Proteins

Product Data Sheet

KY-02327

Cat. No.: HY-124156 CAS No.: 2093407-25-9 Molecular Formula: $C_{20}H_{27}N_{3}O_{4}$ Molecular Weight: 373.45 Target: Wnt

Pathway: Stem Cell/Wnt

Powder -20°C Storage: 3 years

2 years

In solvent -80°C 6 months

> -20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (267.77 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.6777 mL	13.3887 mL	26.7773 mL
	5 mM	0.5355 mL	2.6777 mL	5.3555 mL
	10 mM	0.2678 mL	1.3389 mL	2.6777 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: ≥ 4.24 mg/mL (11.35 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.69 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.69 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

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

In Vitro

KY-02327 (1-10 μ M; 2 days; MC3T3E1 cells, a murine pre-osteoblast cell line) increases β -catenin protein level together with Runx2 and accumulated nuclear β -catenin in a dose-dependent manner^[1].

KY-02327 (1-10 μM) 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) 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.		
	Animal Model:	8-week-old female BL6 mice (ovariectomized (OVX)-induced osteoporosis model mice) $^{[1]}$	
	Dosage:	20 mg/kg	
	Administration:	P.o.; administered for 5 sequential days per week for 4 weeks	
	Result:	Newly formed bones which labeled with calcein were decreased in the femur of vehicle treated OVX mice.	

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.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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