

# **Product** Data Sheet

### **RNPA1000**

Cat. No.: HY-12824 CAS No.: 359600-10-5 Molecular Formula:  $C_{23}H_{18}BrN_3O_3$ 

Molecular Weight: 464.31

Target: Bacterial; Antibiotic

Pathway: Anti-infection

Storage: Powder -20°C 3 years

4°C 2 years

In solvent -80°C 2 years

-20°C 1 year

### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 50 mg/mL (107.69 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.1537 mL	10.7687 mL	21.5373 mL
	5 mM	0.4307 mL	2.1537 mL	4.3075 mL
	10 mM	0.2154 mL	1.0769 mL	2.1537 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 (5.38 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (5.38 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (5.38 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	RNPA1000, an antibiotic, is a potent RnpA inhibitor and inhibits RnpA-mediated cellular RNA degradation. RNPA1000 inhib		
	tRNA maturation with an IC $_{50}$ of 175 $\mu$ M. RNPA1000 displays broad-spectrum antimicrobial activities and inhibits		
	staphylococcal and all Gram-positive bacterial pathogens activity $^{[1][2][3]}$ .		

RNPA1000 displays antimicrobial activity toward Gram-positive bacteria and little or no toxicity toward human cells<sup>[2]</sup>. RNPA1000 limits S. aureus mRNA turnover and growth. RNPA1000 also limits growth of other important Gram-positive bacterial pathogens, exhibits antimicrobial efficacy against biofilm associated S. aureus and protects against the S. aureus

In Vitro

pathogenesis in an animal model of infection<sup>[3]</sup>.

RNPA1000 (IC50= 100-125  $\mu$ M), does not affect the activity of the commercially available E. coli RNase HI, RNase A, RNase I or in-house purified S. aureus RNase J1 at any concentration tested (0-750  $\mu$ M), but does mildly inhibit E. coli RNase III activity (IC50= 500-750  $\mu$ M)[3].

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

#### **REFERENCES**

[1]. Eidem TM, et al. Drug-eluting cements for hard tissue repair: a comparative study using vancomycin and RNPA1000 to inhibit growth of Staphylococcus aureus. J Biomater Appl. 2014 Apr;28(8):1235-46.

[2]. Eidem TM, et al. Small-molecule inhibitors of Staphylococcus aureus RnpA-mediated RNA turnover and tRNA processing. Antimicrob Agents Chemother. 2015 Apr;59(4):2016-28.

[3]. Patrick D Olson, et al. Small molecule inhibitors of Staphylococcus aureus RnpA alter cellular mRNA turnover, exhibit antimicrobial activity, and attenuate pathogenesis. PLoS Pathog. 2011 Feb 10;7(2):e1001287.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA

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