Taesung Park

List of Publications by Year in descending order

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239 papers 8,699 citations

30 h-index 87 g-index

249 all docs 249 docs citations

249 times ranked 16554 citing authors

#	Article	IF	CITATIONS
1	Biological, clinical and population relevance of 95 loci for blood lipids. Nature, 2010, 466, 707-713.	13.7	3,249
2	The genetic architecture of type 2 diabetes. Nature, 2016, 536, 41-47.	13.7	952
3	A large-scale genome-wide association study of Asian populations uncovers genetic factors influencing eight quantitative traits. Nature Genetics, 2009, 41, 527-534.	9.4	937
4	Large-scale genome-wide association studies in east Asians identify new genetic loci influencing metabolic traits. Nature Genetics, 2011, 43, 990-995.	9.4	270
5	Statistical tests for identifying differentially expressed genes in time-course microarray experiments. Bioinformatics, 2003, 19, 694-703.	1.8	152
6	Odds ratio based multifactor-dimensionality reduction method for detecting gene-gene interactions. Bioinformatics, 2007, 23, 71-76.	1.8	144
7	Evaluation of normalization methods for microarray data. BMC Bioinformatics, 2003, 4, 33.	1.2	136
8	HNF4 \hat{l}_{\pm} is a therapeutic target that links AMPK to WNT signalling in early-stage gastric cancer. Gut, 2016, 65, 19-32.	6.1	91
9	Joint Identification of Multiple Genetic Variants via Elasticâ€Net Variable Selection in a Genomeâ€Wide Association Analysis. Annals of Human Genetics, 2010, 74, 416-428.	0.3	84
10	Log-linear model-based multifactor dimensionality reduction method to detect gene–gene interactions. Bioinformatics, 2007, 23, 2589-2595.	1.8	82
11	Personalized identification of altered pathways in cancer using accumulated normal tissue data. Bioinformatics, 2014, 30, i422-i429.	1.8	78
12	A genome-wide scan for signatures of directional selection in domesticated pigs. BMC Genomics, 2015, 16, 130.	1.2	67
13	A novel method to identify high order gene-gene interactions in genome-wide association studies: Gene-based MDR. BMC Bioinformatics, 2012, 13, S5.	1.2	66
14	Clinical implication of serum carcinoembryonic antigen and carbohydrate antigen 19-9 for the prediction of malignancy in intraductal papillary mucinous neoplasm of pancreas. Journal of Hepato-Biliary-Pancreatic Sciences, 2015, 22, 699-707.	1.4	65
15	Comprehensive Metabolomic Search for Biomarkers to Differentiate Early Stage Hepatocellular Carcinoma from Cirrhosis. Cancers, 2019, 11, 1497.	1.7	63
16	Associations of Variants in CHRNA5/A3/B4 Gene Cluster with Smoking Behaviors in a Korean Population. PLoS ONE, 2010, 5, e12183.	1.1	57
17	Enhancing inhibitory synaptic function reverses spatial memory deficits in Shank2 mutant mice. Neuropharmacology, 2017, 112, 104-112.	2.0	56
18	Large-scale genome-wide association study of Asian population reveals genetic factors in FRMD4A and other loci influencing smoking initiation and nicotine dependence. Human Genetics, 2012, 131, 1009-1021.	1.8	52

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19	Molecular subtypes of pancreatic cancer based on miRNA expression profiles have independent prognostic value. Journal of Gastroenterology and Hepatology (Australia), 2016, 31, 1160-1167.	1.4	52
20	Circulating Microbiota-Based Metagenomic Signature for Detection of Hepatocellular Carcinoma. Scientific Reports, 2019, 9, 7536.	1.6	51
21	Association between mutations of critical pathway genes and survival outcomes according to the tumor location in colorectal cancer. Cancer, 2017, 123, 3513-3523.	2.0	50
22	Clinical Characteristics and Outcomes of COVID-19 Cohort Patients in Daegu Metropolitan City Outbreak in 2020. Journal of Korean Medical Science, 2021, 36, e12.	1.1	47
23	β-Caryophyllene potently inhibits solid tumor growth and lymph node metastasis of B16F10 melanoma cells in high-fat diet–induced obese C57BL/6N mice. Carcinogenesis, 2015, 36, 1028-1039.	1.3	44
24	Combining multiple microarrays in the presence of controlling variables. Bioinformatics, 2006, 22, 1682-1689.	1.8	37
25	Serum fibronectin distinguishes the early stages of hepatocellular carcinoma. Scientific Reports, 2017, 7, 9449.	1.6	37
26	Gene–gene interaction analysis for the survival phenotype based on the Cox model. Bioinformatics, 2012, 28, i582-i588.	1.8	36
27	Nonsynonymous Variants in <i>PAX4</i> and <i>GLP1R</i> Are Associated With Type 2 Diabetes in an East Asian Population. Diabetes, 2018, 67, 1892-1902.	0.3	36
28	Powerful p-value combination methods to detect incomplete association. Scientific Reports, 2021, 11, 6980.	1.6	35
29	FARVAT: a family-based rare variant association test. Bioinformatics, 2014, 30, 3197-3205.	1.8	34
30	Definition of Smad3 Phosphorylation Events That Affect Malignant and Metastatic Behaviors in Breast Cancer Cells. Cancer Research, 2014, 74, 6139-6149.	0.4	33
31	Covariance models for nested repeated measures data: analysis of ovarian steroid secretion data. Statistics in Medicine, 2002, 21, 143-164.	0.8	32
32	Relationship between thyroid-stimulating hormone levels and risk of depression among the general population with normal free T4 levels. Psychoneuroendocrinology, 2015, 58, 114-119.	1.3	32
33	Metagenomic Analysis of Serum Microbe-Derived Extracellular Vesicles and Diagnostic Models to Differentiate Ovarian Cancer and Benign Ovarian Tumor. Cancers, 2020, 12, 1309.	1.7	32
34	Pathway-based approach using hierarchical components of collapsed rare variants. Bioinformatics, 2016, 32, i586-i594.	1.8	31
35	Sequence data and association statistics from 12,940 type 2 diabetes cases and controls. Scientific Data, 2017, 4, 170179.	2.4	31
36	Development of Diagnostic Biomarkers for Detecting Diabetic Retinopathy at Early Stages Using Quantitative Proteomics. Journal of Diabetes Research, 2016, 2016, 1-22.	1.0	28

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37	Diagnostic performance enhancement of pancreatic cancer using proteomic multimarker panel. Oncotarget, 2017, 8, 93117-93130.	0.8	28
38	Pathway-Driven Approaches of Interaction between Oxidative Balance and Genetic Polymorphism on Metabolic Syndrome. Oxidative Medicine and Cellular Longevity, 2017, 2017, 1-9.	1.9	28
39	Evaluating the contribution of rare variants to type 2 diabetes and related traits using pedigrees. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 379-384.	3.3	28
40	Association of an IGHV3-66 gene variant with Kawasaki disease. Journal of Human Genetics, 2021, 66, 475-489.	1.1	27
41	An empirical fuzzy multifactor dimensionality reduction method for detecting gene-gene interactions. BMC Genomics, 2017, 18, 115.	1.2	26
42	Ceruloplasmin as a prognostic marker in patients with bile duct cancer. Oncotarget, 2017, 8, 29028-29037.	0.8	26
43	A Modified Entropy-Based Approach for Identifying Gene-Gene Interactions in Case-Control Study. PLoS ONE, 2013, 8, e69321.	1.1	25
44	A unified model based multifactor dimensionality reduction framework for detecting gene–gene interactions. Bioinformatics, 2016, 32, i605-i610.	1.8	24
45	Longitudinal analysis to better characterize Asthmaâ€COPD overlap syndrome: Findings from an adult asthma cohort in Korea (COREA). Clinical and Experimental Allergy, 2019, 49, 603-614.	1.4	23
46	Association between ST8SIA2 and the Risk of Schizophrenia and Bipolar I Disorder across Diagnostic Boundaries. PLoS ONE, 2015, 10, e0139413.	1.1	23
47	Identification of geneâ€gene interactions in the presence of missing data using the multifactor dimensionality reduction method. Genetic Epidemiology, 2009, 33, 646-656.	0.6	22
48	Seasonality and its distinct clinical correlates in bipolar II disorder. Psychiatry Research, 2015, 225, 540-544.	1.7	22
49	Heavy metal accumulation in and food safety of shark meat from Jeju island, Republic of Korea. PLoS ONE, 2019, 14, e0212410.	1.1	22
50	Multivariate generalized multifactor dimensionality reduction to detect gene-gene interactions. BMC Systems Biology, 2013, 7, S15.	3.0	21
51	Multivariate Quantitative Multifactor Dimensionality Reduction for Detecting Gene-Gene Interactions. Human Heredity, 2015, 79, 168-181.	0.4	21
52	Transancestral fine-mapping of four type 2 diabetes susceptibility loci highlights potential causal regulatory mechanisms. Human Molecular Genetics, 2016, 25, 2070-2081.	1.4	21
53	Rare variant association test with multiple phenotypes. Genetic Epidemiology, 2017, 41, 198-209.	0.6	21
54	Effects of government policies on the spread of COVID-19 worldwide. Scientific Reports, 2021, 11, 20495.	1.6	21

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55	Analysis of population-specific pharmacogenomic variants using next-generation sequencing data. Scientific Reports, 2017, 7, 8416.	1.6	20
56	New Common and Rare Variants Influencing Metabolic Syndrome and Its Individual Components in a Korean Population. Scientific Reports, 2018, 8, 5701.	1.6	20
57	Microbiome Markers of Pancreatic Cancer Based on Bacteria-Derived Extracellular Vesicles Acquired from Blood Samples: A Retrospective Propensity Score Matching Analysis. Biology, 2021, 10, 219.	1.3	20
58	Clinical validation of the 2017 international consensus guidelines on intraductal papillary mucinous neoplasm of the pancreas. Annals of Surgical Treatment and Research, 2019, 97, 58.	0.4	20
59	Effects of covariance model assumptions on hypothesis tests for repeated measurements: analysis of ovarian hormone data and pituitary-pteryomaxillary distance data. Statistics in Medicine, 2001, 20, 2441-2453.	0.8	19
60	Cancer survival classification using integrated data sets and intermediate information. Artificial Intelligence in Medicine, 2014, 62, 23-31.	3.8	19
61	Long-term response to mood stabilizer treatment and its clinical correlates in patients with bipolar disorders: a retrospective observational study. International Journal of Bipolar Disorders, 2017, 5, 24.	0.8	19
62	Clinical validation of scoring systems of postoperative pancreatic fistula after pancreatoduodenectomy: applicability to Eastern cohorts?. Hepatobiliary Surgery and Nutrition, 2019, 8, 211-218.	0.7	19
63	Simple pattern-mixture models for longitudinal data with missing observations: analysis of urinary incontinence data., 1999, 18, 2933-2941.		18
64	Improving Disease Prediction by Incorporating Family Disease History in Risk Prediction Models with Large-Scale Genetic Data. Genetics, 2017, 207, 1147-1155.	1.2	18
65	Integration of Traditional and Metabolomics Biomarkers Identifies Prognostic Metabolites for Predicting Responsiveness to Nutritional Intervention against Oxidative Stress and Inflammation. Nutrients, 2017, 9, 233.	1.7	18
66	Hierarchical structural component modeling of microRNA-mRNA integration analysis. BMC Bioinformatics, 2018, 19, 75.	1.2	17
67	Detecting differential DNA methylation from sequencing of bisulfite converted DNA of diverse species. Briefings in Bioinformatics, 2019, 20, 33-46.	3.2	17
68	Systematic approach identifies RHOA as a potential biomarker therapeutic target for Asian gastric cancer. Oncotarget, 2016, 7, 81435-81451.	0.8	17
69	Development of Web-Based Nomograms to Predict Treatment Response and Prognosis of Epithelial Ovarian Cancer. Cancer Research and Treatment, 2019, 51, 1144-1155.	1.3	17
70	Statistical analysis of MMR vaccine adverse events on aseptic meningitis using the case cross-over design. Statistics in Medicine, 2004, 23, 1871-1883.	0.8	16
71	Joint Identification of Genetic Variants for Physical Activity in Korean Population. International Journal of Molecular Sciences, 2014, 15, 12407-12421.	1.8	16
72	A novel fuzzy set based multifactor dimensionality reduction method for detecting gene–gene interaction. Computational Biology and Chemistry, 2016, 65, 193-202.	1.1	16

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73	Development of Machine Learning Models to Predict Platinum Sensitivity of High-Grade Serous Ovarian Carcinoma. Cancers, 2021, 13, 1875.	1.7	16
74	The relationships of present vegetation, bacteria, and soil properties with soil organic matter characteristics in moist acidic tundra in Alaska. Science of the Total Environment, 2021, 772, 145386.	3.9	15
75	22q11-q13 as a hot spot for prediction of disease-free survival in bile duct cancer: integrative analysis of copy number variations. Cancer Genetics, 2014, 207, 57-69.	0.2	14
76	Practical issues in genomeâ€wide association studies for physical activity. Annals of the New York Academy of Sciences, 2011, 1229, 38-44.	1.8	13
77	A new strategy for enhancing imputation quality of rare variants from next-generation sequencing data via combining SNP and exome chip data. BMC Genomics, 2015, 16, 1109.	1.2	13
78	Bis-class: a new classification tool of methylation status using bayes classifier and local methylation information. BMC Genomics, 2014, 15, 608.	1.2	12
79	Novel Trajectories for Identifying Asthma Phenotypes: A Longitudinal Study in Korean Asthma Cohort, COREA. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1850-1857.e4.	2.0	12
80	Prevalence, behavioral manifestations and associated individual and climatic factors of seasonality in the Korean general population. Comprehensive Psychiatry, 2015, 57, 148-154.	1.5	11
81	Calibration of High-Density Lipoprotein Cholesterol Values From the Korea National Health and Nutrition Examination Survey Data, 2008 to 2015. Annals of Laboratory Medicine, 2017, 37, 1-8.	1.2	11
82	WISARD: workbench for integrated superfast association studies for related datasets. BMC Medical Genomics, 2018, 11, 39.	0.7	11
83	Risk prediction for malignant intraductal papillary mucinous neoplasm of the pancreas: logistic regression versus machine learning. Scientific Reports, 2020, 10, 20140.	1.6	11
84	Prediction Models for the Clinical Severity of Patients With COVID-19 in Korea: Retrospective Multicenter Cohort Study. Journal of Medical Internet Research, 2021, 23, e25852.	2.1	11
85	Diagnostic plots for detecting outlying slides in a cDNA microarray experiment. BioTechniques, 2005, 38, 463-471.	0.8	10
86	Patterns of Gene Expression Associated with Pten Deficiency in the Developing Inner Ear. PLoS ONE, 2014, 9, e97544.	1.1	10
87	HisCoM-GGI: Hierarchical structural component analysis of gene–gene interactions. Journal of Bioinformatics and Computational Biology, 2018, 16, 1840026.	0.3	10
88	Proposal of the minimal number of retrieved regional lymph nodes for accurate staging of distal bile duct cancer and clinical validation of the threeâ€tier lymph node staging system (AJCC 8th edition). Journal of Hepato-Biliary-Pancreatic Sciences, 2020, 27, 75-83.	1.4	10
89	Risk Prediction Using Genome-Wide Association Studies on Type 2 Diabetes. Genomics and Informatics, 2016, 14, 138.	0.4	10
90	Joint Modelling of Repeated Measures and Survival Time Data. Biometrical Journal, 2003, 45, 647-658.	0.6	9

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91	Adjusting heterogeneous ascertainment bias for genetic association analysis with extended families. BMC Medical Genetics, 2015, 16, 62.	2.1	9
92	Effects of genetic variants of ST8SIA2 and NCAM1 genes on seasonal mood changes and circadian preference in the general population. Chronobiology International, 2018, 35, 405-415.	0.9	9
93	Psychopathologic structure of bipolar disorders: exploring dimensional phenotypes, their relationships, and their associations with bipolar I and II disorders. Psychological Medicine, 2019, 49, 2177-2185.	2.7	9
94	The influences of DNA methylation and epigenetic clocks, on metabolic disease, in middle-aged Koreans. Clinical Epigenetics, 2020, 12, 148.	1.8	9
95	Integrative Analysis of Multi-Omics Data Based on Blockwise Sparse Principal Components. International Journal of Molecular Sciences, 2020, 21, 8202.	1.8	9
96	Multiâ€biomarker panel prediction model for diagnosis ofÂpancreatic cancer. Journal of Hepato-Biliary-Pancreatic Sciences, 2023, 30, 122-132.	1.4	9
97	The Homocysteine and Metabolic Syndrome: A Mendelian Randomization Study. Nutrients, 2021, 13, 2440.	1.7	9
98	Nonalcoholic fatty liver disease and early prediction of gestational diabetes mellitus using machine learning methods. Clinical and Molecular Hepatology, 2022, 28, 105-116.	4.5	9
99	Error-pooling-based statistical methods for identifying novel temporal replication profiles of human chromosomes observed by DNA tiling arrays. Nucleic Acids Research, 2007, 35, e69.	6.5	8
100	A Comparative Study on Multifactor Dimensionality Reduction Methods for Detecting Gene-Gene Interactions with the Survival Phenotype. BioMed Research International, 2015, 2015, 1-7.	0.9	8
101	Association between the zinc finger protein 804A (<i><scp>ZNF</scp>804A</i>) gene and the risk of schizophrenia and bipolar I disorder across diagnostic boundaries. Bipolar Disorders, 2017, 19, 305-313.	1.1	8
102	Functional conservation of sequence determinants at rapidly evolving regulatory regions across mammals. PLoS Computational Biology, 2018, 14, e1006451.	1.5	8
103	Fuzzy set-based generalized multifactor dimensionality reduction analysis of gene-gene interactions. BMC Medical Genomics, 2018, 11, 32.	0.7	8
104	Statistical Estimation of Effects of Implemented Government Policies on COVID-19 Situation in South Korea. International Journal of Environmental Research and Public Health, 2021, 18, 2144.	1.2	8
105	Estimation of Undetected Asymptomatic COVID-19 Cases in South Korea Using a Probabilistic Model. International Journal of Environmental Research and Public Health, 2021, 18, 4946.	1.2	8
106	Bayesian methods for contingency tables using Gibbs sampling. Statistical Papers, 2004, 45, 33-50.	0.7	7
107	Phenotype prediction from genome-wide association studies: application to smoking behaviors. BMC Systems Biology, 2012, 6, S11.	3.0	7
108	Detecting Genetic Interactions for Quantitative Traits Usingm-Spacing Entropy Measure. BioMed Research International, 2015, 2015, 1-10.	0.9	7

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109	Two simple algorithms on linear combination of multiple biomarkers to maximize partial area under the ROC curve. Computational Statistics and Data Analysis, 2015, 88, 15-27.	0.7	7
110	Gene-set association tests for next-generation sequencing data. Bioinformatics, 2016, 32, i611-i619.	1.8	7
111	Comparing family-based rare variant association tests for dichotomous phenotypes. BMC Proceedings, 2016, 10, 181-186.	1.8	7
112	GxGrare: gene-gene interaction analysis method for rare variants from high-throughput sequencing data. BMC Systems Biology, 2018, 12, 19.	3.0	7
113	Pathway-Based Integrative Analysis of Metabolome and Microbiome Data from Hepatocellular Carcinoma and Liver Cirrhosis Patients. Cancers, 2020, 12, 2705.	1.7	7
114	Diagnostic model for pancreatic cancer using a multi-biomarker panel. Annals of Surgical Treatment and Research, 2021, 100, 144.	0.4	7
115	Development, validation, and comparison of a nomogram based on radiologic findings for predicting malignancy in intraductal papillary mucinous neoplasms of the pancreas: An international multicenter study. Journal of Hepato-Biliary-Pancreatic Sciences, 2023, 30, 133-143.	1.4	7
116	LPEseq: Local-Pooled-Error Test for RNA Sequencing Experiments with a Small Number of Replicates. PLoS ONE, 2016, 11, e0159182.	1.1	7
117	Nuclear receptor and VEGF pathways for gene-blood lead interactions, on bone mineral density, in Korean smokers. PLoS ONE, 2018, 13, e0193323.	1.1	7
118	Gene-Gene Interaction Analysis for the Accelerated Failure Time Model Using a Unified Model-Based Multifactor Dimensionality Reduction Method. Genomics and Informatics, 2016, 14, 166.	0.4	7
119	On the use of working correlation matrices in the gee approach for longitudinal data. Communications in Statistics Part B: Simulation and Computation, 1999, 28, 1011-1029.	0.6	6
120	Risk prediction of type 2 diabetes using common and rare variants. International Journal of Data Mining and Bioinformatics, 2018, 20, 77.	0.1	6
121	Gene-gene interaction analysis for quantitative trait using cluster-based multifactor dimensionality reduction method. International Journal of Data Mining and Bioinformatics, 2018, 20, 1.	0.1	6
122	Multivariate Cluster-Based Multifactor Dimensionality Reduction to Identify Genetic Interactions for Multiple Quantitative Phenotypes. BioMed Research International, 2019, 2019, 1-10.	0.9	6
123	Pathway analysis of rare variants for the clustered phenotypes by using hierarchical structured components analysis. BMC Medical Genomics, 2019, 12, 100.	0.7	6
124	HisCoM-PAGE: Hierarchical Structural Component Models for Pathway Analysis of Gene Expression Data. Genes, 2019, 10, 931.	1.0	6
125	Development and External Validation of Survival Prediction Model for Pancreatic Cancer Using Two Nationwide Databases: Surveillance, Epidemiology and End Results (SEER) and Korea Tumor Registry System-Biliary Pancreas (KOTUS-BP). Gut and Liver, 2021, 15, 912-921.	1.4	6
126	Robust Gene-Gene Interaction Analysis in Genome Wide Association Studies. PLoS ONE, 2015, 10, e0135016.	1.1	6

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127	Prediction of Quantitative Traits Using Common Genetic Variants: Application to Body Mass Index. Genomics and Informatics, 2016, 14, 149.	0.4	6
128	Clinicopathologic and protein markers distinguishing the "polymerase epsilon exonuclease―from the "copy number low―subtype of endometrial cancer. Journal of Gynecologic Oncology, 2022, 33, .	1.0	6
129	<i>FARVATX</i> : Familyâ€Based Rare Variant Association Test for Xâ€Linked Genes. Genetic Epidemiology, 2016, 40, 475-485.	0.6	5
130	Analysis of multiple related phenotypes in genome-wide association studies. Journal of Bioinformatics and Computational Biology, 2016, 14, 1644005.	0.3	5
131	Generalized estimating equations with stabilized working correlation structure. Computational Statistics and Data Analysis, 2017, 106, 1-11.	0.7	5
132	Structural equation modeling for hypertension and type 2 diabetes based on multiple SNPs and multiple phenotypes. PLoS ONE, 2019, 14, e0217189.	1.1	5
133	Gene-Gene Interaction Analysis for the Survival Phenotype Based on the Kaplan-Meier Median Estimate. BioMed Research International, 2020, 2020, 1-10.	0.9	5
134	PATHOME-Drug: a subpathway-based polypharmacology drug-repositioning method. Bioinformatics, 2022, 38, 444-452.	1.8	5
135	Prognostic significance of E-cadherin and ZEB1 expression in intraductal papillary mucinous neoplasm. Oncotarget, 2018, 9, 306-320.	0.8	5
136	CONCORD biomarker prediction for novel drug introduction to different cancer types. Oncotarget, 2018, 9, 1091-1106.	0.8	5
137	Confidence intervals for the COVID-19 neutralizing antibody retention rate in the Korean population. Genomics and Informatics, 2020, 18, e31.	0.4	5
138	Practical Issues in Screening and Variable Selection in Genome-Wide Association Analysis. Cancer Informatics, 2014, 13s7, CIN.S16350.	0.9	4
139	Unified Cox model based multifactor dimensionality reduction method for gene-gene interaction analysis of the survival phenotype. BioData Mining, 2018, 11, 27.	2.2	4
140	Changes in serum fibronectin levels predict tumor recurrence in patients with early hepatocellular carcinoma after curative treatment. Scientific Reports, 2020, 10, 21313.	1.6	4
141	Enhanced Permutation Tests via Multiple Pruning. Frontiers in Genetics, 2020, $11,509$.	1.1	4
142	Do we need to reconsider the CMAM admission and discharge criteria?; an analysis of CMAM data in South Sudan. BMC Public Health, 2020, 20, 511.	1.2	4
143	Forecasting of the COVID-19 pandemic situation of Korea. Genomics and Informatics, 2021, 19, e11.	0.4	4
144	Inclusive Quantification Assay of Serum Desâ€Î³â€Carboxyprothrombin Proteoforms for Hepatocellular Carcinoma Surveillance by Targeted Mass Spectrometry. Hepatology Communications, 2021, 5, 1767-1783.	2.0	4

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145	EFMDR-Fast: An Application of Empirical Fuzzy Multifactor Dimensionality Reduction for Fast Execution. Genomics and Informatics, 2018, 16, e37.	0.4	4
146	Comparison of survival prediction models for pancreatic cancer: Cox model versus machine learning models. Genomics and Informatics, 2022, 20, e23.	0.4	4
147	A chi-square test for detecting multiple joint genetic variants in genome-wide association studies. , $2011, \ldots$		3
148	CARAT-GxG: CUDA-Accelerated Regression Analysis Toolkit for Large-Scale Gene–Gene Interaction with GPU Computing System. Cancer Informatics, 2014, 13s7, CIN.S16349.	0.9	3
149	Pathway-Driven Discovery of Rare Mutational Impact on Cancer. BioMed Research International, 2014, 2014, 1-10.	0.9	3
150	An Efficient Stepwise Statistical Test to Identify Multiple Linked Human Genetic Variants Associated with Specific Phenotypic Traits. PLoS ONE, 2015, 10, e0138700.	1.1	3
151	Using the Generalized Index of Dissimilarity to Detect Gene-Gene Interactions in Multi-Class Phenotypes. PLoS ONE, 2016, 11, e0158668.	1.1	3
152	Multifactor dimensionality reduction analysis of multiple binary traits for gene-gene interaction. International Journal of Data Mining and Bioinformatics, 2016, 14, 293.	0.1	3
153	Correlation estimation with singly truncated bivariate data. Statistics in Medicine, 2017, 36, 1977-1988.	0.8	3
154	Proposed new staging system for ampulla of Vater cancer with greater discriminatory ability: multinational study from eastern and western centers. Journal of Hepato-Biliary-Pancreatic Sciences, 2017, 24, 475-484.	1.4	3
155	PreCimp: Preâ€collapsing imputation approach increases imputation accuracy of rare variants in terms of collapsed variables. Genetic Epidemiology, 2017, 41, 41-50.	0.6	3
156	Statistical methods for metagenomics data analysis. International Journal of Data Mining and Bioinformatics, 2017, 19, 366.	0.1	3
157	Fuzzy heaping mechanism for heaped count data with imprecision. Soft Computing, 2018, 22, 4585-4594.	2.1	3
158	Effect of Interaction between Early Menarche and Genetic Polymorphisms on Triglyceride. Oxidative Medicine and Cellular Longevity, 2019, 2019, 1-9.	1.9	3
159	Develop Nomogram to Predict Malignancy of Intraductal Papillary Mucinous Neoplasm. Methods in Molecular Biology, 2019, 1882, 23-32.	0.4	3
160	Hierarchical structural component model for pathway analysis of common variants. BMC Medical Genomics, 2020, 13, 26.	0.7	3
161	Which National Factors Are Most Influential in the Spread of COVID-19?. International Journal of Environmental Research and Public Health, 2021, 18, 7592.	1.2	3
162	Updated confidence intervals for the COVID-19 antibody retention rate in the Korean population. Genomics and Informatics, 2020, 18, e45.	0.4	3

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163	Spatial rank-based multifactor dimensionality reduction to detect gene–gene interactions for multivariate phenotypes. BMC Bioinformatics, 2021, 22, 480.	1.2	3
164	Analysis of Pharmacogenomic Variants Associated with Population Differentiation. PLoS ONE, 2015, 10, e0119994.	1.1	3
165	Searching susceptibility genes for antipsychotic-induced weight gain: is the 5-HT2C receptor gene a promising candidate?. Personalized Medicine, 2007, 4, 357-361.	0.8	2
166	Estimating cancer gene pathway proximity using network interaction. , 2014, , .		2
167	Biomarker development for pancreatic ductal adenocarcinoma using integrated analysis of mRNA and miRNA expression. , 2014, , .		2
168	VizEpis: A visualization and mapping tool for interpreting epistasis., 2015,,.		2
169	Meta-Qtest: meta-analysis of quadratic test for rare variants. BMC Medical Genomics, 2019, 12, 102.	0.7	2
170	Characterization and Validation of an "Acute Aerobic Exercise Load―as a Tool to Assess Antioxidative and Anti-inflammatory Nutrition in Healthy Subjects Using a Statistically Integrated Approach in a Comprehensive Clinical Trial. Oxidative Medicine and Cellular Longevity, 2019, 2019, 1-14.	1.9	2
171	HisCoM-G×E: Hierarchical Structural Component Analysis of Gene-Based Gene–Environment Interactions. International Journal of Molecular Sciences, 2020, 21, 6724.	1.8	2
172	Editor's introduction to this issue (G&l 19:1, 2021). Genomics and Informatics, 2021, 19, e1.	0.4	2
173	Association between the Arylalkylamine N-Acetyltransferase (AANAT) Gene and Seasonality in Patients with Bipolar Disorder. Psychiatry Investigation, 2021, 18, 453-462.	0.7	2
174	Genome-wide association study identified a novel genetic variation in HLA DRB1 associated with drug hypersensitivity. Annals of Allergy, Asthma and Immunology, $2021, \dots$	0.5	2
175	Kernel-based hierarchical structural component models for pathway analysis. Bioinformatics, 2022, 38, 3078-3086.	1.8	2
176	Mathematical modeling of the impact of Omicron variant on the COVID-19 situation in South Korea. Genomics and Informatics, 2022, 20, e22.	0.4	2
177	Comparative studies for developing protein based cancer prediction model to maximise the ROC-AUC with various variable selection methods. International Journal of Data Mining and Bioinformatics, 2016, 16, 64.	0.1	1
178	Multivariate approach to the analysis of correlated RNA-seq data., 2016,,.		1
179	Risk prediction using common and rare genetic variants: Application to Type 2 diabetes. , 2017, , .		1
180	Cluster-based multifactor dimensionality reduction method to identify gene-gene interactions for quantitative traits in genome-wide studies. , 2017, , .		1

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181	Exact association test for small size sequencing data. BMC Medical Genomics, 2018, 11, 30.	0.7	1
182	Detecting population structures by independent component analysis. , 2018, , .		1
183	Identification of genetic loci affecting body mass index through interaction with multiple environmental factors using structured linear mixed model. Scientific Reports, 2021, 11, 5001.	1.6	1
184	Identifying miRNA-mRNA Integration Set Associated With Survival Time. Frontiers in Genetics, 2021, 12, 634922.	1.1	1
185	Pure additive contribution of genetic variants to a risk prediction model using propensity score matching: application to type 2 diabetes. Genomics and Informatics, 2019, 17, e47.	0.4	1
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