Deoxycorticosterone

Cat. No.:	HY-113414				
CAS No.:	64-85-7				
Molecular Formula:	C ₂₁ H ₃₀ O ₃				
Molecular Weight:	330.46				
Target:	Endogenous Metabolite; Mineralocorticoid Receptor				
Pathway:	Metabolic Enzyme/Protease; Vitamin D Related/Nuclear Receptor				
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 : ≥ 250 mg/mL (756.52 mM) * "≥" means soluble, but saturation unknown.						
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	3.0261 mL	15.1304 mL	30.2609 mL		
		5 mM	0.6052 mL	3.0261 mL	6.0522 mL		
		10 mM	0.3026 mL	1.5130 mL	3.0261 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: ≥ 2.5 mg/mL (7.57 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (7.57 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (7.57 mM); Clear solution						

BIOLOGICAL ACTIVITY Description Deoxycorticosterone is a steroid hormone produced by the adrenal gland that possesses mineralocorticoid activity and acts as an aldosterone precursor. Deoxycorticosterone is an agonist for O. mykiss mineralocorticoid receptor (rtMR) transcription with EC₅₀ of 0.16 nM^[3]. Deoxycorticosterone could acts as an immune stimulator in fish^[4]. IC₅₀ & Target Human Endogenous Metabolite rtMR 0.16 nM (EC50)

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In Vitro	Deoxycorticosterone enhances the rtMR transcriptional activity, involves in the milt fluidity regulation ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.				
In Vivo	Deoxycorticosterone (0.08 mg/kg, i.p, single dosage) increases expressions of C-type lysozyme and apolipoprotein A1 in spleen and gills, which indicates an immune stimulatory effect in Eurasian perch ^[4] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.				
	Animal Model:	Eurasian perch ^[4]			
	Dosage:	0.08 mg/kg			
	Administration:	i.p.			
	Result:	Upregulated mRNA expression of genes coding for 11β-HSD2 and MR in spleen. Unregulated expressions of C type lysozyme mRNA in spleen and Apo A1 mRNA in spleen and gills.			

CUSTOMER VALIDATION

- Proc Natl Acad Sci U S A. 2022 Apr 12;119(15):e2117004119.
- FASEB J. 2023 Apr;37(4):e22869.

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REFERENCES

[1]. Milla S, et al., Plasma 11-deoxycorticosterone (DOC) and mineralocorticoid receptor testicular expression during rainbow trout Oncorhynchus mykiss spermiation: implication with 17alpha, 20beta-dihydroxyprogesterone on the milt fluidity? Reprod Biol Endocrinol. 2008 May 19;6:19.

[2]. Sturm A, et al., 11-deoxycorticosterone is a potent agonist of the rainbow trout (Oncorhynchus mykiss) mineralocorticoid receptor. Endocrinology. 2005 Jan;146(1):47-55.

[3]. Mathieu C, et al., First evidence of the possible implication of the 11-deoxycorticosterone (DOC) in immune activity of Eurasian perch (Perca fluviatilis, L.): comparison with cortisol. Comp Biochem Physiol A Mol Integr Physiol. 2013 Jun;165(2):149-58.

[4]. KAGAWA CM, et al. Action of new steroids in blocking effects of aldosterone and desoxycorticosterone on salt. Science. 1957 Nov 15;126(3281):1015-6.

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

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