

## **Product** Data Sheet

## Cav 2.2 blocker 2

 Cat. No.:
 HY-132268

 CAS No.:
 1204535-44-3

 Molecular Formula:
 C<sub>19</sub>H<sub>20</sub>F<sub>3</sub>N<sub>3</sub>O<sub>3</sub>S

Molecular Weight: 427.44

Target: Calcium Channel

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

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

Analysis.

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (233.95 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.3395 mL	11.6975 mL	23.3951 mL
	5 mM	0.4679 mL	2.3395 mL	4.6790 mL
	10 mM	0.2340 mL	1.1698 mL	2.3395 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: ≥ 2.5 mg/mL (5.85 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (5.85 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (5.85 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	Cav 2.2 blocker 2 is a Cav2.2 calcium channel blocker extracted from patent WO2017046581A1, compound 1. Cav 2.2 blocker 2 can reverses hyperalgesia associated with an injury or inflammation in conjunction with the opioid $^{[1]}$ .
IC <sub>50</sub> & Target	N-type calcium channel
In Vitro	Cav 2.2 blocker 2 (compound 1) (1-10 mg/kg; p.o.) in conjunction with Morphine (0.3 mg/kg) reverses the complete freunds adjuvant (CFA) induced hyperalgesia in rats <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES		
	omprising cav2.2 calcium channel blocker and opioid. WO2017046581A1.	
	Caution: Product has not been fully validated for medical applications. For research use only.	
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