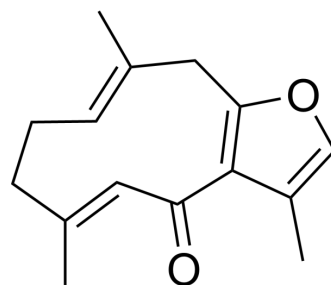


Furanodienone

Cat. No.:	HY-N2184
CAS No.:	24268-41-5
Molecular Formula:	C ₁₅ H ₁₈ O ₂
Molecular Weight:	230.3
Target:	Apoptosis
Pathway:	Apoptosis
Storage:	-20°C, sealed storage, away from moisture and light * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (434.22 mM; Need ultrasonic)																													
	Preparing Stock Solutions	<table border="1"> <thead> <tr> <th>Solvent</th> <th>Mass</th> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> </thead> <tbody> <tr> <td>Concentration</td> <td></td> <td></td> <td></td> <td></td> </tr> <tr> <td>1 mM</td> <td></td> <td>4.3422 mL</td> <td>21.7108 mL</td> <td>43.4216 mL</td> </tr> <tr> <td>5 mM</td> <td></td> <td>0.8684 mL</td> <td>4.3422 mL</td> <td>8.6843 mL</td> </tr> <tr> <td>10 mM</td> <td></td> <td>0.4342 mL</td> <td>2.1711 mL</td> <td>4.3422 mL</td> </tr> </tbody> </table>	Solvent	Mass	1 mg	5 mg	10 mg	Concentration					1 mM		4.3422 mL	21.7108 mL	43.4216 mL	5 mM		0.8684 mL	4.3422 mL	8.6843 mL	10 mM		0.4342 mL	2.1711 mL	4.3422 mL			
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Please refer to the solubility information to select the appropriate solvent.																														
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (10.86 mM); Clear solution																													
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (10.86 mM); Clear solution																													
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (10.86 mM); Clear solution																													

BIOLOGICAL ACTIVITY

Description	Furanodienone is one of the major bioactive constituents derived from Rhizoma Curcumae. Furanodienone induced apoptosis ^[1] .
In Vitro	Furanodienone (0-573.8 μM; 24 hours) exhibits IC ₅₀ values : 56.4 μM (RKO), 73.7 μM (sw480), 251.1 μM (HT-29), 412.5 μM (sw620) and 573.8 μM (LoVo) at 24 hours, while that in 48 h are 51.8 μM (RKO), 44.18 μM (sw480), 168.9 μM (HT-29), 314.2 μM (sw620) and 502.1 μM (LoVo), respectively ^[1] . Furanodienone (0-150 μM; 24 hours) induces apoptosis and shows increase in caspase-9 and -3 activity has been observed in both cells, whereas a relative minor effect on that of caspase-8 in RKO and HT-29 cells ^[1] .

Furanodienone (0-150 μ M; 24 hours) increases the apoptotic rates from 2.34 \pm 0.45% to 19.45 \pm 2.37% and 27.34 \pm 0.79%, in RKO cells at 75 and 150 μ M, whereas 12.4 \pm 1.08 and 20.64 \pm 3.02% apoptosis at the concentration of 75 and 150 μ M are observed in HT-29 cells compared with 2.89 \pm 0.26%^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Western Blot Analysis^[1]

Cell Line:	RKO and HT-29 cells
Concentration:	0 μ M; 50 μ M; 100 μ M; 150 μ M
Incubation Time:	24 hours
Result:	Increased caspase-9 and -3.

Apoptosis Analysis^[1]

Cell Line:	RKO and HT-29 cells
Concentration:	0 μ M; 75 μ M; 150 μ M
Incubation Time:	24 hours
Result:	Induced apoptosis of colorectal cells (RKO and HT-29).

CUSTOMER VALIDATION

- Life. 2022, 12(1), 114.

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REFERENCES

[1]. Li YW, et al. Furanodienone induces cell cycle arrest and apoptosis by suppressing EGFR/HER2 signaling in HER2-overexpressing human breast cancer cells. Cancer Chemother Pharmacol. 2011 Nov;68(5):1315-23.

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

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