

ACC inhibitor alone or co-administered with a DGAT2 in non-alcoholic fatty liver disease: two parallel, placebo- controlled trials

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Why Do So Many Nonalcoholic Steatohepatitis Trials Fail?. <i>Gastroenterology</i> , 2023, 165, 5-10.	1.3	59
2	Phase IIa results for potential NAFLD therapy. <i>Nature Reviews Endocrinology</i> , 2022, 18, 2.	9.6	0
3	Targeting ACC for NASH resolution. <i>Trends in Molecular Medicine</i> , 2022, 28, 5-7.	6.7	9
4	An RNAi therapeutic targeting hepatic DGAT2 in a genetically obese mouse model of nonalcoholic steatohepatitis. <i>Molecular Therapy</i> , 2022, 30, 1329-1342.	8.2	18
5	Lipogenesis inhibitors: therapeutic opportunities and challenges. <i>Nature Reviews Drug Discovery</i> , 2022, 21, 283-305.	46.4	124
6	Pharmacotherapy for Non-Alcoholic Fatty Liver Disease: Emerging Targets and Drug Candidates. <i>Biomedicines</i> , 2022, 10, 274.	3.2	22
7	New avenues for NASH therapy by targeting ACC. <i>Cell Metabolism</i> , 2022, 34, 191-193.	16.2	6
8	Metabolic dysfunction and cancer in HCV: Shared pathways and mutual interactions. <i>Journal of Hepatology</i> , 2022, 77, 219-236.	3.7	16
9	Updates on novel pharmacotherapeutics for the treatment of nonalcoholic steatohepatitis. <i>Acta Pharmacologica Sinica</i> , 2022, 43, 1180-1190.	6.1	22
10	Acetyl-CoA Carboxylases and Diseases. <i>Frontiers in Oncology</i> , 2022, 12, 836058.	2.8	52
11	Lipid alterations in chronic liver disease and liver cancer. <i>JHEP Reports</i> , 2022, 4, 100479.	4.9	69
13	Efficacy and safety of an orally administered DGAT2 inhibitor alone or coadministered with a liver-targeted ACC inhibitor in adults with non-alcoholic steatohepatitis (NASH): rationale and design of the phase II, dose-ranging, dose-finding, randomised, placebo-controlled MIRNA (Metabolic) TJ ETQq1 1 0.784314 1.9fgBT /Overlock 10T	1.9	18
14	Pharmacokinetics, Mass Balance, Metabolism, and Excretion of the Liver-Targeted Acetyl-CoA Carboxylase Inhibitor PF-05221304 (Clesacostat) in Humans. <i>Xenobiotica</i> , 2022, , 1-45.	1.1	1
15	Pathophysiological Mechanisms in Non-Alcoholic Fatty Liver Disease: From Drivers to Targets. <i>Biomedicines</i> , 2022, 10, 46.	3.2	10
16	Quantitative proteomics of HFD-induced fatty liver uncovers novel transcription factors of lipid metabolism. <i>International Journal of Biological Sciences</i> , 2022, 18, 3298-3312.	6.4	6
17	Lipid metabolism in T cell signaling and function. <i>Nature Chemical Biology</i> , 2022, 18, 470-481.	8.0	46
18	Reprogramming of Hepatic Metabolism and Microenvironment in Nonalcoholic Steatohepatitis. <i>Annual Review of Nutrition</i> , 2022, 42, 91-113.	10.1	20
19	Nonalcoholic Steatohepatitis Drug Development Pipeline: An Update. <i>Seminars in Liver Disease</i> , 2022, 42, 379-400.	3.6	17

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20	Impact of NAFLD and its pharmacotherapy on lipid profile and CVD. <i>Atherosclerosis</i> , 2022, 355, 30-44.	0.8	7
21	Effects of Metabolism on Macrophage Polarization Under Different Disease Backgrounds. <i>Frontiers in Immunology</i> , 0, 13, .	4.8	10
22	A Quantitative Systems Pharmacology Model of Liver Lipid Metabolism for Investigation of Non-Alcoholic Fatty Liver Disease. <i>Frontiers in Pharmacology</i> , 0, 13, .	3.5	0
23	Comprehensive molecular mechanisms and clinical therapy in nonalcoholic steatohepatitis: An overview and current perspectives. <i>Metabolism: Clinical and Experimental</i> , 2022, 134, 155264.	3.4	12
24	The postnatal resolution of developmental toxicity induced by pharmacological diacylglycerol acyltransferase 2 (DGAT2) inhibition during gestation in rats. <i>Toxicological Sciences</i> , 0, , .	3.1	0
25	Targeted therapeutics and novel signaling pathways in non-alcohol-associated fatty liver/steatohepatitis (NAFL/NASH). <i>Signal Transduction and Targeted Therapy</i> , 2022, 7, .	17.1	90
26	Paradoxical activation of transcription factor SREBP1c and de novo lipogenesis by hepatocyte-selective ATP-citrate lyase depletion in obese mice. <i>Journal of Biological Chemistry</i> , 2022, 298, 102401.	3.4	5
27	The mitochondrial fission protein Drp1 in liver is required to mitigate NASH and prevents the activation of the mitochondrial ISR. <i>Molecular Metabolism</i> , 2022, 64, 101566.	6.5	14
28	Effectiveness of Lifestyle Interventions for Nonalcoholic Fatty Liver Disease Treatment. , 0, , .		1
29	Progress in research of lipogenesis inhibitors for treatment of nonalcoholic fatty liver disease. <i>World Chinese Journal of Digestology</i> , 2022, 30, 735-742.	0.1	0
30	Association between de novo lipogenesis susceptibility genes and coronary artery disease. <i>Nutrition, Metabolism and Cardiovascular Diseases</i> , 2022, 32, 2883-2889.	2.6	4
31	Molecular mechanisms of metabolic associated fatty liver disease (MAFLD): functional analysis of lipid metabolism pathways. <i>Clinical Science</i> , 2022, 136, 1347-1366.	4.3	56
32	FASN inhibition targets multiple drivers of NASH by reducing steatosis, inflammation and fibrosis in preclinical models. <i>Scientific Reports</i> , 2022, 12, .	3.3	17
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39	Inulin intervention attenuates hepatic steatosis in rats via modulating gut microbiota and maintaining intestinal barrier function. Food Research International, 2023, 163, 112309.	6.2	17
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