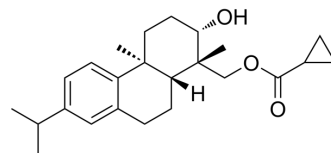


## Antitumor agent-97

<b>Cat. No.:</b>	HY-149829
<b>CAS No.:</b>	2654024-16-3
<b>Molecular Formula:</b>	C <sub>24</sub> H <sub>34</sub> O <sub>3</sub>
<b>Molecular Weight:</b>	370.52
<b>Target:</b>	Apoptosis; Autophagy
<b>Pathway:</b>	Apoptosis; Autophagy
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Antitumor agent-97 (compound 42) is an anticancer agent. Antitumor agent-97 can effectively inhibit the proliferation and autophagy of MGC 803 cells, and induce apoptosis. Antitumor agent-97 also enhances ROS accumulation in MGC 803 cells. Antitumor agent-97 can be used in cancer research <sup>[1]</sup> .																				
<b>In Vitro</b>	<p>Antitumor agent-97 (Compound 42; 0-200 μM; 24 h) inhibits cell viability of MGC 803 cells in a dose-dependent manner, with an IC<sub>50</sub> value of 20.921 μM<sup>[1]</sup>.</p> <p>Antitumor agent-97 (20 μM; 24 h) decreases cell proliferation of MGC 803 cells<sup>[1]</sup>.</p> <p>Antitumor agent-97 (20 μM; 24 h) induces apoptosis and inhibits autophagy in MGC 803 cells<sup>[1]</sup>.</p> <p>Antitumor agent-97 (20 μM; 0-24 h) enhances ROS accumulation in MGC 803 cells<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>MGC 803 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-200 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Suppressed cell viability (IC<sub>50</sub> = 20.921 μM).</td> </tr> </table> <p>Cell Proliferation Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>MGC 803 cells</td> </tr> <tr> <td>Concentration:</td> <td>20 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited cell proliferation.</td> </tr> </table> <p>Western Blot Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>MGC 803 cells</td> </tr> <tr> <td>Concentration:</td> <td>20 μM</td> </tr> </table>	Cell Line:	MGC 803 cells	Concentration:	0-200 μM	Incubation Time:	24 h	Result:	Suppressed cell viability (IC <sub>50</sub> = 20.921 μM).	Cell Line:	MGC 803 cells	Concentration:	20 μM	Incubation Time:	24 h	Result:	Inhibited cell proliferation.	Cell Line:	MGC 803 cells	Concentration:	20 μM
Cell Line:	MGC 803 cells																				
Concentration:	0-200 μM																				
Incubation Time:	24 h																				
Result:	Suppressed cell viability (IC <sub>50</sub> = 20.921 μM).																				
Cell Line:	MGC 803 cells																				
Concentration:	20 μM																				
Incubation Time:	24 h																				
Result:	Inhibited cell proliferation.																				
Cell Line:	MGC 803 cells																				
Concentration:	20 μM																				

Incubation Time:	24 h
Result:	Significantly upregulated the protein expression of LC3B-II and p62 and decreased the protein expression of LC3B-I. Enhanced the protein expression of cleaved caspase-9 and cleaved caspase-3, and decreased the protein expression of B-cell lymphoma-2 (Bcl-2).

## REFERENCES

[1]. Mao XD, et al. Synthesis and Bioactivity Evaluation of Nepetaefolin F and Its Analogues. ACS Omega. 2023 Apr 12;8(16):14830-14840.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA