Product Data Sheet

SR17018

Cat. No.: HY-111454 CAS No.: 2134602-45-0 Molecular Formula: $C_{19}H_{18}Cl_{3}N_{3}O$ Molecular Weight: 410.72

Target: **Opioid Receptor**

Pathway: GPCR/G Protein; Neuronal Signaling

Storage: Powder -20°C 3 years

> 4°C 2 years

In solvent -80°C 2 years

> -20°C 1 year

SOLVENT & SOLUBILITY

In Vitro DMSO: 12.5 mg/mL (30.43 mM; Need ultrasonic)

H₂O: < 0.1 mg/mL (insoluble)

| Preparing Stock Solutions | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg |
|------------------------------|----------------------------|-----------|------------|------------|
| | 1 mM | 2.4347 mL | 12.1737 mL | 24.3475 mL |
| | 5 mM | 0.4869 mL | 2.4347 mL | 4.8695 mL |
| | 10 mM | 0.2435 mL | 1.2174 mL | 2.4347 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: ≥ 1.25 mg/mL (3.04 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1.25 mg/mL (3.04 mM); Clear solution

BIOLOGICAL ACTIVITY

| Description | SR17018 is an mu-opioid-receptor (MOR) agonist, binding with GTP γ S, with an EC $_{50}$ of 97 nM. | | |
|---------------------------|---|--|--|
| IC ₅₀ & Target | EC50: 97 nM (MOR) ^[1] | | |
| In Vitro | SR17018 is an mu-opioid-receptor (MOR) agonist, binding with GTP γ S, with an EC $_{50}$ of 97 nM. SR17018 shows no obvious effect on inducing β arrestin2 recruitment to the MOR at below 10 μ M. SR17018 promotes signaling through G proteins or β arrestin2 ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | | |

CUSTOMER VALIDATION

• Authorea. 2021 Jan 10.

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REFERENCES

[1]. Schmid CL, et al. Bias Factor and Therapeutic Window Correlate to Predict Safer Opioid Analgesics. Cell. 2017 Nov 16;171(5):1165-1175.e13.

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