Proteins



Verosudil

Cat. No.: HY-16758 CAS No.: 1414854-42-4 Molecular Formula: $C_{17}H_{17}N_3O_2S$

Molecular Weight: 327.4 ROCK Target:

Pathway: Cell Cycle/DNA Damage; Cytoskeleton; Stem Cell/Wnt; TGF-beta/Smad

Storage: Powder -20°C 3 years

4°C 2 years -80°C In solvent 6 months -20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 16.67 mg/mL (50.92 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.0544 mL	15.2718 mL	30.5437 mL
	5 mM	0.6109 mL	3.0544 mL	6.1087 mL
	10 mM	0.3054 mL	1.5272 mL	3.0544 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: ≥ 1.67 mg/mL (5.10 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 1.67 mg/mL (5.10 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1.67 mg/mL (5.10 mM); Clear solution

BIOLOGICAL ACTIVITY

Description Verosudil (AR-12286) is a potent, selective Rho-kinase (ROCK) inhibitor with Kis of 2 and 2 nM for ROCK1 and ROCK2, respectively. AR-12286 lowers intraocular pressure (IOP) primarily by increasing aqueous humour outflow through the trabecular meshwork^{[1][2]}.

IC₅₀ & Target ROCK1 ROCK2 PKA MRCKA 69 nM (Ki) 28 nM (Ki) 2 nM (Ki) 2 nM (Ki)

> CAM2A PKC theta

5855 nM (Ki) 9322 nM (Ki)

REFERENCES

[1]. Lin CW, Sherman B, Moore LA, et al. Discovery and Preclinical Development of Netarsudil, a Novel Ocular Hypotensive Agent for the Treatment of Glaucoma. J Ocul Pharmacol Ther. 2018;34(1-2):40-51.

[2]. Kopczynski C, Novack GD, Swearingen D, van Haarlem T. Ocular hypotensive efficacy, safety and systemic absorption of AR-12286 ophthalmic solution in normal volunteers. Br J Ophthalmol. 2013;97(5):567-572.

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

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