



**MK-4074** 

**Solubility:** 

565.62

DMSO: 83.3 mg/mL (147.27 mM; Need ultrasonic)

**Observed Molecular Weight:** 

Catalog No: tcsc0029249

卫	Available Sizes
Size:	5mg
Size:	10mg
Size:	25mg
Size:	50mg
Size:	100mg
	Specifications
<b>CAS</b> 1039	<b>No:</b> 758-22-9
Form	iula: 31 <sup>N</sup> 3 <sup>O</sup> 6
<b>Path</b> Metal	way: polic Enzyme/Protease
<b>Targ</b> Acety	et: ·I-CoA Carboxylase
Purit	y / Grade:



## **Product Description**

MK-4074 is a liver-specific inhibitor of **acetyl-CoA carboxylase** ACC1 and ACC2 with  $IC_{50}$  values of approximately 3 nM.

IC50 & Target: IC50: 3 nM (Acetyl-CoA Carboxylase)[1]

In Vitro: MK-4074 strongly inhibits both ACC1 and ACC2 with  $IC_{50}$  values of approximately 3 nM. MK-4074 is highly liver specific because it is a substrate of organic anion transport protein (OATP) transporters that are present only in hepatocytes, and excretion of MK-4074 from hepatocytes into bile is dependent on the MRP2 efflux transporter<sup>[1]</sup>.

In Vivo: In male KKAy mice, a mouse model of obesity, type 2 diabetes, and fatty liver, a single oral dose of MK-4074 (0.3-3 mg/kg) significantly decreases DNL in a dose-dependent manner with an  $ID_{50}$  value of 0.9 mg/kg 1 hr post-administration. In a time course study, MK-4074 orally at 30 mg/kg reduces hepatic DNL by 83%, 70%, and 51% at 4, 8, and 12 hr post-dose, respectively. Single oral doses of MK-4074 at 30 and 100 mg/kg significantly increases plasma total ketones, a surrogate biomarker for hepatic FAO, by 1.5-fold to 3-fold for up to 8 hr<sup>[1]</sup>.

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!