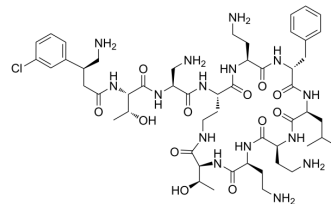


Upleganan

Cat. No.:	HY-128780
CAS No.:	2407717-17-1
Molecular Formula:	C ₅₂ H ₈₂ ClN ₁₅ O ₁₂
Molecular Weight:	1144.75
Target:	Bacterial
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Upleganan (SPR206), a polymyxin analogue, and shows antibiotic activity against multidrug resistant Gram-negative pathogen. The MIC values of Upleganan against <i>Pseudomonas aeruginosa</i> Pa14 and <i>Acinetobacter baumannii</i> NCTC13301 are both 0.125 mg/L ^{[1][2]} .	
IC₅₀ & Target	MIC: 0.125 mg/L (<i>Pseudomonas aeruginosa</i> Pa14) and 0.125 mg/L (<i>Acinetobacter baumannii</i> NCTC13301) ^{[1][2]}	
In Vitro	Upleganan (SPR206) exhibits strong anti-microbial properties against Gram-negative bacteria. The MIC values of SPR206 against <i>E. coli</i> IHMA558090, <i>E. coli</i> ATCC 25922, <i>K. pneumoniae</i> ATCC 13882, <i>P. aeruginosa</i> ATCC 27853, <i>A. baumannii</i> NCTC13424 and <i>A. baumannii</i> ATCC 19003 are 8 mg/L, 0.125 mg/L, 0.125 mg/L, 0.25 mg/L, 0.06 mg/L and 0.125 mg/L, respectively, together with lower cytotoxicity ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
In Vivo	Upleganan (SPR206) (0.125-30 mg/kg; intravenous injection or subcutaneous injection; every 8 hours or every 4 hours; for 16 hours or 24 hours; neutropenic mice) treatment reduces the burden of Pa14 and NCTC13301 in lung tissue and in the thigh model ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	Neutropenic mice infected Pa14 or NCTC13301 ^[2]
	Dosage:	3 mg/kg, 10 mg/kg, 20 mg/kg, 30 mg/kg (intravenous injection); 0.125 mg/kg, 0.5 mg/kg, 1 mg/kg, 4 mg/kg (subcutaneous injection)
	Administration:	Intravenous injection or subcutaneous injection; every 8 hours or every 4 hours; for 16 hours or 24 hours
	Result:	In lung tissue, reduced the burden of Pa14 and NCTC13301 by 1.5 and 3.6 log ₁₀ CFU/mL. In the thigh model, reduced the burden of Ab13301 by 3.4 and 4.3 log ₁₀ CFU/g.

REFERENCES

[1]. Brown P, et al. Design of Next Generation Polymyxins with Lower Toxicity: The Discovery of SPR206. *ACS Infect Dis.* 2019 Oct 11;5(10):1645-1656.

[2]. L. Grosser, et al. In Vivo Efficacy of SPR206 in Murine Lung and Thigh Infection Models Caused by Multidrug Resistant Pathogens *Pseudomonas aeruginosa* and *Acinetobacter baumannii*. Poster-139 ASM ESCMID 2018 Lisbon, Portugal.

[3]. Noushin Akhoundsadegh, et al. Outer Membrane Interaction Kinetics of New Polymyxin B Analogs in Gram-Negative Bacilli. *Antimicrob Agents Chemother.* 2019 Sep 23;63(10):e00935-19.

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

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