

Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

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SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien T. +43(0)1 489 3961-0 F. +43(0)1 489 3961-7 <u>mail@szabo-scandic.com</u> www.szabo-scandic.com

Ethyl pyruvate

Cat. No.:	HY-Y1362		
CAS No.:	617-35-6		
Molecular Formula:	$C_5H_8O_3$		
Molecular Weight:	116.12		
Target:	Autophagy;	Apoptosi	s; Pyroptosis; NF-κB
Pathway:	Autophagy;	Apoptosi	s; Immunology/Inflammation; NF-кВ
Storage:	Pure form	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month

SOLVENT & SOLUBILITY

DMSO:100	mg/mL (861.18 mM; Need ultrasonic)			
	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solu	1 mM	8.6118 mL	43.0589 mL	86.1178 mL
	5 mM	1.7224 mL	8.6118 mL	17.2236 mL
	10 mM	0.8612 mL	4.3059 mL	8.6118 mL

DIOLOGICALACTIV	
Description	Ethyl pyruvate is a simple derivative of the endogenous metabolite pyruvate. Ethyl pyruvate is an HMGB1 release inhibitor. Ethyl pyruvate can induce apoptosis by autophagy. Ethyl pyruvate has anti-inflammatory, antioxidant and anti-tumor activity. Ethyl pyruvate can be used in the study of neurodegenerative diseases such as Alzheimer's and Parkinson's disease [1][2][3][4][5].
In Vitro	Ethyl pyruvate (10 mM, 1 h) has no toxic effect on N9 microglial cells in the range of 1-10 mM. The activation of microglia NLRP3 inflammasome is decreased by inhibiting the HMGB1/ NF-κB /miR-223 signaling pathway ^[2] . Ethyl pyruvate (10-40 mM, 6, 24 h) induces apoptosis in MC38 cells ^[3] . Ethyl pyruvate (5-15 mM, 2 h) has an IC ₅₀ value of 28.83 mM on mouse peritoneal macrophages. Endotoxemia and sepsis are prevented by inhibiting caspase-11-dependent pyroptosis ^[4] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay ^[2]

Product Data Sheet

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	N9 microglial
Concentration:	1-100 mM
Incubation Time:	1 h
Result:	Did not show cytotoxic effects in the range of 1–10 mM.
Western Blot Analysis ^[2]	
Cell Line:	N9 microglial
Concentration:	10 mM
Incubation Time:	1h
Result:	Suppressed LPS (HY-D1056)- and ATP (HY-B2176)-induced IL-1β and IL-18 protein and mRNA levels. Reduced NLRP3, Caspase-1, and ASC Specks. Reduced NF-κB activation and HMGB1 expression level.
Apoptosis Analysis ^[3]	
Cell Line:	MC38
Concentration:	10, 20, 40 mM
Incubation Time:	6, 24 h
Result:	Induced an increase in autophagy and apoptosis in a dose-and time-dependent manne
Result: Ethyl pyruvate (80 mg/k model ^[3] . Ethyl pyruvate (2 or 40 n rat model of paraquat ir MCE has not independer Animal Model:	Induced an increase in autophagy and apoptosis in a dose-and time-dependent manne g intraperitoneal injection for 9 consecutive days) inhibits tumor growth in a mouse liver tumo ng/kg, intraperitoneal injection) has reduced lipid peroxidation and anti-inflammatory effects ntoxication ^[5] . ntly confirmed the accuracy of these methods. They are for reference only.
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In Vivo

Decreased NO concentrations significantly at 6 h and GSH concentrations in the lung.

CUSTOMER VALIDATION

- Cell Death Dis. 2019 Sep 26;10(10):724.
- Life Sci. 2021 Jan 5;118987.
- J Pharm Pharmacol. 2023 Mar 25;rgad021.
- PeerJ. August 4, 2022.
- Research Square Preprint. 2021 Jul.

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REFERENCES

[1]. Liang X, et al. Ethyl pyruvate administration inhibits hepatic tumor growth. J Leukoc Biol. 2009 Sep;86(3):599-607.

[2]. Qiu X, et al. Ethyl pyruvate confers protection against endotoxemia and sepsis by inhibiting caspase-11-dependent cell pyroptosis. Int Immunopharmacol. 2020 Jan;78:106016.

[3]. Lee J, et al. Protective effects of ethyl pyruvate treatment on paraquat-intoxicated rats. Hum Exp Toxicol. 2008 Jan;27(1):49-54.

[4]. Fink MP. Ethyl pyruvate: a novel anti-inflammatory agent. J Intern Med. 2007 Apr;261(4):349-62.

[5]. Olcum M, Tufekci KU, Durur DY, et al. Ethyl Pyruvate Attenuates Microglial NLRP3 Inflammasome Activation via Inhibition of HMGB1/NF-κB/miR-223 Signaling. Antioxidants (Basel). 2021;10(5):745.

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