## CDK7/9-IN-1

Cat. No.: CAS No.: Molecular Formula: Molecular Weight: Target: Pathway: Storage:	HY-145408 2747919-19-1 C <sub>24</sub> H <sub>32</sub> F <sub>3</sub> N <sub>5</sub> O <sub>2</sub> 479.54 CDK Cell Cycle/DNA Damage Please store the product under the recommended conditions in the Certificate of Analysis.	HN HN F F F F
	Analysis.	

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Description	CDK7/9-IN-1 is a cyclin-dependent kinases 7/9 (CDK7/9) inhibitor. CDK7/9-IN-1 selectively inhibits CDK7 over CDK9. CDK7/9-IN-1 inhibits CDK7 with IC <sub>50</sub> s of 0.0656 μM and 0.00574 μM without pre-incubation and after 3 hours pre-incubation, respectively. CDK7/9-IN-1 inhibits CDK9 with an IC <sub>50</sub> of 2.14 μM after 3 hours pre-incubation. CDK7/9-IN-1 can be used for the research of cancer <sup>[1]</sup> .		
IC <sub>50</sub> & Target	CDK7 5.74-65.6 nM (IC <sub>50</sub> )	CDK9 2.14 µM (IC <sub>50</sub> )	
In Vitro	CDK7/9-IN-1 (example 8; 0.001-20 μM; 4 h) inhibits CDK7 with an relative IC <sub>50</sub> of 0.0262 μM for phospho-carboxyl terminal domain (Rbp2) (Ser5) in HCT116 cells <sup>[1]</sup> . CDK7/9-IN-1 (0.001-20 μM; 4 h) inhibits CDK9 with an IC <sub>50</sub> of 2.59 μM for phospho-carboxyl terminal domain (Rbp2) (Ser2) in HCT116 cells <sup>[1]</sup> . CDK7/9-IN-1 (0.001-20 μM; 4 h) inhibits cMyc with an relative IC <sub>50</sub> of 0.0138 μM in HCT116 cells <sup>[1]</sup> . CDK7/9-IN-1 (0.001-20 μM; 4 h) inhibits cMyc with an relative IC <sub>50</sub> of 0.0138 μM in HCT116 cells <sup>[1]</sup> . CDK7/9-IN-1 (20 μM, 2 μM, and 0.2 μM; 1 h) shows excellent selectivity against the 468 protein kinases panel. CDK7/9-IN-1 shows approximately 96% inhibition against CDK7 <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

## REFERENCES

[1]. Maria Carmen FERNANDEZ FIGUEROA, et al. Compounds useful for inhibiting cdk7. WO2021242602 A1.

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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