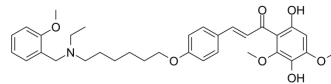


ACHe-IN-12

Cat. No.:	HY-144790
CAS No.:	2764664-52-8
Molecular Formula:	C ₃₃ H ₄₁ NO ₇
Molecular Weight:	563.68
Target:	Amyloid-β; AChE; Monoamine Oxidase
Pathway:	Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	<p>ACHe-IN-12 is a potent and blood-brain barrier (BBB) penetrant acetylcholinesterase (AChE) with IC₅₀s of 0.41 μM and 1.88 μM for rat AChE and electric eel AChE. AChE-IN-12 is also a good antioxidant (ORAC = 3.3 eq), selective metal chelator and hu MAO-B inhibitor (IC₅₀ = 8.8 μM). AChE-IN-12 has remarkable inhibition of self- and Cu²⁺-induced Aβ₁₋₄₂ aggregation, as well as exhibits a good neuroprotective effect. AChE-IN-12 can be used for researching Alzheimer's disease^[1].</p>																
IC₅₀ & Target	IC ₅₀ : 0.41 μM (rat AChE), 1.88 μM (electric eel AChE), 8.8 μM (huMAO-B) ^[1]																
In Vitro	<p>ACHe-IN-12 (compound 17f)(10-100 μM) exhibits no obvious cytotoxicity until the concentration increased up to 50 μM^[1]. AChE-IN-12 (10 and 50 μM) adds up the cell viability of H₂O₂-induced PC12 cells^[1]. 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>PC12 cells^[1]</td> </tr> <tr> <td>Concentration:</td> <td>10, 50 and 100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td></td> </tr> <tr> <td>Result:</td> <td>Did not show obvious cytotoxicity until the concentration increased up to 50 μM.</td> </tr> </table> <p>Cell Viability Assay</p> <table border="1"> <tr> <td>Cell Line:</td> <td>PC12 cells (exposed to 100 μM H₂O₂ for 1 hour to establish the oxidative damage model)^[1]</td> </tr> <tr> <td>Concentration:</td> <td>10 and 50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td></td> </tr> <tr> <td>Result:</td> <td>Added up the cell viability to 65.7% (p < 0.05) and 73.4% (p < 0.05) at 10 and 50 μM, respectively.</td> </tr> </table>	Cell Line:	PC12 cells ^[1]	Concentration:	10, 50 and 100 μM	Incubation Time:		Result:	Did not show obvious cytotoxicity until the concentration increased up to 50 μM.	Cell Line:	PC12 cells (exposed to 100 μM H ₂ O ₂ for 1 hour to establish the oxidative damage model) ^[1]	Concentration:	10 and 50 μM	Incubation Time:		Result:	Added up the cell viability to 65.7% (p < 0.05) and 73.4% (p < 0.05) at 10 and 50 μM, respectively.
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REFERENCES

[1]. Sang Z, Song Q, Cao Z, Deng Y, Zhang L. Design, synthesis, and evaluation of chalcone-Vitamin E-donepezil hybrids as multi-target-directed ligands for the treatment of Alzheimer's disease. *J Enzyme Inhib Med Chem.* 2022;37(1):69-85.

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

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