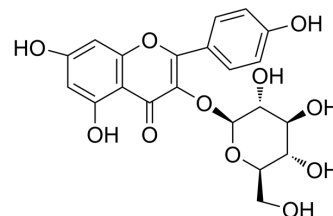


## Astragalin

Cat. No.:	HY-N0015		
CAS No.:	480-10-4		
Molecular Formula:	C <sub>21</sub> H <sub>20</sub> O <sub>11</sub>		
Molecular Weight:	448.38		
Target:	Apoptosis; NF-κB		
Pathway:	Apoptosis; NF-κB		
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 : 100 mg/mL (223.03 mM; Need ultrasonic)

Concentration	Solvent	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.2303 mL	11.1513 mL	22.3025 mL
	5 mM	0.4461 mL	2.2303 mL	4.4605 mL
	10 mM	0.2230 mL	1.1151 mL	2.2303 mL

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

#### In Vivo

- Add each solvent one by one: 1% CMC-Na/saline water  
Solubility: 10 mg/mL (22.30 mM); Suspended solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline  
Solubility: ≥ 2.5 mg/mL (5.58 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)  
Solubility: ≥ 2.5 mg/mL (5.58 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil  
Solubility: ≥ 2.5 mg/mL (5.58 mM); Clear solution

### BIOLOGICAL ACTIVITY

#### Description

Astragalin (Astragaline) a flavonoid with anti-inflammatory, antioxidant, anticancer, bacteriostatic activity. Astragalin inhibits cancer cells proliferation and migration, induces apoptosis. Astragalin is orally active and provides nerve and heart protection, and resistance against osteoporosis<sup>[1]</sup>.

#### IC<sub>50</sub> & Target

p65

**In Vitro**

Astragalin (0-80 µg/mL; 24 h, 48 h, 72 h) inhibits cancer cells viability and migration at 20, 40, 80 µM, and shows no cytotoxicity to normal human colon epithelial cell lines NCM460<sup>[1]</sup>.

Astragalin (80 µg/mL; 4 h, 8 h) interrupts the NF-κB signaling pathway and inhibits NF-κB P65 transcriptional activity in HCT116 cells stimulated by TNF-α<sup>[1]</sup>.

Astragalin (20, 40, 80 µg/mL; 48 h) induces HCT116 cells apoptosis and arrests cell cycle at G0/G1 phase<sup>[1]</sup>.

Astragalin (20, 40, 80 µg/mL; 48 h) increases the level of apoptin in HCT116 and decreases the level of anti-apoptotic proteins in a dose-dependent manner<sup>[1]</sup>.

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

**Cell Viability Assay<sup>[1]</sup>**

Cell Line:	Human colon cancer cell lines (HCT116, LoVo, SW620, SW480, Caco2) and human colon epithelial cell lines NCM460
Concentration:	0, 5, 10, 20, 40, 80 µg/mL
Incubation Time:	24, 48, 72 hours
Result:	Inhibits cancer cells (LoVo, SW620, SW480, Caco2, HCT116) activity at 20, 40, 80 µM. Inhibited HCT116 cells with IC <sub>50</sub> s of 121.845, 87.908, 18.883 µg/mL at 24, 48, 72 hours, respectively. Showed no cytotoxicity with NCM460 cells.

**Cell Cycle Analysis<sup>[1]</sup>**

Cell Line:	Human colon cancer cell HCT116
Concentration:	0, 20, 40, 80 µg/mL
Incubation Time:	48 hours
Result:	Arrested cell cycle at G0/G1.

**Western Blot Analysis<sup>[1]</sup>**

Cell Line:	Human colon cancer cell HCT116
Concentration:	80 µg/mL; pre-treated with 20 ng/mL TNF-α
Incubation Time:	4 or 8 hours
Result:	Increased the expression of pro-apoptotic proteins (caspase 3, caspase 6, caspase 7, caspase 8, caspase 9, P53, and Bax) and decreased the expression of anti-apoptotic proteins (cleaved caspase-3 and Bcl-2) in a dose-dependent manner.

**Cell Migration Assay<sup>[1]</sup>**

Cell Line:	Human colon cancer cell HCT116
Concentration:	0, 20, 40, 80 µg/mL
Incubation Time:	24, 48, 72 hours
Result:	Reduced release of MMP-2 and MMP-9 enzymes. Inhibited cell migration with mobilities of 14.38, 13.44, and 5.29% with 20, 40, and 80 µg/mL concentrations, respectively.

**In Vivo**

Astragalin (25, 50, and 75 mg/kg; p.o.; once every two days; 25 d) reduces the tumor volumes, tumor weight and tumor formation rates and decreases expression of p-NFκB and p-ικκ in nude mouse model with human colon cancer<sup>[1]</sup>.

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

Animal Model:	Male BALB/c nude mice (4-week-old) with HCT116 cells (s.c.) <sup>[1]</sup>
Dosage:	0 mg/kg, 25 mg/kg, 50 mg/kg, 75 mg/kg
Administration:	Oral gavage; once every two days; for 25 days
Result:	Resulted tumor inhibition rates of 17.43%, 34.89% ( $p < 0.01$ ), and 67.06% with 25 mg/kg, 50 mg/kg, 75 mg/kg dose administration. Showed no effect on mouse weight.

## CUSTOMER VALIDATION

- Phytomedicine. 2022 Jul;101:154113.
- Int Immunopharmacol. November 2022, 109278.
- J Cardiovasc Pharmacol. 2021 Feb 1;77(2):217-227.
- Curr Res Food Sci. 2023 Nov 6.
- SSRN. 2023 Feb 7.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

## REFERENCES

[1]. Yang M, et al. Astragalosin Inhibits the Proliferation and Migration of Human Colon Cancer HCT116 Cells by Regulating the NF- $\kappa$ B Signaling Pathway. Front Pharmacol. 2021 Apr 19;12:639256.

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

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