## Markus Scholz

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4339312/publications.pdf

Version: 2024-02-01

160 papers 10,585 citations

38 h-index 92 g-index

166 all docs

166
docs citations

166 times ranked 18577 citing authors

#	Article	IF	CITATIONS
1	A comprehensive 1000 Genomes–based genome-wide association meta-analysis of coronary artery disease. Nature Genetics, 2015, 47, 1121-1130.	9.4	2,054
2	Circular non-coding RNA ANRIL modulates ribosomal RNA maturation and atherosclerosis in humans. Nature Communications, $2016$ , $7$ , $12429$ .	<b>5.</b> 8	859
3	Large-scale cis- and trans-eQTL analyses identify thousands of genetic loci and polygenic scores that regulate blood gene expression. Nature Genetics, 2021, 53, 1300-1310.	9.4	590
4	A catalog of genetic loci associated with kidney function from analyses of a million individuals. Nature Genetics, 2019, 51, 957-972.	9.4	549
5	Study of 300,486 individuals identifies 148 independent genetic loci influencing general cognitive function. Nature Communications, 2018, 9, 2098.	5 <b>.</b> 8	484
6	<i>ANRIL</i> Expression Is Associated With Atherosclerosis Risk at Chromosome 9p21. Arteriosclerosis, Thrombosis, and Vascular Biology, 2010, 30, 620-627.	1.1	402
7	Propionic Acid Shapes the Multiple Sclerosis Disease Course by an Immunomodulatory Mechanism. Cell, 2020, 180, 1067-1080.e16.	13.5	367
8	The power of genetic diversity in genome-wide association studies of lipids. Nature, 2021, 600, 675-679.	13.7	353
9	Alu Elements in ANRIL Non-Coding RNA at Chromosome 9p21 Modulate Atherogenic Cell Functions through Trans-Regulation of Gene Networks. PLoS Genetics, 2013, 9, e1003588.	1.5	323
10	The LIFE-Adult-Study: objectives and design of a population-based cohort study with 10,000 deeply phenotyped adults in Germany. BMC Public Health, 2015, 15, 691.	1.2	287
11	Target genes, variants, tissues and transcriptional pathways influencing human serum urate levels. Nature Genetics, 2019, 51, 1459-1474.	9.4	251
12	Genetic architecture of subcortical brain structures in 38,851 individuals. Nature Genetics, 2019, 51, 1624-1636.	9.4	192
13	Genetic Regulation of Serum Phytosterol Levels and Risk of Coronary Artery Disease. Circulation: Cardiovascular Genetics, 2010, 3, 331-339.	5.1	141
14	Genome-wide association meta-analyses and fine-mapping elucidate pathways influencing albuminuria. Nature Communications, 2019, 10, 4130.	5.8	133
15	GWAS and colocalization analyses implicate carotid intima-media thickness and carotid plaque loci in cardiovascular outcomes. Nature Communications, 2018, 9, 5141.	5.8	119
16	Secretory Phospholipase A2-IIA and Cardiovascular Disease. Journal of the American College of Cardiology, 2013, 62, 1966-1976.	1.2	115
17	Genome-wide analysis identifies novel susceptibility loci for myocardial infarction. European Heart Journal, 2021, 42, 919-933.	1.0	113
18	Identification of Adipokine Clusters Related to Parameters of Fat Mass, Insulin Sensitivity and Inflammation. PLoS ONE, 2014, 9, e99785.	1.1	107

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19	Polymorphisms at <i>PRSS1–PRSS2</i> and <i>CLDN2–MORC4</i> loci associate with alcoholic and non-alcoholic chronic pancreatitis in a European replication study. Gut, 2015, 64, 1426-1433.	6.1	105
20	Genome-wide association study identifies inversion in the <i>CTRB1-CTRB2</i> locus to modify risk for alcoholic and non-alcoholic chronic pancreatitis. Gut, 2018, 67, 1855-1863.	6.1	97
21	Novel loci for childhood body mass index and shared heritability with adult cardiometabolic traits. PLoS Genetics, 2020, 16, e1008718.	1.5	95
22	Dissecting the genetics of the human transcriptome identifies novel trait-related <i>trans </i> eQTLs and corroborates the regulatory relevance of non-protein coding loci. Human Molecular Genetics, 2015, 24, 4746-4763.	1.4	94
23	Relations between lipoprotein(a) concentrations, LPA genetic variants, and the risk of mortality in patients with established coronary heart disease: a molecular and genetic association study. Lancet Diabetes and Endocrinology,the, 2017, 5, 534-543.	5.5	84
24	The novel cystatin C, lactate, interleukin-6, and N-terminal pro-B-type natriuretic peptide (CLIP)-based mortality risk score in cardiogenic shock after acute myocardial infarction. European Heart Journal, 2021, 42, 2344-2352.	1.0	68
25	<i>MB-COMT</i> promoter DNA methylation is associated with working-memory processing in schizophrenia patients and healthy controls. Epigenetics, 2014, 9, 1101-1107.	1.3	65
26	Genetically determined NLRP3 inflammasome activation associates with systemic inflammation and cardiovascular mortality. European Heart Journal, 2021, 42, 1742-1756.	1.0	63
27	Genetic correlations and genome-wide associations of cortical structure in general population samples of 22,824 adults. Nature Communications, 2020, 11, 4796.	5.8	61
28	Genetic variation in the Sorbs of eastern Germany in the context of broader European genetic diversity. European Journal of Human Genetics, 2011, 19, 995-1001.	1.4	59
29	Prognostic and Pathogenic Role of Angiopoietin-1 and -2 in Pneumonia. American Journal of Respiratory and Critical Care Medicine, 2018, 198, 220-231.	2.5	58
30	Relationship Between 12 Adipocytokines and Distinct Components of the Metabolic Syndrome. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 1015-1023.	1.8	55
31	Genetic Factors of the Disease Course after Sepsis: A Genome-Wide Study for 28 Day Mortality. EBioMedicine, 2016, 12, 239-246.	2.7	52
32	Parameters of pulse wave velocity: determinants and reference values assessed in the population-based study LIFE-Adult. Clinical Research in Cardiology, 2018, 107, 1050-1061.	1.5	52
33	fcGENE: A Versatile Tool for Processing and Transforming SNP Datasets. PLoS ONE, 2014, 9, e97589.	1.1	52
34	Modelling Human Granulopoiesis under Poly-chemotherapy with G-CSF Support. Journal of Mathematical Biology, 2005, 50, 397-439.	0.8	50
35	Comparing performance of modern genotype imputation methods in different ethnicities. Scientific Reports, 2016, 6, 34386.	1.6	49
36	A computational model of human granulopoiesis to simulate the hematotoxic effects of multicycle polychemotherapy. Blood, 2004, 104, 2323-2331.	0.6	47

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37	A pre-registered short-term forecasting study of COVID-19 in Germany and Poland during the second wave. Nature Communications, 2021, 12, 5173.	5.8	47
38	Rationale and Design of the Leipzig (LIFE) Heart Study: Phenotyping and Cardiovascular Characteristics of Patients with Coronary Artery Disease. PLoS ONE, 2011, 6, e29070.	1.1	45
39	Sequential organ failure assessment score is an excellent operationalization of disease severity of adult patients with hospitalized community acquired pneumonia $\hat{a} \in \text{``results from the prospective observational PROGRESS study. Critical Care, 2019, 23, 110.}$	2.5	43
40	Meta-analysis uncovers genome-wide significant variants for rapid kidney function decline. Kidney International, 2021, 99, 926-939.	2.6	42
41	Integration of Genome-Wide SNP Data and Gene-Expression Profiles Reveals Six Novel Loci and Regulatory Mechanisms for Amino Acids and Acylcarnitines in Whole Blood. PLoS Genetics, 2015, 11, e1005510.	1.5	41
42	Urine Biomarkers of Tubular Renal Cell Damage for the Prediction of Acute Kidney Injury After Cardiac Surgeryâ€"A Pilot Study. Journal of Cardiothoracic and Vascular Anesthesia, 2017, 31, 2072-2079.	0.6	38
43	Increased Level of Interleukin 6 Associates With Increased 90-Day and 1-Year Mortality in Patients With End-Stage LiverÂDisease. Clinical Gastroenterology and Hepatology, 2018, 16, 730-737.	2.4	38
44	Sepsis-associated acute respiratory distress syndrome in individuals of European ancestry: a genome-wide association study. Lancet Respiratory Medicine, the, 2020, 8, 258-266.	5.2	38
45	Genetic Regulation of PCSK9 (Proprotein Convertase Subtilisin/Kexin Type 9) Plasma Levels and Its Impact on Atherosclerotic Vascular Disease Phenotypes. Circulation Genomic and Precision Medicine, 2018, 11, e001992.	1.6	37
46	Genetic Association Study of Eight Steroid Hormones and Implications for Sexual Dimorphism of Coronary Artery Disease. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 5008-5023.	1.8	37
47	Low-Dose Irradiation Affects Expression of Inflammatory Markers in the Heart of ApoE -/- Mice. PLoS ONE, 2015, 10, e0119661.	1.1	36
48	Ycasd– a tool for capturing and scaling data from graphical representations. BMC Bioinformatics, 2014, 15, 219.	1.2	35
49	Genome-wide meta-analysis identifies novel loci of plaque burden in carotid artery. Atherosclerosis, 2017, 259, 32-40.	0.4	33
50	Pharmacokinetic and -dynamic modelling of G-CSF derivatives in humans. Theoretical Biology and Medical Modelling, 2012, 9, 32.	2.1	32
51	Genome Wide Meta-analysis Highlights the Role of Genetic Variation in RARRES2 in the Regulation of Circulating Serum Chemerin. PLoS Genetics, 2014, 10, e1004854.	1.5	31
52	Genome-wide association study of 23,500 individuals identifies 7 loci associated with brain ventricular volume. Nature Communications, 2018, 9, 3945.	5.8	31
53	FTO Obesity Risk Variants Are Linked to Adipocyte IRX3 Expression and BMI of Children - Relevance of FTO Variants to Defend Body Weight in Lean Children?. PLoS ONE, 2016, 11, e0161739.	1.1	31
54	Modelling Lymphoma Therapy and Outcome. Bulletin of Mathematical Biology, 2014, 76, 401-430.	0.9	29

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55	Associations between DNA methylation and schizophrenia-related intermediate phenotypes — A gene set enrichment analysis. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2015, 59, 31-39.	2.5	29
56	Low sphingosine-1-phosphate plasma levels are predictive for increased mortality in patients with liver cirrhosis. PLoS ONE, 2017, 12, e0174424.	1.1	29
57	Association of metabolic parameters and rs726344 in FNDC5 with serum irisin concentrations. International Journal of Obesity, 2016, 40, 260-265.	1.6	28
58	Plasma levels of apolipoproteins C-III, A-IV, and E are independently associated with stable atherosclerotic cardiovascular disease. Atherosclerosis, 2019, 281, 17-24.	0.4	28
59	The Obesity-Susceptibility Gene TMEM18 Promotes Adipogenesis through Activation of PPARG. Cell Reports, 2020, 33, 108295.	2.9	28
60	Genetic Association of Objective Sleep Phenotypes with a Functional Polymorphism in the Neuropeptide S Receptor Gene. PLoS ONE, 2014, 9, e98789.	1.1	27
61	Population-genetic comparison of the Sorbian isolate population in Germany with the German KORA population using genome-wide SNP arrays. BMC Genetics, 2011, 12, 67.	2.7	26
62	Clinical and lifestyle related factors influencing whole blood metabolite levels – A comparative analysis of three large cohorts. Molecular Metabolism, 2019, 29, 76-85.	3.0	26
63	Different DOACs Control Inflammation in Cardiac Ischemia-Reperfusion Differently. Circulation Research, 2021, 128, 513-529.	2.0	26
64	Proteomics to improve phenotyping in obese patients with heart failure with preserved ejection fraction. European Journal of Heart Failure, 2021, 23, 1633-1644.	2.9	26
65	A biomathematical model of human thrombopoiesis under chemotherapy. Journal of Theoretical Biology, 2010, 264, 287-300.	0.8	25
66	No Association of Coronary Artery Disease with X-Chromosomal Variants in Comprehensive International Meta-Analysis. Scientific Reports, 2016, 6, 35278.	1.6	25
67	Proteomics-Enabled Deep Learning Machine Algorithms Can Enhance Prediction of Mortality. Journal of the American College of Cardiology, 2021, 78, 1621-1631.	1.2	25
68	Impact of genetic similarity on imputation accuracy. BMC Genetics, 2015, 16, 90.	2.7	24
69	Model-based design of chemotherapeutic regimens that account for heterogeneity in leucopoenia. British Journal of Haematology, 2006, 132, 723-735.	1.2	22
70	Genetic correlations reveal the shared genetic architecture of transcription in human peripheral blood. Nature Communications, 2017, 8, 483.	5.8	22
71	Association between lipoprotein(a) level and type 2 diabetes: no evidence for a causal role of lipoprotein(a) and insulin. Acta Diabetologica, 2017, 54, 1031-1038.	1.2	22
72	A Biomathematical Model of Human Erythropoiesis under Erythropoietin and Chemotherapy Administration. PLoS ONE, 2013, 8, e65630.	1.1	21

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73	Genetic Contribution of Variants near SORT1 and APOE on LDL Cholesterol Independent of Obesity in Children. PLoS ONE, 2015, 10, e0138064.	1.1	20
74	Cohort Profile: The Leipzig Research Center for Civilization Diseases–Heart Study (LIFE-Heart). International Journal of Epidemiology, 2020, 49, 1439-1440h.	0.9	19
75	Genome-wide association analysis of pulse wave velocity traits provide new insights into the causal relationship between arterial stiffness and blood pressure. PLoS ONE, 2020, 15, e0237237.	1.1	18
76	A Biomathematical Model of Pneumococcal Lung Infection and Antibiotic Treatment in Mice. PLoS ONE, 2016, 11, e0156047.	1.1	18
77	Peripheral blood RNA biomarkers for cardiovascular disease from bench to bedside: a position paper from the EU-CardioRNA COST action CA17129. Cardiovascular Research, 2022, 118, 3183-3197.	1.8	18
78	Genetic loci and prioritization of genes for kidney function decline derived from a meta-analysis of 62 longitudinal genome-wide association studies. Kidney International, 2022, 102, 624-639.	2.6	18
79	A pharmacokinetic model of filgrastim and pegfilgrastim application in normal mice and those with cyclophosphamideâ€induced granulocytopaenia. Cell Proliferation, 2009, 42, 813-822.	2.4	17
80	Subsequent Event Risk in Individuals With Established Coronary Heart Disease. Circulation Genomic and Precision Medicine, 2019, 12, e002470.	1.6	17
81	Association of plasma trimethylamine N-oxide levels with atherosclerotic cardiovascular disease and factors of the metabolic syndrome. Atherosclerosis, 2021, 335, 62-67.	0.4	17
82	Neutralizing Complement C5a Protects Mice with Pneumococcal Pulmonary Sepsis. Anesthesiology, 2020, 132, 795-807.	1.3	17
83	Genome-wide meta-analysis of phytosterols reveals five novel loci and a detrimental effect on coronary atherosclerosis. Nature Communications, 2022, 13, 143.	5.8	17
84	Differential and shared genetic effects on kidney function between diabetic and non-diabetic individuals. Communications Biology, 2022, 5, .	2.0	17
85	Impact of first- and second-line treatment for Hodgkin's lymphoma on the incidence of AML/MDS and NHL—experience of the German Hodgkin's Lymphoma Study Group analyzed by a parametric model of carcinogenesis. Annals of Oncology, 2011, 22, 681-688.	0.6	16
86	Global and Regional Development of the Human Cerebral Cortex: Molecular Architecture and Occupational Aptitudes. Cerebral Cortex, 2020, 30, 4121-4139.	1.6	16
87	Higher BMI, but not obesity-related genetic polymorphisms, correlates with lower structural connectivity of the reward network in a population-based study. International Journal of Obesity, 2021, 45, 491-501.	1.6	16
88	Pharmacokinetic and pharmacodynamic modelling of the novel human granulocyteâ€fcolonyâ€stimulatingâ€ffactor derivative Maxyâ€G34 and pegfilgrastim in rats. Cell Proliferation, 2009, 42, 823-837.	2.4	15
89	PROGRESS – prospective observational study on hospitalized community acquired pneumonia. BMC Pulmonary Medicine, 2016, 16, 108.	0.8	15
90	The value of noncoronary atherosclerosis for identifying coronary artery disease: results of the Leipzig LIFE Heart Study. Clinical Research in Cardiology, 2016, 105, 172-181.	1.5	15

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91	Genome-wide meta-analysis identifies novel determinants of circulating serum progranulin. Human Molecular Genetics, 2018, 27, 546-558.	1.4	15
92	Modeling individual time courses of thrombopoiesis during multi-cyclic chemotherapy. PLoS Computational Biology, 2019, 15, e1006775.	1.5	15
93	Genome-wide association study identifies an acute myeloid leukemia susceptibility locus near BICRA. Leukemia, 2019, 33, 771-775.	3.3	15
94	HLA Class II Allele Analyses Implicate Common Genetic Components in Type 1 and Non–Insulin-Treated Type 2 Diabetes. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e245-e254.	1.8	15
95	The role of rs2237781 within <i>GRM8</i> in eating behavior. Brain and Behavior, 2013, 3, 495-502.	1.0	14
96	Modelling chemotherapy effects on granulopoiesis. BMC Systems Biology, 2014, 8, 138.	3.0	14
97	A combined model of human erythropoiesis and granulopoiesis under growth factor and chemotherapy treatment. Theoretical Biology and Medical Modelling, 2014, 11, 24.	2.1	14
98	Genome-Wide Association Analysis for Severity of Coronary Artery Disease Using the Gensini Scoring System. Frontiers in Cardiovascular Medicine, 2017, 4, 57.	1.1	14
99	Common variants in the CLDN2-MORC4 and PRSS1-PRSS2 loci confer susceptibility to acute pancreatitis. Pancreatology, 2018, 18, 477-481.	0.5	14
100	Circulating Oxytocin Is Genetically Determined and Associated With Obesity and Impaired Glucose Tolerance. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 5621-5632.	1.8	14
101	Catalyzing Transcriptomics Research in Cardiovascular Disease: The CardioRNA COST Action CA17129. Non-coding RNA, 2019, 5, 31.	1.3	14
102	Genetically Determined Reproductive Aging and Coronary Heart Disease: A Bidirectional 2-sample Mendelian Randomization. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e2952-e2961.	1.8	13
103	DNA methylation patterns reflect individual's lifestyle independent of obesity. Clinical and Translational Medicine, 2022, $12$ , .	1.7	13
104	Growth and Final Height Among Children With Phenylketonuria. Pediatrics, 2017, 140, e20170015.	1.0	12
105	Metabolite-Investigator: an integrated user-friendly workflow for metabolomics multi-study analysis. Bioinformatics, 2021, 37, 2218-2220.	1.8	12
106	Integration of mathematical model predictions into routine workflows to support clinical decision making in haematology. BMC Medical Informatics and Decision Making, 2020, 20, 28.	1.5	12
107	Monocyte subtype counts are associated with 10-year cardiovascular disease risk as determined by the Framingham Risk Score among subjects of the LIFE-Adult study. PLoS ONE, 2021, 16, e0247480.	1.1	12
108	Validity, intra- and inter-observer reliability of automated devices for the assessment of ankle brachial index using photo-plethysmography. BMC Cardiovascular Disorders, 2013, 13, 81.	0.7	10

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109	Relationship Between Determinants of Arterial Stiffness Assessed by Diastolic and Suprasystolic Pulse Oscillometry. Medicine (United States), 2016, 95, e2963.	0.4	10
110	Genetic variants of lipase activity in chronic pancreatitis: TableÂ1. Gut, 2016, 65, 184-185.	6.1	10
111	Cyclin dependent kinase inhibitor $1\mathrm{C}$ is a female-specific marker of left ventricular function after acute myocardial infarction. International Journal of Cardiology, 2019, 274, 319-325.	0.8	10
112	Mendelian randomization analysis does not support causal associations of birth weight with hypertension risk and blood pressure in adulthood. European Journal of Epidemiology, 2020, 35, 685-697.	2.5	9
113	The Effect of FGF21 and Its Genetic Variants on Food and Drug Cravings, Adipokines and Metabolic Traits. Biomedicines, 2021, 9, 345.	1.4	9
114	Meta-GWAS of PCSK9 levels detects two novel loci at <i>APOB</i> and <i>TM6SF2</i> . Human Molecular Genetics, 2022, 31, 999-1011.	1.4	9
115	Comparison and modelling of pegylated or unpegylated G-CSF schedules in CHOP-14 regimen of elderly patients with aggressive B-cell lymphoma. Annals of Hematology, 2013, 92, 1641-1652.	0.8	8
116	Model-based optimization of G-CSF treatment during cytotoxic chemotherapy. Journal of Cancer Research and Clinical Oncology, 2018, 144, 343-358.	1.2	8
117	Circulating Adipokine VASPIN Is Associated with Serum Lipid Profiles in Humans. Lipids, 2019, 54, 203-210.	0.7	8
118	Deep learning detects heart failure with preserved ejection fraction using a baseline electrocardiogram. European Heart Journal Digital Health, 2021, 2, 699-703.	0.7	8
119	Interplay between adipose tissue secreted proteins, eating behavior and obesity. European Journal of Nutrition, 2022, 61, 885-899.	1.8	8
120	Genome-wide analysis of carotid plaque burden suggests a role of IL5 in men. PLoS ONE, 2020, 15, e0233728.	1.1	7
121	Adipocytokines are not associated with gestational diabetes mellitus but with pregnancy status. Cytokine, 2020, 131, 155088.	1.4	7
122	Proteomic profiling of low muscle and high fat mass: a machine learning approach in the KORA S4/FF4 study. Journal of Cachexia, Sarcopenia and Muscle, 2021, 12, 1011-1023.	2.9	7
123	Indications for Potential Parent-of-Origin Effects within the FTO Gene. PLoS ONE, 2015, 10, e0119206.	1.1	7
124	Lineage-Specific Changes in Biomarkers in Great Apes and Humans. PLoS ONE, 2015, 10, e0134548.	1,1	7
125	Sex-Specific Causal Relations between Steroid Hormones and Obesity—A Mendelian Randomization Study. Metabolites, 2021, 11, 738.	1.3	7
126	Effectiveness of cytopenia prophylaxis for different filgrastim and pegfilgrastim schedules in a chemotherapy mouse model. Biologics: Targets and Therapy, 2009, 3, 27-37.	3.0	7

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127	On the Parametrization of Epidemiologic Modelsâ€"Lessons from Modelling COVID-19 Epidemic. Viruses, 2022, 14, 1468.	1.5	7
128	Monitoring Disease Progression and Therapeutic Response in a Disseminated Tumor Model for Non-Hodgkin Lymphoma by Bioluminescence Imaging. Molecular Imaging, 2015, 14, 7290.2015.00010.	0.7	6
129	Dynamics of cytokines, immune cell counts and disease severity in patients with community-acquired pneumonia – Unravelling potential causal relationships. Cytokine, 2020, 136, 155263.	1.4	6
130	Modeling combined chemo- and immunotherapy of high-grade non-Hodgkin lymphoma. Leukemia and Lymphoma, 2016, 57, 1697-1708.	0.6	5
131	A biomathematical model of human erythropoiesis and iron metabolism. Scientific Reports, 2020, 10, 8602.	1.6	5
132	Individual prediction of thrombocytopenia at next chemotherapy cycle: Evaluation of dynamic model performances. British Journal of Clinical Pharmacology, 2021, 87, 3127-3138.	1.1	5
133	Whole Blood Metabolite Profiles Reflect Changes in Energy Metabolism in Heart Failure. Metabolites, 2022, 12, 216.	1.3	5
134	Simultaneous Mass Spectrometry-Based Apolipoprotein Profiling and Apolipoprotein E Phenotyping in Patients with ASCVD and Mild Cognitive Impairment. Nutrients, 2022, 14, 2474.	1.7	5
135	Comparison of scoring methods for the detection of causal genes with or without rare variants. BMC Proceedings, 2011, 5, S49.	1.8	4
136	Genetically programmed changes in transcription of the novel progranulin regulator. Journal of Molecular Medicine, 2020, 98, 1139-1148.	1.7	4
137	Associations of carotid intima media thickness with gene expression in whole blood and genetically predicted gene expression across 48 tissues. Human Molecular Genetics, 2022, 31, 1171-1182.	1.4	4
138	Evaluation of phosphodiesterase 9A as a novel biomarker in heart failure with preserved ejection fraction. ESC Heart Failure, 2021, 8, 1861-1872.	1.4	4
139	Verification of immunology-related genetic associations in BPD supports ABCA3 and five other genes. Pediatric Research, 2022, 92, 190-198.	1.1	4
140	Perioperative Two-Dimensional Left Ventricular Global Longitudinal Strain in Coronary Artery Bypass Surgery: A Prospective Observational Pilot Study. Journal of Cardiothoracic and Vascular Anesthesia, 2022, 36, 166-174.	0.6	4
141	A biomathematical model of immune response and barrier function in mice with pneumococcal lung infection. PLoS ONE, 2020, 15, e0243147.	1.1	4
142	Transcriptome Analyses of Adipose Tissue Samples Identify EGFL6 as a Candidate Gene Involved in Obesity-Related Adipose Tissue Dysfunction in Children. International Journal of Molecular Sciences, 2022, 23, 4349.	1.8	4
143	On the impact of relatedness on SNP association analysis. BMC Genetics, 2017, 18, 104.	2.7	3
144	Markov State Modelling of Disease Courses and Mortality Risks of Patients with Community-Acquired Pneumonia. Journal of Clinical Medicine, 2020, 9, 393.	1.0	3

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145	Prognostic impact of rapid reduction of involved free light chains in multiple myeloma patients under first-line treatment with Bendamustine, Prednisone, and Bortezomib (BPV). Journal of Cancer Research and Clinical Oncology, 2021, 147, 2349-2359.	1.2	3
146	Transcriptomic Research in Heart Failure with Preserved Ejection Fraction: Current State and Future Perspectives. Cardiac Failure Review, 2020, 6, e24.	1.2	3
147	Colocalization analysis of pancreas eQTLs with risk loci from alcoholic and novel non-alcoholic chronic pancreatitis GWAS suggests potential disease causing mechanisms. Pancreatology, 2022, 22, 449-456.	0.5	3
148	An Attempt to Explain the Ascent of Sap in Defoliated Trees. Journal of Mathematical Fluid Mechanics, 2004, 6, 295.	0.4	1
149	Analysis of GPRC6A variants in different pancreatitis etiologies. Pancreatology, 2020, 20, 1262-1267.	0.5	1
150	CD34+ Donor Chimerism and Wilms Tumor Gene 1 (WT1) Expression Provide an Early Indication of Relapse in Patients with Acute Leukemias and MDS after Hematopoietic Cell Transplantation (HCT) with Reduced-Intensity Conditioning Blood, 2006, 108, 548-548.	0.6	1
151	Association of head and neck cancer (HNSCC) subgroups defined by HPV RNA status, gene expression patterns, and TP53 mutations with lymph node metastasis and survival Journal of Clinical Oncology, 2015, 33, 6046-6046.	0.8	1
152	Genetic Regulation of Cytokine Response in Patients with Acute Community-Acquired Pneumonia. Genes, 2022, 13, 111.	1.0	1
153	Impact of medication on blood transcriptome reveals off-target regulations of beta-blockers. PLoS ONE, 2022, 17, e0266897.	1.1	1
154	Genetically regulated gene expression and proteins revealed discordant effects. PLoS ONE, 2022, 17, e0268815.	1,1	1
155	Generalized Solutions of the Capillary Problem. Journal of Mathematical Fluid Mechanics, 2004, 6, 272.	0.4	0
156	P3-086: FTO is Not Related to Imaging Parameters of the Hippocampus: A Volumetric and Diffusion Tensor Imaging Study., 2016, 12, P851-P851.		0
157	Response to Sabour. Pediatric Research, 2017, 82, 177-177.	1.1	0
158	Common variants in glyoxalase I do not increase chronic pancreatitis risk. PLoS ONE, 2019, 14, e0222927.	1.1	0
159	Neuronal hypoxia: protective effects of mononuclear cord blood cells after direct and indirect application. Journal of Stem Cells and Regenerative Medicine, 2007, 2, 16.	2.2	0
160	Integrational approaches for cross-species analysis of lung pathologies at single-cell resolution. Pneumologie, 2022, , .	0.1	0