4-Methylbenzylidene camphor

MedChemExpress

Cat. No.:	HY-17587		
CAS No.:	36861-47-9		
Molecular Formula:	C ₁₈ H ₂₂ O		
Molecular Weight:	254.37		
Target:	Apoptosis; PI3K; Akt; ERK		
Pathway:	Apoptosis; PI3K/Akt/mTOR; MAPK/ERK Pathway; Stem Cell/Wnt		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year

SOLVENT & SOLUBILITY

n Vitro	0, 1	DMSO : 100 mg/mL (393.13 mM; Need ultrasonic) H ₂ O : < 0.1 mg/mL (insoluble)				
		Solvent Mass Concentration	1 mg	5 mg	10 mg	
	Preparing Stock Solutions	1 mM	3.9313 mL	19.6564 mL	39.3128 mL	
		5 mM	0.7863 mL	3.9313 mL	7.8626 mL	
		10 mM	0.3931 mL	1.9656 mL	3.9313 mL	
	Please refer to the sol	Please refer to the solubility information to select the appropriate solvent.				
n Vivo		it one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline g/mL (9.83 mM); Suspended solution; Need ultrasonic				
		2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (9.83 mM); Suspended solution; Need ultrasonic				
		3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (9.83 mM); Clear solution				

BIOLOGICAL ACTIVITY Description 4-Methylbenzylidene camphor (4-MBC) is an endocrine disrupter that produces estrogen-like effects. 4-Methylbenzylidene camphor decreases the proliferation of human trophoblast cells and induces apoptosis. 4-Methylbenzylidene camphor activates PI3K/AKT and ERK1/2 signaling pathways and elevates intracellular ROS production. 4-Methylbenzylidene camphor is a ultraviolet (UV) filter and may hamper normal placental formation during early pregnancy^{[1][2]}. IC₅₀ & Target PI3K Akt ERK1 ERK1

In Vitro

4-Methylbenzylidene camphor (4-MBC; 5-400 μM; for 48 h) inhibits proliferation of HTR8/SVneo cell^[1].

 $\label{eq:2.1} \mbox{4-Methylbenzylidene camphor (10-50 μM; for 48 h) induces apoptotic cell death of human trophoblast cells \mbox{[1]}.$

4-Methylbenzylidene camphor (5-50 μ M; for 48 h) increased the proportion of cells in the SubG1 phase^[1].

4-Methylbenzylidene camphor (50 $\mu\text{M};$ for 48 h) reduces invasion of human trophoblast cells^{[1]}.

4-Methylbenzylidene camphor (50 μ M; 5-120 min) activates PI3K/AKT and ERK1/2 signaling pathways in human trophoblast cells^[1].

4-Methylbenzylidene camphor (20-50 μ M; 24 h) significantly increases the expression of SEMA6 A, GPR56, ITGB4, EPHB4, NRP1^[1].

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

Cell Proliferation Assay^[1]

Cell Line:	HTR8/SVneo cells
Concentration:	0, 5, 10, 20, 50, 100, 200, and 400 μM
Incubation Time:	48 h
Result:	Dose-dependently inhibited cell proliferation of HTR8/SVneo cell.

Apoptosis Analysis^[1]

Cell Line:	HTR8/SVneo cells
Concentration:	10, 20, 50 µM
Incubation Time:	48 h
Result:	Early and late apoptotic cells was significantly increased at 20 μM and 50 $\mu M.$

Cell Cycle Analysis^[1]

Cell Line:	HTR8/SVneo cells
Concentration:	5, 10, 20, 50 μΜ
Incubation Time:	48 h
Result:	Gradually increased the proportion of cells in the SubG1 phase.

Cell Invasion Assay^[1]

Cell Line:	HTR8/SVneo cells	
Concentration:	50 μΜ	
Incubation Time:	48 h	
Result:	Revealed a significant reduction of 81.5% in invasiveness	

Western Blot Analysis^[1]

Cell Line:	HTR8/SVneo cells
Concentration:	50 μΜ
Incubation Time:	0, 5, 15, 30, 60, 120 min
Result:	The phosphorylation of AKT and its downstream kinase protein, P70S6K, peaked at 5 and 15 min, respectively, subsequently decreased after 30 min, and then reactivated at 120

	min
RT-PCR ^[1]	
Cell Line:	HTR8/SVneo cells
Concentration:	20, 50 μM
Incubation Time:	24 h
Result:	Significantly increased the expression of semaphorin 6 A (SEMA6 A), GPR56, integrin subunit beta 4 (ITGB4), EPHB4, neuropilin 1 (NRP1).

mating, during pregnancy and lactation, and to the offspring until adulthood) causes exhibited enhanced proand altered uterine gene expression in Neonates^[2].

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CUSTOMER VALIDATION

• Reprod Toxicol. 2019 Mar;84:49-58.

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REFERENCES

[1]. Changwon Yang, et al. 4-Methylbenzylidene-camphor inhibits proliferation and induces reactive oxygen species-mediated apoptosis of human trophoblast cells. Reprod Toxicol. 2019 Mar:84:49-58.

[2]. Margret Schlumpf, et al. Developmental toxicity of UV filters and environmental exposure: a review. Int J Androl. 2008 Apr;31(2):144-51.

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

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