Esculetin

Cat. No.:	HY-N0284		
CAS No.:	305-01-1		
Molecular Formula:	C ₉ H ₆ O ₄		
Molecular Weight:	178.14		
Target:	PI3K; Akt		
Pathway:	PI3K/Akt/mTOR		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month

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SOLVENT & SOLUBILITY

In Vitro	DMSO : 125 mg/mL (701.70 mM; Need ultrasonic)						
		Solvent Mass Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	5.6136 mL	28.0678 mL	56.1356 mL		
		5 mM	1.1227 mL	5.6136 mL	11.2271 mL		
		10 mM	0.5614 mL	2.8068 mL	5.6136 mL		
	Please refer to the so	lubility information to select the ap	propriate solvent.				
In Vivo		1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (11.68 mM); Clear solution					
		2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (11.68 mM); Clear solution					
		3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (11.68 mM); Clear solution					

BIOLOGICAL ACTIVITY				
Description	Esculetin is an active ingredient extracted mainly from the bark of Fraxinus rhynchophylla. Esculetin inhibits platelet- derived growth factor (PDGF)-induced airway smooth muscle cells (ASMCs) phenotype switching through inhibition of PI3K/Akt pathway. Esculetin has antioxidant, antiinflammatory, and antitumor activities ^[1] .			
IC ₅₀ & Target	РІЗК			
In Vitro	Esculetin (40 μM , 24 h or 48 h) inhibits PDGF-BB-induced ASMC proliferation and migration $^{[1]}.$			

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	Esculetin (100 μM, 36 h) membrane potential by Esculetin (100 μM, 0-12 MCE has not independe	 Esculetin (40 μM, 1 h) inhibits PDGF-BB-induced ECM secretion and activation of PI3K/Akt pathway in ASMC cells^[1]. Esculetin (100 μM, 36 h) induces pancreatic cancer cell (such as PANC-1 cells) apoptosis and loss of mitochondrial membrane potential by activation of caspases 3, 8 and 9^[2]. Esculetin (100 μM, 0-12 h) binds to KEAP1 and inhibits its interaction with Nrf2 in pancreatic cancer cells (in PANC-1 cells)^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis^[2] 		
	Cell Line:	PANC-1 cells		
	Concentration:	100 μΜ		
	Incubation Time:	0-12 h		
	Result:	Increased the amount of phosphorylated form of Nrf2. Increased the levels of Nrf2 in nuclear. Decreased NF-κB protein levels.		
In Vivo	Esculetin (10 or 30 mg/k in an HCT116-implanted Esculetin (oral gavage, 2 mice by increasing antio	g/kg/day, p.o., 7d) ameliorates skin lesion in Imiquimod (HY-B0180) (IMQ)-induced psoriatic mice ^[3] . kg, i.p., three times per week for 28 days) inhibits tumor growth and metastasis via Axin2 suppression d orthotopic mouse model ^[4] . 25 mg/kg for 35 days) inhibits acute restraint stress (ARS)-induced oxidative stress in mature adult oxidant enzyme activities, the GSH/GSSG ratio, and cytochrome c oxidase activity in cortex ^[5] . ently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Imiquimod (HY-B0180) (IMQ)-induced psoriatic mice ^[3]		
	Dosage:	50 and 100 mg/kg/day		
	Administration:	p.o., 7d		
	Result:	Ameliorated the skin lesion and reduced the PASI score. Inhibited epidermal hyperplasia, and CD8 expression in the skin of psoriatic mice.		

CUSTOMER VALIDATION

- Pharmacol Res. 2020 May;155:104751.
- Front Cell Dev Biol. 2021 Nov 10;9:763864.
- J Hepatocell Carcinoma. 2023 Apr 11;10:611-629.

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REFERENCES

[1]. Arora R, et al. Esculetin induces antiproliferative and apoptotic response in pancreatic cancer cells by directly binding to KEAP1. Mol Cancer. 2016 Oct 18;15(1):64.

[2]. Chen Y, et al. Esculetin Ameliorates Psoriasis-Like Skin Disease in Mice by Inducing CD4+Foxp3+ Regulatory T Cells. Front Immunol. 2018 Sep 12;9:2092.

[3]. Kim WK, et al. Esculetin suppresses tumor growth and metastasis by targeting Axin2/E-cadherin axis in colorectal cancer. Biochem Pharmacol. 2018 Jun;152:71-83.

[4]. Martín-Aragón S, et al. Age-dependent effects of esculetin on mood-related behavior and cognition from stressed mice are associated with restoring brain antioxidant status. Prog Neuropsychopharmacol Biol Psychiatry. 2016 Feb 4;65:1-16.

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

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