Antimicrobial agent-8

Cat. No.: HY-151402 Molecular Formula: $C_{39}H_{54}N_{16}$ 746.95 Molecular Weight:

Bacterial Target: Pathway: Anti-infection

Storage: Powder -20°C 3 years

4°C 2 years

In solvent -80°C 6 months

> -20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (133.88 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.3388 mL	6.6939 mL	13.3878 mL
	5 mM	0.2678 mL	1.3388 mL	2.6776 mL
	10 mM	0.1339 mL	0.6694 mL	1.3388 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 (3.35 mM); Clear solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (3.35 mM); Clear solution; Need ultrasonic
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (3.35 mM); Clear solution; Need ultrasonic

BIOLOGICAL ACTIVITY

Description

Antimicrobial agent-8 (Compound 15) is a potent antimicrobial agent, and shows potent antimicrobial activity with an MIC range of 2-8 μg/mL against Gram-negative and Gram-positive bacteria. Antimicrobial agent-8 shows anti-inflammatory activity against lipopolysaccharide-induced inflammation.

In Vitro

Antimicrobial agent-8 (2.8-56.4 μM; 24 h) inhibits Gram-negative bacteria and Gram-positive bacteria growth^[1]. Antimicrobial agent-8 (5 and 20 μg/mL; 18 h) inhibits the production of nitric oxide (NO) and tumor necrosis factor-α (TNF-α) by lipopolysaccharide-stimulated in RAW 264.7 cells^[1].

Antimicrobial agent-8 (1-32 μg/mL, 16 h; 8-128 μg/mL; 24 h) shows potent biofilm inhibitory (MBIC₅₀=1 μg/mL) and eradicating activities (MBEC₅₀=8 μ g/mL) by MDRPA bacteria^[1].

MCE has not independence. Cell Viability Assay ^[1]	ntly confirmed the accuracy of these methods. They are for reference only.	
Cell Line:	E. coli [KCTC 1682], P. aeruginosa [KCTC 1637], S. epidermidis [KCTC 1917] and S. aureus [KCTC1621]	
Concentration:	2.8-56.4 μM	
Incubation Time:	24 hours	
Result:	Inhibited Gram-negative bacteria with MIC values of 5.4 μM for E. coli [KCTC 1682] and P. aeruginosa [KCTC 1637]. Inhibited Gram- positive bacteria with MIC values of 2.7 μM and 5.4 μM for S. epidermidis [KCTC 1917] and S. aureus [KCTC1621], respectively.	
Cell Viability Assay ^[1]		
Cell Line:	RAW 264.7 macrophages	
Concentration:	5 and 20 μg/mL	
Incubation Time:	18 hours	
Result:	Observed LPS-stimulated production of NO with an inhibitory rate of 90.79% at 5 μ g/ml Exhibited inhibitory effects on the LPS-stimulated production of TNF- α with an inhibito rate of 95.4% at 20 μ g/mL.	

REFERENCES

[1]. Dinesh Kumar S, et al. Cationic, amphipathic small molecules based on a triazine-piperazine-triazine scaffold as a new class of antimicrobial agents. Eur J Med Chem. 2022 Sep 8;243:114747.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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