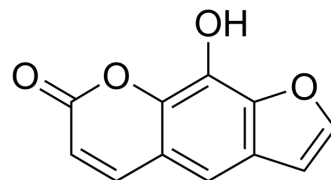


## Xanthotoxol

<b>Cat. No.:</b>	HY-30152		
<b>CAS No.:</b>	2009-24-7		
<b>Molecular Formula:</b>	C <sub>11</sub> H <sub>6</sub> O <sub>4</sub>		
<b>Molecular Weight:</b>	202.16		
<b>Target:</b>	5-HT Receptor; Reactive Oxygen Species; Endogenous Metabolite		
<b>Pathway:</b>	GPCR/G Protein; Neuronal Signaling; Immunology/Inflammation; Metabolic Enzyme/Protease; NF-κB		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 100 mg/mL (494.66 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	<b>Preparing Stock Solutions</b>	1 mM	4.9466 mL	24.7329 mL	49.4658 mL
		5 mM	0.9893 mL	4.9466 mL	9.8932 mL
		10 mM	0.4947 mL	2.4733 mL	4.9466 mL
Please refer to the solubility information to select the appropriate solvent.					
<b>In Vivo</b>	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (12.37 mM); Clear solution  2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (12.37 mM); Clear solution				

### BIOLOGICAL ACTIVITY

<b>Description</b>	Xanthotoxol (8-Hydroxypsoralen) It is a kind of fragrant bean substance, and it is a CYP450 inhibitor. Xanthotoxol has anti-inflammatory, anti-inflammatory, and 5-HT antagonistic and protective effects. Xanthotoxol inhibited CYP3A4 sum CYP1A2 IC50s separation 7.43 μM sum 27.82 μM. Xanthotoxol can pass through MAPK and NF-κB, inhibiting inflammation.	
<b>IC<sub>50</sub> &amp; Target</b>	Human Endogenous Metabolite	serotonin
<b>In Vitro</b>	Xanthotoxol (62.5-250 μM; 20 min) reduces the production of inflammatory factors induced by LPS stimulation in a concentration-dependent manner and inhibits LPS-induced p65 translocation from the cytoplasm to the nucleus in RAW	

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264.7 cells<sup>[3]</sup>.

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

**In Vivo**

Xanthotoxol (5 mg/kg, 10 mg/kg; ip; 2 dose) can attenuate the expression of pro-inflammatory mediators in the ischemia/reperfusion (I/R) rat model, thereby inhibiting the inflammatory response after cerebral ischemia<sup>[4]</sup>.

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

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## CUSTOMER VALIDATION

- Eur J Pharmacol. 2023 Oct 21:176147.
- bioRxiv. 2023 Jun 15.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

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## REFERENCES

[1]. Ma Z, et al. Metabolism and Metabolic Inhibition of Xanthotoxol in Human Liver Microsomes. Evid Based Complement Alternat Med. 2016;2016:5416509.

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

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