Proteins

Product Data Sheet

β-Sheet Breaker Peptide iAβ5

Cat. No.: HY-P1047 CAS No.: 182912-74-9 Molecular Formula: $C_{33}H_{43}N_5O_8$ Molecular Weight: 637.72

Sequence: Leu-Pro-Phe-Phe-Asp

Sequence Shortening: LPFFD Target: Amyloid-β

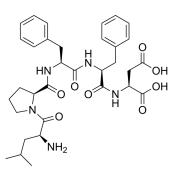
Pathway: **Neuronal Signaling**

Sealed storage, away from moisture and light, under nitrogen Storage:

> Powder -80°C 2 years -20°C 1 year

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture

and light, under nitrogen)



SOLVENT & SOLUBILITY

In Vitro

H₂O: 50 mg/mL (78.40 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.5681 mL	7.8404 mL	15.6809 mL
	5 mM	0.3136 mL	1.5681 mL	3.1362 mL
	10 mM	0.1568 mL	0.7840 mL	1.5681 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description

β-Sheet Breaker Peptide iAβ5 is a potent degrader of cerebral amyloid-beta (Abeta). Abeta deposition is associatied with the Alzheimer disease (AD), due to its related toxicity linked to its beta-sheet conformation and/or aggregation. β-Sheet Breaker Peptide iAβ5 reproducibly induces in vivo disassembly of fibrillar amyloid deposits. Thus, β-Sheet Breaker Peptide iAβ5 prevents and/or reverses neuronal shrinkage caused by Abeta, and reduces the extent of interleukin-1beta positive microglia-like cells that surround the Abeta deposits. β-Sheet Breaker Peptide iAβ5 reduces the size and/or number of cerebral amyloid plaques in AD. β-Sheet Breaker Peptide iAβ5 labeled by hydrophobic benzyl alcohol (HBA) tag, can be used for quantitative assay by showing vivid blue color under acidic conditions^{[1][2][3]}.

In Vitro

β-Sheet Breaker Peptide iAβ5 (1.5 μg/μL; 7 days) incubates with Aβ1-42 (0.5 μg/μL) for 7 days, inhibits amyloid βprotein fibrillogenesis, disassembles preformed fibrils in vitro and prevents neuronal death induced by fibrils in cell culture^[1]. β-Sheet Breaker Peptide iAβ5 (60 μM; 48 h) shows insignificant cytotoxicity in human neuroblastoma (IMR-32) cell treated

	with 50 μ M aggregated A β 1-42 for 2 days ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	β -Sheet Breaker Peptide iA β 5 (100 nmol/rat; Intra-amygdala injection; 7 days) coinjected with A β 1-42 (5 nmol) into in the rat amygdala, blocks A β 1-42 neurotoxicity in tissue culture, and amyloid fibril formation in the rat model ^[1] . β -Sheet Breaker Peptide iA β 5 (100 nmol/rat, 200 nmol/rat; Intra-amygdaloid injection; 7 days) followed by A β 1-42 (5 nmol), induces disassembly of pre-existing A β fibrils in vivo, and leads to a reversal or prevention of A β -induced histopathological changes in rat model ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

- $[1]. Sigurds son EM, et al.\ In\ vivo\ reversal\ of\ amyloid-beta\ lesions\ in\ rat\ brain.\ J\ Neuropathol\ Exp\ Neurol.\ 2000\ Jan; 59(1):11-7.$
- [2]. Okada Y, et al. Acid-Triggered Colorimetric Hydrophobic Benzyl Alcohols for Soluble Tag-Assisted Liquid-Phase Synthesis. Org Lett. 2015 Sep 4;17(17):4264-7.
- [3]. Soto C, et al. Beta-sheet breaker peptides inhibit fibrillogenesis in a rat brain model of amyloidosis: implications for Alzheimer's therapy. Nat Med. 1998 Jul;4(7):822-6.

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

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