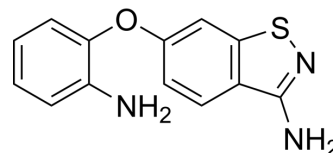


M62812 free base

Cat. No.:	HY-103639
CAS No.:	613262-61-6
Molecular Formula:	C ₁₃ H ₁₁ N ₃ OS
Molecular Weight:	257.31
Target:	Toll-like Receptor (TLR)
Pathway:	Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	M62812 (free base) is a toll-like receptor 4 (TLR4) signal transduction inhibitor. M62812 can suppress endothelial cell and leukocyte activation and prevents lethal septic shock in mice. M62812 can be used for the research of sepsis ^[1] .																
In Vitro	M62812 (0-10 µg/ml) suppresses LPS-induced upregulation of inflammatory cytokines, adhesion molecules and procoagulant activity in human vascular endothelial cells and peripheral mononuclear cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.																
In Vivo	<p>M62812 (i.v.; 10-20 mg/kg; single) protects mice from lethality and reduced inflammatory and coagulatory parameters in a murine d-galactosamine-sensitized endotoxin shock model^[1].</p> <p>M62812 (20 mg/kg/day) prevents mice from lethality in a murine cecal ligation and puncture model^[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>D-galactosamine-sensitized endotoxin shock mouse model^[1]</td> </tr> <tr> <td>Dosage:</td> <td>10-20 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Single intravenous administration</td> </tr> <tr> <td>Result:</td> <td>Suppressed LPS-induced leukocyte and endothelial cell activation in vivo. Prolonged survival in a d-galactosamine-sensitized endotoxin shock mouse model.</td> </tr> </table> <table border="1"> <tr> <td>Animal Model:</td> <td>A cecal ligation and puncture mouse model^[1]</td> </tr> <tr> <td>Dosage:</td> <td>20 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intravenous administration; Everyday</td> </tr> <tr> <td>Result:</td> <td>Reduced mortality in a murine cecal ligation and puncture model.</td> </tr> </table>	Animal Model:	D-galactosamine-sensitized endotoxin shock mouse model ^[1]	Dosage:	10-20 mg/kg	Administration:	Single intravenous administration	Result:	Suppressed LPS-induced leukocyte and endothelial cell activation in vivo. Prolonged survival in a d-galactosamine-sensitized endotoxin shock mouse model.	Animal Model:	A cecal ligation and puncture mouse model ^[1]	Dosage:	20 mg/kg	Administration:	Intravenous administration; Everyday	Result:	Reduced mortality in a murine cecal ligation and puncture model.
Animal Model:	D-galactosamine-sensitized endotoxin shock mouse model ^[1]																
Dosage:	10-20 mg/kg																
Administration:	Single intravenous administration																
Result:	Suppressed LPS-induced leukocyte and endothelial cell activation in vivo. Prolonged survival in a d-galactosamine-sensitized endotoxin shock mouse model.																
Animal Model:	A cecal ligation and puncture mouse model ^[1]																
Dosage:	20 mg/kg																
Administration:	Intravenous administration; Everyday																
Result:	Reduced mortality in a murine cecal ligation and puncture model.																

REFERENCES

[1]. Nakamura M, et al. Toll-like receptor 4 signal transduction inhibitor, M62812, suppresses endothelial cell and leukocyte activation and prevents lethal septic shock in mice. *Eur J Pharmacol.* 2007;569(3):237-243.

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