Product Data Sheet



MM11253

Cat. No.: HY-108530 CAS No.: 345952-44-5 Molecular Formula: $C_{28}H_{30}O_{2}S_{2}$ Molecular Weight: 462.67 Target: RAR/RXR

Pathway: Metabolic Enzyme/Protease Storage: Powder -20°C 3 years

> In solvent -80°C 6 months

-20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 125 mg/mL (270.17 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.1614 mL	10.8068 mL	21.6137 mL
	5 mM	0.4323 mL	2.1614 mL	4.3227 mL
	10 mM	0.2161 mL	1.0807 mL	2.1614 mL

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

BIOLOGICAL ACTIVITY

Description	MM11253 is a potent and selective RAR γ antagonist with an IC $_{50}$ of 44 nM. MM11253 has lower inhibition of RAR α , RAR β and RXR α . MM11253 blocks the growth inhibitory effects of RAR γ -selective agonists ^{[1][3]} .				
IC ₅₀ & Target	RARγ 44 nM (IC ₅₀)	RARγ 44 nM (IC ₅₀)	RARY 44 nM (IC ₅₀)	RARα 1000 nM (IC ₅₀)	
	RARα 1000 nM (IC ₅₀)	RARα 1000 nM (IC ₅₀)	RARβ >1000 nM (IC ₅₀)	RARβ >1000 nM (IC ₅₀)	
	RARβ >1000 nM (IC ₅₀)	RXRα >1000 nM (IC ₅₀)			
In Vitro	MM11253 blocks the ability of MM11254 and MM11389 to inhibit squamous cell carcinoma cell growth ^[2] .				

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

REFERENCES ————————————————————————————————————
[1]. M I Dawson, et al. Retinoic acid (RA) receptor transcriptional activation correlates with inhibition of 12-O-tetradecanoylphorbol-13-acetate-induced ornithine decarboxylase (ODC) activity by retinoids: a potential role for trans-RA-induced ZBP-89 in ODC inhibition. Int J Cancer. 2001 Jan 1;91(1):8-21.
[2]. Q Le, et al. Modulation of retinoic acid receptor function alters the growth inhibitory response of oral SCC cells to retinoids. Oncogene. 2000 Mar 9;19(11):1457-65.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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