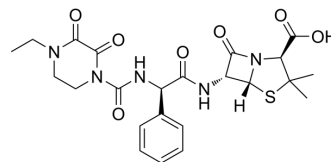


Piperacillin

Cat. No.:	HY-B1923		
CAS No.:	61477-96-1		
Molecular Formula:	C ₂₃ H ₂₇ N ₅ O ₇ S		
Molecular Weight:	517.55		
Target:	Antibiotic; Bacterial; Beta-lactamase; Penicillin-binding protein (PBP)		
Pathway:	Anti-infection		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (193.22 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
	Preparing Stock Solutions		10 mg	
	1 mM	1.9322 mL	9.6609 mL	19.3218 mL
	5 mM	0.3864 mL	1.9322 mL	3.8644 mL
	10 mM	0.1932 mL	0.9661 mL	1.9322 mL
Please refer to the solubility information to select the appropriate solvent.				
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (4.83 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (4.83 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (4.83 mM); Clear solution 			

BIOLOGICAL ACTIVITY

Description	Piperacillin is a semisynthetic broad-spectrum β-lactam antibiotic which exhibits potent bactericidal activity against Gram-negative bacteria as well as select Gram-positive strains through penicillin-binding proteins. Piperacillin is most commonly used in combination with the β-lactamase inhibitor Tazobactam ^{[1][2][3]} .
IC₅₀ & Target	β-lactam
In Vitro	Piperacillin (12.5 μg/mL, 24h) inhibits 92% of isolates of <i>Pseudomonas aeruginosa</i> , 82% of <i>Serratia marcescens</i> , 73% of

Escherichia coli, 61% of Klebsiella spp, and 42% of Enterobacter spp, most Proteus spp. were extremely susceptible. Piperacillin fails to inhibit the growth of gram-negative bacilli when large inocula were used (minimum inhibitory concentration > 25 µg/ml) [2].

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

In Vivo

Piperacillin (100 mg/kg, i.v., 4 times a day, 5 d) in combination with Tazobactam (HY-B1418)(12.5 mg/kg, i.v., 4 times a day, 5 d) prolongs survival in mice with low inoculum of K. pneumoniae[3].

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

Animal Model:	BALB/c low inoculum model of K. pneumoniae, KEN-11 strain [3]
Dosage:	100 mg/kg
Administration:	Intravenous injection (i.v.), 4 times a day, 5 d, in combination with Tazobactam (12.5 mg/kg, i.v., 4 times a day, 5 d)
Result:	Enabled all mice survived whereas all control mice died by 5 d, decreased the number of bacteria in lungs compared with control group. Observed no bacteria in the blood of most mice (except for two mice at the early phase) while bacteria were observed in the blood of control group. Observed few inflammatory cells in the alveoli whereas an influx of numerous inflammatory cells were observed in the control group.

CUSTOMER VALIDATION

- Nat Commun. 2022 Mar 2;13(1):1116.
- Antimicrob Agents Chemother. 2023 May 18;e0160322.
- Microbiol Spectr. 2023 Apr 24;e0069223.
- Microbiol Spectr. 2022 Dec 8;e0303822.
- Biomed Res Int. 2018 Jul 2;2018:3579832.

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REFERENCES

[1]. Fu KP, et al. Piperacillin, a new penicillin active against many bacteria resistant to other penicillins. Antimicrob Agents Chemother. 1978;13(3):358-67.

[2]. Bodey GP, et al. Piperacillin: In Vitro Evaluation. Antimicrob Agents Chemother. 1978;14(1):78-87.

[3]. Harada Y, et al. In vitro and in vivo activities of piperacillin-tazobactam and meropenem at different inoculum sizes of ESBL-producing Klebsiella pneumoniae. Clin Microbiol Infect. 2014;20(11):O831-9.

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

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