

Product Name : COH-SR4 (Mitochondria uncoupler SR4)

Synonyms : —

Cat No. : M22757

CAS Number : 73439-19-7

Molecular Formula : C₁₃H₈Cl₄N₂O

Formula Weight : 350.02

Chemical Name : —

Description : SR4 is a uncoupler of mitochondrial oxidative phosphorylation. SR4 modulating amp-dependent kinase (ampk)-mammalian target of rapamycin (mtor) signaling, and inhibiting proliferation of hepg2 hepatocarcinoma cells SR4 is a novel mitochondrial uncoupler with anti-obesity and anti-diabetic properties. SR4 increased oxygen consumption, dissipated mitochondrial membrane potential, induced mitochondrial swelling, and decreased intracellular ATP in cultured cells and isolated liver mitochondria. Oral feeding of SR4 significantly reduced body weight gain, improved glycemic control and insulin resistance, and prevented dyslipidemia in both high-fat-diet (HFD) induced obese and diabetic db/db mice. SR4 treatment also decreased liver triglycerides and prevented hepatic steatosis in both animal models. Mitochondrial uncoupling of SR4 results to activation of AMP-activated protein kinase (AMPK), leading to the phosphorylation and inhibition of acetyl-CoA carboxylase (ACC). Gene analyses by RT-PCR showed SR4 significantly suppressed the mRNA expression of several lipogenic genes and gluconeogenic genes in the liver of HFD obese mice. RNA sequencing analysis showed that 642 genes were differentially expressed in liver of db/db mice after SR4 treatment (217 upregulated, 425 down-regulated). Gene ontology analysis by DAVID indicated SR4 upregulated amino acid metabolism and down-regulated lipid and fatty acid synthesis and glucose metabolism. These studies demonstrate that SR4 may be a promising compound for treatment of T2DM and obesity.

Pathway : Others

Target : Other Targets

Receptor : Others

Solubility : —

SMILES : Clc1cc(Cl)cc(NC(=O)Nc2cc(Cl)cc(Cl)c2)c1

Storage : (-20°C)

Stability : ≥ 2 years

Reference :

1. Sharad S Singhal, James L Figarola, Jyotsana Singhal, et al. COH-SR4, a Novel Mitochondrial Uncoupler, Improves Metabolic Alterations in Obese and Diabetic Mice.