Nalidixic acid

Cat. No.:	HY-B0398		
CAS No.:	389-08-2		
Molecular Formula:	C ₁₂ H ₁₂ N ₂ O ₃	1	
Molecular Weight:	232.24		
Target:	Bacterial; Antibiotic; Topoisomerase		
Pathway:	Anti-infection; Cell Cycle/DNA Damage		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month

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SOLVENT & SOLUBILITY

		Solvent Mass Concentration	1 mg	5 mg	10 mg	
Preparing Stock Solutions Please refer to the so	Preparing Stock Solutions	1 mM	4.3059 mL	21.5295 mL	43.0589 mL	
	5 mM	0.8612 mL	4.3059 mL	8.6118 mL		
		10 mM	0.4306 mL	2.1529 mL	4.3059 mL	
	Please refer to the solubility information to select the appropriate solvent.					

BIOLOGICAL ACTIVITY				
Description	Nalidixic acid, a quinolone antibiotic, is effective against both gram-positive and gram-negative bacteria. Nalidixic acid acts in a bacteriostatic manner in lower concentrations and is bactericidal in higher concentrations. Nalidixic acid inhibits a subunit of DNA gyrase and topoisomerase IV and reversibly blocks DNA replication in susceptible bacteria ^[1] .			
IC ₅₀ & Target	Quinolone Topoisomerase			
In Vitro	Nalidixic acid is against a variety of microorganisms, it is against with Escherichia coli, Pasteurella spp., Klebsiella pneuiiioniae, Aerobacter aeroyenes, Proteus spp., Salmonella spp., Shigella spp. and Brucella spp. with MIC values of 5.0-12.5 μg/ml, 0.5-2.5 μg/ml, 0.8-25.0 μg/ml, 1.0-25.0 μg/ml, 1.25-30.0 μg/ml, 8-3.2 μg/ml, and 7.5-10.0 μg/ml, respectively ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

Product Data Sheet

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In Vivo

The in vivo activity of Nalidixic acid is most pronounced against Gram-negative bacteria, while Gram-positive organisms are generally more resistant. Maximal activity is observed against systemic infections caused by E. coli, A. aerobacter, Proteus mirabilis, Shigella fkxneri, the ED₅₀ values are 25 mg/kg, 60 mg/kg, 50 mg/kg, and 62 mg/kg, respectively^[1]. The acute toxicity (LD₅₀) of Nalidixic acid in mice following oral and parenteral administration is: oral, 3300 mg/kg; intravenous, 176 mg/kg, and subcutaneous, 500 mg/kg^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

• J Biol Chem. 2021 Dec 29;101554.

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REFERENCES

[1]. antibiotic, DNA gyrase, susceptible bacteria, Escherichia coli, Klebsiella pneuiiioniae, Aerobacter aeroyenes, Proteus spp, Salmonella spp.

[2]. Anna Fàbrega, et al. Mechanism of Action of and Resistance to Quinolones. Microb Biotechnol

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

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