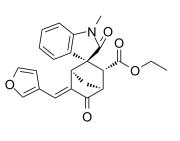
RedChemExpress

PPI-GIT1/ β -Pix interaction-IN-1

Product Data Sheet

Cat. No.:	HY-152095	
CAS No.:	2070916-70-8	
Molecular Formula:	C ₂₃ H ₂₁ NO ₅	
Molecular Weight:	391.42	
Target:	Ras	
Pathway:	GPCR/G Protein; MAPK/ERK Pathway	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	



BIOLOGICAL AC				
Description	PPI-GIT1/β-Pix interaction-IN-1 is a potent and orally active GIT1/β-Pix protein-protein interaction (PPI) inhibitor with a K _D value of 7.7 μM. PPI-GIT1/β-Pix interaction-IN-1 disrupts the GIT/PIX interaction can impact the activation of the downstream Rho GTPase Rac1 and Cdc42. PPI-GIT1/β-Pix interaction-IN-1 inhibits metastasis of gastric cancer ^[1] .			
In Vitro	PPI-GIT1/β-Pix interacti dependent manner and Cdc42 ^[1] . MCE has not independe	 PPI-GIT1/β-Pix interaction-IN-1 (compound 14-5-18; 0-50 μM) inhibits the interaction between GIT1 and β-Pix in living cells^[1] PPI-GIT1/β-Pix interaction-IN-1 (0-50 μM; 24 h; MGC803 cells and MKN45 cells) inhibits gastric cancer cell invasion in a dose-dependent manner and disrupts the GIT/PIX interaction can impact the activation of the downstream Rho GTPase Rac1 and Cdc42^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis^[1] 		
	Cell Line:	MGC803 cells and MKN45 cells		
	Concentration:	0, 5, 20, and 50 μM		
	Incubation Time:	24 hours		
	Result:	Suppressed the expression of GTP-Rac1 and GTP-Cdc42 in a dose-dependent manner.		
In Vivo	nude mice with MGC803	PPI-GIT1/β-Pix interaction-IN-1 (compound 14-5-18; 10 and 30 mg/kg; i.g.; 24 h) inhibits gastric cancer cell invasion in female nude mice with MGC803 xenografts ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Female nude mice with MGC803 xenografts (Four-week-old) $^{[1]}$		
	Dosage:	10 and 30 mg/kg		
	Administration:	oral gavage, daily, for 18 days		
	Result:	Reduced the luminescence intensity in the lungs in a dose-dependent manner.		

REFERENCES

[1]. Gu J, et, al. Construction of a synthetic methodology-based library and its application in identifying a GIT/PIX protein-protein interaction inhibitor. Nat Commun. 2022 Nov 23;13(1):7176.

[2]. Gu J, et, al. Construction of a synthetic methodology-based library and its application in identifying a GIT/PIX protein-protein interaction inhibitor. Nat Commun. 2022 Nov 23;13(1):7176.

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

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