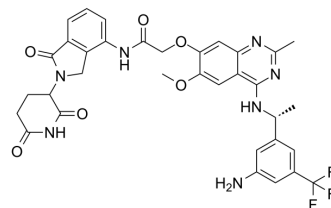


## PROTAC SOS1 degrader-3

<b>Cat. No.:</b>	HY-152145
<b>CAS No.:</b>	3029320-99-5
<b>Molecular Formula:</b>	C <sub>34</sub> H <sub>32</sub> F <sub>3</sub> N <sub>7</sub> O <sub>6</sub>
<b>Molecular Weight:</b>	691.66
<b>Target:</b>	PROTACs; Ras
<b>Pathway:</b>	PROTAC; GPCR/G Protein; MAPK/ERK Pathway
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	PROTAC SOS1 degrader-3 is a potent PROTAC SOS1 degrader. PROTAC SOS1 degrader-3 effectively targeted SOS1 for degradation through the ubiquitin-proteasome system <sup>[1]</sup> .									
<b>IC<sub>50</sub> &amp; Target</b>	Cereblon	SOS1								
<b>In Vitro</b>	<p>PROTAC SOS1 degrader-3 (P7; 0.1-10 μM; 6 hours) induces SOS1 degradation in SW620 cells. In addition to SW620, PROTAC SOS1 degrader-3 also induces SOS1 degradation in other colorectal cancer (CRC) cell lines HCT116, C2BB, and SW1417 in a concentration-dependent manner. The half maximal degradation concentration (DC<sub>50</sub>) for SW620, HCT116, and SW1417 at 24 h is 0.59 μM, 0.75 μM, and 0.19 μM, respectively<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>SW620 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.1 μM, 1 μM, 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>6 hours</td> </tr> <tr> <td>Result:</td> <td>Induced SOS1 degradation.</td> </tr> </table>		Cell Line:	SW620 cells	Concentration:	0.1 μM, 1 μM, 10 μM	Incubation Time:	6 hours	Result:	Induced SOS1 degradation.
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### REFERENCES

[1]. Yujia Bian, et al. Development of SOS1 Inhibitor-Based Degraders to Target KRAS-Mutant Colorectal Cancer. *J Med Chem.* 2022 Dec 22;65(24):16432-16450.

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

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