Proteins

Inhibitors

MS33

Cat. No.: HY-141797 CAS No.: 2407449-11-8 Molecular Formula: $C_{64}H_{84}F_3N_{11}O_7S$

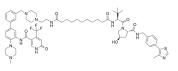
Molecular Weight: 1208.48

Target: Histone Methyltransferase; PROTACs

Pathway: Epigenetics; PROTAC

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

Analysis.



Product Data Sheet

BIOLOGICAL ACTIVITY

Description	MS33 is a potent WDR5 degrader, with K_ds of 870 nM and 120 nM for VCB and WDR5, respectively. MS33 induces WDR5 degradation in an E3 ligase VHL, and proteasome-dependent manner. MS33 can be used for the research of acute myeloid leukemia $^{[1][2][3]}$.		
IC ₅₀ & Target	VHL	WDR5 120 nM (Kd)	VCB 870 nM (Kd)
In Vitro	MS33 (0.05-5 μ M; 18 h) induces WDR5 degradation in a concentration-dependent manner in MV4;11 cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis ^[1]		
	Cell Line:	MV4;11 cells	
	Concentration:	0.05, 0.1, 0.5, 1, 5 μΜ	
	Incubation Time:	1, 2, 4, 8, 16, 24 hours	
	Result:	Concentration- and time-dependently induced WDR5 degradation. The DC $_{50}$ of MS33 was 260 nM, and the maximum degradation of 71%.	

REFERENCES

[1]. Yu X, et, al. A selective WDR5 degrader inhibits acute myeloid leukemia in patient-derived mouse models. Sci Transl Med. 2021 Sep 29;13(613):eabj1578.

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

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