Inhibitors

Product Data Sheet

PPARα/γ agonist 1

Molecular Weight: 281.35

Pathway: Cell Cycle/DNA Damage; Vitamin D Related/Nuclear Receptor

Storage: 4°C, stored under nitrogen, away from moisture

* In solvent : -80°C, 6 months; -20°C, 1 month (stored under nitrogen, away from

moisture)

PPAR

SOLVENT & SOLUBILITY

In Vitro

Target:

DMSO: 100 mg/mL (355.43 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.5543 mL	17.7715 mL	35.5429 mL
	5 mM	0.7109 mL	3.5543 mL	7.1086 mL
	10 mM	0.3554 mL	1.7771 mL	3.5543 mL

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

BIOLOGICAL ACTIVITY

Description	$PPAR\alpha/\gamma\ agonist\ 1\ is\ a\ potent\ and\ dual\ PPAR\alpha/\gamma\ partial\ agonist\ with\ EC_{50}\ values\ of\ 28\ nM\ and\ 69\ nM\ for\ PPAR\alpha\ and\ PPAR\alpha$	
	respectively. $PPAR\alpha/\gamma$ agonist 1 is a promising prototype for dyslipidemia and diabetes research ^[1] .	

 $\begin{array}{ccc} \text{IC}_{\text{50}}\,\&\,\text{Target} & \text{PPAR}\alpha & \text{PPAR}\gamma \\ & & 28\,\text{nM}\,(\text{EC50}) & 69\,\text{nM}\,(\text{EC50}) \end{array}$

REFERENCES

[1]. Luiz A Dutra, et al. Discovery of (E)-4-styrylphenoxy-propanamide: A dual PPARa/y partial agonist that regulates high-density lipoprotein-cholesterol levels, modulates adipogenesis, and improves glucose tolerance in diet-induced obese mice. Bioorg Chem. 2

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

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