
Product Data Sheet

Product Name: δ -Secretase inhibitor 11

Cat. No.: GC66048

Chemical Properties

Cas. No. 842964-18-5

Formula $C_{10}H_{12}N_4O_2$ M.Wt 220.23

Solubility DMSO : 125 mg/mL (567.59 mM; Need ultrasonic) Storage Store at $-20^{\circ}C$

General tips For obtaining a higher solubility , please warm the tube at $37^{\circ}C$ and shake it in the ultrasonic bath for a while. Stock solution can be stored below $-20^{\circ}C$ for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure

Background

Compound 11, a non-toxic and selective δ -secretase inhibitor ($IC_{50}=0.7 \mu M$, in fluorescence-based assay) that interacts with both the active site and allosteric site of δ -secretase in Co-crystal structure analysis. The IC_{50} value of the compound 11 towards δ -secretase in Pala cells is $0.8 \mu M$.

Compound 11 inhibit δ -secretase in Pala cells, with an IC_{50} of $7\mu M$. Compounds 11 is highly permeable assessed by Caco-2 monolayer permeability screen. The human liver microsomal stability screen demonstrates that following 30?min of incubation, 76% of compound 11 remained in human liver microsomes.[1]

Chronic treatment with δ -secretase inhibitor 11 markedly decreases the brain asparaginyl endopeptidase (AEP) activity, reduces the generation of $A\beta_{1-40/42}$ and ameliorates memory loss. The inhibition of AEP with this reagent not only reduces the AEP-cleaved tau fragments and tau hyperphosphorylation, but also attenuates neuroinflammation in the form of microglial activation. Treatment with δ -secretase inhibitor 11 prevents the synaptic loss and alleviated dendritic disruption in senescence-accelerated mouse prone 8 (SAMP8) mouse brain.[2]

[1] Zhang Z, et al. Nat Commun. 2017 Mar 27;8:14740. [2] Wang J, et al. Transl

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

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Neurodegener. 2021 Mar 31;10(1):12.

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