

## **Product** Data Sheet

# alpha-Asarone

Cat. No.:HY-N0700CAS No.:2883-98-9Molecular Formula: $C_{12}H_{16}O_3$ Molecular Weight:208.25

Target: GABA Receptor

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

**Storage:** Powder -20°C 3 years

4°C 2 years -80°C 2 years

In solvent -80°C 2 years

-20°C 1 year

#### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (480.19 mM; Need ultrasonic)

H<sub>2</sub>O: < 0.1 mg/mL (insoluble)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	4.8019 mL	24.0096 mL	48.0192 mL
	5 mM	0.9604 mL	4.8019 mL	9.6038 mL
	10 mM	0.4802 mL	2.4010 mL	4.8019 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 (12.00 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (12.00 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (12.00 mM); Clear solution

### **BIOLOGICAL ACTIVITY**

Description	alpha-Asarone ( $\alpha$ -Asarone) is one of the main psychoactive compounds, and possesses an antidepressant-like activity in mice.
In Vitro	The results indicated that $\alpha$ -asarone significantly attenuated the LPS-stimulated increase in neuroinflammatory responses and suppressed pro-inflammatory cytokine production in BV-2 cells. Mechanistic study revealed that alpha-Asarone ( $\alpha$ -Asarone) inhibited the LPS-stimulated activation via regulation of nuclear factor kappa-B by blocking degradation of

	inhibitor kappa B-alpha signaling in BV-2 microglial cells <sup>[2]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
In Vivo	The present results reveal that the acute treatment of alpha-Asarone ( $\alpha$ -Asarone) elicited biphasic responses on immobility such that the duration of the immobility time is significantly reduced at lower doses (15 and 20 mg/kg, i.p.) but increased at higher doses (50 and 100 mg/kg, i.p.) in the TST. Besides, alpha-Asarone ( $\alpha$ -Asarone) at higher doses (50 and 100 mg/kg, i.p.) significantly decreased the spontaneous locomotor activity <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

#### **REFERENCES**

- [1]. Ranjithkumar Chellian, et al. Biphasic Effects of  $\alpha$ -Asarone on Immobility in the Tail Suspension Test: Evidence for the Involvement of the Noradrenergic and Serotonergic Systems in Its Antidepressant-Like Activity. Front Pharmacol. 2016; 7: 72.
- [2]. Byung-Wook Kim, et al.  $\alpha$ -Asarone attenuates microglia-mediated neuroinflammation by inhibiting NF kappa B activation and mitigates MPTP-induced behavioral deficits in a mouse model of Parkinson's disease. Neuropharmacology, Volume 97, October 2015, Pages 46–57
- [3]. Hye-Jung Park, et al. Effect of  $\alpha$ -asarone on angiogenesis and matrix metalloproteinase. Environmental Toxicology and Pharmacology, Volume 39, Issue 3, May 2015, Pages 1107–1114

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

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