

Product Data Sheet

PAC1R antagonist 1

 Cat. No.:
 HY-147557

 CAS No.:
 2305204-24-2

 Molecular Formula:
 $C_{17}H_{17}ClN_6O_2$

 Molecular Weight:
 372.81

Target: PACAP Receptor
Pathway: GPCR/G Protein

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: 33.33 mg/mL (89.40 mM; ultrasonic and warming and heat to 160°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.6823 mL	13.4117 mL	26.8233 mL
	5 mM	0.5365 mL	2.6823 mL	5.3647 mL
	10 mM	0.2682 mL	1.3412 mL	2.6823 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 (6.71 mM); Clear solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- β -CD in saline) Solubility: 2.5 mg/mL (6.71 mM); Clear solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (6.71 mM); Clear solution; Need ultrasonic

BIOLOGICAL ACTIVITY

Description

PAC1R antagonist 1 (compound 3d) is a potent and orally active antagonist of PAC1 receptor. PAC1R antagonist 1 can inhibit pituitary adenylate cyclase-activating polypeptide (PACAP)- and nerve injury-induced allodynia $^{[1]}$.

REFERENCES

1]. Takasaki I, et, al. Synthesis c .5;186:111902.	of a novel and potent small-mo	olecule antagonist of PAC1 recept	or for the treatment of neuropathic pain. I	Eur J Med Chem. 2020 Jan		
Caution: Product has not been fully validated for medical applications. For research use only.						
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