**Proteins** 

# Inhibitors



# Fobrepodacin disodium

Cat. No.: HY-135655 CAS No.: 1384984-20-6 Molecular Formula:  $C_{21}H_{24}FN_6Na_2O_6P$ 

Molecular Weight: 552.4 Target: Bacterial Pathway: Anti-infection

Storage: 4°C, sealed storage, away from moisture

\* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)

**Product** Data Sheet

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 5 mg/mL (9.05 mM; ultrasonic and adjust pH to 3 with HCl)

	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	1.8103 mL	9.0514 mL	18.1028 mL
	5 mM	0.3621 mL	1.8103 mL	3.6206 mL
	10 mM			

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

### **BIOLOGICAL ACTIVITY**

Description Fobrepodacin (SPR720) disodium is an orally active and potent phosphate proagent of SPR719 (VXc-486; HY-12930). Fobrepodacin disodium has potent bactericidal activities in vivo  $^{\left[1\right]}$ .

In Vivo Fobrepodacin disodium (oral gavage; 10, 30, 100 mg/kg; once per day; 5 times per week for 4 weeks) reduces the mycobacterial burden in a model of chronic tuberculosis infection in mice[1].

> Fobrepodacin disodium (oral; 100 mg/kg; once per day; 5 days per week, for 8 weeks) improves the bactericidal activities of antimycobacterial drugs<sup>[1]</sup>.

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

Animal Model:	Six-week old female BALB/c and C57BL/6 mice uninfected or M. tuberculosis-infected (Erdman) $^{[1]}$
Dosage:	10, 30, 100 mg/kg
Administration:	Oral gavage; once per day; 5 times per week for 4 weeks

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io	d the mycobacterial burden in a model of chronic tuberculosis infectio	Result:

#### **REFERENCES**

[1]. Locher CP, et al. A novel inhibitor of gyrase B is a potent drug candidate for treatment of tuberculosis and nontuberculosis mycobacterial infections. Antimicrob Agents Chemother. 2015 Mar;59(3):1455-65.

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

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Page 2 of 2 www.MedChemExpress.com