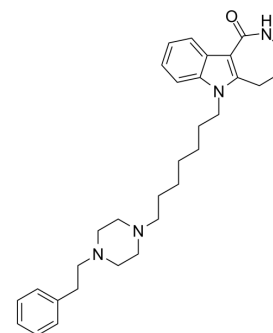


## BChE-IN-31

Cat. No.:	HY-163441
Molecular Formula:	C <sub>31</sub> H <sub>42</sub> N <sub>4</sub> O
Molecular Weight:	486.69
Target:	Cholinesterase (ChE)
Pathway:	Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	BChE-IN-31 (Compound 14d) is a selective BChE inhibitor with an IC <sub>50</sub> of 65 nM. BChE-IN-31 inhibits the self-induced aggregation of neurotoxic amyloid-β (Aβ) peptide <sup>[1]</sup> .									
<b>IC<sub>50</sub> &amp; Target</b>	BChE 65 nM (IC <sub>50</sub> )	AChE 5.9 μM (IC <sub>50</sub> )								
<b>In Vitro</b>	<p>BChE-IN-31 (0-20 μM, 24 h) exhibits a protective effect against NMDA (HY-17551)-induced toxicity and H<sub>2</sub>O<sub>2</sub>-induced oxidation in SH-SY5Y cells at 5 μM<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>SH-SY5Y</td> </tr> <tr> <td>Concentration:</td> <td>0-20 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Improved the cell viability at 5 μM, inhibited cell viability dose-dependently at 5-20 μM.</td> </tr> </table>		Cell Line:	SH-SY5Y	Concentration:	0-20 μM	Incubation Time:	24 h	Result:	Improved the cell viability at 5 μM, inhibited cell viability dose-dependently at 5-20 μM.
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### REFERENCES

[1]. Samarelli F, et al., Novel 6-alkyl-bridged 4-arylalkylpiperazin-1-yl derivatives of azepino [4, 3-b] indol-1 (2H)-one as potent BChE-selective inhibitors showing protective effects against neurodegenerative insults[J]. European Journal of Medicinal Chemistry, 2024: 116353.

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

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