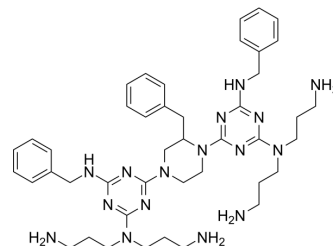


Antimicrobial agent-9

Cat. No.:	HY-151403
CAS No.:	2978694-25-4
Molecular Formula:	C ₄₃ H ₆₂ N ₁₆
Molecular Weight:	803.06
Target:	Bacterial
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Antimicrobial agent-9 (Compound 16) is an antimicrobial agent with an MIC range of 4-8 µg/mL against gram-positive and gram-negative bacteria. Antimicrobial agent-9 also shows anti-inflammatory activity ^[1] .																
In Vitro	<p>Antimicrobial agent-9 (Compound 16) (0-256 µg/mL; 18-24 h) shows antibacterial activity with geometric mean (GM) values of the MICs of 4.5 µg/mL^[1].</p> <p>Antimicrobial agent-9 shows minimum hemolytic concentration (MHC) of >256 µg/mL, the therapeutic index is 113.8^[1].</p> <p>Antimicrobial agent-9 (5 or 20 µg/mL; 18 h) effectively inhibits the release and expression of NO and TNF-α from LPS-stimulated RAW 264.7 cells^[1].</p> <p>Antimicrobial agent-9 is resistant to various physiological salts, human serum, and proteases^[1].</p> <p>Antimicrobial agent-9 exhibits synergistic antimicrobial activity in combination with three conventional antibiotics (Chloramphenicol (HY-B0239), Ciprofloxacin (HY-B0356), and oxacillin) against MDRPA and MRSA, is promising adjuvants in combination with clinically used antibiotics^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td><td>E. coli [KCTC 1682], P. aeruginosa [KCTC 1637], S. epidermidis [KCTC 1917] and S. aureus [KCTC1621]</td></tr> <tr> <td>Concentration:</td><td>0-256 µg/mL</td></tr> <tr> <td>Incubation Time:</td><td>18-24 h</td></tr> <tr> <td>Result:</td><td>Inhibited bacterial growth with MICs of 4, 8, 4 and 2 µg/mL against E. coli [KCTC 1682], P. aeruginosa [KCTC 1637], S. epidermidis [KCTC 1917] and S. aureus [KCTC1621], respectively.</td></tr> </table> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td><td>LPS-stimulated RAW 264.7 macrophages</td></tr> <tr> <td>Concentration:</td><td>5 µg/mL (for NO/iNOS) and 20 µg/mL (for TNF-α)</td></tr> <tr> <td>Incubation Time:</td><td>18 h</td></tr> <tr> <td>Result:</td><td>Effectively inhibited the production and expression of NO and TNF-α from LPS-stimulated</td></tr> </table>	Cell Line:	E. coli [KCTC 1682], P. aeruginosa [KCTC 1637], S. epidermidis [KCTC 1917] and S. aureus [KCTC1621]	Concentration:	0-256 µg/mL	Incubation Time:	18-24 h	Result:	Inhibited bacterial growth with MICs of 4, 8, 4 and 2 µg/mL against E. coli [KCTC 1682], P. aeruginosa [KCTC 1637], S. epidermidis [KCTC 1917] and S. aureus [KCTC1621], respectively.	Cell Line:	LPS-stimulated RAW 264.7 macrophages	Concentration:	5 µg/mL (for NO/iNOS) and 20 µg/mL (for TNF-α)	Incubation Time:	18 h	Result:	Effectively inhibited the production and expression of NO and TNF-α from LPS-stimulated
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RAW 264.7 cells.

REFERENCES

[1]. Dinesh Kumar S, et al. Cationic, amphipathic small molecules based on a triazine-piperazine-triazine scaffold as a new class of antimicrobial agents. Eur J Med Chem. 2022 Sep 8;243:114747.

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

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