# **Product** Data Sheet

## Climbazole

Cat. No.: HY-B1151 CAS No.: 38083-17-9 Molecular Formula:  $\mathsf{C}_{15}\mathsf{H}_{17}\mathsf{CIN}_2\mathsf{O}_2$ 

Molecular Weight: 292.76 Target: Fungal

Pathway: Anti-infection

Storage: Powder -20°C 3 years

2 years

In solvent -80°C 2 years

> -20°C 1 year

### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (341.58 mM; Need ultrasonic)

	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	3.4158 mL	17.0788 mL	34.1577 mL
	5 mM	0.6832 mL	3.4158 mL	6.8315 mL
	10 mM	0.3416 mL	1.7079 mL	3.4158 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 (8.54 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (8.54 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (8.54 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description

Climbazole (BAY-e 6975) is a potent antifungal agent. Climbazole also is a potent inducer of rat hepatic cytochrome P450<sup>[2]</sup>.

In Vitro

Climbazole (20 µM; 48 hours) significantly decreases exosome secretion in aggressive prostate cancer (PCa) cells<sup>[1]</sup>. Climbazole (20 µM) significantly inhibits the protein concentration of Alix, and Rab27a but not nSMase2[1]. Climbazole is a potent inhibitor of exosome biogenesis and/or secretion<sup>[2]</sup>.

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

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[1]. Amrita Datta, et al. High-throughput screening identifie Rep. 2018 May 25;8(1):8161.	d selective inhibitors of exosome	biogenesis and secretion: A drug repurposing strategy for advanced cancer. Sc		
[2]. Y Kobayashi, et al. Climbazole is a new potent inducer of rat hepatic cytochrome P450. J Toxicol Sci. 2001 Aug;26(3):141-50.				
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