IMM-01

Cat. No.: CAS No.: Molecular Formula: Molecular Weight: Target: Pathway: Storage:	HY-124808 218795-74-5 C ₁₂ H ₁₇ N ₃ O ₂ S 267.35 Apoptosis Apoptosis Please store the product under the recommended conditions in the Certificate of Analysis.	N H H H H H H
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BIOLOGICAL ACTIVI			
Description	IMM-01 is a formin agonist that inhibits DID-DAD (diaphanous inhibitory domain-diaphanous autoregulatory domain) binding with an IC ₅₀ 140 nM. IMM-01 acts by disrupting the autoinhibitory bond between the DID and DAD domain and thus activates formins. IMM-01 shows anticancer effects ^[1] .		
In Vitro	IMM-01 (100 μM; 1h) induces microtubule stabilization in SW480 colon carcinoma cells ^[1] . IMM-01 (10 μM) significantly induces LacZ expression in NIH 3T3-SRE-LacZ cells ^[1] . IMM-01 (1-100 μM; 48 hours) induces caspase-3 cleavage during induction of apoptosis in NIH 3T3 cells and SW480 cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Apoptosis Analysis ^[1]		
	Cell Line:	NIH 3T3 cells and SW480 cells	
	Concentration:	1 μM, 10 μM, or 100 μM	
	Incubation Time:	48 hours	
	Result:	Induced caspase-3 cleavage.	
In Vivo	IMM-01 (5-25 mg/kg; i.v.; 2 times a week; for 3 weeks) is able to slow tumor growth in a mouse xenograft model of colon cancer ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Athymic nude female mice (6 to 8 weeks) implanted with SW480 $cells^{[1]}$	
	Dosage:	5 mg/kg, 25 mg/kg	
	Administration:	i.v.; 2 times a week; for 3 weeks	
	Result:	Slowed tumor growth in a dose-dependent manner when administered intravenously via the tail vein.	

REFERENCES

Product Data Sheet



[1]. L Leanne Lash, et al. Small-molecule intramimics of formin autoinhibition: a new strategy to target the cytoskeletal remodeling machinery in cancer cells. Cancer Res. 2013 Nov 15;73(22):6793-803.

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

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