

Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

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Lieferung & Zahlungsart siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
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Rapamycin

Cat. No.:	HY-10219				
CAS No.:	53123-88-9			үн ү	
Molecular Formula:	C ₅₁ H ₇₉ NO ₁₃		-0,		
Molecular Weight:	914.17		HO		
Target:	mTOR; FKB	P; Autop			
Pathway:	PI3K/Akt/m Enzyme/Pr	· ·	O HO		
Storage:	Powder	-20°C	3 years		
		4°C	2 years		
	In solvent	-80°C	6 months		
		-20°C	1 month		

SOLVENT & SOLUBILITY

	Solvent Mass Concentration	1 mg	5 mg	10 mg		
Preparing Stock Solutions	1 mM	1.0939 mL	5.4694 mL	10.9389 mL		
	5 mM	0.2188 mL	1.0939 mL	2.1878 mL		
	10 mM	0.1094 mL	0.5469 mL	1.0939 mL		
2. Add each solvent	 Solubility: ≥ 2.5 mg/mL (2.73 mM); Suspended solution 2. Add each solvent one by one: 10% EtOH >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (2.73 mM); Suspended solution; Need ultrasonic 3. Add each solvent one by one: 10% EtOH >> 90% corn oil Solubility: ≥ 2.5 mg/mL (2.73 mM); Suspended solution 					
3. Add each solvent	one by one: 10% EtOH >> 90% corn					
3. Add each solvent Solubility: ≥ 2.5 m 4. Add each solvent	one by one: 10% EtOH >> 90% corn	ı) >> 45% saline			
3. Add each solvent Solubility: ≥ 2.5 m 4. Add each solvent Solubility: ≥ 2.08 r 5. Add each solvent	one by one: 10% EtOH >> 90% corn g/mL (2.73 mM); Suspended solutior one by one: 10% DMSO >> 40% PEC	n G300 >> 5% Tween-80 % SBE-β-CD in saline)				

RedChemExpress



Product Data Sheet

Description	Rapamycin (Sirolimus; AY 22989) is a potent and specific mTOR inhibitor with an IC ₅₀ of 0.1 nM in HEK293 cells. Rapamycin binds to FKBP12 and specifically acts as an allosteric inhibitor of mTORC1 ^[1] . Rapamycin is an autophagy activator, an immunosuppressant ^[2] .						
IC ₅₀ & Target	mTOR 0.1 nM (IC ₅₀ , in HEK293 cells)	Microbial Metabolite	Autophagy	Human Endogenous Metabolite			
In Vitro	Rapamycin (12.5-100 nM; 24 hours) treatment exerts modest inhibitory effect on lung cancer cell proliferation in a dose- dependent manner in all cell lines (A549, SPC-A-1, 95D and NCI-H446 cells) tested, achieving about 30-40% reduction in cell proliferation at 100 nM vs. ~10% reduction at 12.5 nM ^[3] . Lung cancer cell line 95D cells are exposed to Rapamycin (10 nM, 20 nM) and RP-56976 (1 nM, 10 nM) alone or in combination (Rapamycin 20 nM+ RP-56976 10 nM). After 24 hours exposure to Rapamycin or RP-56976 alone does not significantly alter the level of expression or phosphorylation of ERK1/2, whereas cells treated with the combination of Rapamycin with RP- 56976 exhibit a marked reduction in the phosphorylation levels of ERK1/2 ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay ^[3]						
	Cell Line:	Lung cancer cell lines A549, SPC-A-1, 95D and NCI-H446					
	Concentration:	12.5 nM, 25 nM, 50 nM, 100 nM					
	Incubation Time:	24 hours					
	Result:	Treatment exerted modest inhibitory effect on lung cancer cell proliferation in a dose- dependent manner in all cell lines.					
	Western Blot Analysis ^[3]						
	Cell Line:	95D cells					
	Concentration:	10 nM and 20 nM					
	Incubation Time:	24 hours					
	Result:	Combination treatment with RP-56976 decreased phosphorylation of ERK.					
In Vivo	Rapamycin (2.0 mg/kg; intraperitoneal injection; every other day; 28 days) alone has a moderate inhibitory effect. However, the combination of Metformin and Rapamycin exerts a significantly increased inhibition of tumor growth compared with the control group, the Rapamycin monotherapy group and the Metformin monotherapy group ^[4] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.						
	Animal Model:	24 male nu/nu mice aged 4-5 week old (15-20 g) ^[4]					
	Dosage:	2.0 mg/kg					
	Administration:	Intraperitoneal injection; every other day; 28 days					
	Result:	Had a moderate inhibitory effect in monotherapy group. The combination with Metformin exerted a significantly increased inhibition of tumor growth.					

CUSTOMER VALIDATION

- Nature. 2021 Jun;594(7862):271-276.
- Nature. 2018 Jun;558(7711):540-546.
- Nature. 2016 Dec 1;540(7631):119-123.
- Cell. 2023 Jun 22;186(13):2802-2822.e22.
- Cancer Cell. 2021 Mar 8;39(3):380-393.e8.

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REFERENCES

[1]. Edwards SR, et al. The rapamycin-binding domain of the protein kinase mammalian target of rapamycin is a destabilizing domain. J Biol Chem, 2007, 282(18), 13395-13401.

[2]. Rangaraju S, et al. Rapamycin activates autophagy and improves myelination in explant cultures from neuropathicmice. J Neurosci. 2010 Aug 25;30(34):11388-97.

[3]. Niu H, et al. Rapamycin potentiates cytotoxicity by RP-56976 possibly through downregulation of Survivin in lung cancer cells. J Exp Clin Cancer Res. 2011 Mar 10;30:28.

[4]. Zhang JW, et al. Metformin synergizes with rapamycin to inhibit the growth of pancreatic cancer in vitro and in vivo. Oncol Lett. 2018 Feb;15(2):1811-1816.

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

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