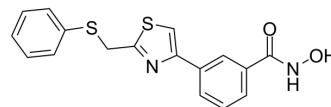


## HDAC8-IN-4

<b>Cat. No.:</b>	HY-153583
<b>CAS No.:</b>	1600528-05-9
<b>Molecular Formula:</b>	C <sub>17</sub> H <sub>14</sub> N <sub>2</sub> O <sub>2</sub> S <sub>2</sub>
<b>Molecular Weight:</b>	342.44
<b>Target:</b>	HDAC
<b>Pathway:</b>	Cell Cycle/DNA Damage; Epigenetics
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	HDAC8-IN-4 is a selective inhibitor of HDAC8. HDAC8-IN-4 inhibits HDAC8 and HDAC3 with IC <sub>50</sub> s of 0.15 and 12 μM <sup>[1]</sup> .																	
<b>IC<sub>50</sub> &amp; Target</b>	HDAC8 0.15 μM (IC <sub>50</sub> )	HDAC3 12 μM (IC <sub>50</sub> )																
<b>In Vitro</b>	<p>HDAC8-IN-4 (Compound 9) (1 to 100 μM, 5-6 days) inhibits HDAC8 protein level in HeLa cells<sup>[1]</sup>.          HDAC8-IN-4 (0-50 μM approximately, 72 h) inhibits T-cell lymphoma cells (Jurkat, HH, MT4, HUT78 cells) growth<sup>[1]</sup>.          MCE has not independently confirmed the accuracy of these methods. They are for reference only.          Cell Proliferation Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>Jurkat, HH, MT4, HUT78 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-50 μM approximately</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited cell growth with IC<sub>50</sub>s of 2, 7.4, 5.8, 27 μM).</td> </tr> </table> <p>Western Blot Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>HeLa cells</td> </tr> <tr> <td>Concentration:</td> <td>0-40 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>4 h</td> </tr> <tr> <td>Result:</td> <td>Selectively inhibited HDAC8 over other HDACs in cells.</td> </tr> </table>		Cell Line:	Jurkat, HH, MT4, HUT78 cells	Concentration:	0-50 μM approximately	Incubation Time:	72 h	Result:	Inhibited cell growth with IC <sub>50</sub> s of 2, 7.4, 5.8, 27 μM).	Cell Line:	HeLa cells	Concentration:	0-40 μM	Incubation Time:	4 h	Result:	Selectively inhibited HDAC8 over other HDACs in cells.
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### REFERENCES

[1]. Suzuki T, et al. Design, synthesis, and biological activity of NCC149 derivatives as histone deacetylase 8-selective inhibitors. ChemMedChem. 2014 Mar;9(3):657-64.

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

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