# **Product** Data Sheet

# Dirithromycin

Cat. No.:HY-B0643CAS No.:62013-04-1Molecular Formula: $C_{42}H_{78}N_2O_{14}$ Molecular Weight:835.07

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 Ethanol : ≥ 50 mg/mL (59.88 mM)

DMSO: 33.33 mg/mL (39.91 mM; Need ultrasonic)

\* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.1975 mL	5.9875 mL	11.9750 mL
	5 mM	0.2395 mL	1.1975 mL	2.3950 mL
	10 mM	0.1198 mL	0.5988 mL	1.1975 mL

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

In Vivo

- 1. Add each solvent one by one: 10% EtOH >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility:  $\geq$  2.5 mg/mL (2.99 mM); Clear solution
- 2. Add each solvent one by one: 10% EtOH >> 90% (20% SBE- $\beta$ -CD in saline) Solubility:  $\geq$  2.5 mg/mL (2.99 mM); Clear solution
- 3. Add each solvent one by one: 10% EtOH >> 90% corn oil Solubility:  $\geq$  2.5 mg/mL (2.99 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	Dirithromycin (LY237216), a derivative of Erythromycin, is a potent and orally active semi-synthetic macrolide antibiotic.  Dirithromycin is active against gram-positive bacteria, Legionella spp., Helicobacter pylori, and Chlamydia trachomatis <sup>[1][2]</sup> .
IC., & Target	Macrolide

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In Vitro	Dirithromycin possesses an in vitro spectrum of antimicrobial activity which is similar to that of Erythromycin <sup>[2]</sup> . Dirithromycin exhibits excellent in vitro activity against several strains of Legionella, with MICs of ~1.0 and <0.25 $\mu$ g/mL at pH=7.1 and 7.4, respectively <sup>[2]</sup> . Dirithromycin demonstrates potent activity against several strains of Helicobacter pytori, with MICs of <0.5 $\mu$ g/mL <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	Dirithromycin (s.c. for 2 times) is effective against experimental infections caused by S. aureus, S. pyogenes, and S. pneumoniae in mice, with $ED_{50}$ s of 1.0, 0.6, and <0.6 mg/kg <sup>[2]</sup> . Dirithromycin (p.o. for 2 times) is effective against experimental infections caused by S. aureus, S. pyogenes, and S. pneumoniae in mice, with $ED_{50}$ s of 27, 34, and 23 mg/kg <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **CUSTOMER VALIDATION**

• Cell Prolif. 2021 Jan;54(1):e12953.

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#### **REFERENCES**

[1]. Wasilewski MM, et, al. Five-day dirithromycin therapy is as effective as seven-day erythromycin therapy for acute exacerbations of chronic bronchitis. J Antimicrob Chemother. 1999 Apr;43(4):541-8.

[2]. Counter FT, et, al. Synthesis and antimicrobial evaluation of dirithromycin (AS-E 136; LY237216), a new macrolide antibiotic derived from erythromycin. Antimicrob Agents Chemother. 1991 Jun;35(6):1116-26.

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

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