## **Product** Data Sheet

## cis-(Z)-Flupentixol dihydrochloride

Cat. No.: HY-15856 CAS No.: 51529-01-2

Molecular Formula:  $C_{23}H_{27}Cl_{2}F_{3}N_{2}OS$ 

Molecular Weight: 507.44

Target: Dopamine Receptor; 5-HT Receptor Pathway: GPCR/G Protein; Neuronal Signaling

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 62.5 mg/mL (123.17 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.9707 mL	9.8534 mL	19.7068 mL
	5 mM	0.3941 mL	1.9707 mL	3.9414 mL
	10 mM	0.1971 mL	0.9853 mL	1.9707 mL

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

## **BIOLOGICAL ACTIVITY**

Description cis-(Z)-Flupentixol dihydrochloride is a potent and selective DA D1/D2 receptor antagonist, with  $K_i$  values of 0.38 nM and 7 nM for D2 receptor and 5-HT $_{2A}$ , respectively  $^{[1][2]}$ .

IC<sub>50</sub> & Target D<sub>2</sub> Receptor 5-HT<sub>2A</sub> Receptor

0.38 nM (Ki) 7 nM (Ki)

cis-(Z)-Flupentixol (0.25, or 0.5 mg/kg, i.p.) pretreatment dose-dependently reduces cocaine-induced activity<sup>[1]</sup>. In Vivo

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	$Rats^{[1]}.$
Dosage:	0.125, 0.25, or 0.5 mg/kg.
Administration:	IP.
Result:	Reduced cocaine-induced activity.

REFERENCES
NEI ENERGES
[1]. Jennifer M Wenzel, et al. The dopamine antagonist cis-flupenthixol blocks the expression of the conditioned positive but not the negative effects of cocaine in rats. Pharmacol Biochem Behav. 2013 Dec;114-115:90-6.
[2]. Philip Seeman, et al. Atypical antipsychotics: mechanism of action. Can J Psychiatry. 2002 Feb;47(1):27-38.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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