Zymostenol

Cat. No.:	HY-113345		
CAS No.:	566-97-2		
Molecular Formula:	C ₂₇ H ₄₆ O		
Molecular Weight:	386.65		
Target:	Endogenou	s Metabo	lite; ROR
Pathway:	Metabolic E	nzyme/P	rotease; Vitamin D Related/Nuclear Receptor
Storage:	Powder In solvent	-20°C -80°C	3 years 6 months
		-20°C	1 month

SOLVENT & SOLUBILITY

		Solvent Mass Concentration	1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.5863 mL	12.9316 mL	25.8632 mL
		5 mM	0.5173 mL	2.5863 mL	5.1726 mL
		10 mM			

BIOLOGICAL ACTI	VITY		
Description	Zymostenol (5a-Cholest-8-en-3b-ol) is a late-stage precursor in the biosynthesis of cholesterol. Zymostenol is a RORy agonist $(EC_{50}: 1 \ \mu M)^{[1][2][3]}$.		
IC ₅₀ & Target	Human Endogenous Metabolite	RORγ 1 μM (EC50)	
In Vitro	Zymostenol (12-104 μM) enhances the formation of myelin basic protein-positive (MBP+) oligodendrocytes from mous epiblast stem cell-derived oligodendrocyte progenitor cells ^[2] . Zymostenol (20 μM, 3 days) arrest MCF-7 cell cycle in the G0-G1 phase ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Cycle Analysis ^[2]		
	Cell Line:	MCF-7 cells	
	Concentration:	20 µM	

HO

Incubation Time:	3 days
Result:	Induced 67 \pm 1% accumulation of cells in the G0-G1 phase of the cell cycle.

REFERENCES

[1]. Hu X, et al. Sterol metabolism controls T(H)17 differentiation by generating endogenous RORy agonists. Nat Chem Biol. 2015 Feb;11(2):141-7.

[2]. Hubler Z, et al. Accumulation of 8,9-unsaturated sterols drives oligodendrocyte formation and remyelination. Nature. 2018 Aug;560(7718):372-376.

[3]. Payré B, et al. Microsomal antiestrogen-binding site ligands induce growth control and differentiation of human breast cancer cells through the modulation of cholesterol metabolism. Mol Cancer Ther. 2008 Dec;7(12):3707-18.

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

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