Anti-MRSA agent 7

®

Cat. No.:	HY-149271	
Molecular Formula:	$C_{22}H_{20}BrF_2N_3O_4$	F
Molecular Weight:	508.31	Br
Target:	Bacterial; DNA/RNA Synthesis; Topoisomerase	
Pathway:	Anti-infection; Cell Cycle/DNA Damage	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	



BIOLOGICAL ACTIV	1 Y								
Description	Anti-MRSA agent DNA gyrase, S. au	: 7 (Compound ureus topo IV a	ا 12) is a potent a and E. coli topo ۱	antibacterial age V with IC ₅₀ s of 0.	ent. Anti-MRSA aş .185, 0.365, 0.34	gent 7 inhibits \$ L and 0.059 μM,	5. aureus DNA g , respectively ^[1] .	yrase, E. coli	
IC ₅₀ & Target	TOPO IV 0.059 μΜ (IC ₅₀ , E. coli)		TOPO IV 0.341 μM (IC ₅₀ , S. aureus)		DNA gyrase 0.185 μΜ (IC ₅₀ , S. aureus)		DNA gyrase 0.365 μΜ (IC ₅₀ , E. coli)		
In Vitro	Anti-MRSA agent 7 (Compound 12) shows a dose-dependent killing efficacy achieving bactericidal effect against plankto methicillin-resistant S. aureus (ATCC 43300) at 8 × MIC after 8 h of treatment, after which re-growth occurs ^[1] . Antimicrobial activity of Anti-MRSA agent 7 (Compound 12) against a panel of Gram-positive and Gram-negative bacteri pathogens ^[1]						nst planktonic ive bacterial		
	Strain	S. aureus (ATCC 29213)	MRSAQA-12.1	<i>E. coli</i> N43 (CGSC# 5583)	MRSA QA-11.7	MRSA QA-11.2	<i>E. faecalis</i> DRK 057	<i>E. coli</i> (ATCC 25922)	
	MIC (µM)	0.03	0.03	0.06	0.06	0.124	4.07	252	
	Strain	E. coli D22	A. baumannii	E. coli ESBL QA:11.3	K. pneumoniae	S. alachua RDK 030c	P. aeruginosa RDK 184		
	MIC (µM)	252	252	>252	>252	>252	>252		
	MCE has not inde	ependently co	nfirmed the accu	aracy of these m	ethods. They are	e for reference of	only.		
In Vivo	Anti-MRSA agent neutropenic mor MCE has not inde	7 (Compound use thigh infec ependently co	l 12; 20 and 40 m tion model ^[1] . nfirmed the accu	g/kg; i.p.; QID fo aracy of these m	or 1 day) demons ethods. They are	trates high in v	ivo efficacy in M only.	IRSA	
	Animal Model:		CD-1 female mice, MRSA neutropenic mouse thigh infection $model^{[1]}$						
	Dosage:		20 and 40 mg/kg						

Administration:	IP/QID (four times a day) at 2, 8, 14, and 20 h post infection
Result:	Demonstrated inhibition of bacterial growth in a dose dependent manner.

REFERENCES

[1]. Kokot M, et al. Amide containing NBTI antibacterials with reduced hERG inhibition, retained antimicrobial activity against gram-positive bacteria and in vivo efficacy. Eur J Med Chem. 2023 Mar 15;250:115160.

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

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