Product Data Sheet

KY-02327 acetate

Cat. No.: HY-124156A Molecular Formula: $C_{22}H_{31}N_3O_6$

433.5 Molecular Weight: Wnt

Pathway: Stem Cell/Wnt

Storage: 4°C, sealed storage, away from moisture and light

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture

and light)

SOLVENT & SOLUBILITY

In Vitro

Target:

 $H_2O : \ge 160 \text{ mg/mL} (369.09 \text{ mM})$

DMSO: 140 mg/mL (322.95 mM; ultrasonic and warming and heat to 80°C)

* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.3068 mL	11.5340 mL	23.0681 mL
	5 mM	0.4614 mL	2.3068 mL	4.6136 mL
	10 mM	0.2307 mL	1.1534 mL	2.3068 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 (5.77 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (5.77 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (5.77 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

KY-02327 acetate, a metabolically stabilized KY-02061 analog, is a potent Dishevelled (Dvl)-CXXC5 interaction inhibitor. KY-02327 acetate shows an activating effect on the Wnt/ β -catenin pathway, resulting in promotion of osteoblast differentiation [1]

In Vitro

KY-02327 (1-10 μM; 2 days; MC3T3E1 cells, a murine pre-osteoblast cell line) acetate increases β-catenin protein level together with Runx2 and accumulated nuclear β -catenin in a dose-dependent manner [1]. KY-02327 (1-10 μM) acetate increases the mRNA levels of osteoblast differentiation markers collagen 1a (Col1a) and

	osteocalcin (OCN) $^{[1]}$. MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	KY-02327 (20 mg/kg; p.o.; 5 sequential days per week for 4 weeks) acetate successfully rescues bone loss in the ovariectomized (OVX) mouse model ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Kim HY, et al. Small molecule inhibitors of the Dishevelled-CXXC5 interaction are new drug candidates for bone anabolic osteoporosis therapy. EMBO Mol Med. 2016;8(4):375-387.

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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