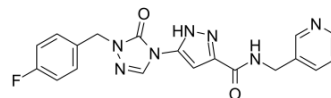


## SCD1 inhibitor-3

<b>Cat. No.:</b>	HY-139077
<b>CAS No.:</b>	1282606-48-7
<b>Molecular Formula:</b>	C <sub>19</sub> H <sub>16</sub> FN <sub>7</sub> O <sub>2</sub>
<b>Molecular Weight:</b>	393.37
<b>Target:</b>	Stearoyl-CoA Desaturase (SCD)
<b>Pathway:</b>	Metabolic Enzyme/Protease
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	SCD1 inhibitor-3 is a safe, potent and orally active SCD1 inhibitor. SCD1 inhibitor-3 can be used for the research of metabolic diseases such as obesity, type II diabetes and dyslipidemia, as well as skin diseases, acne and cancer <sup>[1]</sup> .																
<b>IC<sub>50</sub> &amp; Target</b>	SCD1 <sup>[1]</sup>																
<b>In Vivo</b>	<p>SCD1 inhibitor-3 (5 mg/kg; p.o.; 4 hours) reduces the plasma C16:1/C16:0 triglycerides desaturation index by 54 %<sup>[1]</sup>.            SCD1 inhibitor-3 (2~10 mg/kg; p.o.; 4 hours) makes a dose-responsive reduction of plasma triglycerides desaturation index<sup>[1]</sup>.</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>Lewis rats</td> </tr> <tr> <td>Dosage:</td> <td>5 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>P.o.; 4 hours</td> </tr> <tr> <td>Result:</td> <td>Reduced the plasma C16:1/C16:0 triglycerides desaturation index by 54 %.</td> </tr> </table> <table border="1"> <tr> <td>Animal Model:</td> <td>Lewis rats</td> </tr> <tr> <td>Dosage:</td> <td>2~10 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>P.o.; 4 hours</td> </tr> <tr> <td>Result:</td> <td>A dose-responsive reduction of plasma triglycerides desaturation index.</td> </tr> </table>	Animal Model:	Lewis rats	Dosage:	5 mg/kg	Administration:	P.o.; 4 hours	Result:	Reduced the plasma C16:1/C16:0 triglycerides desaturation index by 54 %.	Animal Model:	Lewis rats	Dosage:	2~10 mg/kg	Administration:	P.o.; 4 hours	Result:	A dose-responsive reduction of plasma triglycerides desaturation index.
Animal Model:	Lewis rats																
Dosage:	5 mg/kg																
Administration:	P.o.; 4 hours																
Result:	Reduced the plasma C16:1/C16:0 triglycerides desaturation index by 54 %.																
Animal Model:	Lewis rats																
Dosage:	2~10 mg/kg																
Administration:	P.o.; 4 hours																
Result:	A dose-responsive reduction of plasma triglycerides desaturation index.																

### REFERENCES

[1]. Sun S, et al. Discovery of triazolone derivatives as novel, potent stearoyl-CoA desaturase-1 (SCD1) inhibitors. *Bioorg Med Chem.* 2015;23(3):455-465.

---

**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](mailto:tech@MedChemExpress.com)

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