Honghuang Lin

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8637910/publications.pdf

Version: 2024-02-01

173 papers 13,689 citations

51 h-index

36271

27389 106 g-index

204 all docs

204 docs citations

times ranked

204

23688 citing authors

#	Article	IF	CITATIONS
1	Genetic meta-analysis of diagnosed Alzheimer's disease identifies new risk loci and implicates Aβ, tau, immunity and lipid processing. Nature Genetics, 2019, 51, 414-430.	9.4	1,962
2	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. Nature Genetics, 2017, 49, 1373-1384.	9.4	783
3	Multi-ethnic genome-wide association study for atrial fibrillation. Nature Genetics, 2018, 50, 1225-1233.	9.4	552
4	Rare and low-frequency coding variants alter human adult height. Nature, 2017, 542, 186-190.	13.7	544
5	Meta-analysis identifies six new susceptibility loci for atrial fibrillation. Nature Genetics, 2012, 44, 670-675.	9.4	533
6	Genome-wide association and Mendelian randomisation analysis provide insights into the pathogenesis of heart failure. Nature Communications, 2020, 11, 163.	5.8	466
7	Genetic associations at 53 loci highlight cell types and biological pathways relevant for kidney function. Nature Communications, 2016, 7, 10023.	5.8	412
8	The genetics of blood pressure regulation and its target organs from association studies in 342,415 individuals. Nature Genetics, 2016, 48, 1171-1184.	9.4	362
9	Genome Analyses of >200,000 Individuals Identify 58 Loci for Chronic Inflammation and Highlight Pathways that Link Inflammation and Complex Disorders. American Journal of Human Genetics, 2018, 103, 691-706.	2.6	326
10	Protein-altering variants associated with body mass index implicate pathways that control energy intake and expenditure in obesity. Nature Genetics, 2018, 50, 26-41.	9.4	286
11	Large-scale analyses of common and rare variants identify 12 new loci associated with atrial fibrillation. Nature Genetics, 2017, 49, 946-952.	9.4	279
12	PROFEAT: a web server for computing structural and physicochemical features of proteins and peptides from amino acid sequence. Nucleic Acids Research, 2006, 34, W32-W37.	6.5	270
13	Trans-ancestry meta-analyses identify rare and common variants associated with blood pressure and hypertension. Nature Genetics, 2016, 48, 1151-1161.	9.4	261
14	DNA methylation signatures of chronic low-grade inflammation are associated with complex diseases. Genome Biology, 2016, 17, 255.	3.8	251
15	Atrial Fibrillation. Circulation, 2011, 124, 1982-1993.	1.6	225
16	Evaluation of MHC class I peptide binding prediction servers: Applications for vaccine research. BMC Immunology, 2008, 9, 8.	0.9	207
17	Genetic Predisposition, Clinical Risk Factor Burden, and Lifetime Risk of Atrial Fibrillation. Circulation, 2018, 137, 1027-1038.	1.6	196
18	Whole-Exome Sequencing Identifies Rare and Low-Frequency Coding Variants Associated with LDL Cholesterol. American Journal of Human Genetics, 2014, 94, 233-245.	2.6	193

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19	Genetic architecture of subcortical brain structures in 38,851 individuals. Nature Genetics, 2019, 51, 1624-1636.	9.4	192
20	Whole exome sequencing study identifies novel rare and common Alzheimer's-Associated variants involved in immune response and transcriptional regulation. Molecular Psychiatry, 2020, 25, 1859-1875.	4.1	191
21	Evaluation of MHC-II peptide binding prediction servers: applications for vaccine research. BMC Bioinformatics, 2008, 9, S22.	1.2	187
22	Integrating Genetic, Transcriptional, and Functional Analyses to Identify 5 Novel Genes for Atrial Fibrillation. Circulation, 2014, 130, 1225-1235.	1.6	183
23	Association Between Titin Loss-of-Function Variants and Early-Onset Atrial Fibrillation. JAMA - Journal of the American Medical Association, 2018, 320, 2354.	3.8	144
24	Higher dietary anthocyanin and flavonol intakes are associated with anti-inflammatory effects in a population of US adults. American Journal of Clinical Nutrition, 2015, 102, 172-181.	2.2	143
25	Loci influencing blood pressure identified using a cardiovascular gene-centric array. Human Molecular Genetics, 2013, 22, 1663-1678.	1.4	141
26	Novel Genetic Markers Associate With Atrial Fibrillation Risk in Europeans and Japanese. Journal of the American College of Cardiology, 2014, 63, 1200-1210.	1.2	127
27	Shortâ€Term Exposure to Air Pollution and Biomarkers of Oxidative Stress: The Framingham Heart Study. Journal of the American Heart Association, 2016, 5, .	1.6	109
28	Pleiotropic genes for metabolic syndrome and inflammation. Molecular Genetics and Metabolism, 2014, 112, 317-338.	0.5	107
29	Large-scale genome-wide analysis identifies genetic variants associated with cardiac structure and function. Journal of Clinical Investigation, 2017, 127, 1798-1812.	3.9	106
30	Plasma microRNAs are associated with atrial fibrillation and change after catheter ablation (the) Tj ETQq0 0 0 rgB1	Г <u>(O</u> yerloc	k 10 Tf 50 30
31	ROBO4 variants predispose individuals to bicuspid aortic valve and thoracic aortic aneurysm. Nature Genetics, 2019, 51, 42-50.	9.4	101
32	Common genetic variation at the IL1RL1 locus regulates IL-33/ST2 signaling. Journal of Clinical Investigation, 2013, 123, 4208-4218.	3.9	101
33	Genetic Obesity and the Risk of Atrial Fibrillation. Circulation, 2017, 135, 741-754.	1.6	96
34	Deep learning enables genetic analysis of the human thoracic aorta. Nature Genetics, 2022, 54, 40-51.	9.4	90
35	Relations between circulating microRNAs and atrial fibrillation: Data from the Framingham Offspring Study. Heart Rhythm, 2014, 11, 663-669.	0.3	80
36	A support vector machines approach for virtual screening of active compounds of single and multiple mechanisms from large libraries at an improved hit-rate and enrichment factor. Journal of Molecular Graphics and Modelling, 2008, 26, 1276-1286.	1.3	76

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37	A Whole-Blood Transcriptome Meta-Analysis Identifies Gene Expression Signatures of Cigarette Smoking. Human Molecular Genetics, 2016, 25, ddw288.	1.4	76
38	Recent progresses in the application of machine learning approach for predicting protein functional class independent of sequence similarity. Proteomics, 2006, 6, 4023-4037.	1.3	72
39	PR interval genome-wide association meta-analysis identifies 50 loci associated with atrial and atrioventricular electrical activity. Nature Communications, 2018, 9, 2904.	5.8	71
40	Support vector machines approach for predicting druggable proteins: recent progress in its exploration and investigation of its usefulness. Drug Discovery Today, 2007, 12, 304-313.	3.2	69
41	Efficacy of different protein descriptors in predicting protein functional families. BMC Bioinformatics, 2007, 8, 300.	1.2	66
42	TANTIGEN: a comprehensive database of tumor T cell antigens. Cancer Immunology, Immunotherapy, 2017, 66, 731-735.	2.0	66
43	PEARLS:Â Program for Energetic Analysis of Receptorâ°Ligand System. Journal of Chemical Information and Modeling, 2006, 46, 445-450.	2.5	64
44	Assessment of the Relationship Between Genetic Determinants of Thyroid Function and Atrial Fibrillation. JAMA Cardiology, 2019, 4, 144.	3.0	64
45	Novel Mutation in <i>FLNC</i> (Filamin C) Causes Familial Restrictive Cardiomyopathy. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	62
46	Prediction of transporter family from protein sequence by support vector machine approach. Proteins: Structure, Function and Bioinformatics, 2005, 62, 218-231.	1.5	61
47	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
48	Prediction of MHC-binding peptides of flexible lengths from sequence-derived structural and physicochemical properties. Molecular Immunology, 2007, 44, 866-877.	1.0	60
49	Multi-ancestry GWAS of the electrocardiographic PR interval identifies 202 loci underlying cardiac conduction. Nature Communications, 2020, 11, 2542.	5.8	59
50	Measures of Biologic Age in a Community Sample Predict Mortality and Age-Related Disease: The Framingham Offspring Study. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2018, 73, 757-762.	1.7	59
51	Computer prediction of drug resistance mutations in proteins. Drug Discovery Today, 2005, 10, 521-529.	3.2	58
52	Prediction of the functional class of metal-binding proteins from sequence derived physicochemical properties by support vector machine approach. BMC Bioinformatics, 2006, 7, S13.	1.2	58
53	Next Steps in Cardiovascular Disease Genomic Research—Sequencing, Epigenetics, and Transcriptomics. Clinical Chemistry, 2012, 58, 113-126.	1.5	55
54	Machine learning approaches for predicting compounds that interact with therapeutic and ADMET related proteins. Journal of Pharmaceutical Sciences, 2007, 96, 2838-2860.	1.6	54

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55	A Global Characterization and Identification of Multifunctional Enzymes. PLoS ONE, 2012, 7, e38979.	1.1	53
56	Gene expression and genetic variation in human atria. Heart Rhythm, 2014, 11, 266-271.	0.3	48
57	Methylome-wide Association Study of Atrial Fibrillation in Framingham Heart Study. Scientific Reports, 2017, 7, 40377.	1.6	48
58	Exome-chip meta-analysis identifies novel loci associated with cardiac conduction, including ADAMTS6. Genome Biology, 2018, 19, 87.	3.8	47
59	Genome-Wide Association Study of Cardiac Structure and Systolic Function in African Americans. Circulation: Cardiovascular Genetics, 2013, 6, 37-46.	5.1	46
60	MicroRNA Signature of Cigarette Smoking and Evidence for a Putative Causal Role of MicroRNAs in Smoking-Related Inflammation and Target Organ Damage. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	45
61	Fresh frozen plasma transfusion in acute variceal haemorrhage: Results from a multicentre cohort study. Liver International, 2021, 41, 1901-1908.	1.9	45
62	Proteomics Profiling and Risk of Newâ€Onset Atrial Fibrillation: Framingham Heart Study. Journal of the American Heart Association, 2019, 8, e010976.	1.6	42
63	Integrative Omics Approach to Identifying Genes Associated With Atrial Fibrillation. Circulation Research, 2020, 126, 350-360.	2.0	41
64	Prediction of the functional class of lipid binding proteins from sequence-derived properties irrespective of sequence similarity. Journal of Lipid Research, 2006, 47, 824-831.	2.0	40
65	Association of exome sequences with plasma C-reactive protein levels in >9000 participants. Human Molecular Genetics, 2015, 24, 559-571.	1.4	36
66	Association of Habitual Physical Activity With Cardiovascular Disease Risk. Circulation Research, 2020, 127, 1253-1260.	2.0	36
67	Epigenetic Age and the Risk of Incident Atrial Fibrillation. Circulation, 2021, 144, 1899-1911.	1.6	35
68	Whole Exome Sequencing in Atrial Fibrillation. PLoS Genetics, 2016, 12, e1006284.	1.5	35
69	Genetic analysis of right heart structure and function in 40,000 people. Nature Genetics, 2022, 54, 792-803.	9.4	34
70	Shared genetic susceptibility of vascular-related biomarkers with ischemic and recurrent stroke. Neurology, 2016, 86, 351-359.	1.5	33
71	Diminished <i>PRRX1</i> Expression Is Associated With Increased Risk of Atrial Fibrillation and Shortening of the Cardiac Action Potential. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	33
72	Association of Methylation Signals With Incident Coronary Heart Disease in an Epigenome-Wide Assessment of Circulating Tumor Necrosis Factor α. JAMA Cardiology, 2018, 3, 463.	3.0	33

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73	Dana-Farber repository for machine learning in immunology. Journal of Immunological Methods, 2011, 374, 18-25.	0.6	32
74	Genome-Wide Association Analysis of Plasma B–Type Natriuretic Peptide in Blacks. Circulation: Cardiovascular Genetics, 2015, 8, 122-130.	5.1	32
75	Protein Biomarkers and Risk of Atrial Fibrillation. Circulation: Arrhythmia and Electrophysiology, 2020, 13, e007607.	2.1	31
76	Discovery of novel heart rate-associated loci using the Exome Chip. Human Molecular Genetics, 2017, 26, 2346-2363.	1.4	29
77	Genome sequencing unveils a regulatory landscape of platelet reactivity. Nature Communications, 2021, 12, 3626.	5.8	29
78	Prediction of factor Xa inhibitors by machine learning methods. Journal of Molecular Graphics and Modelling, 2007, 26, 505-518.	1.3	28
79	Whole blood gene expression and white matter Hyperintensities. Molecular Neurodegeneration, 2017, 12, 67.	4.4	28
80	MHC-BPS: MHC-binder prediction server for identifying peptides of flexible lengths from sequence-derived physicochemical properties. Immunogenetics, 2006, 58, 607-613.	1.2	27
81	Metabolomic Profiling in Relation to New-Onset Atrial Fibrillation (from the Framingham Heart) Tj ETQq1 1 0.784	4314.rgBT 0.7	Oyerlock 10
82	ExomeChip-Wide Analysis of 95 626 Individuals Identifies 10 Novel Loci Associated With QT and JT Intervals. Circulation Genomic and Precision Medicine, 2018, 11, e001758.	1.6	27
83	Recent exposure to particle radioactivity and biomarkers of oxidative stress and inflammation: The Framingham Heart Study. Environment International, 2018, 121, 1210-1216.	4.8	27
84	Common Coding Variants in <i>SCN10A</i> Are Associated With the Nav1.8 Late Current and Cardiac Conduction. Circulation Genomic and Precision Medicine, 2018, 11, e001663.	1.6	26
85	Quality control and integration of genotypes from two calling pipelines for whole genome sequence data in the Alzheimer's disease sequencing project. Genomics, 2019, 111, 808-818.	1.3	26
86	Aptamer-Based Proteomic Platform Identifies Novel Protein Predictors of Incident Heart Failure and Echocardiographic Traits. Circulation: Heart Failure, 2020, 13, e006749.	1.6	26
87	Derivation of Stable Microarray Cancer-Differentiating Signatures Using Consensus Scoring of Multiple Random Sampling and Gene-Ranking Consistency Evaluation. Cancer Research, 2007, 67, 9996-10003.	0.4	25
88	Assessment of the Mid-Life Demographic and Lifestyle Risk Factors of Dementia Using Data from the Framingham Heart Study Offspring Cohort. Journal of Alzheimer's Disease, 2018, 63, 1119-1127.	1,2	25
89	Whole blood gene expression and interleukin-6 levels. Genomics, 2014, 104, 490-495.	1.3	24
90	Targeted sequencing in candidate genes for atrial fibrillation: The Cohorts for Heart and Aging Research in Genomic Epidemiology (CHARGE) Targeted Sequencing Study. Heart Rhythm, 2014, 11, 452-457.	0.3	24

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91	Genome-Wide Meta-Analyses of Plasma Renin Activity and Concentration Reveal Association With the Kininogen 1 and Prekallikrein Genes. Circulation: Cardiovascular Genetics, 2015, 8, 131-140.	5.1	24
92	Gain-of-function mutations in GATA6 lead to atrial fibrillation. Heart Rhythm, 2017, 14, 284-291.	0.3	24
93	Whole Blood Gene Expression and Atrial Fibrillation: The Framingham Heart Study. PLoS ONE, 2014, 9, e96794.	1.1	23
94	Gene expression markers of age-related inflammation in two human cohorts. Experimental Gerontology, 2015, 70, 37-45.	1.2	23
95	Genetic invalidation of Lp-PLA2 as a therapeutic target: Large-scale study of five functional Lp-PLA2-lowering alleles. European Journal of Preventive Cardiology, 2017, 24, 492-504.	0.8	22
96	Variants in angiopoietin-2 (<i>ANGPT2</i>) contribute to variation in nocturnal oxyhaemoglobin saturation level. Human Molecular Genetics, 2016, 25, ddw324.	1.4	21
97	Homology-Free Prediction of Functional Class of Proteins and Peptides by Support Vector Machines. Current Protein and Peptide Science, 2008, 9, 70-95.	0.7	19
98	Common and Rare Coding Genetic Variation Underlying the Electrocardiographic PR Interval. Circulation Genomic and Precision Medicine, 2018, 11, e002037.	1.6	19
99	Whole Blood Gene Expression Associated With Clinical Biological Age. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2019, 74, 81-88.	1.7	19
100	Whole blood microRNA expression associated with stroke: Results from the Framingham Heart Study. PLoS ONE, 2019, 14, e0219261.	1.1	19
101	An Artificial Intelligence-Assisted Method for Dementia Detection Using Images from the Clock Drawing Test. Journal of Alzheimer's Disease, 2021, 83, 581-589.	1.2	19
102	Identification of digital voice biomarkers for cognitive health. Exploration of Medicine, 2020, 1, 406-417.	1.5	19
103	Sequencing of 2 Subclinical Atherosclerosis Candidate Regions in 3669 Individuals. Circulation: Cardiovascular Genetics, 2014, 7, 359-364.	5.1	18
104	Strategies to Design and Analyze Targeted Sequencing Data. Circulation: Cardiovascular Genetics, 2014, 7, 335-343.	5.1	18
105	Integrated Multiomics Approach to Identify Genetic Underpinnings of Heart Failure and Its Echocardiographic Precursors. Circulation Genomic and Precision Medicine, 2019, 12, e002489.	1.6	18
106	Validation of Polygenic Scores for QT Interval in Clinical Populations. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	17
107	Genetic Determinants of Electrocardiographic P-Wave Duration and Relation to Atrial Fibrillation. Circulation Genomic and Precision Medicine, 2020, 13, 387-395.	1.6	16
108	FAM13A Represses AMPK Activity and Regulates Hepatic Glucose and Lipid Metabolism. IScience, 2020, 23, 100928.	1.9	16

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109	Association Between the Digital Clock Drawing Test and Neuropsychological Test Performance: Large Community-Based Prospective Cohort (Framingham Heart Study). Journal of Medical Internet Research, 2021, 23, e27407.	2.1	16
110	Comparison of On-Site Versus Remote Mobile Device Support in the Framingham Heart Study Using the Health eHeart Study for Digital Follow-up: Randomized Pilot Study Set Within an Observational Study Design. JMIR MHealth and UHealth, 2019, 7, e13238.	1.8	16
111	Gene-gene Interaction Analyses for Atrial Fibrillation. Scientific Reports, 2016, 6, 35371.	1.6	15
112	Prefrontal cortex eQTLs/mQTLs enriched in genetic variants associated with alcohol use disorder and other diseases. Epigenomics, 2020, 12, 789-800.	1.0	15
113	MoViES: molecular vibrations evaluation server for analysis of fluctuational dynamics of proteins and nucleic acids. Nucleic Acids Research, 2004, 32, W679-W685.	6.5	14
114	Mutation of a common amino acid in NKX2.5 results in dilated cardiomyopathy in two large families. BMC Medical Genetics, 2016, 17, 83.	2.1	14
115	Epigenetic Analyses of Human Left Atrial Tissue Identifies Gene Networks Underlying Atrial Fibrillation. Circulation Genomic and Precision Medicine, 2020, 13, e003085.	1.6	14
116	Novel Risk Modeling Approach of Atrial Fibrillation With Restricted Mean Survival Times. Circulation: Cardiovascular Quality and Outcomes, 2020, 13, e005918.	0.9	14
117	Proteomic Signatures of Lifestyle Risk Factors for Cardiovascular Disease: A Crossâ€6ectional Analysis of the Plasma Proteome in the Framingham Heart Study. Journal of the American Heart Association, 2021, 10, e018020.	1.6	14
118	Prediction of functional class of novel plant proteins by a statistical learning method. New Phytologist, 2005, 168, 109-121.	3.5	13
119	Circuit Performance Classification With Active Learning Guided Sampling for Support Vector Machines. IEEE Transactions on Computer-Aided Design of Integrated Circuits and Systems, 2015, 34, 1467-1480.	1.9	13
120	Crossâ€sectional relations of wholeâ€blood mi <scp>RNA</scp> expression levels and hand grip strength in a community sample. Aging Cell, 2017, 16, 888-894.	3.0	13
121	Adherence of Mobile App-Based Surveys and Comparison With Traditional Surveys: eCohort Study. Journal of Medical Internet Research, 2021, 23, e24773.	2.1	13
122	A comparison of whole genome sequencing with exome sequencing for family-based association studies. BMC Proceedings, 2014, 8, S38.	1.8	12
123	Sequencing of <i>SCN5A</i> Identifies Rare and Common Variants Associated With Cardiac Conduction: Cohorts for Heart and Aging Research in Genomic Epidemiology (CHARGE) Consortium. Circulation: Cardiovascular Genetics, 2014, 7, 365-373.	5.1	12
124	Using data science to diagnose and characterize heterogeneity of Alzheimer's disease. Alzheimer's and Dementia: Translational Research and Clinical Interventions, 2019, 5, 264-271.	1.8	12
125	New biomarkers from multiomics approaches: improving risk prediction of atrial fibrillation. Cardiovascular Research, 2021, 117, 1632-1644.	1.8	12
126	Transcriptome-wide association study of inflammatory biologic age. Aging, 2017, 9, 2288-2301.	1.4	12

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127	Genome-Wide Association Study for Endothelial Growth Factors. Circulation: Cardiovascular Genetics, 2015, 8, 389-397.	5.1	11
128	Whole exome sequencing in the Framingham Heart Study identifies rare variation in HYAL2 that influences platelet aggregation. Thrombosis and Haemostasis, 2017, 117, 1083-1092.	1.8	11
129	Epigenome-Wide Association Study of Soluble Tumor Necrosis Factor Receptor 2 Levels in the Framingham Heart Study. Frontiers in Pharmacology, 2018, 9, 207.	1.6	11
130	Healthy diet is associated with gene expression in blood: the Framingham Heart Study. American Journal of Clinical Nutrition, 2019, 110, 742-749.	2.2	11
131	The genomics of heart failure: design and rationale of the HERMES consortium. ESC Heart Failure, 2021, 8, 5531-5541.	1.4	11
132	Genetic Reduction in Left Ventricular Protein Kinase C- \hat{l}_{\pm} and Adverse Ventricular Remodeling in Human Subjects. Circulation Genomic and Precision Medicine, 2018, 11, e001901.	1.6	10
133	Relations between plasma microRNAs, echocardiographic markers of atrial remodeling, and atrial fibrillation: Data from the Framingham Offspring study. PLoS ONE, 2020, 15, e0236960.	1.1	10
134	Automatic synchronization and distribution of biological databases and software over low-bandwidth networks among developing countries. Bioinformatics, 2008, 24, 299-301.	1.8	9
135	Association of Habitual Physical Activity With Home Blood Pressure in the Electronic Framingham Heart Study (eFHS): Cross-sectional Study. Journal of Medical Internet Research, 2021, 23, e25591.	2.1	9
136	EDEM3 Modulates Plasma Triglyceride Level through Its Regulation of LRP1 Expression. IScience, 2020, 23, 100973.	1.9	8
137	Advances in Exploration of Machine Learning Methods for Predicting Functional Class and Interaction Profiles of Proteins and Peptides Irrespective of Sequence Homology. Current Bioinformatics, 2007, 2, 95-112.	0.7	7
138	A Lowâ€Frequency Variant in MAPK14 Provides Mechanistic Evidence of a Link With Myeloperoxidase: A Prognostic Cardiovascular Risk Marker. Journal of the American Heart Association, 2014, 3, .	1.6	7
139	Low oxygen saturation during sleep reduces CD1D and RAB20 expressions that are reversed by CPAP therapy. EBioMedicine, 2020, 56, 102803.	2.7	7
140	Rare Coding Variants Associated With Electrocardiographic Intervals Identify Monogenic Arrhythmia Susceptibility Genes: A Multi-Ancestry Analysis. Circulation Genomic and Precision Medicine, 2021, 14, e003300.	1.6	7
141	Identifying Blood Biomarkers for Dementia Using Machine Learning Methods in the Framingham Heart Study. Cells, 2022, 11, 1506.	1.8	7
142	Design and Preliminary Findings of Adherence to the Self-Testing for Our Protection From COVID-19 (STOP COVID-19) Risk-Based Testing Protocol: Prospective Digital Study. JMIR Formative Research, 2022, 6, e38113.	0.7	7
143	Familial clustering of hypertensive target organ damage in the community. Journal of Hypertension, 2018, 36, 1086-1093.	0.3	6
144	Tissue-specific Network Analysis of Genetic Variants Associated with Coronary Artery Disease. Scientific Reports, 2018, 8, 11492.	1.6	6

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145	Network Analysis of Depression-Related Transcriptomic Profiles. NeuroMolecular Medicine, 2019, 21, 143-149.	1.8	6
146	P-wave signal-averaged electrocardiography: Reference values, clinical correlates, and heritability in the Framingham Heart Study. Heart Rhythm, 2021, 18, 1500-1507.	0.3	6
147	Functional analysis of HapMap SNPs. Gene, 2012, 511, 358-363.	1.0	5
148	Rare genetic variant analysis on blood pressure in related samples. BMC Proceedings, 2014, 8, S35.	1.8	5
149	Relevance vector and feature machine for statistical analog circuit characterization and built-in self-test optimization. , $2016, \ldots$		5
150	Association of Genetic Variation in Coronary Artery Disease–Related Loci With the Risk of Heart Failure With Preserved Versus Reduced Ejection Fraction. Circulation, 2018, 137, 1290-1292.	1.6	5
151	Exploring the Hierarchical Influence of Cognitive Functions for Alzheimer Disease: The Framingham Heart Study. Journal of Medical Internet Research, 2020, 22, e15376.	2.1	5
152	Neck Circumference and Risk of Incident Atrial Fibrillation in the Framingham Heart Study. Journal of the American Heart Association, 2022, 11, e022340.	1.6	5
153	Lymphocyte activation gene-3-associated protein networks are associated with HDL-cholesterol and mortality in the Trans-omics for Precision Medicine program. Communications Biology, 2022, 5, 362.	2.0	5
154	Transcriptomic Heterogeneity of Alzheimer's Disease Associated with Lipid Genetic Risk. NeuroMolecular Medicine, 2020, 22, 534-541.	1.8	4
155	Human Leukocyte Antigen Typing Using a Knowledge Base Coupled with a High-Throughput Oligonucleotide Probe Array Analysis. Frontiers in Immunology, 2014, 5, 597.	2.2	3
156	Relations Between BMI Trajectories and Habitual Physical Activity Measured by a Smartwatch in the Electronic Cohort of the Framingham Heart Study: Cohort Study. JMIR Cardio, 2022, 6, e32348.	0.7	3
157	Associations Between the Digital Clock Drawing Test and Brain Volume: Large Community-Based Prospective Cohort (Framingham Heart Study). Journal of Medical Internet Research, 2022, 24, e34513.	2.1	3
158	Comparing Bowtie and BWA to Align Short Reads from a RNA-Seq Experiment. Advances in Intelligent and Soft Computing, 2012, , 197-207.	0.2	2
159	Shared Genetic and Environmental Architecture of Cardiac Phenotypes Assessed via Echocardiography. Circulation Genomic and Precision Medicine, 2021, 14, e003244.	1.6	2
160	Comparison of Daily Routines Between Middle-aged and Older Participants With and Those Without Diabetes in the Electronic Framingham Heart Study: Cohort Study. JMIR Diabetes, 2022, 7, e29107.	0.9	2
161	The association between social network index, atrial fibrillation, and mortality in the Framingham Heart Study. Scientific Reports, 2022, 12, 3958.	1.6	2
162	Prediction of antibiotic resistance proteins from sequence-derived properties irrespective of sequence similarity. International Journal of Antimicrobial Agents, 2008, 32, 221-226.	1.1	1

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163	Genomic profiling by machine learning. , 2011, , .		1
164	Loci influencing blood pressure identified using a cardiovascular gene-centric array. Human Molecular Genetics, 2013, 22, 3394-3395.	1.4	1
165	Article Commentary: Microarray Data Analysis of Gene Expression Evolution. Gene Regulation and Systems Biology, 2009, 3, GRSB.S2997.	2.3	0
166	Association of genetic variations and gene expression in a family-based study. BMC Proceedings, 2016, 10, 109-112.	1.8	0
167	0021 Lower Oxygen Saturation During Sleep Is Associated With Reduced Expressions Of Cd1d And Rab20 That Is Potentially Reversed By CPAP Therapy. Sleep, 2019, 42, A8-A9.	0.6	0
168	Comparative transâ€ethnic metaâ€analysis of whole exome sequencing variation for Alzheimer's disease (AD) in 18,402 individuals of the Alzheimer's Disease Sequencing Project (ADSP). Alzheimer's and Dementia, 2020, 16, e041583.	0.4	0
169	Assessing whole genome sequencing variation for Alzheimer's disease in 4707 individuals from the Alzheimer's Disease Sequencing Project (ADSP). Alzheimer's and Dementia, 2020, 16, e045548.	0.4	0
170	Frequency of familial Alzheimer's disease gene mutations within the Alzheimer Disease Sequencing Project (ADSP). Alzheimer's and Dementia, 2020, 16, e046203.	0.4	0
171	2010 Dietary Guidelines for Americans and decreased inflammation. FASEB Journal, 2013, 27, lb397.	0.2	0
172	The 2010 Dietary Guidelines for Americans food groups associated with decreased inflammation (370.3). FASEB Journal, 2014, 28, 370.3.	0.2	0
173	Sparse Relevance Kernel Machine-Based Performance Dependency Analysis of Analog and Mixed-Signal Circuits. , 2019, , 423-447.		0