## **Product** Data Sheet

## Aβ42-IN-1

 $\begin{array}{lll} \textbf{Cat. No.:} & \textbf{HY-130609} \\ \textbf{CAS No.:} & 2582757-69-3 \\ \textbf{Molecular Formula:} & \textbf{C}_{29}\textbf{H}_{27}\textbf{ClN}_4\textbf{O}_2 \\ \end{array}$ 

Molecular Weight: 499

**Target:** γ-secretase

Pathway: Neuronal Signaling; Stem Cell/Wnt

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

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	Aβ42-IN-1, compound 1v, is a novel, potent and orally active γ-secretase modulator (GSM). Aβ42-IN-1 potently reduced Aβ42 levels with an IC <sub>50</sub> value of 0.091 μM without CYP3A4 inhibition. Aβ42-IN-1 shows a sustained pharmacokinetic profile.	
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IC <sub>50</sub> & Target	IC50: $0.091  \mu \text{M}$ (γ-secretase modulator) $^{[1]}$	
In Vivo	Aβ42-IN-1 (orally adminstation; 30 mg/kg)shows a good pharmacokinetic profile in mice, significantly reduces Aβ42 levels in the brain and plasma, by 36% and 66%, respectively. while the brain/plasma ratio of 1v is slightly lower than that of 1a <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	8- week-old male ddY mice
	Dosage:	30 mg/kg
	Administration:	Orally adminstation; 30 mg/kg
		Reduced Aβ42 levels in the brain and plasma, by 36% and 66%, respectively.

## **REFERENCES**

[1]. Sekioka R, et al. Discovery of N-ethylpyridine-2-carboxamide derivatives as a novel scaffold for orally active γ-secretase modulators. Bioorg Med Chem. 2019 Nov 6:115132.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA