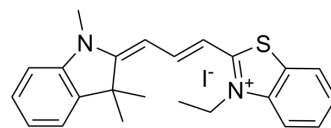


## AC-93253

<b>Cat. No.:</b>	HY-118343
<b>CAS No.:</b>	108527-83-9
<b>Molecular Formula:</b>	C <sub>23</sub> H <sub>25</sub> IN <sub>2</sub> S
<b>Molecular Weight:</b>	488.43
<b>Target:</b>	Sirtuin
<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>	AC-93253 is a selective, potent SIRT2 inhibitor. AC93253 can inhibit SIRT2 with an IC <sub>50</sub> value of 6 μM. AC93253 can be used for the research of tumors <sup>[1]</sup> .										
<b>IC<sub>50</sub> &amp; Target</b>	SIRT1 45.3 μM (IC <sub>50</sub> )	SIRT2 6 μM (IC <sub>50</sub> )	SIRT3 24.6 μM (IC <sub>50</sub> )								
<b>In Vitro</b>	<p>AC93253 can inhibit SIRT1, SIRT2 and SIRT3 with IC<sub>50</sub> values of 45.3 μM, 6 μM and 24.6 μM, respectively<sup>[1]</sup>.            AC-93253 (0, 2, 5, 10 μM; 16 h) significantly enhanced the acetylation of tubulin, p53, and histone H4<sup>[1]</sup>.            AC-93253 exhibits selective cytotoxicity towards four tumor cell lines in a single agent with IC<sub>50</sub> values ranging from 10 to 100 nM<sup>[1]</sup>.            AC-93253 significantly triggered apoptosis<sup>[1]</sup>.            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>NCI-H460 cells, HeLa cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 2, 5, 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>16 h</td> </tr> <tr> <td>Result:</td> <td>Increased the acetylation levels of α-tubulin in a dose-dependent manner. Increased the level of histone protein and p53.</td> </tr> </table>			Cell Line:	NCI-H460 cells, HeLa cells	Concentration:	0, 2, 5, 10 μM	Incubation Time:	16 h	Result:	Increased the acetylation levels of α-tubulin in a dose-dependent manner. Increased the level of histone protein and p53.
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Concentration:	0, 2, 5, 10 μM										
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Result:	Increased the acetylation levels of α-tubulin in a dose-dependent manner. Increased the level of histone protein and p53.										

### REFERENCES

[1]. Zhang, Yingjia et al. Identification of a small molecule SIRT2 inhibitor with selective tumor cytotoxicity. Biochemical and biophysical research communications vol. 386,4 (2009): 729-33.

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

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