## Multi-kinase-IN-6

Cat. No.:	HY-156470
CAS No.:	3009081-73-3
Molecular Formula:	C <sub>17</sub> H <sub>15</sub> N <sub>5</sub> O <sub>2</sub>
Molecular Weight:	321.33
Target:	Trk Receptor; Anaplastic lymphoma kinase (ALK); c-Kit; EGFR; Pim; Casein Kinase; N-O Checkpoint Kinase (Chk); CDK; Apoptosis
Pathway:	Neuronal Signaling; Protein Tyrosine Kinase/RTK; JAK/STAT Signaling; Cell Cycle/DNA Damage; Stem Cell/Wnt; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

BIOLOGICAL ACTIVITY					
Description	Multi-kinase-IN-6 (compound 10e) is a multikinase inhibitor that shows good enzyme inhibitory activity against TrkA, ALK2, c-KIT, EGFR, PIM1, CK2α, CHK1, and CDK2. Multi-kinase-IN-6 reveals antiproliferative activity against MCF7, HCT116 and EKVX with IC <sub>50</sub> values of 3.36 μM, 1.40 μM and 3.49 μM, respectively. Multi-kinase-IN-6 shows cell cycle arrest at the G1/S phase and G1 phase in MCF7 and HCT116 cells with good apoptotic effect <sup>[1]</sup> .				
IC₅o & Target	TrkA 0.33 μΜ (IC <sub>50</sub> )	CDK2 0.71 μΜ (IC <sub>50</sub> )	CK2 0.09 μΜ (IC <sub>50</sub> )	ΡΙΜ1 0.42 μΜ (IC <sub>50</sub> )	
	Chk1 0.15 μΜ (IC <sub>50</sub> )	EGFR 0.1 μΜ (IC <sub>50</sub> )	c-KIT 0.05 μΜ (IC <sub>50</sub> )	ALK2 0.03 μΜ (IC <sub>50</sub> )	
	RET 0.53 μΜ (IC <sub>50</sub> )	FLT3 0.34 μΜ (IC <sub>50</sub> )	ROS1 0.75 μΜ (IC <sub>50</sub> )		

## REFERENCES

[1]. Mustafa A. Al-Qadhi, et al. Design and synthesis of certain 7-Aryl-2-Methyl-3-Substituted Pyrazolo{1,5-a}Pyrimidines as multikinase inhibitors. European Journal of Medicinal Chemistry. Volume 262, 15 December 2023, 115918.

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

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**Product** Data Sheet



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