Amphomycin

MedChemExpress

Cat. No.:	HY-P3078	
CAS No.:	1402-82-0	
Target:	Bacterial; Antibiotic	
Pathway:	Anti-infection	Amphomycin
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

Product Data Sheet

Description	Amphomycin is a lipopeptide antibiotic that inhibits peptidoglycan synthesis and blocks cell wall development. Amphomycin exhibits potent antibacterial activities against methicillin-resistant S. aureus (MRSA), vancomycin-resistant enterococci (VRE), penicillin-gentamicin-erythromycin-resistant S. pneumonia, and linezolid-quinupristin-dalfopristin- resistant enterococci ^{[1][2][3]} .		
IC_{50} & Target	Lipopeptide		
In Vitro	Amphomycin (40 μ g/mL; 60 min) leads S. aureus to exhibit cell wall thinning and Park's nucleotide accumulation ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay ^[1] Cell Line: S. aureus whole cells Concentration: 40 μ g/mL Incubation Time: 60 min Result: Resulted in the thinning of cell wall and the accumulation of Park's nucleotide.		
In Vivo	Amphomycin (50 mg/kg; p.o.; single) shows low oral availability and (5-10mg/kg; i.v.; single) exhibits a long half-life (5.2-8.0 h in mice and 4.6-7.1 h in rats) ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

REFERENCES

[1]. Singh M, et al. Solid-state NMR characterization of amphomycin effects on peptidoglycan and wall teichoic acid biosyntheses in Staphylococcus aureus. Sci Rep. 2016 Aug 19;6:31757.

[2]. Pasetka CJ, et al. Novel antimicrobial lipopeptides with long in vivo half-lives. Int J Antimicrob Agents. 2010 Feb;35(2):182-5.

[3]. Tanaka H, et al. Amphomycin inhibits phospho-N-acetylmuramyl-pentapeptide translocase in peptidoglycan synthesis of Bacillus. Biochem Biophys Res Commun. 1979 Feb 14;86(3):902-8.

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

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