**Proteins** 

# Inhibitors

# **SEN12333**

Cat. No.: HY-107678 CAS No.: 874450-44-9 Molecular Formula:  $C_{20}H_{25}N_3O_2$ Molecular Weight: 339.43

Target: nAChR

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

Storage: Powder -20°C 3 years

In solvent

4°C 2 years -80°C 6 months

-20°C 1 month

**Product** Data Sheet

### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (294.61 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.9461 mL	14.7306 mL	29.4612 mL
	5 mM	0.5892 mL	2.9461 mL	5.8922 mL
	10 mM	0.2946 mL	1.4731 mL	2.9461 mL

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

In Vivo

1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (7.37 mM); Clear solution

# **BIOLOGICAL ACTIVITY**

Description	SEN 12333 (WAY-317538) is a potent, selective and orally active $\alpha$ 7 nAChR agonist. SEN12333 displays high affinity for the rat $\alpha$ 7 nAChRs expressed in GH4C1 cells ( $K_{>i}$ =260 nM) and acts as full agonist in functional Ca <sup>2+</sup> flux studies (EC <sub>50</sub> =1.6 $\mu$ M). SEN 12333 is used for AD and schizophrenia research <sup>[1]</sup> .
In Vitro	In whole-cell patch-clamp recordings, SEN12333 activates peak currents and maximal total charges similar to acetylcholine (EC $_{50}$ = 12 $\mu$ M). SEN12333 displays functional antagonism at histamine H3 receptors (IC $_{50}$ =103 nM) and weak agonist activity at human ganglionic $\alpha$ 3 nAChRs (IC $_{50}$ =8.5 $\mu$ M)[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	SEN12333 (intraperitoneal injection; 3 mg/kg) improves episodic memory in a novel object recognition task in rats in conditions of spontaneous forgetting as well as cognitive disruptions induced via glutamatergic or cholinergic mechanisms. SEN12333 also prevents a scopolamine-induced deficit in a passive avoidance task <sup>[1]</sup> .

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

### **REFERENCES**

[1]. Corinne Beinat, et al. The Recent Development of  $\alpha 7$  Nicotinic Acetylcholine Receptor (nAChR) Ligands as Therapeutic Candidates for the Treatment of Central Nervous System (CNS) Diseases. Curr Pharm Des. 2016;22(14):2134-51.

[2]. Renza Roncarati, et al. Procognitive and neuroprotective activity of a novel alpha7 nicotinic acetylcholine receptor agonist for treatment of neurodegenerative and cognitive disorders. J Pharmacol Exp Ther. 2009 May;329(2):459-68.

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

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