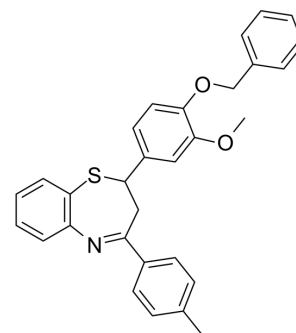


## α-Glucosidase-IN-17

<b>Cat. No.:</b>	HY-151141
<b>CAS No.:</b>	2820424-84-6
<b>Molecular Formula:</b>	C <sub>30</sub> H <sub>27</sub> NO <sub>2</sub> S
<b>Molecular Weight:</b>	465.61
<b>Target:</b>	Glucosidase
<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>	α-Glucosidase-IN-17 (Compound 12B) is a potent, orally active α-glucosidase inhibitor with an IC <sub>50</sub> of 3.79 μM. α-Glucosidase-IN-17 shows antidiabetic activity <sup>[1]</sup> .								
<b>IC<sub>50</sub> &amp; Target</b>	IC <sub>50</sub> : 3.79 μM (α-glucosidase) <sup>[1]</sup>								
<b>In Vivo</b>	<p>α-Glucosidase-IN-17 (Compound 12B) (10 and 20 mg/kg; p.o.; b.w. for 4 weeks) shows antidiabetic activity in <a href="#">Streptozocin</a> (HY-13753)-induced diabetic rats<sup>[1]</sup>.</p> <p>α-Glucosidase-IN-17 (10 and 20 mg/kg; p.o.; once) significantly decreases the serum glucose level after the administration of glucose (3 g/kg, oral) in rats<sup>[1]</sup>.</p> <p>α-Glucosidase-IN-17 (2000mg/kg; p.o.; b.w. for 2 weeks) demonstrates no mortality in mice<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>Male Wistar albino rats (170–200 g), Streptozotocin-induced diabetes model<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>10 and 20 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Oral administration, b.w. for 4 weeks</td> </tr> <tr> <td>Result:</td> <td>Decreased the level of blood glucose, reversed Streptozocin-induced body weight loss. Showed antihyperlipidemic effects on Streptozotocin-induced diabetes, reduced to a significant level of serum biomarkers.</td> </tr> </table>	Animal Model:	Male Wistar albino rats (170–200 g), Streptozotocin-induced diabetes model <sup>[1]</sup>	Dosage:	10 and 20 mg/kg	Administration:	Oral administration, b.w. for 4 weeks	Result:	Decreased the level of blood glucose, reversed Streptozocin-induced body weight loss. Showed antihyperlipidemic effects on Streptozotocin-induced diabetes, reduced to a significant level of serum biomarkers.
Animal Model:	Male Wistar albino rats (170–200 g), Streptozotocin-induced diabetes model <sup>[1]</sup>								
Dosage:	10 and 20 mg/kg								
Administration:	Oral administration, b.w. for 4 weeks								
Result:	Decreased the level of blood glucose, reversed Streptozocin-induced body weight loss. Showed antihyperlipidemic effects on Streptozotocin-induced diabetes, reduced to a significant level of serum biomarkers.								

### REFERENCES

[1]. Mehmood R, et al. Synthesis of Novel 2, 3-Dihydro-1, 5-Benzothiazepines as α-Glucosidase Inhibitors: In Vitro, In Vivo, Kinetic, SAR, Molecular Docking, and QSAR Studies. ACS Omega, 2022.

---

**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