

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

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Product Data Sheet

4-Hydroxycoumarin

Cat. No.: HY-N6856 CAS No.: 1076-38-6 Molecular Formula: $C_9H_6O_3$ **Molecular Weight:** 162.14

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 (616.75 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	6.1675 mL	30.8375 mL	61.6751 mL
	5 mM	1.2335 mL	6.1675 mL	12.3350 mL
	10 mM	0.6168 mL	3.0838 mL	6.1675 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 (12.83 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (12.83 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (12.83 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

4-Hydroxycoumarin is an orally active coumarin derivative, one of the most versatile heterocyclic scaffolds, often used in the synthesis of various organic compounds. 4-Hydroxycoumarin possesses both electrophilic and nucleophilic properties. 4-Hydroxycoumarin is an HIV protease inhibitor and tyrosine kinase inhibitor. 4-Hydroxycoumarin has anti-inflammatory, antibacterial and anti-tumor effects^{[1][2][3][4]}.

In Vitro

4-Hydroxycoumarin (0, 50, 160, 500 μ M, 24 h) disrupts the actin cytoskeleton in B16-F10 melanoma cells but not in B82 fibroblasts. However, they can reduce their adhesion and motility to extracellular matrix proteins^[1].

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

Cell Viability Assay^[2]

Cell Line:	B16-F10, B82	
Concentration:	0, 50, 160, 500 μΜ	
Incubation Time:	24 h	
Result:	Had no appreciable effect on cell viability.	
Western Blot Analysis [2]		
Cell Line:	B16-F10	
Concentration:	0, 50, 160, 500 μΜ	
Incubation Time:	24 h	
Result:	Reduced the adhesion to ECM proteins and the tyrosine phosphorylation of several proteins in a concentration-dependent manner. Inhibited the migration at 500 μ M.	

In Vivo

- 4-Hydroxycoumarin (10, 20 or 40 mg/kg/day, oral gavage) can effectively inhibit tumor growth and improve survival rate in melanoma mice $^{[3]}$.
- 4-Hydroxycoumarin (5, 10, 25, 50 mg/kg/ day, Oral gavage) can significantly reduce symptoms in a rat model of trinitrobenzene sulfonic acid-induced colitis [4].

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

Animal Model:	Mouse melanoma model $^{[3]}$	
Dosage:	10, 20 or 40 mg/kg	
Administration:	i.g.	
Result:	Reduced >85% of the number of pulmonary tumors. Diminished the tumor size from day 22 and increased survival time at 10 mg/kg/day.	
Animal Model:	Rat colitis model ^[4]	
Dosage:	5, 10, 25, 50 mg/kg	
Administration:	p.o.	
Result:	Reduced damage score and extension of the lesion at doses of 10 and 25 mg/kg. Counteracted GSH content and reduced AP activity at a dose of 5 mg/kg. Showed a good recovery of the intestinal cytoarchitecture at doses of 5 and 25 mg/kg.	

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

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