# **Screening Libraries**



# **JPD447**

Cat. No.: HY-139628 CAS No.: 2883235-86-5 Molecular Formula:  $C_{20}H_{23}FN_{4}$ Molecular Weight: 338.42 Bacterial Target: Pathway: Anti-infection

Storage: Powder -20°C 3 years 2 years

-80°C In solvent 6 months

> -20°C 1 month

**Product** Data Sheet

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (295.49 mM; ultrasonic and warming and heat to 80°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.9549 mL	14.7745 mL	29.5491 mL
	5 mM	0.5910 mL	2.9549 mL	5.9098 mL
	10 mM	0.2955 mL	1.4775 mL	2.9549 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 (7.39 mM); Clear solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (7.39 mM); Clear solution; Need ultrasonic

## **BIOLOGICAL ACTIVITY**

Description

JPD447, a MAC-0547630 derivative, is a novel class of UppS inhibitor to potentiate  $\beta$ -lactam antibiotics.

### **REFERENCES**

[1]. Workman SD, et al. Structural Insights into the Inhibition of Undecaprenyl Pyrophosphate Synthase from Gram-Positive Bacteria. J Med Chem. 2021 Sep 23;64(18):13540-13550.

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

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

Page 2 of 2 www.MedChemExpress.com