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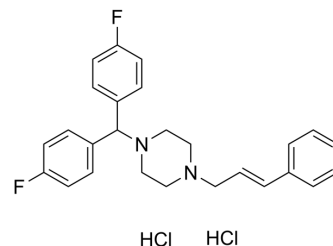
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Flunarizine dihydrochloride

Cat. No.:	HY-B0358A
CAS No.:	30484-77-6
Molecular Formula:	C ₂₆ H ₂₈ Cl ₂ F ₂ N ₂
Molecular Weight:	477.42
Target:	Calcium Channel; Sodium Channel; Dopamine Receptor
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling; GPCR/G Protein
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	DMSO : 50 mg/mL (104.73 mM; Need ultrasonic) H ₂ O : 1 mg/mL (2.09 mM; Need ultrasonic)				
	Preparing Stock Solutions	Mass Solvent Concentration	1 mg	5 mg	10 mg
		1 mM	2.0946 mL	10.4730 mL	20.9459 mL
		5 mM	0.4189 mL	2.0946 mL	4.1892 mL
		10 mM	0.2095 mL	1.0473 mL	2.0946 mL
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: PBS Solubility: 10 mg/mL (20.95 mM); Clear solution; Need ultrasonic				
	2. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (5.24 mM); Clear solution				
	3. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (5.24 mM); Clear solution				
	4. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (5.24 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	Flunarizine dihydrochloride is a potent dual Na ⁺ /Ca ²⁺ channel (T-type) blocker. Flunarizine dihydrochloride is a D ₂ dopamine receptor antagonist. Flunarizine dihydrochloride shows anticonvulsive and antimigraine activity, and peripheral vasodilator effects ^{[1][2][3][4][5]} .	
IC ₅₀ & Target	T-type calcium channel	D ₂ Receptor

In Vitro

Flunarizine blocks sodium currents (I_{Na}) and calcium currents (I_{Ca}) with IC_{50} values of 0.94 μ M and 1.77 μ M in cultured rat cortical neurons, respectively^[2].

Flunarizine (10 and 30 μ M; 24 h) shows cytotoxic effects to chromaffin cells^[4].

Flunarizine (1-30 μ M) causes clear cytoprotection of chromaffin cell at concentrations of 3-10 μ M^[4].

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

Cell Viability Assay^[4]

Cell Line:	Chromaffin cells
Concentration:	10 and 30 μ M
Incubation Time:	24 hours
Result:	Showed a tendency to increase cell death at the concentration of 10 μ M, and showed near 100% cell loss at the concentration of 30 μ M.

In Vivo

Flunarizine (intraperitoneal injection; 30 mg/kg; once) protects mice from lipopolysaccharide- (LPS-) induced acute lung injury (ALI)^[5].

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

Animal Model:	Male BALB/c mice (6-8 weeks old) with acute lung injury induced by lipopolysaccharide ^[5]
Dosage:	30 mg/kg
Administration:	Intraperitoneal injection; 30 mg/kg; once
Result:	Suppressed the LPS-induced cell influx, protein leakage, and inflammatory cytokines release. Inhibited the pulmonary inflammation.

CUSTOMER VALIDATION

- Sci Rep. 2018 Nov 16;8(1):16932.
- J Leukoc Biol. 2021 Sep 17.
- J Ethnopharmacol. 2020 May 23;254:112727.

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REFERENCES

- [1]. Novalbos J, et al. Effects of dotarizine and flunarizine on chromaffin cell viability and cytosolic Ca^{2+} . Eur J Pharmacol. 1999 Feb 5;366(2-3):309-17.
- [2]. Wan L, et al. Mibefradil and Flunarizine, Two T-Type Calcium Channel Inhibitors, Protect Mice against Lipopolysaccharide-Induced Acute Lung Injury. Mediators Inflamm. 2020 Nov 10;2020:3691701.
- [3]. Hong-Seob So, et al. Protective effect of T-type calcium channel blocker flunarizine on cisplatin-induced death of auditory cells. Hear Res. 2005 Jun;204(1-2):127-39.
- [4]. Qing Ye, et al. Flunarizine blocks voltage-gated Na^{+} and Ca^{2+} currents in cultured rat cortical neurons: A possible locus of action in the prevention of migraine. Neurosci Lett. 2011 Jan 10;487(3):394-9.
- [5]. Celia M Santi, et al. Differential inhibition of T-type calcium channels by neuroleptics. J Neurosci. 2002 Jan 15;22(2):396-403.

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

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