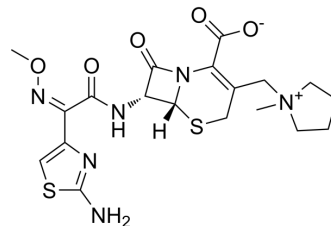


Cefepime

Cat. No.:	HY-B0692		
CAS No.:	88040-23-7		
Molecular Formula:	C ₁₉ H ₂₄ N ₆ O ₅ S ₂		
Molecular Weight:	480.56		
Target:	Antibiotic; Bacterial; 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 : 125 mg/mL (260.11 mM; Need ultrasonic)
 H₂O : 100 mg/mL (208.09 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
	1 mM	2.0809 mL	10.4045 mL	20.8091 mL
	5 mM	0.4162 mL	2.0809 mL	4.1618 mL
	10 mM	0.2081 mL	1.0405 mL	2.0809 mL

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

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
 Solubility: ≥ 2.08 mg/mL (4.33 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
 Solubility: ≥ 2.08 mg/mL (4.33 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
 Solubility: ≥ 2.08 mg/mL (4.33 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Cefepime (BMY-28142) is a broad-spectrum and cross the blood-brain barrier cephalosporin. Cefepime shows antibacterial effects against both Gram-positive and Gram-negative aerobic bacteria. Cefepime induces neurotoxicity^{[1][2][3][4]}.

IC₅₀ & Target

β-lactam

In Vitro

Cefepime chloride exerts its antibacterial effects by binding to penicillin-binding proteins^[2].

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

In Vivo

Cefepime (80 mg/kg; i.p.) significantly increases the half-life and mice survived^[4].

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

Animal Model:	Male CD-1 mice ^[4]
Dosage:	80 mg/kg
Administration:	I.p.
Result:	Significantly prolonged the half-life of cefepime and all mice survived at 18-22 mg/kg cisplatin, and when pretreatment with 26 mg/kg cisplatin significantly decreased survival with the half-life of cefepime was not significantly longer than of 18 mg/kg cisplatin.

CUSTOMER VALIDATION

- Emerg Microbes Infect. 2024 Dec;13(1):2321981.
- Pharmaceutics. 2023 Nov 30, 15(12), 2705.
- Vet Microbiol. 2024 May, 292, 110046.
- J Antibiot (Tokyo). 2023 Feb 1.
- Animal Diseases. 02 November 2021.

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REFERENCES

- [1]. Yahav D, et al. Efficacy and safety of cefepime: a systematic review and meta-analysis. Lancet Infect Dis. 2007 May;7(5):338-48.
- [2]. Barradell LB, Bryson HM. Cefepime. A review of its antibacterial activity, pharmacokinetic properties and therapeutic use. Drugs. 1994;47(3):471-505.
- [3]. Payne LE, et al. Cefepime-induced neurotoxicity: a systematic review. Crit Care. 2017 Nov 14;21(1):276. doi: 10.1186/s13054-017-1856-1.
- [4]. Máthé A, et al. The effect of different doses of cisplatin on the pharmacokinetic parameters of cefepime in mice. Lab Anim. 2006 Jul;40(3):296-300.

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

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