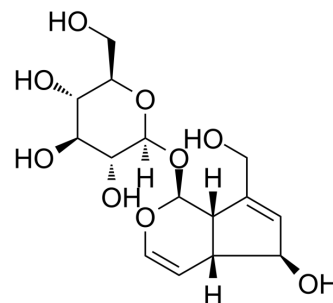


Aucubin

Cat. No.:	HY-N0664		
CAS No.:	479-98-1		
Molecular Formula:	C ₁₅ H ₂₂ O ₉		
Molecular Weight:	346.33		
Target:	Bacterial		
Pathway:	Anti-infection		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (288.74 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg
		Concentration				
		1 mM		2.8874 mL	14.4371 mL	28.8742 mL
5 mM			0.5775 mL	2.8874 mL	5.7748 mL	
	10 mM		0.2887 mL	1.4437 mL	2.8874 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 (7.22 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (7.22 mM); Clear solution					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (7.22 mM); Suspended solution; Need ultrasonic					

BIOLOGICAL ACTIVITY

Description	Aucubin, an iridoid glucoside, is isolated from <i>Plantago asiatica</i> , <i>Eucommia ulmoides</i> , the leaves of <i>Aucuba japonica</i> and more recently from butterfly larva. Aucubin has many biological activities, such as antioxidant, anti-aging, anti-inflammatory, antimicrobial, anti-fibrotic, anti-cancer, hepatoprotective, neuroprotective and osteoprotective effects ^{[1][2][3]} .
In Vitro	Aucubin (0.001-1 μg/mL; pretreated for 30 min) dose-dependently inhibits IgE-induced TNF-α and IL-6 production and expression in RBL-2H3 cells, with IC ₅₀ s of 0.101 and 0.19 μg/mL, respectively ^[2] .

Aucubin (0.01 µg/mL; pretreated for 30 min) inhibits IgE-induced nuclear translocation of p65 subunit of NF-κB and degradation of IκBα in RBL-2H3 cells^[2].
Aucubin (0.001-1 mM; 12 h) increases PC12 cellular viability and markedly inhibits H₂O₂-induced apoptotic cell death^[4].
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

Aucubin (5 mg/kg; i.p. for 15 d) has antioxidant and pancreas-protective effects on rats with streptozotocin-induced diabetes ^[1].
Aucubin (40-200 mg/kg; a single i.p.) exhibits significant protective activity against α-amanitin intoxication in mice^[5].
Aucubin (5 mg/kg/day; i.p. for 21 d) decreases the breathing frequency, increases the lung dynamic compliance, alleviates lung parenchymal fibrotic changes, and reduces the intrapulmonary collagen disposition and inflammatory injury of BLM-stimulated mice^[6].
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male Wistar rats (200-230 g) induced diabetes by a injection of streptozotocin ^[1]
Dosage:	5 mg/kg
Administration:	I.p. twice daily for the first 5 days, followed by single injections daily for the last 10 days
Result:	Increased the body weight of streptozotocin-diabetic rats. Lowered the blood glucose level. Decreased the level of lipid peroxidation and increased the activities of antioxidant enzymes. Increased in insulin immunoreactivity and the number of immunoreactive β cells compared with untreated diabetic rats.

CUSTOMER VALIDATION

- Research Square Preprint. 2021 Feb.

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REFERENCES

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

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