Proteins

Inhibitors



QR-0217

Cat. No.: HY-153416 CAS No.: 1027786-12-4 Molecular Formula: C₁₉H₁₃NO₃ Molecular Weight: 303.31

Target: Amyloid-β

Pathway: **Neuronal Signaling**

Storage: Powder -20°C 3 years

4°C 2 years

In solvent -80°C 6 months

> -20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (329.70 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.2970 mL	16.4848 mL	32.9696 mL
	5 mM	0.6594 mL	3.2970 mL	6.5939 mL
	10 mM	0.3297 mL	1.6485 mL	3.2970 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 (8.24 mM); Clear solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- β -CD in saline) Solubility: 2.5 mg/mL (8.24 mM); Clear solution; Need ultrasonic

BIOLOGICAL ACTIVITY

Description	QR-0217 is a potent A β 1-40 aggregation inhibitor with an IC $_{50}$ value of 7.5 μ M. QR-0217 inhibits α -synuclein aggregation. QR-0217 reduces memory impairments caused by A β neurotoxicity ^[1] .		
IC ₅₀ & Target	IC_{50} :7.5 μM (Aβ1-40) ^[1]		
In Vitro	QR-0217 (20, 100 μ M; 96 h) inhibits α -synuclein (4 μ M) aggregation ^[1] . QR-0217 (50 μ M) significantly rescues impairment of LTP in the APP/PS1 transgenic mice hippocampal slices ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

Caution: Product has not been fully validated for medical applications. For research use only. Tel: 609-228-6998 Fax: 609-228-5909 E-mail: tech@MedChemExpress.com Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA	REFERENCES					
Tel: 609-228-6898 Fax: 609-228-5909 E-mail: tech@MedChemExpress.com	[1]. Michael D. Carter, et al. Compounds and methods for treating protein folding disorders. US8362066B2.					
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