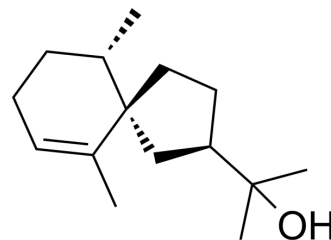


(-)-Hinesol

Cat. No.:	HY-N1930
CAS No.:	23811-08-7
Molecular Formula:	C ₁₅ H ₂₆ O
Molecular Weight:	222.37
Target:	Apoptosis
Pathway:	Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	(-)-Hinesol (Hinesol) is a potent anticancer agent. (-)-Hinesol induces apoptosis and cell cycle arrest at G0/G1 phase. (-)-Hinesol downregulates MEK/ERK pathway and NF-κB pathway and mediates the expression of cyclin D1, Bax and Bcl-2. (-)-Hinesol has the potential for the research of non-small cell lung cancer ^[1] .																
In Vitro	<p>(-)-Hinesol (0-25 μg/ml; 24, 48 h) shows antiproliferative activity of the A549 and NCI-H1299 cells in a dose- and time-dependent manner^[1].</p> <p>(-)-Hinesol (0, 2, 8 μg/ml; 24 h) induces apoptosis and cell cycle arrest at G0/G1 phase with increases the expression of Bax and decreases the expression of Bcl-2 and cyclin D1^[1].</p> <p>(-)-Hinesol (0, 2, 8 μg/ml; 24 h) decreases the expression of phosphor-ERK1/2, phosphor-MEK1/2, phosphor-IκBα and -p65 level, and shows no change for the total protein levels of IκBα and p65 in A549 cells^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay^[1]</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 30%;">Cell Line:</td> <td>A549, NCI-H1299 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-25 μg/ml</td> </tr> <tr> <td>Incubation Time:</td> <td>24, 48 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited the proliferation of the A549 and NCI-H1299 cells in a dose- and time-dependent manner.</td> </tr> </table> <p>Apoptosis Analysis^[1]</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 30%;">Cell Line:</td> <td>A549 cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 2, 8 μg/ml</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Induced apoptosis with the apoptotic cells was increased to 21.2 ± 0.96% and 36 ± 1.04% after treatment with hinesol at 2 and 8 μg/mL, respectively.</td> </tr> </table> <p>Western Blot Analysis^[1]</p>	Cell Line:	A549, NCI-H1299 cells	Concentration:	0-25 μg/ml	Incubation Time:	24, 48 h	Result:	Inhibited the proliferation of the A549 and NCI-H1299 cells in a dose- and time-dependent manner.	Cell Line:	A549 cells	Concentration:	0, 2, 8 μg/ml	Incubation Time:	24 h	Result:	Induced apoptosis with the apoptotic cells was increased to 21.2 ± 0.96% and 36 ± 1.04% after treatment with hinesol at 2 and 8 μg/mL, respectively.
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Cell Line:	A549 cells
Concentration:	0, 2, 8 µg/ml
Incubation Time:	24 h
Result:	Increased the protein expression of Bax and decreased the expression of Bcl-2.

Cell Cycle Analysis^[1]

Cell Line:	A549 cells
Concentration:	0, 2, 8 µg/ml
Incubation Time:	24 h
Result:	Showed concentration-dependent increase in the percentage of cell in the G0/G1 phase, and a decrease of the percentage in G2/M phase.

REFERENCES

[1]. Guo W, et al. The antitumor effect of hinesol, extract from *Atractylodes lancea* (Thunb.) DC. by proliferation, inhibition, and apoptosis induction via MEK/ERK and NF-κB pathway in non-small cell lung cancer cell lines A549 and NCI-H1299. *J Cell Biochem.* 2019 Nov;120(11):18600-18607.

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

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