

Shaoqi Rao

List of Publications by Year in descending order

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Version: 2024-02-01

78
papers

2,896
citations

236833

25
h-index

168321

53
g-index

80
all docs

80
docs citations

80
times ranked

3919
citing authors

#	ARTICLE	IF	CITATIONS
1	Stratified meta-analysis by ethnicity revealed that ADRB3 Trp64Arg polymorphism was associated with coronary artery disease in Asians, but not in Caucasians. <i>Medicine (United States)</i> , 2020, 99, e18914.	0.4	7
2	A Seven-Long Non-coding RNA Signature Improves Prognosis Prediction of Lung Adenocarcinoma: An Integrated Competing Endogenous RNA Network Analysis. <i>Frontiers in Genetics</i> , 2020, 11, 625977.	1.1	5
3	Genome-Wide Association and Functional Studies Identify <i>SCML4</i> and <i>THSD7A</i> as Novel Susceptibility Genes for Coronary Artery Disease. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2018, 38, 964-975.	1.1	32
4	Complex Segregation Analysis Provides Evidence for Autosomal Dominant Transmission in the Chinese Han Families with Ankylosing Spondylitis. <i>BioMed Research International</i> , 2017, 2017, 1-6.	0.9	1
5	Association between apolipoprotein B EcoRI polymorphisms and coronary heart disease. <i>Wiener Klinische Wochenschrift</i> , 2016, 128, 890-897.	1.0	2
6	Association Between Apolipoprotein B XbaI Polymorphism and Coronary Heart Disease in Han Chinese Population: A Meta-Analysis. <i>Genetic Testing and Molecular Biomarkers</i> , 2016, 20, 304-311.	0.3	6
7	Molecular Basis of Gene-Gene Interaction: Cyclic Cross-Regulation of Gene Expression and Post-GWAS Gene-Gene Interaction Involved in Atrial Fibrillation. <i>PLoS Genetics</i> , 2015, 11, e1005393.	1.5	47
8	Association of <i>DRD3</i> , <i>COMT</i> , and <i>SLC6A4</i> Gene Polymorphisms with Type 2 Diabetes in Southern Chinese: A Hospital-Based Case-Control Study. <i>Diabetes Technology and Therapeutics</i> , 2015, 17, 580-586.	2.4	24
9	Knowledge-based analysis of genetic associations of rheumatoid arthritis to inform studies searching for pleiotropic genes: a literature review and network analysis. <i>Arthritis Research and Therapy</i> , 2015, 17, 202.	1.6	14
10	Genome-Wide Linkage Scan Identifies Two Novel Genetic Loci for Coronary Artery Disease: In GeneQuest Families. <i>PLoS ONE</i> , 2014, 9, e113935.	1.1	8
11	Pathway-based Analysis of the Hidden Genetic Heterogeneities in Cancers. <i>Genomics, Proteomics and Bioinformatics</i> , 2014, 12, 31-38.	3.0	6
12	A Novel Molecular Diagnostic Marker for Familial and Early-Onset Coronary Artery Disease and Myocardial Infarction in the <i>LRP8</i> Gene. <i>Circulation: Cardiovascular Genetics</i> , 2014, 7, 514-520.	5.1	21
13	Identifying functional modules for coronary artery disease by a prior knowledge-based approach. <i>Gene</i> , 2014, 537, 260-268.	1.0	11
14	Genomic Scan Reveals Loci under Altitude Adaptation in Tibetan and Dahe Pigs. <i>PLoS ONE</i> , 2014, 9, e110520.	1.1	42
15	Association between rs10118757(A/G) in methylthioadenosine phosphorylase gene and coronary artery disease in Chinese Hans. <i>Gene</i> , 2013, 526, 344-346.	1.0	3
16	Atorvastatin Treatment of Rats with Ischemia-Reperfusion Injury Improves Adipose-Derived Mesenchymal Stem Cell Migration and Survival via the SDF-1 α /CXCR-4 Axis. <i>PLoS ONE</i> , 2013, 8, e79100.	1.1	27
17	To Control False Positives in Gene-Gene Interaction Analysis: Two Novel Conditional Entropy-Based Approaches. <i>PLoS ONE</i> , 2013, 8, e81984.	1.1	3
18	Joint Effects of Genetic Variants in Multiple Loci on the Risk of Coronary Artery Disease in Chinese Han Subjects. <i>Circulation Journal</i> , 2012, 76, 1987-1992.	0.7	21

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19	SDF-1 α upregulation by atorvastatin in rats with acute myocardial infarction via nitric oxide production confers anti-inflammatory and anti-apoptotic effects. <i>Journal of Biomedical Science</i> , 2012, 19, 99.	2.6	24
20	Lack of association between four SNPs in the SLC22A3-LPAL2-LPA gene cluster and coronary artery disease in a Chinese Han population: a case control study. <i>Lipids in Health and Disease</i> , 2012, 11, 128.	1.2	15
21	Phenotype-Genotype Correlation in 295 Chinese Deaf Subjects with Biallelic Causative Mutations in the GJB2 Gene. <i>Genetic Testing and Molecular Biomarkers</i> , 2011, 15, 619-625.	0.3	10
22	Adaptation of the Audit of Diabetes-Dependent Quality of Life questionnaire to people with diabetes in China. <i>Diabetes Research and Clinical Practice</i> , 2011, 94, 45-52.	1.1	31
23	Disease embryo development network reveals the relationship between disease genes and embryo development genes. <i>Journal of Theoretical Biology</i> , 2011, 287, 100-108.	0.8	9
24	Short-term Outcomes of Induction Therapy With Tacrolimus Versus Cyclophosphamide for Active Lupus Nephritis: A Multicenter Randomized Clinical Trial. <i>American Journal of Kidney Diseases</i> , 2011, 57, 235-244.	2.1	169
25	DOSim: An R package for similarity between diseases based on Disease Ontology. <i>BMC Bioinformatics</i> , 2011, 12, 266.	1.2	88
26	Brief communication: Y-chromosome haplogroup analysis indicates that Chinese Tuvans share distinctive affinity with Siberian Tuvans. <i>American Journal of Physical Anthropology</i> , 2011, 144, 492-497.	2.1	11
27	Efficacy of Atorvastatin combined with adipose-derived mesenchymal stem cell transplantation on cardiac function in rats with acute myocardial infarction. <i>Acta Biochimica Et Biophysica Sinica</i> , 2011, 43, 857-866.	0.9	26
28	Ala499Val (C > T) and Lys939Gln (A > C) polymorphisms of the XPC gene: their correlation with the risk of primary gallbladder adenocarcinoma—a case-control study in China. <i>Carcinogenesis</i> , 2011, 32, 496-501.	1.3	18
29	Functional Homogeneity in microRNA Target Heterogeneity—a New Sight into Human microRNomics. <i>OMICS A Journal of Integrative Biology</i> , 2011, 15, 25-35.	1.0	10
30	Plasma Oxidized Low-Density Lipoprotein Is an Independent Risk Factor in Young Patients with Coronary Artery Disease. <i>Disease Markers</i> , 2011, 31, 295-301.	0.6	14
31	A Novel Evolution-Based Method for Detecting Gene-Gene Interactions. <i>PLoS ONE</i> , 2011, 6, e26435.	1.1	1
32	Plasma oxidized low-density lipoprotein is an independent risk factor in young patients with coronary artery disease. <i>Disease Markers</i> , 2011, 31, 295-301.	0.6	17
33	Psychometric properties of the Chinese version of the Pediatric Quality of Life Inventory, 4.0 generic core scales. <i>Quality of Life Research</i> , 2010, 19, 1229-1233.	1.5	68
34	A genome-wide linkage scan identifies multiple quantitative trait loci for HDL-cholesterol levels in families with premature CAD and MI. <i>Journal of Lipid Research</i> , 2010, 51, 1442-1451.	2.0	11
35	Mapping and characterization of two relevance networks from SNP and gene levels. <i>Progress in Natural Science: Materials International</i> , 2009, 19, 653-657.	1.8	1
36	Behavioural development of school-aged children who live around a multi-metal sulphide mine in Guangdong province, China: a cross-sectional study. <i>BMC Public Health</i> , 2009, 9, 217.	1.2	69

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37	Effects of cutoff thresholds for minor allele frequencies on HapMap resolution: A real dataset-based evaluation of the Chinese Han and Tibetan populations. <i>Science Bulletin</i> , 2009, 54, 2069-2075.	4.3	1
38	Association between four SNPs on chromosome 9p21 and myocardial infarction is replicated in an Italian population. <i>Journal of Human Genetics</i> , 2008, 53, 144-150.	1.1	112
39	Constructing disease-specific gene networks using pair-wise relevance metric: Application to colon cancer identifies interleukin 8, desmin and enolase 1 as the central elements. <i>BMC Systems Biology</i> , 2008, 2, 72.	3.0	64
40	Cell cycle-dependent gene networks relevant to cancer. <i>Progress in Natural Science: Materials International</i> , 2008, 18, 945-952.	1.8	3
41	A feature ensemble technology to identify molecular mechanisms for distinction between multiple subtypes of lymphoma. <i>Progress in Natural Science: Materials International</i> , 2008, 18, 1491-1500.	1.8	1
42	A systematic method for mapping multiple loci: An application to construct a genetic network for rheumatoid arthritis. <i>Gene</i> , 2008, 408, 104-111.	1.0	13
43	Four SNPs on Chromosome 9p21 in a South Korean Population Implicate a Genetic Locus That Confers High Cross-Race Risk for Development of Coronary Artery Disease. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2008, 28, 360-365.	1.1	183
44	Edge-based scoring and searching method for identifying condition-responsive protein protein interaction sub-network. <i>Bioinformatics</i> , 2007, 23, 2121-2128.	1.8	139
45	An LRP8 Variant Is Associated with Familial and Premature Coronary Artery Disease and Myocardial Infarction. <i>American Journal of Human Genetics</i> , 2007, 81, 780-791.	2.6	77
46	Data mining of RNA expression and DNA genotype data: Presentation Group 5 contributions to Genetic Analysis Workshop 15. <i>Genetic Epidemiology</i> , 2007, 31, S43-S50.	0.6	1
47	Unravelling the hidden heterogeneities of diffuse large B-cell lymphoma based on coupled two-way clustering. <i>BMC Genomics</i> , 2007, 8, 332.	1.2	7
48	Widely predicting specific protein functions based on protein-protein interaction data and gene expression profile. <i>Science in China Series C: Life Sciences</i> , 2007, 50, 125-134.	1.3	10
49	A novel model-free approach for reconstruction of time-delayed gene regulatory networks. <i>Science in China Series C: Life Sciences</i> , 2006, 49, 190-200.	1.3	2
50	Identifying disease feature genes based on cellular localized gene functional modules and regulation networks. <i>Science Bulletin</i> , 2006, 51, 1848-1856.	1.7	3
51	Discovery of Time-Delayed Gene Regulatory Networks based on temporal gene expression profiling. <i>BMC Bioinformatics</i> , 2006, 7, 26.	1.2	60
52	Effects of replacing the unreliable cDNA microarray measurements on the disease classification based on gene expression profiles and functional modules. <i>Bioinformatics</i> , 2006, 22, 2883-2889.	1.8	36
53	SAGE Programs. <i>Methods in Molecular Medicine</i> , 2006, , 61-89.	0.8	3
54	Analysis of Sib-Pair IBD Profiles Using Ensemble Decision Tree Approach: Application to Alcoholism. <i>Lecture Notes in Computer Science</i> , 2006, , 774-779.	1.0	0

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55	Association Research on Potassium Channel Subtypes and Functional Sites. Lecture Notes in Computer Science, 2006, , 780-785.	1.0	0
56	SAGE Programs: Model-Free Linkage Analysis for Complex Cardiovascular Phenotypes. , 2006, 128, 61-90.		2
57	Towards precise classification of cancers based on robust gene functional expression profiles. BMC Bioinformatics, 2005, 6, 58.	1.2	146
58	Reply to Ray and Weeks: Linkage for Restless Legs Syndrome on Chromosome 9p Is Significant. American Journal of Human Genetics, 2005, 76, 707-710.	2.6	6
59	A robust hybrid between genetic algorithm and support vector machine for extracting an optimal feature gene subset. Genomics, 2005, 85, 16-23.	1.3	112
60	Miscues on the "lack of MEF2A mutations" in coronary artery disease. Journal of Clinical Investigation, 2005, 115, 1399-1400.	3.9	14
61	Analysis of Sib-Pair IBD Profiles and Genomic Context for Identification of the Relevant Molecular Signatures for Alcoholism. Lecture Notes in Computer Science, 2005, , 845-851.	1.0	1
62	A Novel Feature Ensemble Technology to Improve Prediction Performance of Multiple Heterogeneous Phenotypes Based on Microarray Data. Lecture Notes in Computer Science, 2005, , 869-879.	1.0	0
63	Large-Scale Ensemble Decision Analysis of Sib-Pair IBD Profiles for Identification of the Relevant Molecular Signatures for Alcoholism. Lecture Notes in Computer Science, 2005, , 1184-1189.	1.0	1
64	A Novel Ensemble Decision Tree Approach for Mining Genes Coding Ion Channels for Cardiopathy Subtype. Lecture Notes in Computer Science, 2005, , 852-860.	1.0	1
65	Gene mining: a novel and powerful ensemble decision approach to hunting for disease genes using microarray expression profiling. Nucleic Acids Research, 2004, 32, 2685-2694.	6.5	85
66	Genome-Wide Linkage Scan Identifies a Novel Genetic Locus on Chromosome 5p13 for Neonatal Atrial Fibrillation Associated With Sudden Death and Variable Cardiomyopathy. Circulation, 2004, 110, 3753-3759.	1.6	114
67	Identification of an angiogenic factor that when mutated causes susceptibility to Klippelâ€™Trenaunay syndrome. Nature, 2004, 427, 640-645.	13.7	289
68	Premature Myocardial Infarction Novel Susceptibility Locus on Chromosome 1P34-36 Identified by Genomewide Linkage Analysis. American Journal of Human Genetics, 2004, 74, 262-271.	2.6	195
69	Genomewide Linkage Scan Identifies a Novel Susceptibility Locus for Restless Legs Syndrome on Chromosome 9p. American Journal of Human Genetics, 2004, 74, 876-885.	2.6	186
70	On the Significance of Linkage Studies of Complex Traits: Reply to Newton-Cheh et al.. American Journal of Human Genetics, 2004, 75, 152-154.	2.6	4
71	An ensemble method for gene discovery based on DNA mi-croarray data. Science in China Series C: Life Sciences, 2004, 47, 396.	1.3	6
72	Longitudinal data analysis in pedigree studies. Genetic Epidemiology, 2003, 25, S18-S28.	0.6	32

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73	Genetic linkage analysis of longitudinal hypertension phenotypes using three summary measures. BMC Genetics, 2003, 4, S24.	2.7	4
74	Multivariate sib-pair linkage analysis of longitudinal phenotypes by three step-wise analysis approaches. BMC Genetics, 2003, 4, S68.	2.7	6
75	Locating the Genes Underlying a Simulated Complex Disease by Discriminant Analysis. Genetic Epidemiology, 2001, 21, S516-S521.	0.6	10
76	Strategies for genetic mapping of categorical traits. Genetica, 2000, 109, 183-197.	0.5	20
77	Linkage of chromosome 1 markers to alcoholism-related phenotypes by sib pair linkage analysis of principal components. Genetic Epidemiology, 1999, 17, S271-6.	0.6	16
78	Mapping quantitative trait loci for ordered categorical traits in four-way crosses. Heredity, 1998, 81, 214-224.	1.2	58