

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Data Sheet (Cat.No.T1051)



Retinoic acid

Chemical Properties

CAS No.: 302-79-4

Formula: C20H28O2

Molecular Weight: 300.44

Appearance: no data available

Storage: keep away from direct sunlight

Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Biological Description

Description	Retinoic acid (Tretinoin), a metabolite of vitamin A, is a natural agonist of the retinoic acid receptor RAR and inhibits RAR $\alpha/\beta/\gamma$ (IC50=14 nM). Retinoic acid induces cellular differentiation, reduces cellular proliferation, and inhibits tumorigenesis.				
Targets(IC50)	Retinoid Receptor, Endogenous Metabolite, PPAR, Autophagy				
In vitro	Tretinoin prevents skin atrophy induced by corticosteroids in hairless mice. When coadministered with miquimod in guinea pigs, tretinoin induces tattoo fading and moderate pigment clearance histopathologically. Applications of tretinoin on incisions in the skin of 45 CD-1 mice increase fibroblast differentiation and reduce collagen production. In aged male Fischer 344 rats treated with tretinoin, renal cortex protein content is 30% lower compared to controls, potentially due to suppressed expression of tumor necrosis factor-β1 and osteopontin.				
In vivo	In studies evaluating the impact on glutathione levels and catalase activity, Tretinoin increased both metrics in a time- and dose-dependent manner, offering protective and mitigating effects against H2O2 cytotoxicity in human renal mesangial cells. Treatment with Tretinoin resulted in elevated mRNA levels of catalase and γ -glutamylcysteine synthetase (the catalytic subunit responsible for the rate-limiting step in reduced glutathione synthesis) in cultured mesangial cells. Additionally, Tretinoin upregulated matrix metalloproteinase-8/13 in human keloid-derived fibroblasts.				
Cell Research	Retinoic acid is dissolved in DMSO and stored, and then diluted with appropriate medium before use[3]. P19 cell are induced to undergo neuronal differentiation according to established procedures. Briefly, cells are cultured on 1% agarose-coated 10 cm dishes at 3×10 5 cells/mL in α-minimal essential medium supplemented with 10% FBS. Differentiation is induced by addition of Retinoic acid (1 μM) and medium containing Retinoic acid replaced 2 days later. On day 4, cell aggregates are collected by centrifugation, separated to single cells by trypsin/EDTA treatment, replated onto poly-L-lysine-coated plates, and cultured in α-minimal essential medium supplemented with 10% FBS. On day 6, medium is replaced with neurobasal medium containing B27 supplement and 2 mM GlutaMAX. Medium is replaced every 2 days for an additional week[3].				

Solubility Information

A DRUG SCREENING EXPERT

Solubility	DMSO: 45 mg/mL (149.78 mM), Ethanol: 6 mg/mL (19.97 mM), H2O: <1
	mg/mL (insoluble or slightly soluble), (< 1 mg/ml refers to the product slightly
	soluble or insoluble)

Preparing Stock Solutions

	1mg	5mg	10mg	
1 mM	3.3285 mL	16.6423 mL	33.2845 mL	
5 mM	0.6657 mL	3.3285 mL	6.6569 mL	
10 mM	0.3328 mL	1.6642 mL	3.3285 mL	
50 mM	0.0666 mL	0.3328 mL	0.6657 mL	

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

Chen Q, et al. Retinoic acid regulates cell cycle progression and cell differentiation in human monocytic THP-1 cells. Exp Cell Res. 2004 Jul 1;297(1):68-81.

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