

Product Data Sheet

MCE MedChemExpress

Multitarget AD inhibitor-1

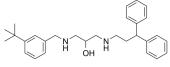
Cat. No.:HY-136813CAS No.:2205015-77-4Molecular Formula: $C_{29}H_{38}N_2O$ Molecular Weight:430.62

Target: Beta-secretase; Amyloid-β; Cholinesterase (ChE)

Pathway: Neuronal Signaling

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.



BIOLOGICAL ACTIVITY

Description	Multitarget AD inhibitor-1 is a selective and reversible butyrylcholinesterase (BuChE) inhibitor with IC $_{50}$ s of 7.22 μ M and 1.55 μ M for hBuChE and eqBuChE (BuChE from equine serum), respectively. Multitarget AD inhibitor-1 inhibits β -secretase (IC $_{50\text{hBACE-1}}$ =41.60 μ M), amyloid β aggregation (IC $_{50\text{A}\beta}$ =3.09 μ M), tau aggregation. Multitarget AD inhibitor-1, a diphenylpropylamine derivative, has the potential for multifunctional disease-modifying anti-Alzheimer's research[1].
In Vitro	Multitarget AD inhibitor-1 (compound 10) selectively inhibits BuChE from equine serum (eqBuChE) and hBuChE over AChE from electric eel (eeAChE; 21.78% at 10 μ M)[1]. Multitarget AD inhibitor-1 (10 μ M) inhibits full-length tau aggregation with 44.4%[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Dawid Panek, et al. Design, Synthesis, and Biological Evaluation of 1-Benzylamino-2-hydroxyalkyl Derivatives as New Potential Disease-Modifying Multifunctional Anti-Alzheimer's Agents. ACS Chem Neurosci. 2018 May 16;9(5):1074-1094.

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

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