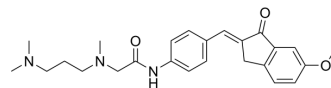


## ACHe/BChE/MAO-B-IN-3

<b>Cat. No.:</b>	HY-152113
<b>Molecular Formula:</b>	C <sub>25</sub> H <sub>31</sub> N <sub>3</sub> O <sub>3</sub>
<b>Molecular Weight:</b>	421.53
<b>Target:</b>	Monoamine Oxidase; Cholinesterase (ChE)
<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>	ACHe/BChE/MAO-B-IN-3, an indan-1-one derivative, is a potent MAO-B inhibitor with an IC <sub>50</sub> of 0.0359 μM for human MAO-B. AChE/BChE/MAO-B-IN-3 is a potent AChE and BChE enzyme inhibitor, with IC <sub>50</sub> s of 0.0473 μM and 0.0782 μM for human AChE and BChE enzyme, respectively. AChE/BChE/MAO-B-IN-3 shows significant antioxidant activity and has the potential for Alzheimer's disease (AD) research <sup>[1]</sup> .		
<b>IC<sub>50</sub> &amp; Target</b>	hMAO-B 0.0359 μM (IC <sub>50</sub> )	hAChE 0.0473 μM (IC <sub>50</sub> )	hBChE 0.0782 μM (IC <sub>50</sub> )
<b>In Vitro</b>	ACHe/BChE/MAO-B-IN-3 (compound D38) has the IC <sub>50</sub> value of 6.8118 μM on the NIH/3T3 fibroblast cell line <sup>[1]</sup> . AChE/BChE/MAO-B-IN-3 (1 mM) demonstrates strong human AChE and BChE enzyme inhibition profiles by generating at 93% activity, respectively <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

### REFERENCES

[1]. Begüm Nurpelin Sağlık, et al. Design, Synthesis, and In Vitro and In Silico Approaches of Novel Indanone Derivatives as Multifunctional Anti-Alzheimer Agents. ACS Omega. 2022 Dec 7;7(50):47378-47404.

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

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