

Product Data Sheet

A2AR-agonist-1

Cat. No.: HY-18776

CAS No.: 41552-95-8Molecular Formula: $C_{20}H_{22}N_6O_4$ Molecular Weight: 410.43

Target: Adenosine Receptor
Pathway: GPCR/G Protein

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: 100 mg/mL (243.65 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.4365 mL	12.1823 mL	24.3647 mL
	5 mM	0.4873 mL	2.4365 mL	4.8729 mL
	10 mM	0.2436 mL	1.2182 mL	2.4365 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 (6.09 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.09 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.09 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

A2AR-agonist-1 is a potent A2AR and ENT1 agonist with Ki of 4.39 and 3.47 for A2AR and ENT1.IC50 value: 4.39 and 3.47 (Ki) [1]Target: A2AR and ENT1A2AR-agonist-1 is a novel dual-action compound, targeting the Adenosine A2A Receptor and Adenosine Transporter for Neuroprotection.[1]

CUSTOMER VALIDATION

• Int Immunopharmacol. 2022 Dec 16;114:109567.

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REFERENCES

- [1]. Chen JB, et al. Design and synthesis of novel dual-action compounds targeting the adenosine A(2A) receptor and adenosine transporter for neuroprotection. ChemMedChem. 2011 Aug 1;6(8):1390-1400.
- [2]. Lin Yun-Lian, et al. Dual-action compounds targeting adenosine A2A receptor and adenosine transporter for prevention and treatment of neurodegenerative diseases. From PCT Int. Appl. (2012), WO 2012064340 A1 20120518.
- [3]. Chen, Chih-Cheng, et al. Methods and compositions for treating pain. From PCT Int. Appl. (2013), WO 2013120078 A1 20130815.

Caution: Product has not been fully validated for medical applications. For research use only.

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