

# C16-Ceramide

Cat. No.: HY-100354 CAS No.: 24696-26-2 Molecular Formula:  $C_{34}H_{67}NO_3$ Molecular Weight: 537.9

Target: Endogenous Metabolite; MDM-2/p53; Apoptosis

Pathway: Metabolic Enzyme/Protease; Apoptosis

Storage: 4°C, protect from light

\* In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)

**Product** Data Sheet

#### **SOLVENT & SOLUBILITY**

In Vitro DMF: 20 mg/mL (37.18 mM; ultrasonic and warming and heat to 60°C)

> Ethanol: 20 mg/mL (37.18 mM; warming and heat to 60°C) DMSO: < 1 mg/mL (ultrasonic) (insoluble or slightly soluble)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.8591 mL	9.2954 mL	18.5908 mL
	5 mM	0.3718 mL	1.8591 mL	3.7182 mL
	10 mM	0.1859 mL	0.9295 mL	1.8591 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

1. Add each solvent one by one: 10% EtOH >> 90% corn oil Solubility: ≥ 2 mg/mL (3.72 mM); Clear solution

### **BIOLOGICAL ACTIVITY**

Description	${\tt C16-Ceramide\ is\ a\ natural\ small\ molecule\ activating\ p53\ through\ the\ direct\ and\ selective\ binding}^{[1]}.$
IC <sub>50</sub> & Target	p53 <sup>[1]</sup> , Apoptosis <sup>[2]</sup>
In Vitro	C16-Ceramide interacts with p53 within its core domain. p53 forms complex with natural C16-Ceramide in the cell <sup>[1]</sup> . C16-Ceramide (2.5-50 $\mu$ M; 0-48 h) strongly decreased HCT116 cell viability in a time- and concentration-dependent manner <sup>[2]</sup> . C16-Ceramide (12 $\mu$ M; 48 h) induces apoptosis through Btf (Bcl-2-associated transcription factor) in HCT116 cells <sup>[2]</sup> . C16-ceramide (12 $\mu$ M; 0-6 h) and Btf expression up-regulate p53 and BAX expression. C16-ceramide down-regulates Mdm2 expression via Btf <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay <sup>[2]</sup>

Cell Line:	HCT116 cells	
Concentration:	$2.5, 5, 10, 12, 20, 50 \mu\text{M}$	
Incubation Time:	0-48 h	
Result:	Strongly decreased cell viability in a time- and concentration-dependent manner.	
Western Blot Analysis <sup>[2]</sup>		
Cell Line:	HCT116 cells	
Concentration:	12 μΜ	
Incubation Time:	1, 3 and 6 h	
Result:	Increased PARP cleavage, decreased pro-caspase 3. Decreased the levels of stratifin an stathmin, increased the expression of prohibitin and Btf. RNAi-mediated Btf depletion alsopartially inhibited BAX expression after the treatment. Significantly decreased luciferase activity and Mdm2 protein expression levels.	

## **CUSTOMER VALIDATION**

• ACS Nano. 2023 Jul 13.

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#### **REFERENCES**

[1]. Fekry B, et al. C16-ceramide is a natural regulatory ligand of p53 in cellular stress response. Nat Commun. 2018 Oct 8;9(1):4149.

[2]. Rénert AF, et al. The proapoptotic C16-ceramide-dependent pathway requires the death-promoting factor Btf in colon adenocarcinoma cells. J Proteome Res. 2009 Oct;8(10):4810-22.

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

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