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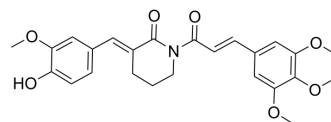
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Apoptosis inducer 17

Cat. No.:	HY-163451
Molecular Formula:	C ₂₅ H ₂₇ NO ₇
Molecular Weight:	453.48
Target:	Apoptosis
Pathway:	Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Apoptosis inducer 17 is a Curcumin-Piperlongumine hybrid molecule that increases cell cycle arrest and apoptosis through activating JNK signaling pathway in lung cancer cells ^[1] .																				
In Vitro	<p>Apoptosis inducer 17 (compound CP; 0.027-60 μM; 72 h) inhibits cell proliferation, with IC₅₀ values of 0.021 μM, 0.027 μM, and 0.73 μM for H446, H1299, and SBC-2 cells, respectively^[1].</p> <p>Apoptosis inducer 17 (compound CP; 0.03-0.3 μM; 24 h) displays superior inhibition of colony formation. Apoptosis inducer 17 induces dose-dependent G2/M arrest in H446 and H1299 cells^[1].</p> <p>Apoptosis inducer 17 (compound CP; 0.03-0.3 μM; 24 h) induces apoptosis of H446 and H1299 cells is mediated by caspase activity. And also activates JNK signaling cascade^[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"> <tr> <td>Cell Line:</td> <td>H446, H1299, and SBC-2 cells</td> </tr> <tr> <td>Concentration:</td> <td>60 μM, 20 μM, 6.67 μM, 2.22 μM, 0.74 μM, 0.25 μM, 0.082 μM, and 0.027 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Effectively inhibited cell proliferation.</td> </tr> </table> <p>Cell Cycle Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>H446, H1299 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.03 μM, 0.1 μM, 0.3 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Induced dose-dependent G2/M arrest.</td> </tr> </table> <p>Apoptosis Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>H446, H1299 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.03 μM, 0.1 μM, 0.3 μM</td> </tr> </table>	Cell Line:	H446, H1299, and SBC-2 cells	Concentration:	60 μM, 20 μM, 6.67 μM, 2.22 μM, 0.74 μM, 0.25 μM, 0.082 μM, and 0.027 μM	Incubation Time:	72 h	Result:	Effectively inhibited cell proliferation.	Cell Line:	H446, H1299 cells	Concentration:	0.03 μM, 0.1 μM, 0.3 μM	Incubation Time:	24 h	Result:	Induced dose-dependent G2/M arrest.	Cell Line:	H446, H1299 cells	Concentration:	0.03 μM, 0.1 μM, 0.3 μM
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	Incubation Time:	24 h
	Result:	Induced apoptosis of H446 and H1299 cells.
	Western Blot Analysis ^[1]	
	Cell Line:	H446, H1299 cells
	Concentration:	0.03 μ M, 0.1 μ M, 0.3 μ M
	Incubation Time:	24 h
	Result:	Led to a dose-dependent increase in the phosphorylation of JNK, c-Jun and activating transcription factor-2 (ATF-2) in both H446 and H1299 cells.
In Vivo	Apoptosis inducer 17 (compound CP; 10 mg/kg; ip; twice a day; for 15 days) shows a significant reduction in H446 cell growth, without any significant changes in body weight ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	BALB/c nude mice (4-6 weeks old) injected with H446 cells ^[1]
	Dosage:	10 mg/kg
	Administration:	ip; twice a day; for 15 days
	Result:	Prevented growth of lung cancer in xenograft tumors.

REFERENCES

[1]. Curcumin-Piperlongumine Hybrid Molecule Increases Cell Cycle Arrest and Apoptosis in Lung Cancer through JNK/c-Jun Signaling Pathway

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

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