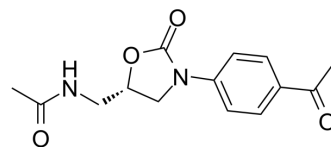


Dup-721

Cat. No.:	HY-139618		
CAS No.:	104421-21-8		
Molecular Formula:	C ₁₄ H ₁₆ N ₂ O ₄		
Molecular Weight:	276.29		
Target:	Bacterial; Antibiotic		
Pathway:	Anti-infection		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



BIOLOGICAL ACTIVITY

Description	DuP-721 is a broad spectrum and orally active antibacterial agent against a variety of clinically susceptible and resistant bacteria, especially <i>M. tuberculosis</i> ^[1] .
In Vitro	DuP-721 (1.5-4 µg/ml) inhibits equally the strains of <i>Mycobacterium tuberculosis</i> susceptible and resistant to conventional antituberculosis drug. And it does not show cross resistance to any of the anti-tuberculosis drugs tested ^[1] . DuP-721 is inactive against <i>M. avium</i> and <i>M. intracellulare</i> at 250 µg/ml. It inhibits <i>M. gordonae</i> and <i>M. fortuitum</i> at 3.9 µg/ml and <i>M. kansasii</i> and <i>M. scrofulaceum</i> at 1.95 µg/ml and 15.6 µg/ml, respectively ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	DuP-721 (oral gavage; 50-160 mg/kg) is protective against <i>M. tuberculosis</i> infection in mice. DuP-721 protects 100% of the infected animals at 50 mg/kg p.o. dose when administered daily for 17 days, and the same effect is observed at 160 mg/kg dose when the drug is administered only on day 11 and 12 post infection ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Affiliatio, et al. Antimycobacterial activities of oxazolidinones: a review. *Infect Disord Drug Targets*. 2006 Dec;6(4):343-54.

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