**Proteins** 

# **Product** Data Sheet

CRPRLCKHCRPRLC (Disulfide bridge:Cvs1-Cvs6; Cvs9-Cvs14)

## **MM 54**

Cat. No.: HY-P2271 CAS No.: 1313027-43-8 Molecular Formula:  $C_{70}H_{121}N_{29}O_{15}S_4$ 

Molecular Weight: 1737.17

**Sequence Shortening:** CRPRLCKHCRPRLC (Disulfide bridge:Cys1-Cys6; Cys9-Cys14)

Target: Apelin Receptor (APJ) Pathway: GPCR/G Protein

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

> Powder -80°C 2 years 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

 $H_2O : \ge 100 \text{ mg/mL} (57.56 \text{ mM})$ 

\* "≥" means soluble, but saturation unknown.

| Preparing<br>Stock Solutions | Solvent Mass<br>Concentration | 1 mg      | 5 mg      | 10 mg     |
|------------------------------|-------------------------------|-----------|-----------|-----------|
|                              | 1 mM                          | 0.5756 mL | 2.8782 mL | 5.7565 mL |
|                              | 5 mM                          | 0.1151 mL | 0.5756 mL | 1.1513 mL |
|                              | 10 mM                         | 0.0576 mL | 0.2878 mL | 0.5756 mL |

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

### **BIOLOGICAL ACTIVITY**

| Description | MM 54 (compound 5) is a competitive antagonist at APJ, with an IC <sub>50</sub> of 93 nM. MM 54 behaves as a potent and selective inhibitor of apelin binding and APLNR activation <sup>[1][2]</sup> .   |
|-------------|--|
| In Vitro    | MM 54 inhibits more than 95% of apelin binding to APLNR at the dose of 10 $\mu$ M $^{[2]}$ . MCE has not independently confirmed the accuracy of these methods. They are for reference only.   |
| In Vivo     | MM 54 (2 mg/kg, ip, bi-weekly for 4 weeks) possesses anti-tumor activity in glioblastoma models with no obvious toxicity <sup>[2]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.  Animal Model:  Tumour-bearing nude-mice <sup>[2]</sup> . |

Page 1 of 2 www.MedChemExpress.com

| Dosage:         | 2 mg/kg.  |
|-----------------|---|
| Administration: | Intraperitoneal injection, bi-weekly for 4 weeks. |
| Result:         | Reduced tumor progression (glioblastoma).         |

#### **REFERENCES**

[1]. N J Maximilian Macaluso, et al. Discovery of a competitive apelin receptor (APJ) antagonist. ChemMedChem. 2011 Jun 6;6(6):1017-23.

[2]. Elizabeth Harford-Wright, et al. Pharmacological targeting of apelin impairs glioblastoma growth. Brain. 2017 Nov 1;140(11):2939-2954.

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

Page 2 of 2 www.MedChemExpress.com