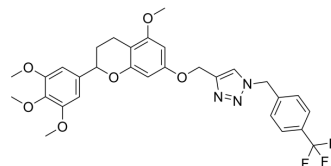


AChE/BuChE-IN-3

Cat. No.:	HY-147939
CAS No.:	2742707-47-5
Molecular Formula:	C ₃₀ H ₃₀ F ₃ N ₃ O ₆
Molecular Weight:	585.57
Target:	Cholinesterase (ChE); Amyloid-β
Pathway:	Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	AChE/BuChE-IN-3 is a potent and blood-brain barrier (BBB) penetrant AChE and BuChE dual inhibitor with IC ₅₀ s of 0.65 μM and 5.77 μM for AChE and BuChE. AChE/BuChE-IN-3 also inhibits Aβ ₁₋₄₂ aggregation. AChE/BuChE-IN-3 has effectively neuroprotective activities and nearly no toxicity on SH-SY5Y cells. AChE/BuChE-IN-3 can be used for researching Alzheimer's disease ^[1] .									
IC₅₀ & Target	AChE 0.65 μM (IC ₅₀)	BChE 5.77 μM (IC ₅₀)								
In Vitro	<p>AChE/BuChE-IN-3 (compound C4) (12.5-100 μM; 48 hours) exhibits no cytotoxicity on the human neuroblastoma cell line SH-SY5Y, and slightly increases cell viabilities^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>SH-SY5Y</td> </tr> <tr> <td>Concentration:</td> <td>12.5 μM, 25 μM, 50 μM and 100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> <tr> <td>Result:</td> <td>Exhibited no cytotoxicity on the human neuroblastoma cell line SH-SY5Y, and slightly increased cell viabilities.</td> </tr> </table>		Cell Line:	SH-SY5Y	Concentration:	12.5 μM, 25 μM, 50 μM and 100 μM	Incubation Time:	48 hours	Result:	Exhibited no cytotoxicity on the human neuroblastoma cell line SH-SY5Y, and slightly increased cell viabilities.
Cell Line:	SH-SY5Y									
Concentration:	12.5 μM, 25 μM, 50 μM and 100 μM									
Incubation Time:	48 hours									
Result:	Exhibited no cytotoxicity on the human neuroblastoma cell line SH-SY5Y, and slightly increased cell viabilities.									

REFERENCES

[1]. Shi S, et al. Semi-synthesis and biological evaluation of flavone hybrids as multifunctional agents for the potential treatment of Alzheimer's disease. *Bioorg Chem.* 2020 Jul;100:103917.

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

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA