

Product Data Sheet

tau/Aβ40 aggregation-IN-1

Cat. No.: HY-149272

Molecular Formula: $C_{23}H_{28}N_2O_2S$ Molecular Weight: 396.55

Target: Tau Protein; Amyloid-β; Cholinesterase (ChE)

Pathway: Neuronal Signaling

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

Analysis.

BIOLOGICAL ACTIVITY

Description	tau/A β 40 aggregation-IN-1 (C respectively ^[1] .	au/A eta 40 aggregation-IN-1 (Compound 20) is a tau and A eta_{40} aggregation inhibitor with IC $_{50}$ s of 1.8 μ M and 1.3 μ M, espectively ^[1] .		1.8 μM and 1.3 μM,
IC ₅₀ & Target	Aβ ₄₀ aggregation	tau aggregation	equine serum BChE	hAChE

IC₅₀ & Target Aβ₄₀ aggregation tau aggregation equine serum BChE hAChE $1.3 \,\mu\text{M} \,(\text{IC}_{50})$ $1.8 \,\mu\text{M} \,(\text{IC}_{50})$ $0.96 \,\mu\text{M} \,(\text{IC}_{50})$ $16.10 \,\mu\text{M} \,(\text{IC}_{50})$

In Vitro tau/A β 40 aggregation-IN-1 (Compound 20; 0.1-10 μ M; 24 h) shows neuroprotective effect against Okadaic acid (HY-N6785)-induced tau toxicity in SH-SY5Y cells^[1].

tau/A β 40 aggregation-IN-1 inhibits electric eel AChE, human AChE and equine serum BChE with IC₅₀s of 4.4, 16.10 and 0.96 μ M, respectively. tau/A β 40 aggregation-IN-1 shows 16% inhibition against human BChE at 10 μ M^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

 ${\sf Cell\ Viability\ Assay}^{[1]}$

Cell Line:	Neuroblastoma cell line SH-SY5Y	
Concentration:	0.1, 1, 5, 10 and 50 μM	
Incubation Time:	24 h with Okadaic acid (100 nM)	
Result:	Exhibited a potent neuroprotective effect from 0.1 to 10 μM .	

REFERENCES

[1]. Tonelli M, et al. Thioxanthenone-based derivatives as multitarget therapeutic leads for Alzheimer's disease. Eur J Med Chem. 2023 Mar 15;250:115169.

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

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