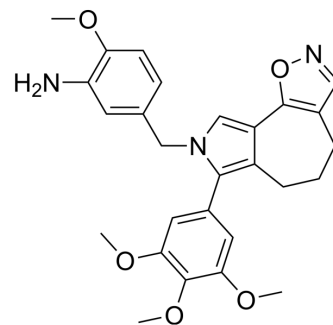


## Apoptosis inducer 12

Cat. No.:	HY-149222
CAS No.:	1971096-01-1
Molecular Formula:	C <sub>27</sub> H <sub>29</sub> N <sub>3</sub> O <sub>5</sub>
Molecular Weight:	475.54
Target:	Apoptosis
Pathway:	Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Apoptosis inducer 12 (Compound 3z) is an apoptosis inducer that induces apoptosis through the mitochondrial pathway. Apoptosis inducer 12 can be used for the research of cancer <sup>[1]</sup> .																
<b>In Vitro</b>	<p>Apoptosis inducer 12 (Compound 3z; 72 h) shows antiproliferative activity against nine human cancer cell types (leukemia, non-small-cell lung, colon, central nervous system, melanoma, ovarian, renal, prostate and breast) with GI<sub>50</sub> of 0.24-96.6 μM. leukemia and prostate cell lines are particularly responsive to treatment with Apoptosis inducer 12, with GI<sub>50</sub> values of 0.30-0.65 μM and 0.43-0.84 μM, respectively<sup>[1]</sup>.</p> <p>Apoptosis inducer 12 (0.5-10 μM; 24 h) blocks the cell cycle in metaphase in A549, CCRF-CEM and VL51 cells<sup>[1]</sup>.</p> <p>Apoptosis inducer 12 (0.5-5 μM; 48 h) induces apoptosis in A549, CCRF-CEM and VL51 cells<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cycle Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>A549, CCRF-CEM and VL51</td> </tr> <tr> <td>Concentration:</td> <td>0.5, 1, 5 and 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Induced a block in G2/M accompanied by a strong decrease in S phase cells.</td> </tr> </table> <p>Apoptosis Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>A549, CCRF-CEM and VL51</td> </tr> <tr> <td>Concentration:</td> <td>0.5, 1 and 5 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Induced massive apoptosis in a dose-dependent manner.</td> </tr> </table>	Cell Line:	A549, CCRF-CEM and VL51	Concentration:	0.5, 1, 5 and 10 μM	Incubation Time:	24 h	Result:	Induced a block in G2/M accompanied by a strong decrease in S phase cells.	Cell Line:	A549, CCRF-CEM and VL51	Concentration:	0.5, 1 and 5 μM	Incubation Time:	48 h	Result:	Induced massive apoptosis in a dose-dependent manner.
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### REFERENCES

[1]. Barreca M, et al. Identification of pyrrolo[3',4':3,4]cyclohepta[1,2-d][1,2]oxazoles as promising new candidates for the treatment of lymphomas. Eur J Med Chem. 2023

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

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