# Ziyuglycoside I

Cat. No.: HY-N0331 CAS No.: 35286-58-9  $C_{41}H_{66}O_{13}$ Molecular Formula: Molecular Weight: 766.96

Target: MDM-2/p53; Apoptosis

Pathway: **Apoptosis** 

Storage: -20°C, protect from light

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

**Product** Data Sheet

### **SOLVENT & SOLUBILITY**

In Vitro

Methanol: 125 mg/mL (162.98 mM; Need ultrasonic) DMSO: 100 mg/mL (130.38 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.3038 mL	6.5192 mL	13.0385 mL
	5 mM	0.2608 mL	1.3038 mL	2.6077 mL
	10 mM	0.1304 mL	0.6519 mL	1.3038 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 (2.71 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (2.71 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (2.71 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	Ziyuglycoside I isolated from S. officinalis root, has anti-wrinkle activity, and increases the expression of type I collagen. Ziyuglycoside I could be used as an active ingredient for cosmetics <sup>[1]</sup> . Ziyuglycoside I triggers cell cycle arrest and apoptosis mediated by p53, it can be a potential agent candidate for treating triple-negative breast cancer (TNBC) <sup>[2]</sup> .
In Vitro	Ziyuglycoside I (5-160 $\mu$ M; 24 hours) reveals a marked anti-proliferation activity with an IC <sub>50</sub> value of 13.96 $\mu$ M in MDA-MB-231 cells <sup>[2]</sup> . Ziyuglycoside I (5-20 $\mu$ M; 24 hours) induces MDA-MB-231 cell apoptosis by increasing the percentage of apoptotic cells from 2.43% to 44.76% at 20 $\mu$ M <sup>[2]</sup> .

Ziyuglycoside I (5-20  $\mu$ M; 24 hours) induces the cleavage of caspas-8, caspase-9, and caspase-3, up-regulates the proapoptotic protein Bax, and down-regulates anti-apoptotic protein Bal-2, dose-dependently reduces the level of mito-cyto c expression<sup>[2]</sup>.

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

 $\operatorname{Cell Viability} \operatorname{Assay}^{[1]}$ 

Cell Line:	MDA-MB-231 cells		
Concentration:	5 μM, 10 μM, 20 μM, 40 μM, 80 μM, and 160 μM		
Incubation Time:	24 hours		
Result:	Showed cytotoxicity on MDA-MB-231 cells.		
Cell Cycle Analysis <sup>[1]</sup>			
Cell Line:	MDA-MB-231 cells		
Concentration:	5 μM, 10 μM, and 20 μM		
Incubation Time:	24 hours		
Result:	Induced G2/M phase arrest and apoptosis on MDA-MB-231 Cells		
Western Blot Analysis <sup>[1]</sup>			
Cell Line:	MDA-MB-231 cells		
Concentration:	5 μM, 10 μM, and 20 μM		
Incubation Time:	24 hours		
Result:	Induced apoptosis in MDA-MB-231 Cells through Intrinsic and Extrinsic Pathways.		

#### **REFERENCES**

[1]. Kim YH, et al. Anti-wrinkle activity of ziyuglycoside I isolated from a Sanguisorba officinalis root extract and its application as a cosmeceutical ingredient. Biosci Biotechnol Biochem. 2008 Feb;72(2):303-11. Epub 2008 Feb 7.

[2]. Zhu X, et al. Ziyuglycoside I Inhibits the Proliferation of MDA-MB-231 Breast Carcinoma Cells through Inducing p53-Mediated G2/M Cell Cycle Arrest and Intrinsic/Extrinsic Apoptosis. Int J Mol Sci. 2016 Nov 22;17(11). pii: E1903

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA