

# Shuo Jiao

## List of Publications by Year in descending order

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Version: 2024-02-01

31  
papers

1,991  
citations

361413

20  
h-index

434195

31  
g-index

31  
all docs

31  
docs citations

31  
times ranked

5082  
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of Genetic Susceptibility Loci for Colorectal Tumors in a Genome-Wide Meta-analysis. <i>Gastroenterology</i> , 2013, 144, 799-807.e24.	1.3	292
2	Whole-Exome Sequencing Identifies Rare and Low-Frequency Coding Variants Associated with LDL Cholesterol. <i>American Journal of Human Genetics</i> , 2014, 94, 233-245.	6.2	193
3	Meta-analysis of new genome-wide association studies of colorectal cancer risk. <i>Human Genetics</i> , 2012, 131, 217-234.	3.8	183
4	Association of Aspirin and NSAID Use With Risk of Colorectal Cancer According to Genetic Variants. <i>JAMA - Journal of the American Medical Association</i> , 2015, 313, 1133.	7.4	171
5	Characterization of Gene-Environment Interactions for Colorectal Cancer Susceptibility Loci. <i>Cancer Research</i> , 2012, 72, 2036-2044.	0.9	140
6	Genome-wide association study of colorectal cancer identifies six new susceptibility loci. <i>Nature Communications</i> , 2015, 6, 7138.	12.8	138
7	Estimating the heritability of colorectal cancer. <i>Human Molecular Genetics</i> , 2014, 23, 3898-3905.	2.9	114
8	Genome-Wide Diet-Gene Interaction Analyses for Risk of Colorectal Cancer. <i>PLoS Genetics</i> , 2014, 10, e1004228.	3.5	81
9	Powerful Cocktail Methods for Detecting Genome-Wide Gene-Environment Interaction. <i>Genetic Epidemiology</i> , 2012, 36, 183-194.	1.3	80
10	Common and rare von Willebrand factor (VWF) coding variants, VWF levels, and factor VIII levels in African Americans: the NHLBI Exome Sequencing Project. <i>Blood</i> , 2013, 122, 590-597.	1.4	70
11	A Systematic Mapping Approach of 16q12.2/FTO and BMI in More Than 20,000 African Americans Narrows in on the Underlying Functional Variation: Results from the Population Architecture using Genomics and Epidemiology (PAGE) Study. <i>PLoS Genetics</i> , 2013, 9, e1003171.	3.5	63
12	A Pooled Analysis of Smoking and Colorectal Cancer: Timing of Exposure and Interactions with Environmental Factors. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2012, 21, 1974-1985.	2.5	54
13	Gene-Environment Interaction Involving Recently Identified Colorectal Cancer Susceptibility Loci. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2014, 23, 1824-1833.	2.5	48
14	SBERIA: Set-Based Gene-Environment Interaction Test for Rare and Common Variants in Complex Diseases. <i>Genetic Epidemiology</i> , 2013, 37, 452-464.	1.3	39
15	Genome-Wide Interaction Analyses between Genetic Variants and Alcohol Consumption and Smoking for Risk of Colorectal Cancer. <i>PLoS Genetics</i> , 2016, 12, e1006296.	3.5	38
16	Imputation of coding variants in African Americans: better performance using data from the exome sequencing project. <i>Bioinformatics</i> , 2013, 29, 2744-2749.	4.1	36
17	Pleiotropic effects of genetic risk variants for other cancers on colorectal cancer risk: PAGE, GECCO and CCFR consortia. <i>Gut</i> , 2014, 63, 800-807.	12.1	35
18	Genome-Wide Search for Gene-Gene Interactions in Colorectal Cancer. <i>PLoS ONE</i> , 2012, 7, e52535.	2.5	35

#	ARTICLE	IF	CITATIONS
19	Identification of a common variant with potential pleiotropic effect on risk of inflammatory bowel disease and colorectal cancer. <i>Carcinogenesis</i> , 2015, 36, 999-1007.	2.8	28
20	The use of imputed values in the meta-analysis of genome-wide association studies. <i>Genetic Epidemiology</i> , 2011, 35, 597-605.	1.3	27
21	CYP24A1 variant modifies the association between use of oestrogen plus progestogen therapy and colorectal cancer risk. <i>British Journal of Cancer</i> , 2016, 114, 221-229.	6.4	18
22	Genetic variants of adiponectin and risk of colorectal cancer. <i>International Journal of Cancer</i> , 2015, 137, 154-164.	5.1	16
23	Powerful Set-Based Gene-Environment Interaction Testing Framework for Complex Diseases. <i>Genetic Epidemiology</i> , 2015, 39, 609-618.	1.3	15
24	Whole-exome imputation of sequence variants identified two novel alleles associated with adult body height in African Americans. <i>Human Molecular Genetics</i> , 2014, 23, 6607-6615.	2.9	14
25	A Variational Bayes Discrete Mixture Test for Rare Variant Association. <i>Genetic Epidemiology</i> , 2014, 38, 21-30.	1.3	12
26	Evaluation of the colorectal cancer risk conferred by rare <i>UNC5C</i> alleles. <i>World Journal of Gastroenterology</i> , 2014, 20, 204.	3.3	11
27	The t-mixture model approach for detecting differentially expressed genes in microarrays. <i>Functional and Integrative Genomics</i> , 2008, 8, 181-186.	3.5	10
28	On correcting the overestimation of the permutation-based false discovery rate estimator. <i>Bioinformatics</i> , 2008, 24, 1655-1661.	4.1	10
29	No Evidence of Gene-Calcium Interactions from Genome-Wide Analysis of Colorectal Cancer Risk. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2014, 23, 2971-2976.	2.5	9
30	Fine-Mapping of Common Genetic Variants Associated with Colorectal Tumor Risk Identified Potential Functional Variants. <i>PLoS ONE</i> , 2016, 11, e0157521.	2.5	8
31	Estimating the Proportion of Equivalently Expressed Genes in Microarray Data Based on Transformed Test Statistics. <i>Journal of Computational Biology</i> , 2010, 17, 177-187.	1.6	3