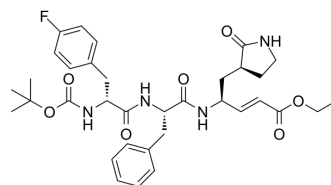


SARS-CoV-2 Mpro-IN-5

Cat. No.:	HY-151901
CAS No.:	3023276-79-8
Molecular Formula:	C ₃₄ H ₄₃ FN ₄ O ₇
Molecular Weight:	638.73
Target:	SARS-CoV
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	SARS-CoV-2 Mpro-IN-5 is a dual Inhibitor of Main Protease (M ^{Pro}) and Cathepsin L (CatL), with IC ₅₀ s of 1800 nM and 145 nM respectively. SARS-CoV-2 Mpro-IN-5 has antiviral activity against SARS-CoV2. SARS-CoV-2 Mpro-IN-5 blocks SARS-CoV2 replication in hACE2 expressing A549 cells with IC ₅₀ value of 14.7 nM ^[1] .								
IC₅₀ & Target	MPro/CatL ^[1]								
In Vitro	<p>SARS-CoV-2 Mpro-IN-5 (SM142) blocks SARS-CoV2 replication in A549-hACE2 cells with IC₅₀ value of 14.7 nM^[1].</p> <p>SARS-CoV-2 Mpro-IN-5 (0-50 μM, 24 h) cause cytotoxicity in A549-hACE2 cells^[1].</p> <p>SARS-CoV-2 Mpro-IN-5 inhibits OC-43 virus mRNA expression A549 cells^[1].</p> <p>SARS-CoV-2 Mpro-IN-5 inhibits SARS-CoV2 infection by inhibiting both MPro and CatL^[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>A549-hACE2 cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 0.1, 0.2, 2, 20, 50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Dose-dependently inhibited cell viability, reduce by 15% at 50 μM.</td> </tr> </table>	Cell Line:	A549-hACE2 cells	Concentration:	0, 0.1, 0.2, 2, 20, 50 μM	Incubation Time:	24 h	Result:	Dose-dependently inhibited cell viability, reduce by 15% at 50 μM.
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In Vivo	<p>SARS-CoV-2 Mpro-IN-5 (SM142) (10 mg/kg for i.n. or 25 mg/kg for i.p.) protects K18-hACE2 mice from SARS-CoV2-induced weight loss and lethality^[1].</p> <p>SARS-CoV-2 Mpro-IN-5 (3 mg/kg, i.v.) shows a half-life of 2.1 h and high clearance of 18490 mL/min/kg in in male C57Bl/6 mice^[1].</p> <p>SARS-CoV-2 Mpro-IN-5 (10 mg/kg, p.o.) shows oral bioavailability of 37.5%^[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>K18-hACE2 transgenic mice^[1]</td> </tr> <tr> <td>Dosage:</td> <td>10 mg/kg (i.n.) or 25 mg/kg (i.p.)</td> </tr> <tr> <td>Administration:</td> <td>Intranasal inhalation (i.n.), once daily for 3 days, prior to the infection; or Intraperitoneal</td> </tr> </table>	Animal Model:	K18-hACE2 transgenic mice ^[1]	Dosage:	10 mg/kg (i.n.) or 25 mg/kg (i.p.)	Administration:	Intranasal inhalation (i.n.), once daily for 3 days, prior to the infection; or Intraperitoneal		
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Dosage:	10 mg/kg (i.n.) or 25 mg/kg (i.p.)								
Administration:	Intranasal inhalation (i.n.), once daily for 3 days, prior to the infection; or Intraperitoneal								

	injection (i.p.), twice daily for 5 days, postinfection administration.
Result:	Prevented weight loss and prolonged survival.

REFERENCES

[1]. Mondal S, et al. Dual Inhibitors of Main Protease (MPro) and Cathepsin L as Potent Antivirals against SARS-CoV2. J Am Chem Soc. 2022 Nov 23;144(46):21035-21045.

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

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