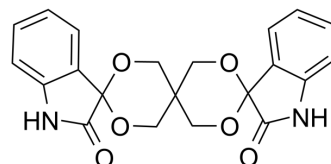


JW67

Cat. No.:	HY-108442		
CAS No.:	442644-28-2		
Molecular Formula:	C ₂₁ H ₁₈ N ₂ O ₆		
Molecular Weight:	394		
Target:	Wnt		
Pathway:	Stem Cell/Wnt		
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 : 50 mg/mL (126.90 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.5381 mL	12.6904 mL	25.3807 mL
		5 mM	0.5076 mL	2.5381 mL	5.0761 mL
10 mM		0.2538 mL	1.2690 mL	2.5381 mL	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 2.5 mg/mL (6.35 mM); Suspended solution; Need ultrasonic Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.35 mM); Clear solution 				

BIOLOGICAL ACTIVITY

Description	JW67 inhibits the canonical Wnt signaling with an IC ₅₀ of 1.17 μM ^[1] . JW67 affects the multiprotein complex consisting of β-catenin/GSK-3β/AXIN/APC/CK1 that rapidly reduces active β-catenin with a subsequent downregulation of Wnt target genes. JW67 also inhibits colorectal cancer cell growth ^[1] .
IC ₅₀ & Target	IC ₅₀ : 1.17 μM (Wnt signaling) ^[1]
In Vitro	<p>JW67 (1 μM; 24 hours) increases concentration of AXIN2 and decreases the active form of β-catenin^[1].</p> <p>JW67 affects the expression of Wnt target genes and reduces cell growth of colon cancer cell lines^[1].</p> <p>JW67 shows a concentration dependent reduction in proliferation with a GI₅₀ value of 7.8 μM^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>

Western Blot Analysis^[1]

Cell Line:	SW480 cells
Concentration:	1 μ M
Incubation Time:	24 hours
Result:	Increased the AXIN2 expression.

RT-PCR^[1]

Cell Line:	SW480 cells
Concentration:	10 or 25 μ M
Incubation Time:	72 hours
Result:	Reduced growth of SW480 CRC cells in vitro by inhibiting cell-cycle progression at the G1/S.

REFERENCES

[1]. Waaler J, et al. Novel synthetic antagonists of canonical Wnt signaling inhibit colorectal cancer cell growth. *Cancer Res.* 2011;71(1):197-205.

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