Product Data Sheet



Camobucol

Cat. No.: HY-14916 CAS No.: 216167-92-9 Molecular Formula: $C_{33}H_{50}O_4S_2$

Molecular Weight: 575 Target: Others Pathway: Others

Storage: Powder -20°C 3 years

In solvent

2 years -80°C 6 months

-20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (173.91 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.7391 mL	8.6957 mL	17.3913 mL
	5 mM	0.3478 mL	1.7391 mL	3.4783 mL
	10 mM	0.1739 mL	0.8696 mL	1.7391 mL

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

BIOLOGICAL ACTIVITY

Description

Camobucol (AGIX 4207) is an orally active, phenolic antioxidant and anti-inflammatory compound with antirheumatic properties.

In Vitro

Camobucol exhibits potent antioxidant activity toward lipid peroxides in vitro and displays enhanced cellular uptake. $Camobucol \ selectively \ inhibits \ tumor \ necrosis \ factor \ (TNF)-\alpha-inducible \ levels \ of \ the \ redox-sensitive \ genes, \ vascular \ cell$ adhesion molecule-1 and monocyte chemoattractant protein-1, with less inhibition of E-selectin, and no effect on intracellular adhesion molecule-1 expression in endothelial cells. In addition, Camobucol inhibits cytokine-induced levels of monocyte chemoattractant protein-1, interleukin (IL)-6, and IL-8 from endothelial cells and human fibroblast-like synoviocytes as well as lipopolysaccharide-induced release of TNF-α, IL-1β, and IL-6 from human peripheral blood mononuclear cells. Camobucol does not inhibit TNF- α -induced nuclear translocation of nuclear factor of the κ -enhancer in B cells (NF-кВ), suggesting that the mechanism of action is independent of this redox-sensitive transcription factor^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Kunsch C, et al. AGIX-4207 [2- antioxidant and anti-inflammato			ethylethyl]thio]-2,6-bis(1,1-dimethyle n	ethyl)phenoxy]acetic acid], a novel	
			ical applications. For research u		
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