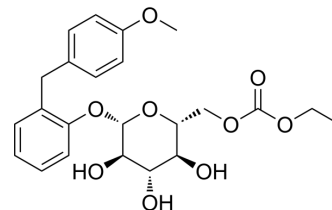


Sergliflozin etabonate

Cat. No.:	HY-12611
CAS No.:	408504-26-7
Molecular Formula:	C ₂₃ H ₂₈ O ₉
Molecular Weight:	448.46
Target:	SGLT
Pathway:	Membrane Transporter/Ion Channel
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Sergliflozin etabonate (GW-869682X) is a potent and orally active sodium glucose cotransporter (SGLT2) inhibitor. Sergliflozin etabonate shows antidiabetic and antihyperglycemic effects. Sergliflozin etabonate significantly reduces non-fasting blood glucose levels in diabetic mice. Sergliflozin etabonate has the potential for the research of diabetes ^[1] .																
IC₅₀ & Target	SGLT2																
In Vivo	<p>Sergliflozin etabonate (1, 3, 10, 30 mg/kg; oral; once) significantly reduces non-fasting blood glucose levels in KK-Ay mice in a dose-dependent manner^[1].</p> <p>Sergliflozin etabonate (1.7, 8.4, 45.7 mg/kg in the 0.001, 0.005, and 0.025% sergliflozin etabonate; fed; daily for 9 weeks) reduces non-fasting blood glucose level and decreases triglyceride content in the liver in a dose-dependent manner in KK-Ay mice^[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>14 weeks, Female KK-Ay mice^[1]</td> </tr> <tr> <td>Dosage:</td> <td>1, 3, 10, 30 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Oral; once</td> </tr> <tr> <td>Result:</td> <td>Significantly reduced non-fasting blood glucose level with the reduction in blood glucose level at 2 h post dosing in the 1 mg/kg group, 3 mg/kg group, 10 mg/kg group, and 30 mg/kg group was 12, 15, 28, and 39%, respectively.</td> </tr> </table> <table border="1"> <tr> <td>Animal Model:</td> <td>4 weeks, Female KK-Ay mice^[1]</td> </tr> <tr> <td>Dosage:</td> <td>1.7, 8.4, 45.7 mg/kg in the 0.001, 0.005, and 0.025% sergliflozin etabonate</td> </tr> <tr> <td>Administration:</td> <td>Fed, daily for 9 weeks</td> </tr> <tr> <td>Result:</td> <td>Reduced non-fasting blood glucose level and decreased triglyceride content in the liver in a dose-dependent manner in diabetic control KK-Ay mice.</td> </tr> </table>	Animal Model:	14 weeks, Female KK-Ay mice ^[1]	Dosage:	1, 3, 10, 30 mg/kg	Administration:	Oral; once	Result:	Significantly reduced non-fasting blood glucose level with the reduction in blood glucose level at 2 h post dosing in the 1 mg/kg group, 3 mg/kg group, 10 mg/kg group, and 30 mg/kg group was 12, 15, 28, and 39%, respectively.	Animal Model:	4 weeks, Female KK-Ay mice ^[1]	Dosage:	1.7, 8.4, 45.7 mg/kg in the 0.001, 0.005, and 0.025% sergliflozin etabonate	Administration:	Fed, daily for 9 weeks	Result:	Reduced non-fasting blood glucose level and decreased triglyceride content in the liver in a dose-dependent manner in diabetic control KK-Ay mice.
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REFERENCES

[1]. Katsuno K, et al. Long-term treatment with sergliflozin etabonate improves disturbed glucose metabolism in KK-A(y) mice. Eur J Pharmacol. 2009 Sep 15;618(1-3):98-104.

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

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