

# Dihydrocapsaicin

Cat. No.: HY-N0361 CAS No.: 19408-84-5 Molecular Formula:  $C_{18}H_{29}NO_3$ 

Molecular Weight: 307.43

TRP Channel; Reactive Oxygen Species; Apoptosis; Caspase; Bcl-2 Family; Akt; PI3K Target:

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling; Immunology/Inflammation;

Metabolic Enzyme/Protease; NF-κB; Apoptosis; PI3K/Akt/mTOR

Storage: 4°C, protect from light

\* In solvent: -80°C, 6 months; -20°C, 1 month (protect from light)

**Product** Data Sheet

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (325.28 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.2528 mL	16.2639 mL	32.5277 mL
	5 mM	0.6506 mL	3.2528 mL	6.5055 mL
	10 mM	0.3253 mL	1.6264 mL	3.2528 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 (8.13 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (8.13 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (8.13 mM); Clear solution

# **BIOLOGICAL ACTIVITY**

Description Dihydrocapsaicin, a capsaicin, is a potent and selective TRPV1 (transient receptor potential vanilloid channel 1) agonist. Dihydrocapsaicin reduces AIF, Bax, and Caspase-3 expressions, and increased Bcl-2, Bcl-xL and p-Akt levels.

Dihydrocapsaicin enhances the hypothermia-induced neuroprotection following ischemic stroke via PI3K/Akt regulation in

rat[1][2][3]

PI3K IC<sub>50</sub> & Target TRPV1 Caspase 3 Bax

> Bcl-2 Bcl-xL

In Vitro	Dihydrocapsaicin (0-100 $\mu$ M) inhibits platelet aggregation and the activity of clotting factors VIII:C (6.26-100 $\mu$ M) and IX (25-100 $\mu$ M)[ $^{3}$ ]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
In Vivo	Dihydrocapsaicin (0.5 mg/kg, IP, once) exhibits hypothermic effect and neuroprotection in rat MCAO models <sup>[2]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Sprague-Dawley rats (adult, male, 300-340 g, subjected to right middle cerebral artery occlusion (MCAO) <sup>[2]</sup>	
	Dosage:	0.5 mg/kg	
	Administration:	IP, once	
	Result:	Exhibits hypothermic effect, rectal temperature dropped to approximately 35.0 ⋈ at 30 min, stayed at equal or below 35.0 ⋈ for approximately 20 min, and then gradually returned to approximately 36.5 ⋈ at 120 min. Significantly reduced Ischemia-reperfusion induced infarct volume (36.2% ± 2.5%). Reduces ROS levels at 24 h, and reduced ischemia-reperfusion induced a high level of cell death.	

# **CUSTOMER VALIDATION**

• Chem Eng J. 2023 Apr 15.

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## **REFERENCES**

- [1]. Adams MJ, et al. Effect of capsaicin and dihydrocapsaicin on in vitro blood coagulation and platelet aggregation. Thromb Res. 2009 Dec;124(6):721-3.
- [2]. Gao F, et al. Impairment in function and expression of transient receptor potential vanilloid type 4 in Dahl salt-sensitive rats: significance and mechanism. Hypertension. 2010 Apr;55(4):1018-25.
- [3]. Dihydrocapsaicin, et al. Dihydrocapsaicin (DHC) enhances the hypothermia-induced neuroprotection following ischemic stroke via PI3K/Akt regulation in rat. Brain Res. 2017 Sep 15;1671:18-25.

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

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