

## PROTAC CDK9 degrader-6

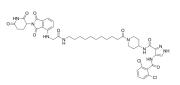
Cat. No.: HY-149963 CAS No.: 2935587-91-8 Molecular Formula:  $C_{42}H_{49}Cl_{2}N_{9}O_{8}$ 

Molecular Weight: 878.8 CDK Target:

Pathway: Cell Cycle/DNA Damage

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.



## **BIOLOGICAL ACTIVITY**

Description	PROTAC CDK9 degrader-6 is a PROTAC targeting to CDK9 sepcifically. PROTAC CDK9 degrader-6 mediates CDK9 degradation
	via the proteasome. PROTAC CDK9 degrader-6 degrades CDK9 with DC $_{50}$ s of 0.10 $\mu$ M and 0.14 $\mu$ M for the CDK9 $_{42}$ and CDK9 $_{55}$
	isoforms, respectively $^{[1]}$ .

IC<sub>50</sub> & Target CDK9<sub>42</sub> CDK9<sub>55</sub>

0.03 μM (DC50) 0.05 μM (DC50)

PROTAC CDK9 degrader-6 (compound 16) (1  $\mu$ M; 6 h) decreases the protein level of MCL2, and competely degrades CDK9 in In Vitro MV411 cells<sup>[1]</sup>.

> PROTAC CDK9 degrader-6 (1 µM; 1-6 h) time-dependently decreases the protein level of MCL2, and CDK9, and remaines suppression for  $24 h^{[1]}$ .

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Western Blot Analysis<sup>[1]</sup>

Cell Line:	MV411 cells
Concentration:	1 μΜ
Incubation Time:	1 h, 2 h, 4 h, 6 h
Result:	Degraded CDK9 starting at 2 h incubation and reaching a plateau at 4 h.  Remained suppressed for 24 h with some recurrence at 48 h after finished treatment.

## **REFERENCES**

[1]. Tokarski RJ 2nd, et al. Bifunctional degraders of cyclin dependent kinase 9 (CDK9): Probing the relationship between linker length, properties, and selective protein degradation. Eur J Med Chem. 2023 Jun 5;254:115342.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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