## **Product** Data Sheet

## HHS-0701

 Cat. No.:
 HY-138665

 CAS No.:
 2851993-91-2

 Molecular Formula:
  $C_{20}H_{20}N_4O_3S$  

 Molecular Weight:
 396.46

Target: Prostaglandin Receptor

Pathway: GPCR/G Protein

Storage: Powder -20°C 3 years

4°C 2 years

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## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (252.23 mM; Need ultrasonic)

	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.5223 mL	12.6116 mL	25.2232 mL
Stock Solutions	5 mM	0.5045 mL	2.5223 mL	5.0446 mL
	10 mM	0.2522 mL	1.2612 mL	2.5223 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: ≥ 2.08 mg/mL (5.25 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (5.25 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (5.25 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description HHS-0701, a sulfur-triazole exchange (SuTEx) ligand, is a potent tyrosine-reactive prostaglandin reductase 2 (PTGR2) inhibitor. HHS-0701 blocks PTGR2 metabolism of the lipid substrate 15-Keto-PGE2<sup>[1]</sup>.

In Vitro HHS-0701 (0.1, 0.25, 0.5, 1, 2.5, 5, 10, 25  $\mu$ M; 2 hours) results concentration-dependent blockade of probe labeling in treatment of recombinant PTGR2 overexpressing HEK293T cells<sup>[1]</sup>.

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

Page 1 of 2

<sup>\*</sup> The compound is unstable in solutions, freshly prepared is recommended.

REFERENCES
[1]. Emmanuel K Toroitich, et al.Discovery of a Cell-Active SuTEx Ligand of Prostaglandin Reductase 2. Chembiochem. 2021 Jun 15;22(12):2134-2139.
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

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