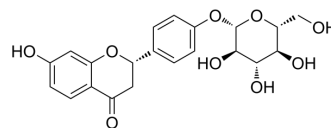


Liquiritin

Cat. No.:	HY-N0376												
CAS No.:	551-15-5												
Molecular Formula:	C ₂₁ H ₂₂ O ₉												
Molecular Weight:	418.39												
Target:	Reactive Oxygen Species												
Pathway:	Immunology/Inflammation; Metabolic Enzyme/Protease; NF-κB												
Storage:	<table border="0"> <tr> <td>Powder</td> <td>-20°C</td> <td>3 years</td> </tr> <tr> <td></td> <td>4°C</td> <td>2 years</td> </tr> <tr> <td>In solvent</td> <td>-80°C</td> <td>2 years</td> </tr> <tr> <td></td> <td>-20°C</td> <td>1 year</td> </tr> </table>	Powder	-20°C	3 years		4°C	2 years	In solvent	-80°C	2 years		-20°C	1 year
Powder	-20°C	3 years											
	4°C	2 years											
In solvent	-80°C	2 years											
	-20°C	1 year											



SOLVENT & SOLUBILITY

In Vitro

DMSO : 150 mg/mL (358.52 mM; Need ultrasonic)
 H₂O : < 0.1 mg/mL (ultrasonic) (insoluble)

	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.3901 mL	11.9506 mL	23.9011 mL
	5 mM	0.4780 mL	2.3901 mL	4.7802 mL
	10 mM	0.2390 mL	1.1951 mL	2.3901 mL

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

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
 Solubility: ≥ 2.5 mg/mL (5.98 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
 Solubility: ≥ 2.5 mg/mL (5.98 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
 Solubility: ≥ 2.5 mg/mL (5.98 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Liquiritin, a flavonoid isolated from *Glycyrrhiza uralensis*, is a potent and competitive AKR1C1 inhibitor with IC₅₀s of 0.62 μM, 0.61 μM, and 3.72 μM for AKR1C1, AKR1C2 and AKR1C3, respectively. Liquiritin efficiently inhibits progesterone metabolism mediated by AKR1C1 in vivo^[1]. Liquiritin acts as an antioxidant and has neuroprotective, anti-cancer and anti-inflammatory activity^[2].

IC₅₀ & Target

IC₅₀: 0.62 μM (AKR1C1), 0.61 μM (AKR1C2) and 3.72 μM (AKR1C3)^[1]

In Vitro

Liquiritin can target the residues Ala-27, Val-29, Ala-25, and Asn-56 of AKR1C1^[1].

Liquiritin (50 μ M; 6 hours) results in 85.00% of reduction in progesterone metabolism, which is mediated by Aldo-keto reductase family 1 member C1 (AKR1C1) enzymatic activity in HEC-1-B cells^[1].

Liquiritin (100 μ M) increases glucose-6-phosphate dehydrogenase expression on B65 neuroblastoma cells^[2].

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

CUSTOMER VALIDATION

- Acta Biomater. 2022 Aug 11;S1742-7061(22)00454-8.
- Int Immunopharmacol. December 2021, 108283.
- Drug Des Devel Ther. 2023 Oct 3;17:3047-3060.
- Plant Direct. 2022 Sep 2;6(9):e442.
- Planta Med. 2021 Feb 5.

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REFERENCES

[1]. Nakatani Y, et al. Neuroprotective effect of liquiritin as an antioxidant via an increase in glucose-6-phosphate dehydrogenase expression on B65 neuroblastoma cells. Eur J Pharmacol. 2017 Nov 15;815:381-390.

[2]. Zeng C, et al. Liquiritin, as a Natural Inhibitor of AKR1C1, Could Interfere With the Progesterone Metabolism. Front Physiol. 2019 Jul 3;10:833.

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

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