PI4KIIIbeta-IN-11

MedChemExpress

HY-148075	/N
2365241-79-6	N
$C_{_{33}}H_{_{39}}N_7O_3$	
581.71	NH
РІ4К	N-N
PI3K/Akt/mTOR	HOTIN
Please store the product under the recommended conditions in the Certificate of Analysis.	
	2365241-79-6 C ₃₃ H ₃₉ N ₇ O ₃ 581.71 PI4K PI3K/Akt/mTOR Please store the product under the recommended conditions in the Certificate of

BIOLOGICAL ACTIVITY			
Description	PI4KIIIbeta-IN-11 is an inhibitor of PI4KIIIβ, with a mean pIC ₅₀ value of at least 9.1. PI4KIIIβ plays a key role in diseases research of RNA viruses and Plasmodium falciparum ^{[1][2]} .		
IC ₅₀ & Target	ΡΙ4ΚΙΙΙβ 9.1 (pIC ₅₀)		
In Vitro	PI4KIIIβ may be a relevant target protein for inhibition of Hedgehog signaling, and regulates trafficking from the Golgi system to the plasma membrane ^[1] . PI4KIIIbeta-IN-11 (compound 22) (0.5 μM; 0-45 min) shows metabolic stability in human microsomes, with the intrinsic clearance of 33.7 mL/min/g ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
In Vivo	PI4KIIIbeta-IN-11 (compound 22) (1 mg/kg; i.v.; last over 1 h) displays a low spleen concentration in rats ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Male Wistar Han Rats ^[2]	
	Dosage:	1 mg/kg	
	Administration:	Intravenous injection, formulated in 2% DMSO in Kleptose (aq, 10% w/v); over 1 hour; euthanised at 12 h	
	Result:	Showed a low concentration in spleen of 5.54 ng/g.	

REFERENCES

[1]. Kremer L, et al. Discovery of the Hedgehog Pathway Inhibitor Pipinib that Targets PI4KIIIß. Angew Chem Int Ed Engl. 2019 Nov 11;58(46):16617-16628.

[2]. Barton Nicholas Paul, et al. Preparation of pyrazolopyrimidinamines derivatives as PI4kIII-beta inhibitors: World Intellectual Property Organization, WO2019141694. 2019-07-25.

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

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