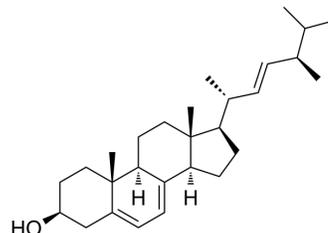


## Ergosterol

Cat. No.:	HY-N0181
CAS No.:	57-87-4
Molecular Formula:	C <sub>28</sub> H <sub>44</sub> O
Molecular Weight:	396.65
Target:	Endogenous Metabolite
Pathway:	Metabolic Enzyme/Protease
Storage:	4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



### SOLVENT & SOLUBILITY

In Vitro	DMSO : 3.57 mg/mL (9.00 mM; ultrasonic and warming and heat to 60°C)				
	Ethanol : 2.6 mg/mL (6.55 mM; ultrasonic and warming and heat to 80°C)				
	Preparing Stock Solutions	Solvent	1 mg	5 mg	10 mg
		Concentration			
		1 mM	2.5211 mL	12.6056 mL	25.2111 mL
5 mM		0.5042 mL	2.5211 mL	5.0422 mL	
	10 mM	---	---	---	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 0.5% CMC-Na >> 0.5% Tween-80 Solubility: 2.5 mg/mL (6.30 mM); Suspended solution; Need ultrasonic				

### BIOLOGICAL ACTIVITY

Description	Ergosterol is the primary sterol found in fungi, with antioxidative, anti-proliferative, and anti-inflammatory effects.
In Vivo	Ergosterol (25, 50 mg/kg, p.o.) significantly mitigates the reduced cardiac performance in rats induced by LPS, increases SOD activity and decreases the formation of MDA, CK-MB, and LDH in LPS-induced sepsis rats <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### PROTOCOL

Animal Administration <sup>[2]</sup>	Rats <sup>[2]</sup> Experimental myocardial injury in rats is performed by LPS injection (15 mg/kg). Dexmedetomidine (Dex) is used as a positive control. The experimental animals are randomly divided into five groups (n = 10) as follows: Control group, rats
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receive 2% gum acacia suspension orally at a dose of 2 mL/kg for 5 days, followed by normal saline injected intraperitoneally on day 5; LPS group, rats receive 2% gum acacia suspension at dose of 2 mL/kg for 5 days with LPS simultaneously injected intraperitoneally day 5; LPS+ Dex group, rats are treated with 2 mg/kg Dex suspension followed by LPS injection on day 5; LPS + Ergosterol (25 mg/kg, 50 mg/kg) groups, 25 or 50 mg/kg Ergosterol are given to rats orally for 5 consecutive days, and LPS is injected on day 5. Twelve hours after LPS treatment, blood samples are collected through the retro-orbital plexus. The serum specimens are centrifugated at  $4,000 \times g$  for 15 min and stored at  $-80^{\circ}\text{C}$  until needed. Thereafter, rats are anesthetized and sacrificed. Heart tissues are removed and homogenized in ice-cold phosphate buffered saline (50 mM, pH 7.4). Heart tissue homogenates from different groups are centrifuged at  $12,000 \times g$  for 45 min at  $4^{\circ}\text{C}$  and the supernatants retained for further biochemical evaluations<sup>[2]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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## CUSTOMER VALIDATION

- Commun Biol. 2023 Jan 3;6(1):1.

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## REFERENCES

[1]. Xu J, et al. Ergosterol Attenuates LPS-Induced Myocardial Injury by Modulating Oxidative Stress and Apoptosis in Rats. Cell Physiol Biochem. 2018;48(2):583-592.

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

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