CP21R7

Cat. No.:	HY-100207		
CAS No.:	125314-13-	8	
Molecular Formula:	$C_{19}H_{15}N_{3}O_{2}$		
Molecular Weight:	317.34		
Target:	GSK-3		
Pathway:	PI3K/Akt/m	TOR; Ste	m Cell/Wnt
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year

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SOLVENT & SOLUBILITY

In Vitro	DMSO : ≥ 32 mg/mL (100.84 mM) * "≥" means soluble, but saturation unknown.						
Preparing Stock Solutions		Solvent Mass Concentration	1 mg	5 mg	10 mg		
	1 mM	3.1512 mL	15.7560 mL	31.5119 mL			
		5 mM	0.6302 mL	3.1512 mL	6.3024 mL		
		10 mM	0.3151 mL	1.5756 mL	3.1512 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 (7.88 mM); Clear solution						

BIOLOGICAL ACTIVITY				
Description	CP21R7 is potent GSK-3β inhibitor, with an IC ₅₀ of 1.8 nM; CP21R7 also shows inhibitory activitiy against PKCα, with an IC ₅₀ of 1900 nM.			
IC ₅₀ & Target	GSK-3β 1.8 nM (IC ₅₀)	ΡΚCα 1900 nM (IC ₅₀)		
In Vitro	CP21R7 (Compound 9) is a selective inhibitor of GSK-3β, with an IC ₅₀ of 1.8 nM; the IC ₅₀ of CP21R7 against PKCα is 1900 nM ^[1] . CP21R7 (CP21, 3 μM) potently activates canonical Wnt signaling with highest activity. CP21 significantly increases total levels of intracellular β-catenin. CP21 combined with BMP4 induces commitment of hPSCs towards mesoderm ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

Product Data Sheet

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NH₂

CUSTOMER VALIDATION

- Small. 2023 Jan 12;e2207194.
- J Mol Cell Cardiol. 2023 Jan 16;176:21-32.
- J Mol Cell Cardiol. 2021 Oct 16;S0022-2828(21)00197-8.
- STAR Protoc. 2022 Apr 13;3(2):101296.

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REFERENCES

[1]. Gong L, et al. Discovery of potent and bioavailable GSK-3beta inhibitors. Bioorg Med Chem Lett. 2010 Mar 1;20(5):1693-6.

[2]. Patsch C, et al. Generation of vascular endothelial and smooth muscle cells from human pluripotent stem cells. Nat Cell Biol. 2015 Aug;17(8):994-1003.

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

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