

# **Product** Data Sheet

# **Felodipine**

Cat. No.: HY-B0309 CAS No.: 72509-76-3 Molecular Formula:  $C_{18}H_{19}Cl_2NO_4$ 

Molecular Weight: 384.25

Target: Calcium Channel; Autophagy

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling; Autophagy

**Storage:** Powder -20°C 3 years

In solvent

4°C 2 years -80°C 1 year

-20°C 6 months

### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (260.25 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.6025 mL	13.0124 mL	26.0247 mL
	5 mM	0.5205 mL	2.6025 mL	5.2049 mL
	10 mM	0.2602 mL	1.3012 mL	2.6025 mL

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility:  $\geq$  2.5 mg/mL (6.51 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.51 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	Felodipine, a dihydropyridine, is a potent, vasoselective calcium channel antagonist. Felodipine lowers blood pressure (BP) by selective action on vascular smooth muscle, especially in the resistance vessels. Felodipine, an anti-hypertensive agent, induces autophagy. Felodipine can cross the blood-brain barrier <sup>[1][2][3]</sup> .	
IC <sub>50</sub> & Target	L-type calcium channel	
In Vitro	Felodipine, a dihydropyridine calcium antagonist, is the most potent in relaxing porcine coronary arteries (IC <sub>50</sub> =0.15 nM) <sup>[1]</sup> . Felodipine, an L-type calcium channel blocker, induces autophagy and clears diverse aggregate-prone, neurodegenerative disease-associated proteins <sup>[2]</sup> . Felodipine blocks the muscarinic receptor-mediated (carbachol) Ca2+-dependent contraction of guinea pig ileum	

longitudinal smooth muscle (GPILSM) with an  $IC_{50}$  of 1.45 nM<sup>[3]</sup>.

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

### **CUSTOMER VALIDATION**

• Int J Pharm. 2017 Jan 30;517(1-2):19-24.

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#### **REFERENCES**

[1]. Johnson JD, et al. Calcium and calmodulin antagonists binding to calmodulin and relaxation of coronary segments. J Pharmacol Exp Ther. 1983;226(2):330-334.

[2]. Siddiqi FH, et al. Felodipine induces autophagy in mouse brains with pharmacokinetics amenable to repurposing [published correction appears in Nat Commun. 2019 Jun 4;10(1):2530]. Nat Commun. 2019;10(1):1817. Published 2019 Apr 18.

[3]. Yiu, S. and E.E. Knaus, Synthesis, biological evaluation, calcium channel antagonist activity, and anticonvulsant activity of felodipine coupled to a dihydropyridine-pyridinium salt redox chemical delivery system. J Med Chem, 1996. 39(23): p. 4576-82.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA