## HPK1-IN-35

Cat. No.:	HY-149232	N
CAS No.:	2935903-77-6	N
Molecular Formula:	C <sub>30</sub> H <sub>32</sub> N <sub>8</sub> O <sub>3</sub> S	
Molecular Weight:	584.69	
Target:	МАР4К	Ň <sub>Ŷ</sub> Ň
Pathway:	MAPK/ERK Pathway	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

BIOLOGICAL ACTIV			
Description	HPK1-IN-35 is a potent and selective HPK1 inhibitor with an IC <sub>50</sub> value of 3.5 nM. HPK1-IN-35 decreases the expression of p-SLP76 and promotes IL-2 secretion <sup>[1]</sup> .		
IC₅₀ & Target	HPK1 3.5 nM (IC <sub>50</sub> )		
In Vitro	HPK1-IN-35 (compound 31) (0.1, 0.3, 1, 3 μM; 1 h) decreases the expression of p-SLP76 stimulated by anti-CD3 antibody (1 μg/mL) in a dose-dependent manner <sup>[1]</sup> . HPK1-IN-35 (0-10 μM; 24 h) promotes IL-2 secretion in Jurkat cells in a dose-dependent manner with EC50 values of 1.19 μM <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis <sup>[1]</sup>		
	Cell Line:	Jurkat cells	
	Concentration:	0.1, 0.3, 1, 3 μΜ	
	Incubation Time:	1 h	
	Result:	Dose-dependently suppressed the phosphorylation of SLP76 stimulated by anti-CD3 antibody in Jurkat cells with IC50 values of 1.04 $\mu M.$	

## REFERENCES

[1]. Feifei Wu, et al. Discovery of 7H-Pyrrolo[2,3-d] pyrimidine Derivatives as potent hematopoietic progenitor kinase 1 (HPK1) inhibitors. European Journal of Medicinal Chemistry. 2023, 254: 115355.

## **Product** Data Sheet



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

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