

Raffaella Rametta

List of Publications by Citations

Source: <https://exaly.com/author-pdf/7224896/raffaella-rametta-publications-by-citations.pdf>

Version: 2024-04-26

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

62

papers

3,951

citations

35

h-index

62

g-index

62

ext. papers

4,680

ext. citations

5.7

avg, IF

4.68

L-index

#	Paper	IF	Citations
62	Homozygosity for the patatin-like phospholipase-3/adiponutrin I148M polymorphism influences liver fibrosis in patients with nonalcoholic fatty liver disease. <i>Hepatology</i> , 2010 , 51, 1209-17	11.2	445
61	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	13.3	347
60	Transmembrane 6 superfamily member 2 gene variant disentangles nonalcoholic steatohepatitis from cardiovascular disease. <i>Hepatology</i> , 2015 , 61, 506-14	11.2	311
59	PNPLA3 has retinyl-palmitate lipase activity in human hepatic stellate cells. <i>Human Molecular Genetics</i> , 2014 , 23, 4077-85	5.6	230
58	Statin use and non-alcoholic steatohepatitis in at risk individuals. <i>Journal of Hepatology</i> , 2015 , 63, 705-12	13.4	227
57	Patatin-like phospholipase domain-containing 3 I148M polymorphism, steatosis, and liver damage in chronic hepatitis C. <i>Hepatology</i> , 2011 , 53, 791-9	11.2	199
56	Increased expression and activity of the transcription factor FOXO1 in nonalcoholic steatohepatitis. <i>Diabetes</i> , 2008 , 57, 1355-62	0.9	128
55	Genetic variants regulating insulin receptor signalling are associated with the severity of liver damage in patients with non-alcoholic fatty liver disease. <i>Gut</i> , 2010 , 59, 267-73	19.2	117
54	DJ-1 modulates alpha-synuclein aggregation state in a cellular model of oxidative stress: relevance for Parkinson's disease and involvement of HSP70. <i>PLoS ONE</i> , 2008 , 3, e1884	3.7	104
53	Dietary iron overload induces visceral adipose tissue insulin resistance. <i>American Journal of Pathology</i> , 2013 , 182, 2254-63	5.8	101
52	MERTK rs4374383 polymorphism affects the severity of fibrosis in non-alcoholic fatty liver disease. <i>Journal of Hepatology</i> , 2016 , 64, 682-90	13.4	79
51	Alpha 1-antitrypsin mutations in NAFLD: high prevalence and association with altered iron metabolism but not with liver damage. <i>Hepatology</i> , 2006 , 44, 857-64	11.2	74
50	Protective effect of TAT-delivered alpha-synuclein: relevance of the C-terminal domain and involvement of HSP70. <i>FASEB Journal</i> , 2004 , 18, 1713-5	0.9	72
49	The rs2294918 E434K variant modulates patatin-like phospholipase domain-containing 3 expression and liver damage. <i>Hepatology</i> , 2016 , 63, 787-98	11.2	70
48	Serum hepcidin and macrophage iron correlate with MCP-1 release and vascular damage in patients with metabolic syndrome alterations. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2011 , 31, 683-90	9.4	67
47	A randomized trial of iron depletion in patients with nonalcoholic fatty liver disease and hyperferritinemia. <i>World Journal of Gastroenterology</i> , 2014 , 20, 3002-10	5.6	66
46	Hepatic notch signaling correlates with insulin resistance and nonalcoholic fatty liver disease. <i>Diabetes</i> , 2013 , 62, 4052-62	0.9	65

45	The APOC3 T-455C and C-482T promoter region polymorphisms are not associated with the severity of liver damage independently of PNPLA3 I148M genotype in patients with nonalcoholic fatty liver. <i>Journal of Hepatology</i> , 2011 , 55, 1409-14	13.4	65
44	Serum ferritin levels are associated with vascular damage in patients with nonalcoholic fatty liver disease. <i>Nutrition, Metabolism and Cardiovascular Diseases</i> , 2011 , 21, 568-75	4.5	62
43	PNPLA3 I148M Variant Influences Circulating Retinol in Adults with Nonalcoholic Fatty Liver Disease or Obesity. <i>Journal of Nutrition</i> , 2015 , 145, 1687-91	4.1	59
42	Paradoxical dissociation between hepatic fat content and de novo lipogenesis due to PNPLA3 sequence variant. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015 , 100, E821-5	5.6	55
41	Iron-dependent regulation of MDM2 influences p53 activity and hepatic carcinogenesis. <i>American Journal of Pathology</i> , 2010 , 176, 1006-17	5.8	54
40	Beta-globin mutations are associated with parenchymal siderosis and fibrosis in patients with non-alcoholic fatty liver disease. <i>Journal of Hepatology</i> , 2010 , 53, 927-33	13.4	51
39	The I148M PNPLA3 polymorphism influences serum adiponectin in patients with fatty liver and healthy controls. <i>BMC Gastroenterology</i> , 2012 , 12, 111	3	50
38	LPIN1 rs13412852 polymorphism in pediatric nonalcoholic fatty liver disease. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2012 , 54, 588-93	2.8	46
37	Lack of association between peroxisome proliferator-activated receptors alpha and gamma2 polymorphisms and progressive liver damage in patients with non-alcoholic fatty liver disease: a case control study. <i>BMC Gastroenterology</i> , 2010 , 10, 102	3	46
36	The TM6SF2 E167K genetic variant induces lipid biosynthesis and reduces apolipoprotein B secretion in human hepatic 3D spheroids. <i>Scientific Reports</i> , 2019 , 9, 11585	4.9	44
35	Genetic and Epigenetic Modifiers of Alcoholic Liver Disease. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	44
34	Prevalence and Risk Factors of Significant Fibrosis in Patients With Nonalcoholic Fatty Liver Without Steatohepatitis. <i>Clinical Gastroenterology and Hepatology</i> , 2019 , 17, 2310-2319.e6	6.9	42
33	Patatin-like phospholipase domain containing-3 gene I148M polymorphism, steatosis, and liver damage in hereditary hemochromatosis. <i>World Journal of Gastroenterology</i> , 2012 , 18, 2813-20	5.6	42
32	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	6.5	38
31	The A736V Tmprss6 polymorphism influences hepatic iron overload in nonalcoholic fatty liver disease. <i>PLoS ONE</i> , 2012 , 7, e48804	3.7	37
30	Mboat7 down-regulation by hyper-insulinemia induces fat accumulation in hepatocytes. <i>EBioMedicine</i> , 2020 , 52, 102658	8.8	36
29	PNPLA3 I148M polymorphism, clinical presentation, and survival in patients with hepatocellular carcinoma. <i>PLoS ONE</i> , 2013 , 8, e75982	3.7	36
28	The UCP2 -866G>A promoter region polymorphism is associated with nonalcoholic steatohepatitis. <i>Liver International</i> , 2015 , 35, 1574-80	7.9	35

27	Liver transcriptomics highlights interleukin-32 as novel NAFLD-related cytokine and candidate biomarker. <i>Gut</i> , 2020 , 69, 1855-1866	19.2	34
26	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, 229-42	4.2	32
25	High fat diet subverts hepatocellular iron uptake determining dysmetabolic iron overload. <i>PLoS ONE</i> , 2015 , 10, e0116855	3.7	30
24	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	6	30
23	gene variation bridges atherogenic dyslipidemia with hepatic inflammation in NAFLD patients. <i>Journal of Lipid Research</i> , 2019 , 60, 1144-1153	6.3	27
22	Effect of the A736V TMPRSS6 polymorphism on the penetrance and clinical expression of hereditary hemochromatosis. <i>Journal of Hepatology</i> , 2012 , 57, 1319-25	13.4	27
21	HFE mutations modulate the effect of iron on serum hepcidin-25 in chronic hemodialysis patients. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2009 , 4, 1331-7	6.9	24
20	Hepcidin resistance in dysmetabolic iron overload. <i>Liver International</i> , 2016 , 36, 1540-8	7.9	22
19	A tetra-primer amplification refractory mutation system polymerase chain reaction for the evaluation of rs12979860 IL28B genotype. <i>Journal of Viral Hepatitis</i> , 2011 , 18, 628-30	3.4	22
18	Hepatic steatosis and PNPLA3 I148M variant are associated with serum Fetuin-A independently of insulin resistance. <i>European Journal of Clinical Investigation</i> , 2014 , 44, 627-33	4.6	21
17	HFE gene mutations and oxidative stress influence serum ferritin, associated with vascular damage, in hemodialysis patients. <i>American Journal of Nephrology</i> , 2007 , 27, 101-7	4.6	19
16	The A736V TMPRSS6 polymorphism influences hepcidin and iron metabolism in chronic hemodialysis patients: TMPRSS6 and hepcidin in hemodialysis. <i>BMC Nephrology</i> , 2013 , 14, 48	2.7	17
15	Liver transplantation for hepatocellular carcinoma in a patient with a novel telomerase mutation and steatosis. <i>Journal of Hepatology</i> , 2013 , 58, 399-401	13.4	11
14	GNPAT rs11558492 is not a Major Modifier of Iron Status: Study of Italian Hemochromatosis Patients and Blood Donors. <i>Annals of Hepatology</i> , 2017 , 16, 451-456	3.1	11
13	HFE genotype influences erythropoiesis support requirement in hemodialysis patients: a prospective study. <i>American Journal of Nephrology</i> , 2008 , 28, 311-6	4.6	11
12	PCSK9 rs11591147 R46L loss-of-function variant protects against liver damage in individuals with NAFLD. <i>Liver International</i> , 2021 , 41, 321-332	7.9	10
11	GNPAT rs11558492 is not a Major Modifier of Iron Status: Study of Italian Hemochromatosis Patients and Blood Donors. <i>Annals of Hepatology</i> , 2017 , 16, 451-456	3.1	9
10	Proprotein convertase 7 rs236918 associated with liver fibrosis in Italian patients with HFE-related hemochromatosis. <i>Journal of Gastroenterology and Hepatology (Australia)</i> , 2016 , 31, 1342-8	4	9

9	Ceruloplasmin gene variants are associated with hyperferritinemia and increased liver iron in patients with NAFLD. <i>Journal of Hepatology</i> , 2021 , 75, 506-513	13.4	8
8	Dysmetabolic Hyperferritinemia and Dysmetabolic Iron Overload Syndrome (DIOS): Two Related Conditions or Different Entities?. <i>Current Pharmaceutical Design</i> , 2020 , 26, 1025-1035	3.3	7
7	A novel alpha1-antitrypsin null variant (PiQ0Milano). <i>World Journal of Hepatology</i> , 2013 , 5, 458-61	3.4	7
6	Increased circulating adiponectin in males with chronic HCV hepatitis. <i>European Journal of Internal Medicine</i> , 2015 , 26, 635-9	3.9	5
5	From Environment to Genome and Back: A Lesson from Mutations. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	5
4	GNPAT p.D519G variant and iron metabolism during oral iron tolerance test. <i>Hepatology</i> , 2017 , 65, 384-385	3	3
3	Adipocyte PHLPP2 inhibition prevents obesity-induced fatty liver. <i>Nature Communications</i> , 2021 , 12, 1822	17.4	3
2	Impact of natural neuromedin-B receptor variants on iron metabolism. <i>American Journal of Hematology</i> , 2020 , 95, 167-177	7.1	2
1	Juvenile hemochromatosis associated with heterozygosity for novel hemojuvelin mutations and with unknown cofactors. <i>Annals of Hepatology</i> , 2014 , 13, 568-71	3.1	1