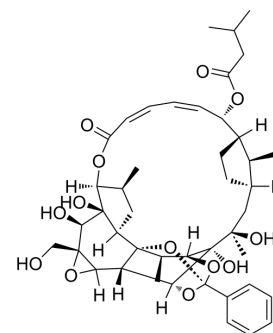


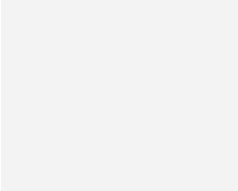
Trigothysoid N

Cat. No.:	HY-135744
CAS No.:	1501943-08-3
Molecular Formula:	C ₄₄ H ₅₈ O ₁₃
Molecular Weight:	794.92
Target:	Apoptosis; STAT; MMP
Pathway:	Apoptosis; JAK/STAT Signaling; Stem Cell/Wnt; Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Trigothysoid N is a daphnane diterpenoid with anticancer activity. Trigothysoid N inhibits tumor proliferation and migration by targeting mitochondria, regulating the STAT3/FAK signal pathway, and suppressing angiogenesis. Trigothysoid N also induce apoptosis, can be used for research of non-small cell lung cancer (NSCLC) ^[1] .								
In Vitro	<p>Trigothysoid N (5, 15, and 45 μM; 48 h) suppresses tumor cell growth in a dose-dependent manner^[1].</p> <p>Trigothysoid N (5, 10, and 20 μM; 48 h) induces apoptosis and arrests the cell cycle at G0/G1 phase^[1].</p> <p>Trigothysoid N (5, 10, and 20 μM; 48 h) induces MMP depolarization, and increased cellular ROS production, to stimulate A549 cell apoptosis^[1].</p> <p>Trigothysoid N induces apoptosis by mitochondria-dependent signaling, and inhibits A549 cell metastasis via regulating FAK signal pathway^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Apoptosis Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>A549 cells</td> </tr> <tr> <td>Concentration:</td> <td>5, 10, and 20 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Resulted the apoptotic cell percentages increased from 9.9% (control) to 12.6% (5.0 μM), 15.4% (10.0 μM), and 59.0% (20.0 μM).</td> </tr> </table>	Cell Line:	A549 cells	Concentration:	5, 10, and 20 μM	Incubation Time:	48 h	Result:	Resulted the apoptotic cell percentages increased from 9.9% (control) to 12.6% (5.0 μM), 15.4% (10.0 μM), and 59.0% (20.0 μM).
Cell Line:	A549 cells								
Concentration:	5, 10, and 20 μM								
Incubation Time:	48 h								
Result:	Resulted the apoptotic cell percentages increased from 9.9% (control) to 12.6% (5.0 μM), 15.4% (10.0 μM), and 59.0% (20.0 μM).								
In Vivo	<p>Trigothysoid N (0.025, 0.05, and 0.1 μM; 4 h) shows antiangiogenic activity and antitumor activity against A549 in a transgenic zebrafish model^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Transgenic zebrafish model transplanted with CM-Dil-stained A549 cells^[1]</td> </tr> <tr> <td>Dosage:</td> <td>0.025, 0.05, and 0.1 μM</td> </tr> <tr> <td>Administration:</td> <td></td> </tr> <tr> <td>Result:</td> <td>Inhibited angiogenesis. Disrupted intersegmental blood vessels (ISVs) and dorsal</td> </tr> </table>	Animal Model:	Transgenic zebrafish model transplanted with CM-Dil-stained A549 cells ^[1]	Dosage:	0.025, 0.05, and 0.1 μM	Administration:		Result:	Inhibited angiogenesis. Disrupted intersegmental blood vessels (ISVs) and dorsal
Animal Model:	Transgenic zebrafish model transplanted with CM-Dil-stained A549 cells ^[1]								
Dosage:	0.025, 0.05, and 0.1 μM								
Administration:									
Result:	Inhibited angiogenesis. Disrupted intersegmental blood vessels (ISVs) and dorsal								



longitudinal anastomotic vessels (DLAVs) in zebrafish.
Inhibited tumor cell proliferation with inhibitory rates of 5.7% (0.025 μM), 14.8% (0.05 μM)
and 56.9% (0.1 μM), respectively.

REFERENCES

[1]. Li Y, et al. Trigothysoid N inhibits tumor proliferation and migration by targeting mitochondria and the STAT3/FAK pathway[J]. Arabian Journal of Chemistry, 2023, 16(8): 104930.

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

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