ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Y von Coburg, et al. Potential utility of histamine H3 receptor antagonist pharmacophore in antipsychotics. *Bioorg Med Chem Lett*. 2009 Jan 15;19(2):538-42.

[2]. B L Roth, et al. Binding of typical and atypical antipsychotic agents to 5-hydroxytryptamine-6 and 5-hydroxytryptamine-7 receptors. *J Pharmacol Exp Ther*. 1994 Mar;268(3):1403-10.

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

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