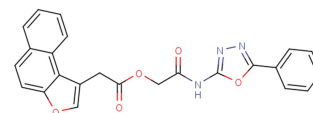


LtaS-IN-1

Chemical Properties

CAS No.: 877950-01-1
Formula: C₂₄H₁₇N₃O₅
Molecular Weight: 427.41
Appearance: N/A
Storage: 0-4°C for short term (days to weeks), or -20°C for long term (months).

**Biological Description**

Description	LtaS-IN-1 alone inhibits Enterococcus.spp 28 strains with varying MIC values ranging from 0.5 µg/mL to 64 µg/mL. LtaS-IN-1 combination with antibiotics abolishes multidrug-resistant E. faecium growth almost completely. LtaS-IN-1 is a potent small-molecule inhibitor of Lipoteichoic acid (LTA) synthesis in multidrug-resistant (MDR) E. faecium and by altering the cell wall morphology.
Targets(IC ₅₀)	strain E1630: (MIC)0.5 µg/mL strain E1590: 0.5 µg/mL
In vitro	LtaS-IN-1 (20 µM) combines with either ampicillin (20 µg/mL), gentamicin (10 µg/mL), linezolid (5 µg/mL), daptomycin (10 µg/mL+50 µg/mL calcium chloride) or vancomycin (20 µg/mL) can inhibit strains E7128 and E7130 growth by 97-100%, while LtaS-IN-1 alone only gives 73% (strain E7128) and 8% (strain E7130) of growth inhibition, respectively[1].LtaS-IN-1 (0-100 µM) inhibits strain E745 growth as a concentration-dependent manner. At the concentration 10 µM leads to an 60% reduction in the final OD600 for this strain. Meanwhile, LtaS-IN-1 does not affect E. faecium growth in control group[1]. LtaS-IN-1 is against Enterococcus spp 28 strains with varying MIC values ranging from 0.5 µg/mL to 64 µg/mL. LtaS-IN-1 inhibits strain E1630 and E1590 with the MIC values of 0.5µg/mL[1].

Solubility Information

Solubility	< 1 mg/ml refers to the product slightly soluble or insoluble
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.34 mL	11.698 mL	23.397 mL
5 mM	0.468 mL	2.34 mL	4.679 mL
10 mM	0.234 mL	1.17 mL	2.34 mL
50 mM	0.047 mL	0.234 mL	0.468 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. The storage conditions and period of the stock solution: - 80 °C for 6 months; - 20 °C for 1 month. Please use it as soon as possible.

Reference

1. Paganelli FL, et al. Lipoteichoic acid synthesis inhibition in combination with antibiotics abrogates growth of multidrug-resistant Enterococcus faecium.Int J Antimicrob Agents. 2017 Mar;49(3):355-363.

Inhibitors · Natural Compounds · Compound Libraries

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