Miclxin

Cat. No.:	HY-138301		
CAS No.:	2494198-61-5		
Molecular Formula:	C ₂₆ H ₂₇ N ₅ O ₂		
Molecular Weight:	441.52		
Target:	β-catenin		
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

Preparing Stock Solutio		Solvent Mass Concentration	1 mg	5 mg	10 mg	
	Preparing Stock Solutions	1 mM	2.2649 mL	11.3245 mL	22.6490 mL	
		5 mM	0.4530 mL	2.2649 mL	4.5298 mL	
		10 mM	0.2265 mL	1.1325 mL	2.2649 mL	
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	 Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 1 mg/mL (2.26 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1 mg/mL (2.26 mM); Clear solution 					

DIOLOGICAL ACTIV	
Description	Miclxin (DS37262926) is a potent inhibitor of mutant β-catenin, involving in Wnt signaling pathway. Miclxin induces β- catenin-dependent apoptosis, leads to severe mitochondrial damage with the loss of mitochondrial membrane. Miclxin kills tumor via targeting to MIC60, a major components of the mitochondrial contact site and cristae organizing system (MICOS) complex ^[1] .
In Vitro	Miclxin (0-15 μM; 48 h) inhibits the growth of β-catenin-mutated HCT116 cells and isogenic HCT116 (CTNNB1 Δ45/-) cells, and induces (10 μM; 24 h) apoptosis in HCT116 cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay ^[1]

Product Data Sheet

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Cell Line:	HCT116 (CTNNB1 +/–) and (CTNNB1 △45/–)
Concentration:	0, 5, 8, 10, 12, and 15 μM
Incubation Time:	48 hours
Result:	Inhibited cell growth with inhibition rate of 100% at 15 $\mu\text{M}.$

REFERENCES

[1]. Ikeda H, et al. Miclxin, a Novel MIC60 Inhibitor, Induces Apoptosis via Mitochondrial Stress in β-Catenin Mutant Tumor Cells. ACS Chem Biol. 2020 Aug 21;15(8):2195-2204.

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

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