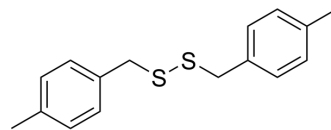


IITR08367

Cat. No.:	HY-163473
Molecular Formula:	C ₁₆ H ₁₈ S ₂
Molecular Weight:	274.44
Target:	Bacterial
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	IITR08367 is a potent effector pump <i>Acinetobacter baumannii</i> Fosfomycin Efflux pump (AbaF) inhibitor for enhancing the antimicrobial activity of Fosfomycin (HY-B1075A) against <i>Acinetobacter baumannii</i> . IITR08367 acts by interfering with Fosfomycin/H ⁺ reverse transporter activity. ^[1]								
In Vitro	<p>IITR08367 (0-50 μM) concentration-dependently enhances the activity of Fosfomycin against AbaF-expressing strain of <i>Escherichia coli</i>. A concentration of 50 μM can increase the postantibiotic effect of Fosfomycin by 30 min^[1].</p> <p>IITR08367 (25 μM; 16 min) disrupts the H⁺ gradient across the membrane, ultimately inhibiting the H⁺ gradient-driven efflux pump without causing membrane damage^[1].</p> <p>IITR08367 (100 μM; 12 h) and Fosfomycin (64 mg/L) in combination can inhibit <i>A. baumannii</i> RPTC-15 growth^[1].</p> <p>IITR08367 (100 μM) eliminates the Biofilm Forming Ability of <i>A. baumannii</i> with and without the addition of Fosfomycin^[1].</p> <p>IITR08367 (12.5-200 μM) is not toxic to erythrocytes^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>RBCs</td> </tr> <tr> <td>Concentration:</td> <td>12.5; 25; 50; 100; 200 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>6 h</td> </tr> <tr> <td>Result:</td> <td>Showed less than 40% cytotoxicity till 200 μM</td> </tr> </table>	Cell Line:	RBCs	Concentration:	12.5; 25; 50; 100; 200 μM	Incubation Time:	6 h	Result:	Showed less than 40% cytotoxicity till 200 μM
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In Vivo	<p>IITR08367 (i.p.; 30 mg/kg; every 12 h for 54 h) co-administered with Fosfomycin to Urinary tract infections mice shows significant improvement in the damage caused by <i>A. baumannii</i> RPTC-15, with tissue ultrastructure similar to that of uninfected mice^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Urinary tract infections mice^[1]</td> </tr> <tr> <td>Dosage:</td> <td>30 mg/kg; every 12 h for 54 h</td> </tr> <tr> <td>Administration:</td> <td>i.p.</td> </tr> </table>	Animal Model:	Urinary tract infections mice ^[1]	Dosage:	30 mg/kg; every 12 h for 54 h	Administration:	i.p.		
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Result:	Reduced the load of <i>A. baumannii</i> RPTC-15 in the kidney and bladder by approximately $3\log_{10}$.
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REFERENCES

[1]. Saini M, et al. Small Molecule IITR08367 Potentiates Antibacterial Efficacy of Fosfomycin against *Acinetobacter baumannii* by Efflux Pump Inhibition. *ACS Infect Dis.* 2024 Apr 1.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA