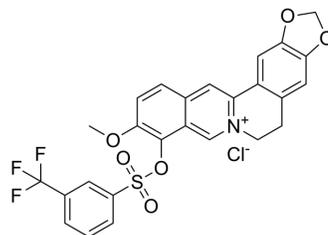


Lipid-lowering agent-1

Cat. No.:	HY-115914
CAS No.:	2304859-34-3
Molecular Formula:	C ₂₆ H ₁₉ ClF ₃ NO ₆ S
Molecular Weight:	565.95
Target:	Others
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Lipid-lowering agent-1 is a potent lipid-lowering agent. Lipid-lowering agent-1 has significant pharmacological effects on the inhibition of low-density lipoprotein cholesterol (LDLC) and promotion of high-density lipoprotein cholesterol (HDL) production. Lipid-lowering agent-1 shows potent hypolipidemic effect in high-fat diet rats ^[1] .																
In Vitro	<p>Lipid-lowering agent-1 (compound 1m) (12.5 μM; 72 hours) does not exhibit significant toxicity to both HepG2 and 3T3-L1 cells at a high concentration^[1].</p> <p>Lipid-lowering agent-1 (10 μM; 24 hours) has significant pharmacological effects on the inhibition of LDLC and promotion of HDLC production^[1].</p> <p>Lipid-lowering agent-1 (10 μM; 24 hours) inhibits the conversion of 3T3-L1 cells from pre-adipose to adipose-like with the inhibition of triglycerides (TG) by 70.4%^[1].</p> <p>Lipid-lowering agent-1 (10 μM; 10 days) decreases yellow lipid droplets effectively in 3T3-L1 cells^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HepG2 and 3T3-L1^[1]</td> </tr> <tr> <td>Concentration:</td> <td>12.5 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 hours</td> </tr> <tr> <td>Result:</td> <td>Did not exhibit significant toxicity to both HepG2 and 3T3-L1 cells at a high concentration.</td> </tr> </table> <p>Cell Differentiation Assay</p> <table border="1"> <tr> <td>Cell Line:</td> <td>3T3-L1 cells^[1]</td> </tr> <tr> <td>Concentration:</td> <td>10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibited the conversion of 3T3-L1 cells from pre-adipose to adipose-like with the inhibition of triglycerides (TG) by 70.4%</td> </tr> </table>	Cell Line:	HepG2 and 3T3-L1 ^[1]	Concentration:	12.5 μM	Incubation Time:	72 hours	Result:	Did not exhibit significant toxicity to both HepG2 and 3T3-L1 cells at a high concentration.	Cell Line:	3T3-L1 cells ^[1]	Concentration:	10 μM	Incubation Time:	24 hours	Result:	Inhibited the conversion of 3T3-L1 cells from pre-adipose to adipose-like with the inhibition of triglycerides (TG) by 70.4%
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In Vivo	Lipid-lowering agent-1 (15 mg/kg; i.g., daily for 40 or 80 days) reduces blood cholesterolby, triglycerides and LDLC by 19.6%, 34.52%, and 41.49% after dosing 40 days, moreover the three indexes is still better than that of berberine after dosing 80																

days^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	High-fat diet (HFD) rats ^[1]
Dosage:	15 mg/kg
Administration:	i.g., daily for 40 or 80 days
Result:	Reduced blood cholesterolby, triglycerides and LDLC by 19.6%, 34.52%, and 41.49% after dosing 40 days, moreover the three indexes was still better than that of berberine after dosing 80 days.

REFERENCES

[1]. Kong Y, et al. Discovery and structural optimization of 9-O-phenylsulfonyl-berberines as new lipid-lowering agents. *Bioorg Chem.* 2022;121:105665.

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

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