# Data Sheet (Cat.No.T15748)



#### LG100754

# **Chemical Properties**

CAS No.: 180713-37-5

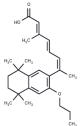
Formula: C26H36O3

Molecular Weight: 396.56

Appearance: no data available

Storage: Storage: 2005 for 2 years

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



# **Biological Description**

Description	LG100754 (UVI 2112) is an insulin sensitizer that functions through RXR. LG100754 is an RXR dimer modulator. LG100754 acts as an RXR: RXR homodimer antagonist, but functions as an agonist towards RXR: PPAR $\alpha$ and RXR: PPAR $\gamma$ heterodimers.		
Targets(IC50)	PPAR		
In vitro	LG100754 functions as an activator of endogenous RXR heterodimers, specifically facilitating the activity of RXR: PPARγ heterodimers. It plays a crucial role in regulating insulin-dependent signaling pathways. Additionally, LG100754 has been shown to mitigate TNFα-mediated insulin resistance in mature adipocytes[1].		
In vivo	LG100754 (100 mg/kg) fully blocks the enhance in glucose levels. It also shows that LG100754 can improve insulin resistance in vivo[1].		

# **Solubility Information**

Solubility	DMSO: 90.0 mg/mL (227.0 mM), Sonication is recommended.
	(< 1 mg/ml refers to the product slightly soluble or insoluble)

### **Preparing Stock Solutions**

	1mg	5mg	10mg
1 mM	2.5217 mL	12.6084 mL	25.2169 mL
5 mM	0.5043 mL	2.5217 mL	5.0434 mL
10 mM	0.2522 mL	1.2608 mL	2.5217 mL
50 mM	0.0504 mL	0.2522 mL	0.5043 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.

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#### Reference

Cesario RM, et al. The rexinoid LG100754 is a novel RXR:PPARgamma agonist and decreases glucose levels in vivo. Mol Endocrinol. 2001 Aug;15(8):1360-9.



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