# ATP synthase inhibitor 2

Cat. No.: HY-150983 CAS No.: 2814540-76-4 Molecular Formula:  $C_{21}H_{22}N_2O_3S$ Molecular Weight: 382.48

Target: ATP Synthase; Bacterial

Pathway: Membrane Transporter/Ion Channel; Anti-infection

Storage: Powder -20°C 3 years

2 years

In solvent -80°C 6 months

> -20°C 1 month

**Product** Data Sheet

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 250 mg/mL (653.63 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.6145 mL	13.0726 mL	26.1452 mL
	5 mM	0.5229 mL	2.6145 mL	5.2290 mL
	10 mM	0.2615 mL	1.3073 mL	2.6145 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)

Solubility: ≥ 2.08 mg/mL (5.44 mM); Clear solution

2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (5.44 mM); Clear solution

### **BIOLOGICAL ACTIVITY**

Description	can inhibit Pseudomonas aeruginosa (PA) ATP synthase inhibitor ( $C_{50}$ –10 µg/mL). ATP synthase inhibitor 2 can inhibit Pseudomonas aeruginosa (PA) ATP synthasis activity completely at 128 µg/mL <sup>[1]</sup> .
IC <sub>50</sub> & Target	IC50: 10 μg/mL (Pseudomonas aeruginosa ATP synthase) <sup>[1]</sup>
In Vitro	ATP synthase inhibitor 2 (Compound 22) (0-256 µg/mL; 12-15 h) treatment inhibits ATP synthase but not shows whole cell antibiotic activity against Pseudomonas aeruginosa <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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Cell Viability Assay<sup>[1]</sup>

Cell Line:	MSSA and WTPA	
Concentration:	0-256 μg/mL	
Incubation Time:	12-15 hours	
Result:	Showed no activities against MSSA and WTPA (MICs>128 $\mu$ g/mL and >256 $\mu$ g/mL, respectively).	

## **REFERENCES**

[1]. John F. Ciprich, et al. Synthesis and Evaluation of Pseudomonas aeruginosa ATP Synthase Inhibitors. ACS Omega.

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

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