

Marica Meroni

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4388290/publications.pdf>

Version: 2024-02-01

51
papers

3,122
citations

218592

26
h-index

214721

47
g-index

58
all docs

58
docs citations

58
times ranked

3918
citing authors

#	ARTICLE	IF	CITATIONS
1	The MBOAT7-TMC4 Variant rs641738 Increases Risk of Nonalcoholic Fatty Liver Disease in Individuals of European Descent. <i>Gastroenterology</i> , 2016, 150, 1219-1230.e6.	0.6	506
2	Genome-wide association study of non-alcoholic fatty liver and steatohepatitis in a histologically characterised cohort. <i>Journal of Hepatology</i> , 2020, 73, 505-515.	1.8	279
3	MBOAT7 rs641738 variant and hepatocellular carcinoma in non-cirrhotic individuals. <i>Scientific Reports</i> , 2017, 7, 4492.	1.6	193
4	Non-invasive stratification of hepatocellular carcinoma risk in non-alcoholic fatty liver using polygenic risk scores. <i>Journal of Hepatology</i> , 2021, 74, 775-782.	1.8	193
5	Hepatocyte Notch activation induces liver fibrosis in nonalcoholic steatohepatitis. <i>Science Translational Medicine</i> , 2018, 10, .	5.8	151
6	Macrophage MerTK Promotes Liver Fibrosis in Nonalcoholic Steatohepatitis. <i>Cell Metabolism</i> , 2020, 31, 406-421.e7.	7.2	141
7	Liver fat accumulation is associated with circulating PCSK9. <i>Annals of Medicine</i> , 2016, 48, 384-391.	1.5	119
8	Alcohol or Gut Microbiota: Who Is the Guilty?. <i>International Journal of Molecular Sciences</i> , 2019, 20, 4568.	1.8	106
9	miRNA Signature in NAFLD: A Turning Point for a Non-Invasive Diagnosis. <i>International Journal of Molecular Sciences</i> , 2018, 19, 3966.	1.8	98
10	The rs2294918 E434K variant modulates patatin-like phospholipase domain-containing 3 expression and liver damage. <i>Hepatology</i> , 2016, 63, 787-798.	3.6	93
11	PNPLA3 overexpression results in reduction of proteins predisposing to fibrosis. <i>Human Molecular Genetics</i> , 2016, 25, ddd341.	1.4	86
12	Rare Pathogenic Variants Predispose to Hepatocellular Carcinoma in Nonalcoholic Fatty Liver Disease. <i>Scientific Reports</i> , 2019, 9, 3682.	1.6	85
13	The Role of Probiotics in Nonalcoholic Fatty Liver Disease: A New Insight into Therapeutic Strategies. <i>Nutrients</i> , 2019, 11, 2642.	1.7	81
14	Genetic and Epigenetic Modifiers of Alcoholic Liver Disease. <i>International Journal of Molecular Sciences</i> , 2018, 19, 3857.	1.8	75
15	Liver transcriptomics highlights interleukin-32 as novel NAFLD-related cytokine and candidate biomarker. <i>Gut</i> , 2020, 69, 1855-1866.	6.1	75
16	Mboat7 down-regulation by hyper-insulinemia induces fat accumulation in hepatocytes. <i>EBioMedicine</i> , 2020, 52, 102658.	2.7	71
17	Insulin resistance promotes Lysyl Oxidase Like 2 induction and fibrosis accumulation in non-alcoholic fatty liver disease. <i>Clinical Science</i> , 2017, 131, 1301-1315.	1.8	64
18	Nutrition and Genetics in NAFLD: The Perfect Binomium. <i>International Journal of Molecular Sciences</i> , 2020, 21, 2986.	1.8	60

#	ARTICLE	IF	CITATIONS
19	Mitochondrial dynamics and nonalcoholic fatty liver disease (NAFLD): new perspectives for a fairy-tale ending?. <i>Metabolism: Clinical and Experimental</i> , 2021, 117, 154708.	1.5	59
20	Î²-Klotho gene variation is associated with liver damage in children with NAFLD. <i>Journal of Hepatology</i> , 2020, 72, 411-419.	1.8	48
21	The role of insulin resistance in nonalcoholic steatohepatitis and liver disease development – a potential therapeutic target?. <i>Expert Review of Gastroenterology and Hepatology</i> , 2016, 10, 229-242.	1.4	44
22	TM6SF2/PNPLA3/MBOAT7 Loss-of-Function Genetic Variants Impact on NAFLD Development and Progression Both in Patients and in InÂVtro Models. <i>Cellular and Molecular Gastroenterology and Hepatology</i> , 2022, 13, 759-788.	2.3	44
23	Fibronectin Type III Domain-Containing Protein 5 rs3480 A>G Polymorphism, Irisin, and Liver Fibrosis in Patients With Nonalcoholic Fatty Liver Disease. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017, 102, 2660-2669.	1.8	42
24	PCSK7 gene variation bridges atherogenic dyslipidemia with hepatic inflammation in NAFLD patients. <i>Journal of Lipid Research</i> , 2019, 60, 1144-1153.	2.0	42
25	Protein phosphatase 1 regulatory subunit 3B gene variation protects against hepatic fat accumulation and fibrosis in individuals at high risk of nonalcoholic fatty liver disease. <i>Hepatology Communications</i> , 2018, 2, 666-675.	2.0	38
26	MBOAT7 down-regulation by genetic and environmental factors predisposes to MAFLD. <i>EBioMedicine</i> , 2020, 57, 102866.	2.7	38
27	Genetics Is of the Essence to Face NAFLD. <i>Biomedicines</i> , 2021, 9, 1359.	1.4	30
28	mir-101-3p Downregulation Promotes Fibrogenesis by Facilitating Hepatic Stellate Cell Transdifferentiation During Insulin Resistance. <i>Nutrients</i> , 2019, 11, 2597.	1.7	24
29	MAFLD in COVID-19 patients: an insidious enemy. <i>Expert Review of Gastroenterology and Hepatology</i> , 2020, 14, 867-872.	1.4	23
30	Remodeling of Mitochondrial Plasticity: The Key Switch from NAFLD/NASH to HCC. <i>International Journal of Molecular Sciences</i> , 2021, 22, 4173.	1.8	23
31	Notch signaling and progenitor/ductular reaction in steatohepatitis. <i>PLoS ONE</i> , 2017, 12, e0187384.	1.1	18
32	Low Lipoprotein(a) Levels Predict Hepatic Fibrosis in Patients With Nonalcoholic Fatty Liver Disease. <i>Hepatology Communications</i> , 2022, 6, 535-549.	2.0	18
33	The rs599839 A>G Variant Disentangles Cardiovascular Risk and Hepatocellular Carcinoma in NAFLD Patients. <i>Cancers</i> , 2021, 13, 1783.	1.7	16
34	Impact of Sarcopenia and Myosteatosi s in Non-Cirrhotic Stages of Liver Diseases: Similarities and Differences across Aetiologies and Possible Therapeutic Strategies. <i>Biomedicines</i> , 2022, 10, 182.	1.4	15
35	NDP-MSH treatment recovers marginal lungs during ex vivo lung perfusion (EVL P). <i>Peptides</i> , 2021, 141, 170552.	1.2	12
36	The KLB rs17618244 gene variant is associated with fibrosing MAFLD by promoting hepatic stellate cell activation. <i>EBioMedicine</i> , 2021, 65, 103249.	2.7	11

#	ARTICLE	IF	CITATIONS
37	Hepatic IRF3 fuels dysglycemia in obesity through direct regulation of <i>Ppp2r1b</i> . <i>Science Translational Medicine</i> , 2022, 14, eabh3831.	5.8	11
38	Neurotensin up-regulation is associated with advanced fibrosis and hepatocellular carcinoma in patients with MAFLD. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2020, 1865, 158765.	1.2	10
39	Genetics, Immunity and Nutrition Boost the Switching from NASH to HCC. <i>Biomedicines</i> , 2021, 9, 1524.	1.4	10
40	PD-1/PD-L1 Immuno-Mediated Therapy in NAFLD: Advantages and Obstacles in the Treatment of Advanced Disease. <i>International Journal of Molecular Sciences</i> , 2022, 23, 2707.	1.8	9
41	MAFLD definition underestimates the risk to develop HCC in genetically predisposed patients. <i>Journal of Internal Medicine</i> , 2022, 291, 374-376.	2.7	8
42	Impact of natural neuromedin-B receptor variants on iron metabolism. <i>American Journal of Hematology</i> , 2020, 95, 167-177.	2.0	7
43	From Environment to Genome and Back: A Lesson from HFE Mutations. <i>International Journal of Molecular Sciences</i> , 2020, 21, 3505.	1.8	7
44	Cutting-Edge Therapies and Novel Strategies for Acute Intermittent Porphyria: Step-by-Step towards the Solution. <i>Biomedicines</i> , 2022, 10, 648.	1.4	7
45	Î±-Lipoic Acid Improves Hepatic Metabolic Dysfunctions in Acute Intermittent Porphyria: A Proof-of-Concept Study. <i>Diagnostics</i> , 2021, 11, 1628.	1.3	5
46	Genetic and metabolic factors: the perfect combination to treat metabolic associated fatty liver disease. <i>Exploration of Medicine</i> , 2020, 1, 218-243.	1.5	4
47	PS-005-Evaluation of neuromedin-B receptor variants effect on iron metabolism and liver disease. <i>Journal of Hepatology</i> , 2019, 70, e7-e8.	1.8	0
48	PS-006-MBOAT7 downregulation induces hepatic lipid accumulation. <i>Journal of Hepatology</i> , 2019, 70, e8.	1.8	0
49	THU-323-Impact of genetic polymorphisms associated with NAFLD on hepatic and vascular complications in diabetes. <i>Journal of Hepatology</i> , 2019, 70, e302.	1.8	0
50	FRI-320-TM6SF2 silencing impairs lipid metabolism and trafficking in HepG2 cells carrying the I148M PNPLA3 variant and MBOAT7 deletion. <i>Journal of Hepatology</i> , 2019, 70, e536-e537.	1.8	0
51	FRI-333-ATG7 genetic variant and defective autophagy: A novel risk factor for non-alcoholic fatty liver disease progression in patients with type 2 diabetes mellitus. <i>Journal of Hepatology</i> , 2019, 70, e542.	1.8	0