



**SZABO
SCANDIC**

Part of Europa Biosite

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic



Silymarin

Cat. No.:	HY-N7073		
CAS No.:	65666-07-1		
Target:	SARS-CoV		
Pathway:	Anti-infection		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month

Silymarin

SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (Need ultrasonic)
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 3 mg/mL (Infinity mM); Clear solution

BIOLOGICAL ACTIVITY

Description	Silymarin is an extract of the milk thistle (<i>Silybum marianum</i>). Silymarin is an effective SARS-CoV-2 main protease (M ^{PRO}) inhibitor. Silymarin can significantly reduce tumor cell proliferation, angiogenesis as well as insulin resistance. Silymarin has the chemopreventive effect on hepatocellular carcinoma (HCC). Silymarin has the potential for COVID-19 research ^{[1][2][4]} .									
In Vitro	<p>Silymarin (0-120 µg/ml; 24 hours) inhibits AGS cell viability, the viability of AGS cells is 77.9% in the presence of 20 µg/ml silymarin, 71.5% at 40 µg/ml, 59.8% at 60 µg/ml, 44.5% at 80 µg/ml, 35.3% at 100 µg/ml and 33.9% at 120 µg/ml^[1]. Silymarin (40-80 µg/ml; 24 hours) inhibits the migration of the AGS cells in a concentration-dependent manner. It inhibit migration of AGS cells 59.4% at 40 µg/ml and 21.7% at 80 µg/ml,respectively^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>AGS cells</td> </tr> <tr> <td>Concentration:</td> <td>20 µg/ml, 40 µg/ml, 80 µg/ml, 100 µg/ml and 120 µg/ml</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Indicated a significant concentration-dependent inhibitory effect on AGS cells starting at 20 µg/ml.</td> </tr> </table>		Cell Line:	AGS cells	Concentration:	20 µg/ml, 40 µg/ml, 80 µg/ml, 100 µg/ml and 120 µg/ml	Incubation Time:	24 hours	Result:	Indicated a significant concentration-dependent inhibitory effect on AGS cells starting at 20 µg/ml.
Cell Line:	AGS cells									
Concentration:	20 µg/ml, 40 µg/ml, 80 µg/ml, 100 µg/ml and 120 µg/ml									
Incubation Time:	24 hours									
Result:	Indicated a significant concentration-dependent inhibitory effect on AGS cells starting at 20 µg/ml.									
In Vivo	Silymarin (oral gavage; 10, 20, 50, 100, and 200 mg/kg) decreases the immobility time in a dose-dependent manner in forced swimming test (FST). It also lowers the immobility measure dose-dependently in tail suspension test (TST). Additionally, 50% of maximum response (ED ₅₀) of Silymarin is around 10 mg/kg. The dose 100 mg/kg proved the most effective dose in both the tests ^[3] .									

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

CUSTOMER VALIDATION

- Molecules. 2021, 26(5), 1409.
- Naunyn Schmiedebergs Arch Pharmacol. 2023 Apr 13.
- Biomed Chromatogr. 2022 Jun 16;e5427.
- Indian J Exp Biol. 2022 Dec.

See more customer validations on www.MedChemExpress.com

REFERENCES

- [1]. Féher J, et al. Silymarin in the prevention and treatment of liver diseases and primary liver cancer. Curr Pharm Biotechnol. 2012 Jan;13(1):210-7.
- [2]. Sung-Hyun Kim, et al. Silymarin induces inhibition of growth and apoptosis through modulation of the MAPK signaling pathway in AGS human gastric cancer cells. Oncol Rep. 2019 Nov;42(5):1904-1914.
- [3]. Mina Khoshnoodi, et al. Possible involvement of nitric oxide in antidepressant-like effect of silymarin in male mice. Pharm Biol. 2015 May;53(5):739-45.
- [4]. Anna Maria Sardanelli, et al. SARS-CoV-2 Main Protease Active Site Ligands in the Human Metabolome. Molecules 2021, 26(5), 1409.

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