MCH(human, mouse, rat) TFA

Cat. No.: HY-P1205A

Molecular Formula: $C_{107}H_{161}F_{3}N_{30}O_{28}S_{4}$

Molecular Weight:

Sequence Shortening: DFDMLRCMLGRVYRPCWQV (Disulfide bridge:Cys7-Cys16)

DFDMLRCMLGRVYRPCWQV (Disulfide bridge:Cvs7-Cvs16) (TFA salt)

Target: MCHR1 (GPR24)

Pathway: GPCR/G Protein; Neuronal Signaling Storage: Sealed storage, away from moisture

> Powder -80°C 2 years

> > -20°C 1 year

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

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (39.99 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	0.3999 mL	1.9993 mL	3.9986 mL
	5 mM	0.0800 mL	0.3999 mL	0.7997 mL
	10 mM	0.0400 mL	0.1999 mL	0.3999 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 (1.00 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (1.00 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (1.00 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

MCH (human, mouse, rat) TFA is a potent peptide agonist of MCH-R and exhibits binding IC₅₀ values of 0.3nM and 1.5 nM for MCH1R and MCH2R, respectively. MCH (human, mouse, rat) is a highly sensitive to MCH-2R in a CHO cell line and monitoring mobilization of intracellular calcium with FLIPR, exhibits functional activation EC_{50} values of 3.9 nM and 0.1nM for human MCH-1R and MCH-2R, respectively^[1].

IC₅₀ & Target

Binding IC50: 0.3 nM (MCH1R) IC50: 1.5 nM (MCH1R)[1]

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REFERENCES				
[1]. A W Sailer, et al. Identifica	tion and characterization of	a second melanin-concentrating	g hormone receptor, MCH-2R. Proc Natl Aca	d Sci U S A. 2001 Jun 19;98(13):7564-9
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