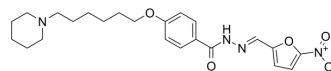


TGFβ1-IN-3

Cat. No.:	HY-151955
CAS No.:	2883813-58-7
Molecular Formula:	C ₂₃ H ₃₀ N ₄ O ₅
Molecular Weight:	442.51
Target:	TGF-β Receptor
Pathway:	TGF-beta/Smad
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	TGFβ1-IN-3 is a diarylacylhydrazones derivative that effectively suppresses the activation and proliferation of fibroblasts. TGFβ1-IN-3 can be used for idiopathic pulmonary fibrosis (IPF) research ^[1] .								
In Vitro	<p>TGFβ1-IN-3 (compound 44) shows inhibitory effect against NIH-3T3 cells with an IC₅₀ of 0.57 μM^[1].</p> <p>TGFβ1-IN-3 (compound 44; 10 μM; for 24 h) inhibits TGF-β1-induced abnormal activation of NIH-3T3 and A549 cells, as well as migration and epithelial-mesenchymal transition (EMT) of A549 cells^[1].</p> <p>TGFβ1-IN-3 (compound 44) could bind to STAT3, and able to interact with Ile659, and the hydrophilic group piperidine formed intermolecular forces with Ser636, Arg609, and Pro639. The nitrofuran of TGFβ1-IN-3 could interact with Lys658^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>NIH-3T3 or A549 cells</td> </tr> <tr> <td>Concentration:</td> <td>10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Were able to inhibit TGF-β1-induced activation of fibroblasts in vitro.</td> </tr> </table>	Cell Line:	NIH-3T3 or A549 cells	Concentration:	10 μM	Incubation Time:	24 h	Result:	Were able to inhibit TGF-β1-induced activation of fibroblasts in vitro.
Cell Line:	NIH-3T3 or A549 cells								
Concentration:	10 μM								
Incubation Time:	24 h								
Result:	Were able to inhibit TGF-β1-induced activation of fibroblasts in vitro.								
In Vivo	<p>TGFβ1-IN-3 (compound 44; 30-60 mg/kg; p.o; once daily; for 22 days) improves mouse lung function and slows the progression of IPF. TGFβ1-IN-3 could reverse the pulmonary fibrosis in treatment model^[1].</p> <p>Pharmacokinetic parameters of TGFβ1-IN-3 (compound 44) in rats^[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>C57BL/6 mice (6-10 weeks) injected with Bleomycin for establishing pulmonary fibrosis model^[1].</td> </tr> <tr> <td>Dosage:</td> <td>30 mg/kg, 60 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Orally administration; once daily; for 22 days</td> </tr> <tr> <td>Result:</td> <td>Could reverse the pulmonary fibrosis in treatment model.</td> </tr> </table>	Animal Model:	C57BL/6 mice (6-10 weeks) injected with Bleomycin for establishing pulmonary fibrosis model ^[1] .	Dosage:	30 mg/kg, 60 mg/kg	Administration:	Orally administration; once daily; for 22 days	Result:	Could reverse the pulmonary fibrosis in treatment model.
Animal Model:	C57BL/6 mice (6-10 weeks) injected with Bleomycin for establishing pulmonary fibrosis model ^[1] .								
Dosage:	30 mg/kg, 60 mg/kg								
Administration:	Orally administration; once daily; for 22 days								
Result:	Could reverse the pulmonary fibrosis in treatment model.								

REFERENCES

[1]. Xingping Su, et al. Design, synthesis and biological evaluation of novel diarylacylhydrazones derivatives for the efficient treatment of idiopathic pulmonary fibrosis. Eur J Med Chem. 2023 Jan 5;245(Pt 2):114918.

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