# **Product** Data Sheet



## **URAT1** inhibitor 1

Cat. No.: HY-114309 CAS No.: 2268720-62-1 Molecular Formula:  $C_{19}H_{15}Br_2N_5O_2S_2$ 

Molecular Weight: 569.29 URAT1 Target:

Pathway: Membrane Transporter/Ion Channel

Storage: Powder -20°C 3 years 4°C 2 years

In solvent -80°C 6 months

> -20°C 1 month

#### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 125 mg/mL (219.57 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.7566 mL	8.7829 mL	17.5657 mL
	5 mM	0.3513 mL	1.7566 mL	3.5131 mL
	10 mM	0.1757 mL	0.8783 mL	1.7566 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 (3.65 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (3.65 mM); Clear solution

### **BIOLOGICAL ACTIVITY**

Description	URAT1 inhibitor 1 (1g) is a uric acid transporter 1 (URAT1) inhibitor, with an IC <sub>50</sub> of 32 nM. URAT1 inhibitor 1 has potential to treat hyperuricemia associated with gout <sup>[1]</sup> .
IC <sub>50</sub> & Target	IC50: 32 nM (URAT1) <sup>[1]</sup> .

#### **REFERENCES**

[1]. Wu JW, et al. Synthesis, biological evaluation and 3D-QSAR studies of 1,2,4-triazole-5-substituted carboxylic acid bioisosteres as uric acid transporter 1 (URAT1)

nhibitors for the treatment of h	hyperuricemia associated v	with gout. Bioorg Med Chem Lett.	2019 Feb 1;29(3):383-388.			
Caution: Product has not been fully validated for medical applications. For research use only.						
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