Proteins



Product Data Sheet

5-Hydroxydecanoate sodium

Cat. No.: HY-136615 CAS No.: 71186-53-3 Molecular Formula: C₁₀H₁₉NaO₃ Molecular Weight: 210.25

Potassium Channel Target:

Pathway: Membrane Transporter/Ion Channel 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: 12.5 mg/mL (59.45 mM; Need ultrasonic) H₂O: 10 mg/mL (47.56 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	4.7562 mL	23.7812 mL	47.5624 mL
	5 mM	0.9512 mL	4.7562 mL	9.5125 mL
	10 mM	0.4756 mL	2.3781 mL	4.7562 mL

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

In Vivo

- 1. Add each solvent one by one: PBS Solubility: 6.25 mg/mL (29.73 mM); Clear solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 1.25 mg/mL (5.95 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 1.25 mg/mL (5.95 mM); Suspended solution; Need ultrasonic
- 4. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1.25 mg/mL (5.95 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	5-Hydroxydecanoate sodium is a selective ATP-sensitive K^+ (K_{ATP}) channel blocker (IC_{50} of ~30 μ M). 5-Hydroxydecanoate sodium is a substrate for mitochondrial outer membrane acyl-CoA synthetase and has antioxidant activity ^{[1][2]} .
IC ₅₀ & Target	IC50: ~30 μM (K _{ATP})
In Vitro	5-Hydroxydecanoate (5-HD) treatment abolishs the beneficial effects of penehyclidine hydrochloride (PHC) preconditioning

	in anoxia/reoxygenation (A/R) ainduced injury in H9c2 cells. 5-Hydroxydecanoate blocks the inhibitory effect of PHC on Ca ²⁺ overload and ROS production. 5-Hydroxydecanoate promotes the release of Cyt-C from mitochondria into cytoplasm. 5-Hydroxydecanoate attenuats the anti-apoptotic effect of PHC. PHC treatment shows remarkably decreases levels of Bax and cleaved caspase-3, and increases levels of Bcl-2. 5-Hydroxydecanoate pretreatment reverses the effects of PHC on their expression levels. 5-Hydroxydecanoate blocks the effects of PHC on K _{ATP} channels ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	5-Hydroxydecanoate (100 μM) treatment abolishes the effects of ischemic preconditioning (IPC) on the contractile recovery and does not affect its effect on the contracture, lactate production, glycogenolysis and viable tissue in rats ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

- [1]. Congna Zi, et al. Penehyclidine hydrochloride protects against anoxia/reoxygenation injury in cardiomyocytes through ATP-sensitive potassium channels, and the $Akt/GSK-3\beta$ and Akt/mTOR signaling pathways. Cell Biol Int. 2020 Jun;44(6):1353-1362.
- [2]. Xiantao Li, et al. 5-Hydroxydecanoate and coenzyme A are inhibitors of native sarcolemmal KATP channels in inside-out patches. Biochim Biophys Acta. 2010 Mar;1800(3):385-91.
- [3]. M G Marina Prendes, et al. Effects of 5-hydroxydecanoate and ischemic preconditioning on the ischemic-reperfused heart of fed and fasted rats. J Physiol Biochem. 2005 Sep;61(3):447-56.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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