



# SZABO SCANDIC

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## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

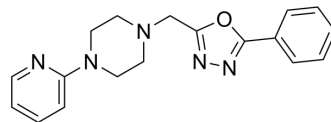
[mail@szabo-scandic.com](mailto:mail@szabo-scandic.com)

[www.szabo-scandic.com](http://www.szabo-scandic.com)

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

## AChE-IN-63

<b>Cat. No.:</b>	HY-158261
<b>CAS No.:</b>	876685-78-8
<b>Molecular Formula:</b>	C <sub>18</sub> H <sub>19</sub> N <sub>5</sub> O
<b>Molecular Weight:</b>	321.38
<b>Target:</b>	Cholinesterase (ChE); Amyloid-β; Beta-secretase
<b>Pathway:</b>	Neuronal Signaling
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	<p>AChE-IN-63 (Compound 5AD) is a selective inhibitor of hAChE (IC<sub>50</sub>=0.103 μM). AChE-IN-63 also inhibits hBChE and hBACE-1 (IC<sub>50</sub>= 10 μM (hBChE); 1.342 μM (hBACE-1)). AChE-IN-63 inhibits Aβ aggregation, preventing the formation and deposition of Aβ<sub>1-42</sub>. AChE-IN-63 can effectively penetrate the blood-brain barrier and is orally effective. It is primarily used in Alzheimer's disease research<sup>[1]</sup>.</p>									
<b>IC<sub>50</sub> &amp; Target</b>	<p>hAChE 0.103 μM (IC<sub>50</sub>)</p>	<p>hBChE 10 μM (IC<sub>50</sub>)</p>								
<b>In Vitro</b>	<p>AChE-IN-63 (5-20 μM; 48 h) inhibits Aβ<sub>1-42</sub> aggregation in a concentration-dependent manner<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>									
<b>In Vivo</b>	<p>AChE-IN-63 (p.o.; 100 mg/kg; single dose) exhibits no toxicity or abnormal reactions in Swiss albino mice<sup>[1]</sup>. AChE-IN-63 (p.o.; 5-20 mg/kg; single dose) improves learning and memory in cognitive dysfunction mice in the Scopolamine (HY-N0296) Y-Maze Model and demonstrates good antioxidant properties<sup>[1]</sup>. 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>Scopolamine Y-Maze Model <sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>5; 10 ;20 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>p.o.; single dose</td> </tr> <tr> <td>Result:</td> <td>Showed significant improvement in spontaneous alternation behavior. Reduced AChE and MDA levels and increased CAT levels.</td> </tr> </table>		Animal Model:	Scopolamine Y-Maze Model <sup>[1]</sup>	Dosage:	5; 10 ;20 mg/kg	Administration:	p.o.; single dose	Result:	Showed significant improvement in spontaneous alternation behavior. Reduced AChE and MDA levels and increased CAT levels.
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### REFERENCES

[1]. Singh A, et al. Structure-Guided Design, Synthesis, and Biological Evaluation of Peripheral Anionic Site Selective and Brain Permeable Novel Oxadiazole-Piperazine Conjugates against Alzheimer's Disease with Antioxidant Potential. ACS Omega. 2024 Apr 11;9(16):18169-18182.

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**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