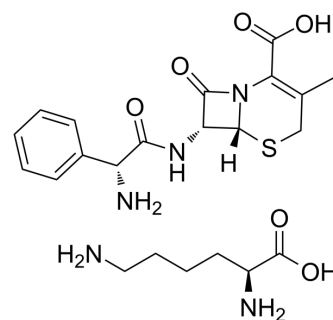


Cephalexin (lysine)

Cat. No.:	HY-B0200D
CAS No.:	53950-14-4
Molecular Formula:	C ₂₂ H ₃₁ N ₅ O ₆ S
Molecular Weight:	493.58
Target:	Bacterial; Antibiotic
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Cephalexin (Cefalexin) lysine is a potent, orally active new semisynthetic cephalosporin antibiotic with a broad antibacterial spectrum. Cephalexin lysine has antibacterial activity against a wide variety of gram-positive and gram-negative bacteria. Cephalexin lysine targets penicillin-binding proteins (PBPs) to inhibit bacterial cell wall assembly. Cephalexin lysine is used for the research of pneumonia, strep throat, and bacterial endocarditis, et al ^{[1][2]} .								
IC₅₀ & Target	Bacteria ^[1]								
In Vitro	<p>Cephalexin lysine (10 µg/mL) disrupts polymer peptidoglycan (PG) biogenesis by inactivating enzymes called penicillin-binding proteins (PBPs)^[1].</p> <p>Cephalexin lysine inhibits a broad spectrum of gram-positive and gram-negative organisms with MIC values of 2, 2, 2, 4, 4, 4 and 5.7 µg/mL for <i>Bacillus anthracis</i>, <i>Edwardsiella taFda</i>, <i>Vibrio cholera</i>, <i>Pasteurella multocida</i>, <i>Edwardsiella tarda</i>, <i>Alcaligenes sp</i> and <i>Proteus rettgeri</i>, respectively^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>								
In Vivo	<p>Cephalexin lysine (0-50 mg/kg; p.o.; for 3.5 hours) has antibacterial activity in male Swiss-Webster mice with infected bacterial^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 30%;">Animal Model:</td> <td>Male Swiss-Webster mice with infected bacterial^[2]</td> </tr> <tr> <td>Dosage:</td> <td>0-50 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Oral administration; for 3.5 hours</td> </tr> <tr> <td>Result:</td> <td>Had antibacterial activity against <i>Streptococcus pyogenes</i>, <i>Streptococcus pneumoniae</i>, <i>Staphylococcus aureus</i> and several gram-negative species mice.</td> </tr> </table>	Animal Model:	Male Swiss-Webster mice with infected bacterial ^[2]	Dosage:	0-50 mg/kg	Administration:	Oral administration; for 3.5 hours	Result:	Had antibacterial activity against <i>Streptococcus pyogenes</i> , <i>Streptococcus pneumoniae</i> , <i>Staphylococcus aureus</i> and several gram-negative species mice.
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CUSTOMER VALIDATION

- Theranostics. 2022 Jan 1;12(3):1187-1203.

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- J Med Chem. 2021 Sep 21.
 - Chemosphere. 2021, 131417.
 - Chemosphere. 2019 Jun;225:378-387.
 - Infect Immun. 2018 May 22;86(6). pii: e00090-18.

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REFERENCES

- [1]. Cho H, et, al. Beta-lactam antibiotics induce a lethal malfunctioning of the bacterial cell wall synthesis machinery. Cell. 2014 Dec 4;159(6):1300-11.
- [2]. Buck RE, et, al. Cefadroxil, a new broad-spectrum cephalosporin. Antimicrob Agents Chemother. 1977 Feb;11(2):324-30.
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Caution: Product has not been fully validated for medical applications. For research use only.

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