**Proteins** 

# Inhibitors



# **Anti-MRSA agent 5**

Cat. No.: HY-149013 CAS No.: 2490154-44-2 Molecular Formula:  $C_{25}H_{22}N_4O_4$ Molecular Weight: 442.47

Target: Bacterial Pathway: Anti-infection

Please store the product under the recommended conditions in the Certificate of Storage:

Analysis.

**Product** Data Sheet

# **BIOLOGICAL ACTIVITY**

Description Anti-MRSA agent 5 (B14) is a potent

agent with  $MIC_{50}$  values of 0.38  $\mu$ g/mL and has low hERG activity with an  $IC_{50}$  values of >40  $\mu$ M. Anti-MRSA agent 5 (B14) also has low cytotoxicity to mammal cells and unlikely to acquire bacterial resistance [1].

IC<sub>50</sub> & Target MIC: 0.38 μg/mL (MRSA); IC50: >40 Mm (hERG)

In Vitro

Anti-MRSA agent 5 (B14) is highly selective toxic in bacterial cells. Anti-MRSA agent has marginal inhibitory activities with an IC<sub>50</sub> values of exceeding 40 μM. Anti-MRSA agent 5 (B14) has strong antibacterial activity for all five clinical isolates of MRSA (NRS-1, NRS-70, NRS-100, NRS-108, NRS-271) with MIC values between < 0.17 and 0.69 μg/mL.Anti-MRSA agent 5 (B14) inhibits all three clinical isolates of fluoroquinolone-resistant MRSA (19-25, 19-26, 19-27) and exhibits good potency against  $VISA^{[1]}$ .

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

Cell Cytotoxicity Assay

Cell Line:	A549, MDA-MB-231, PC-3, ABAE.
Concentration:	0–400 μΜ
Incubation Time:	48h
Result:	Showed marginal inhibitory activities with anIC $_{50}$ exceeding 40 $\mu\text{M}$ and displayed highly selective toxicity toward bacterial cells.

## **REFERENCES**

[1]. Runzhe Songa, et al, Design and synthesis of novel desfluoroquinolone-aminopyrimidine hybrids as potent anti-MRSA agents with low hERG activity, Bioorg Chem. 2020 Oct:103:104176.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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