## **Product** Data Sheet

# **Eliprodil**

Cat. No.: HY-12881 CAS No.: 119431-25-3 Molecular Formula: C<sub>20</sub>H<sub>23</sub>ClFNO

Molecular Weight: 347.85 iGluR Target:

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

-20°C Storage: Powder 3 years

> 4°C 2 years

-80°C In solvent 2 years

> -20°C 1 year

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 14.29 mg/mL (41.08 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.8748 mL	14.3740 mL	28.7480 mL
	5 mM	0.5750 mL	2.8748 mL	5.7496 mL
	10 mM	0.2875 mL	1.4374 mL	2.8748 mL

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

In Vivo

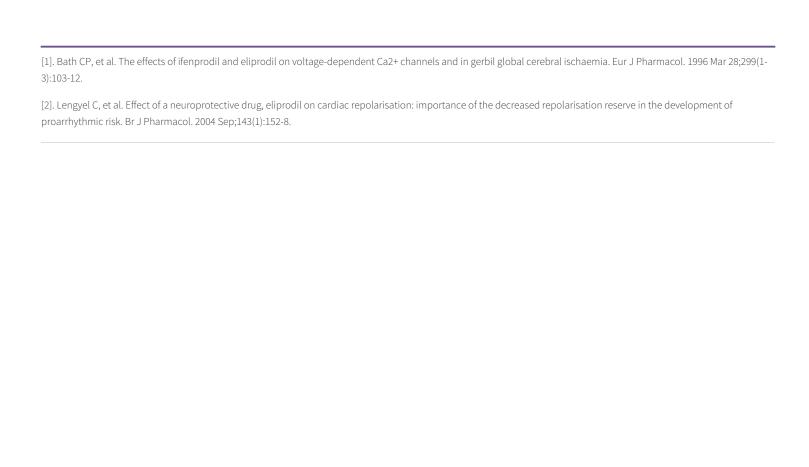
- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 1.43 mg/mL (4.11 mM); Suspended solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1.43 mg/mL (4.11 mM); Clear solution

### **BIOLOGICAL ACTIVITY**

Description

Eliprodil(SL-820715) is a non-competitive NR2B-NMDA receptor antagonist(IC50=1 uM), less potent for NR2A- and NR2Ccontaining receptors(IC50> 100 uM).IC50 value:Target: NR2B-NMDA antagonistHuman N-type Ca2+ channel currents were inhibited by ifenprodil and eliprodil with IC50 values of 50 microM and 10 microM respectively whereas P-type Ca2+ channel currents were inhibited reversibly by ifenprodil and eliprodil with approximate IC50 values of 60 microM and 9 microM respectively. eliprodil (1 microm) produced a moderate reverse rate-dependent prolongation of the action potential duration (7.4+/-1.5, 8.9+/-2.1 and 9.9+/-1.8% at cycle lengths of 300, 1000 and 5000 ms, respectively; n=9).

#### **REFERENCES**



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

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