

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

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- Trockeneiszuschlag
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4'-Hydroxychalcone

Cat. No.: HY-N7056 CAS No.: 2657-25-2 Molecular Formula: $C_{15}H_{12}O_{2}$ Molecular Weight: 224.25

Target: Proteasome; Oxidative Phosphorylation

Pathway: Metabolic Enzyme/Protease Powder

2 years

-80°C In solvent 6 months -20°C 1 month

-20°C

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

Storage:

DMSO: 250 mg/mL (1114.83 mM; Need ultrasonic)

3 years

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	4.4593 mL	22.2965 mL	44.5931 mL
	5 mM	0.8919 mL	4.4593 mL	8.9186 mL
	10 mM	0.4459 mL	2.2297 mL	4.4593 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: ≥ 2.08 mg/mL (9.28 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (9.28 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (9.28 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

4'-Hydroxychalcone is a chalcone isolated from licorice root, with hepatoprotective activity. 4'-Hydroxychalcone inhibits $TNF\alpha-induced\ NF-\kappa B\ activation\ via\ proteasome\ inhibition.\ 4'-Hydroxychalcone\ induces\ a\ rapid\ potassium\ release\ from$ mitochondrial vesicles and causes deterioration of respiratory control and oxidative phosphorylation of isolated rat liver mitochondria[1][2][3].

In Vitro

4'-Hydroxychalcone (20-40 μM; 2 hours) inhibits TNFα-induced (20 ng/mL; 6 hours) NF-kB pathway activation in a dosedependent manner^[3].

- 4'-Hydroxychalcone (0.1-25 μ M ; 8 hours) inhibits proteasome activity in a dose-dependent manner but has no effect on IKK activity^[3].
- 4'-Hydroxychalcone inhibits TNF α -dependent degradation of IkB α and subsequently prevents p50/p65 nuclear translocation leading to 4'-Hydroxychalcone-inhibited expression of NF-kB target genes^[3].
- 4'-Hydroxychalcone affects cancer cell viability but has no significant effect on non-transformed cell viability [3].

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

Cell Viability Assay^[3]

Cell Line:	K562 cells, Jurkat cells, U937 cells, PBMCs	
Concentration:	5 μΜ, 10 μΜ, 20 μΜ, 24 μΜ, 28 μΜ, 32 μΜ, 40 μΜ, 60 μΜ	
Incubation Time:	24 hours	
Result:	Affected cancer cell viability but has no significant effect on non-transformed cell viability.	

Western Blot Analysis^[3]

Cell Line:	Jurkat cells	
Concentration:	60 μM (followed by TNFα 20 ng/mL)	
Incubation Time:	2 hours	
Result:	Inhibited TNF α -dependent degradation of IkB α and prevents p50/p65 nuclear translocation.	

In Vivo

4'-Hydroxychalcone has hepatoprotective activity against Acetaminophen induced hepatotoxicity in mice $^{[2]}$. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male albino mice(5-30 g) ^[2]	
Dosage:	25 mg/kg, 50 mg/kg, 100 mg/kg	
Administration:	Oral administration, 12h intervals, 4 doses	
Result:	Significantly reduced the mortality rate induced by Acetaminophen (1 g/kg).	

REFERENCES

- [1]. Inoue B, et al. The effects of echinatin and its related compounds on the mitochondrial energy transfer reaction. J Toxicol Sci. 1982 Nov;7(4):245-54.
- [2]. Ali Reza, et al. Hepatoprotective activity of phloretin and hydroxychalcones against Acetaminophen Induced hepatotoxicity in mice. Iranian Journal of Pharmaceutical Sciences. 2011 Jan; 7(2).
- [3]. Orlikova B, et al. The aromatic ketone 4'-hydroxychalcone inhibits TNF α -induced NF- κ B activation via proteasome inhibition. Biochem Pharmacol. 2011 Sep 15;82(6):620-31.

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

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