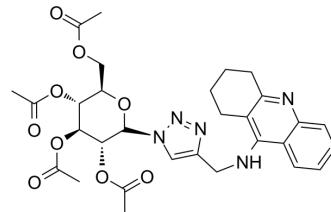


AChe-IN-9

| | |
|--------------------|---|
| Cat. No.: | HY-143291 |
| Molecular Formula: | C ₃₀ H ₃₅ N ₅ O ₉ |
| Molecular Weight: | 609.63 |
| Target: | Cholinesterase (ChE) |
| Pathway: | Neuronal Signaling |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |



BIOLOGICAL ACTIVITY

| | | | | | | | | | |
|-------------------------------------|---|------------|-------------|----------------|----------------------------------|------------------|----------|---------|---|
| Description | AChe-IN-9 is a Tacrine (HY-111338) glycoconjugate tethered with acetylated β-Glucose. AChE-IN-9 is also an AChE inhibitor with an IC ₅₀ value of 0.4 μM, with lower hepatotoxicity on healthy cells. Tacrine is used in Alzheimer's research ^[1] . | | | | | | | | |
| IC₅₀ & Target | Acetylcholinesterase 0.4 μM (IC ₅₀) | | | | | | | | |
| In Vitro | <p>Tacrine (HY-111338) is mainly metabolized by CYP 1A2, but its glycoconjugates have shown more affinity towards CYP 3A4 rather than CYP 1A2^[1].</p> <p>AChe-IN-9 (compound A-1) (200 μM; 24 h) is non-toxic on HepG2 cell line with 100% cell viability, and shows lower hepatotoxicity than Tacrine (HY-111338) on healthy HepG2 cells^[1].</p> <p>AChe-IN-9 (10 μM; 24 h) inhibits AChE activity with inhibition rate of 96.6%^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HepG2 cells</td> </tr> <tr> <td>Concentration:</td> <td>3.125, 6.25, 25, 50, 100, 200 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Showed much lower toxic effect on HepG2 cell than Tacrine (HY-111338) in a dose dependent manner, showing the inflexion point at the concentration of 100 μM.</td> </tr> </table> | Cell Line: | HepG2 cells | Concentration: | 3.125, 6.25, 25, 50, 100, 200 μM | Incubation Time: | 24 hours | Result: | Showed much lower toxic effect on HepG2 cell than Tacrine (HY-111338) in a dose dependent manner, showing the inflexion point at the concentration of 100 μM. |
| Cell Line: | HepG2 cells | | | | | | | | |
| Concentration: | 3.125, 6.25, 25, 50, 100, 200 μM | | | | | | | | |
| Incubation Time: | 24 hours | | | | | | | | |
| Result: | Showed much lower toxic effect on HepG2 cell than Tacrine (HY-111338) in a dose dependent manner, showing the inflexion point at the concentration of 100 μM. | | | | | | | | |

REFERENCES

[1]. Kaur Gulati H, et al. Design, Synthesis, biological investigations and molecular interactions of triazole linked tacrine glycoconjugates as Acetylcholinesterase inhibitors with reduced hepatotoxicity. *Bioorg Chem.* 2022 Jan;118:105479.

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

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