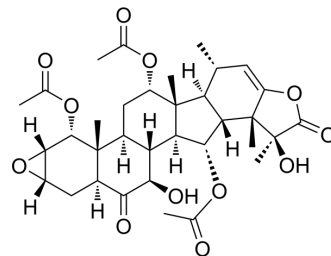


Taccalonolide E

Cat. No.:	HY-N11067
CAS No.:	134954-57-7
Molecular Formula:	C ₃₄ H ₄₄ O ₁₂
Molecular Weight:	644.71
Target:	Apoptosis
Pathway:	Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Taccalonolide E is a microtubule stabilizer and induces cancer cell apoptosis.																
In Vitro	<p>Taccalonolide E (5 or 10 μM; 18 h) causes an increased density of cellular microtubules in interphase cells and the formation of thick bundles of microtubules in A-10 cells^[1].</p> <p>Taccalonolide E (1 or 5 μM; 18 h) causes the appearance of abnormal multipolar mitotic spindles in A-10 and HeLa cells^[1].</p> <p>Taccalonolide E (5 μM; 6-24 h) arrests cell cycle at G2-M phase^[1].</p> <p>Taccalonolide E (0.5-5 μM; 18 h) initiates micronucleation in interphase A-10 cells^[1].</p> <p>Taccalonolide E (48 h) inhibits cancer cell proliferation^[1].</p> <p>Taccalonolide E (10 μM; 1 h) causes polymerization of tubulin in MDA-MB-435 cells^[1].</p> <p>Taccalonolide E (5 μM; 4-30 h) initiates Bcl-2 phosphorylation, MAPK activation, and apoptosis^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cycle Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MDA-MB-435 cells</td> </tr> <tr> <td>Concentration:</td> <td>5 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>6, 12, 18, or 24 h</td> </tr> <tr> <td>Result:</td> <td>Caused cells to accumulate in the G2-M phase of the cell cycle.</td> </tr> </table> <p>Cell Proliferation Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>SK-OV-3, MDA-MB-435, NCI/ADR, 1 A9, PTX 10, PTX 22 and 1A9/A8 cells</td> </tr> <tr> <td>Concentration:</td> <td></td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited the proliferation with IC₅₀s of 0.99 ± 0.08, 0.78 ± 0.17, 21.1 ± 0.46, 0.34 ± 0.04, 1.64 ± 0.25, 4.01 ± 0.20 and 1.42 ± 0.30 μM against SK-OV-3, MDA-MB-435, NCI/ADR, 1 A9, PTX 10, PTX 22 and 1A9/A8 cells, respectively.</td> </tr> </table> <p>Western Blot Analysis^[1]</p>	Cell Line:	MDA-MB-435 cells	Concentration:	5 μM	Incubation Time:	6, 12, 18, or 24 h	Result:	Caused cells to accumulate in the G2-M phase of the cell cycle.	Cell Line:	SK-OV-3, MDA-MB-435, NCI/ADR, 1 A9, PTX 10, PTX 22 and 1A9/A8 cells	Concentration:		Incubation Time:	48 h	Result:	Inhibited the proliferation with IC ₅₀ s of 0.99 ± 0.08, 0.78 ± 0.17, 21.1 ± 0.46, 0.34 ± 0.04, 1.64 ± 0.25, 4.01 ± 0.20 and 1.42 ± 0.30 μM against SK-OV-3, MDA-MB-435, NCI/ADR, 1 A9, PTX 10, PTX 22 and 1A9/A8 cells, respectively.
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Cell Line:	MDA-MB-435 cells
Concentration:	5 μ M
Incubation Time:	4, 6, 12, 24, or 30 h
Result:	Activated ERK1/2 and increased PARP cleavage.

REFERENCES

[1]. Tinley TL, et al. Taccalonolides E and A: Plant-derived steroids with microtubule-stabilizing activity. *Cancer Res.* 2003 Jun 15;63(12):3211-20.

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

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