(Rac)-Indoximod

Cat. No.:	HY-133897	
CAS No.:	26988-72-7	
Molecular Formula:	$C_{12}H_{14}N_2O_2$	
Molecular Weight:	218.25	
Target:	Indoleamine 2,3-Dioxygenase (IDO); Apoptosis	
Pathway:	Metabolic Enzyme/Protease; Apoptosis	
Storage:	4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)	

SOLVENT & SOLUBILITY

		Solvent Mass Concentration	1 mg	5 mg	10 mg			
	Preparing Stock Solutions	1 mM	4.5819 mL	22.9095 mL	45.8190 mL			
		5 mM	0.9164 mL	4.5819 mL	9.1638 mL			
		10 mM	0.4582 mL	2.2910 mL	4.5819 mL			

BIOLOGICAL ACTIVITY					
Description	(Rac)-Indoximod (1-Methyl-DL-tryptophan) is an indoleamine 2,3-dioxygenase (IDO) inhibitor. Co-treatment with IFN-γ and (Rac)-Indoximod markedly reduces the activity of human cardiac myofibroblasts (hCMs) expressing α-SMA and induces apoptosis through up-regulating the IRF-1, Fas, and FasL genes ^[1] .				
IC ₅₀ & Target	IDO				
In Vitro	(Rac)-Indoximod (1-Methyl-DL-tryptophan; 0.5 mM; 3 days) inhibits tryptophan depletion, thereby partially reversing the growth-inhibitory activity of IFN-γ, but eventually induces cell death regardless of tryptophan depletion ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay ^[1]				
	Cell Line:	Human cardiac myofibroblasts (hCMs)			
	Concentration:	0.5 mM			
	Incubation Time:	3 days			



Product Data Sheet

Result:

Growth retardation by IFN- γ was partially reversed on day 2, but cell viability was further reduced on day 3.

CUSTOMER VALIDATION

• Research Square Preprint. 2023 May 22.

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REFERENCES

[1]. Jun-Won Lee, et al. Co-treatment with interferon-γ and 1-methyl tryptophan ameliorates cardiac fibrosis through cardiac myofibroblasts apoptosis. Mol Cell Biochem. 2019 Aug; 458(1-2):197-205.

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

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