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Diagnostik & molekulare Diagnostik



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Proteins

Product Data Sheet

DIM-C-pPhOH

Cat. No.: HY-112055 CAS No.: 151358-47-3 Molecular Formula: $C_{23}H_{18}N_{2}O$ Molecular Weight: 338.4

Apoptosis; Nuclear Hormone Receptor 4A/NR4A Target: Pathway: Apoptosis; Vitamin D Related/Nuclear Receptor

Storage: Powder -20°C 3 years

4°C 2 years

In solvent -80°C 2 years

-20°C 1 year

SOLVENT & SOLUBILITY

In Vitro

DMSO: 125 mg/mL (369.39 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.9551 mL	14.7754 mL	29.5508 mL
	5 mM	0.5910 mL	2.9551 mL	5.9102 mL
	10 mM	0.2955 mL	1.4775 mL	2.9551 mL

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: ≥ 2.08 mg/mL (6.15 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (6.15 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (6.15 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	DIM-C-pPhOH is a nuclear receptor 4A1 (NR4A1) antagonist. DIM-C-pPhOH inhibits cancer cell growth and mTOR signaling, induce apoptosis and cellular stress. DIM-C-pPhOH reduces cell proliferation with IC50 values of 13.6 μM and 13.0 μM for ACHN cells and 786-O cells, respectively ^[1] .
IC ₅₀ & Target	Nur77/NR4A1
In Vitro	DIM-C-pPhOH (7.5-20 μ M; 24 hours; ACHN and 786-O cells) treatment significantly decreases cell proliferation [1].

DIM-C-pPhOH (20 μ M; 24 hours; ACHN and 786-O cells) treatment induces Annexin V staining in ACHN and 786-O cells, confirming that DIM-C-pPhOH induce apoptosis, and also induces cleavage of caspases 7 and 8^[1].

DIM-C-pPhOH (15-20 μ M; 24 hours; ACHN and 786-O cells) treatment inhibits NR4A1-regulated expression of survivin, bcl-2 and EGFR in ACHN and 786-O cells. And also induces sestrin 2, activates AMPK α and inhibits activation of mTOR and downstream kinases^[1].

DIM-C-pPhOH decreases expression of $\beta1$ -integrin protein and mRNA and $\beta1$ -integrin-dependent responses in MCF7, MDA-MB-231, and SKBR3 cells and also inhibits migration of the latter two cell lines^[2].

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

Cell Proliferation Assay^[1]

Cell Line:	ACHN and 786-O cells	
Concentration:	7.5 μΜ, 15 μΜ, 20 μΜ	
Incubation Time:	24 hours	
Result:	Significantly decreased cell proliferation.	
Apoptosis Analysis ^[1]		
Cell Line:	ACHN and 786-O cells	
Concentration:	20 μΜ	
Incubation Time:	24 hours	
Result:	Induced apoptosis in ACHN and 786-O cells.	
Western Blot Analysis ^[1]		
Cell Line:	ACHN and 786-O cells	
Concentration:	15 μΜ, 20 μΜ	
Incubation Time:	24 hours	
Result:	Inhibited NR4A1-regulated expression of survivin, bcl-2 and EGFR in ACHN and 786-O cells. And also induced sestrin 2, activated AMPK α and inhibited activation of mTOR and downstream kinases.	

In Vivo

DIM-C-pPhOH (30 mg/kg; oral gavage; daily; for 50 days; male athymic nude mice) treatment results in a significant inhibition of tumor growth $^{[1]}$.

 $\label{eq:mce} \mbox{MCE has not independently confirmed the accuracy of these methods. They are for reference only.}$

Animal Model:	Male athymic nude mice (aged 6-7 weeks) injected with ACHN $\operatorname{cells}^{[1]}$	
Dosage:	30 mg/kg/day	
Administration:	Oral gavage; daily; for 50 days	
Result:	Resulted in a significant inhibition of tumor growth.	

CUSTOMER VALIDATION

- Redox Biol. 2021 Jan;38:101807.
- J Ethnopharmacol. 2024 Jan 7:117690.
- J Cardiovasc Transl Res. 2023 May 30.

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REFERENCES

- [1]. Hedrick E, et al. Nuclear Receptor 4A1 (NR4A1) as a Drug Target for Renal Cell Adenocarcinoma. PLoS One. 2015 Jun 2;10(6):e0128308.
- $[2]. \ Hedrick\ E, et\ al.\ NR4A1\ Antagonists\ Inhibit\ \beta1-Integrin-Dependent\ Breast\ Cancer\ Cell\ Migration.\ Mol\ Cell\ Biol.\ 2016\ Apr\ 15;36(9):1383-94.$

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

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