Yongyong Shi

List of Publications by Year in descending order

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		147566	123241
107	4,241	31	61
papers	citations	h-index	g-index
111	111	111	8854
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	A partition-ligation-combination-subdivision EM algorithm for haplotype inference with multiallelic markers: update of the SHEsis (http://analysis.bio-x.cn). Cell Research, 2009, 19, 519-523.	5.7	706
2	Genome-wide association study identifies eight new risk loci for polycystic ovary syndrome. Nature Genetics, 2012, 44, 1020-1025.	9.4	505
3	The OncoArray Consortium: A Network for Understanding the Genetic Architecture of Common Cancers. Cancer Epidemiology Biomarkers and Prevention, 2017, 26, 126-135.	1.1	278
4	A genome-wide association study identifies new susceptibility loci for non-cardia gastric cancer at 3q13.31 and 5p13.1. Nature Genetics, 2011, 43, 1215-1218.	9.4	250
5	Common variants on 8p12 and 1q24.2 confer risk of schizophrenia. Nature Genetics, 2011, 43, 1224-1227.	9.4	224
6	Proteogenomic characterization identifies clinically relevant subgroups of intrahepatic cholangiocarcinoma. Cancer Cell, 2022, 40, 70-87.e15.	7.7	120
7	Identification of recurrent USP48 and BRAF mutations in Cushing's disease. Nature Communications, 2018, 9, 3171.	5.8	106
8	Genome-wide association analysis identifies three new risk loci for gout arthritis in Han Chinese. Nature Communications, 2015, 6, 7041.	5.8	88
9	SHEsisPlus, a toolset for genetic studies on polyploid species. Scientific Reports, 2016, 6, 24095.	1.6	77
10	Consensus paper of the WFSBP Task Force on Genetics: Genetics, epigenetics and gene expression markers of major depressive disorder and antidepressant response. World Journal of Biological Psychiatry, 2017, 18, 5-28.	1.3	75
11	DNA origami-based shape IDs for single-molecule nanomechanical genotyping. Nature Communications, 2017, 8, 14738.	5.8	73
12	Identification of new susceptibility loci for gastric non-cardia adenocarcinoma: pooled results from two Chinese genome-wide association studies. Gut, 2017, 66, 581-587.	6.1	68
13	An international genome-wide meta-analysis of primary biliary cholangitis: Novel risk loci and candidate drugs. Journal of Hepatology, 2021, 75, 572-581.	1.8	62
14	Exome Array Analysis Identifies Variants in SPOCD1 and BTN3A2 That Affect Risk for Gastric Cancer. Gastroenterology, 2017, 152, 2011-2021.	0.6	58
15	Germline Mutations in CDH23, Encoding Cadherin-Related 23, Are Associated with Both Familial and Sporadic Pituitary Adenomas. American Journal of Human Genetics, 2017, 100, 817-823.	2.6	57
16	Susceptibility loci for metabolic syndrome and metabolic components identified in Han Chinese: a multiâ€stage genomeâ€wide association study. Journal of Cellular and Molecular Medicine, 2017, 21, 1106-1116.	1.6	56
17	Genome-wide Analysis of the Role of Copy Number Variation in Schizophrenia Risk in Chinese. Biological Psychiatry, 2016, 80, 331-337.	0.7	55
18	Wholeâ€exome sequencing of oral mucosal melanoma reveals mutational profile and therapeutic targets. Journal of Pathology, 2018, 244, 358-366.	2.1	52

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19	Genetic risk of extranodal natural killer T-cell lymphoma: a genome-wide association study in multiple populations. Lancet Oncology, The, 2020, 21, 306-316.	5.1	49
20	Genetics of schizophrenia: A consensus paper of the WFSBP Task Force on Genetics. World Journal of Biological Psychiatry, 2017, 18, 492-505.	1.3	48
21	A genome-wide gene-environment interaction analysis for tobacco smoke and lung cancer susceptibility. Carcinogenesis, 2014, 35, 1528-1535.	1.3	47
22	A Genome-Wide Association Study Identifies a Locus on TERT for Mean Telomere Length in Han Chinese. PLoS ONE, 2014, 9, e85043.	1.1	46
23	Genome-Wide Association Study of Bladder Cancer in a Chinese Cohort Reveals a New Susceptibility Locus at 5q12.3. Cancer Research, 2016, 76, 3277-3284.	0.4	46
24	Common variants at 10p12.31, 10q21.1 and 13q12.13 are associated with sporadic pituitary adenoma. Nature Genetics, 2015, 47, 793-797.	9.4	43
25	Low-Frequency Coding Variants at 6p21.33 and 20q11.21 Are Associated with Lung Cancer Risk in Chinese Populations. American Journal of Human Genetics, 2015, 96, 832-840.	2.6	41
26	MicroRNA-137 Inhibits EFNB2 Expression Affected by a Genetic Variant and Is Expressed Aberrantly in Peripheral Blood of Schizophrenia Patients. EBioMedicine, 2016, 12, 133-142.	2.7	41
27	Genomic dissection of 43 serum urate-associated loci provides multiple insights into molecular mechanisms of urate control. Human Molecular Genetics, 2020, 29, 923-943.	1.4	40
28	Distinct severity stages of obstructive sleep apnoea are correlated with unique dyslipidaemia: large-scale observational study. Thorax, 2016, 71, 347-355.	2.7	38
29	Significant association of GRM7 and GRM8 genes with schizophrenia and major depressive disorder in the Han Chinese population. European Neuropsychopharmacology, 2016, 26, 136-146.	0.3	35
30	MiRNAs of peripheral blood as the biomarker of schizophrenia. Hereditas, 2018, 155, 9.	0.5	35
31	The GSK3B gene confers risk for both major depressive disorder and schizophrenia in the Han Chinese population. Journal of Affective Disorders, 2015, 185, 149-155.	2.0	34
32	Glucose and Insulin-Related Traits, Type 2 Diabetes and Risk of Schizophrenia: A Mendelian Randomization Study. EBioMedicine, 2018, 34, 182-188.	2.7	34
33	Fine mapping the MHC region identified four independent variants modifying susceptibility to chronic hepatitis B in Han Chinese. Human Molecular Genetics, 2016, 25, 1225-1232.	1.4	33
34	Loci with genome-wide associations with schizophrenia in the Han Chinese population. British Journal of Psychiatry, 2015, 207, 490-494.	1.7	29
35	Mutational landscape of penile squamous cell carcinoma in a Chinese population. International Journal of Cancer, 2019, 145, 1280-1289.	2.3	28
36	Genome-wide association studies identify susceptibility loci for epithelial ovarian cancer in east Asian women. Gynecologic Oncology, 2019, 153, 343-355.	0.6	28

3

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37	X Chromosome Contribution to the Genetic Architecture of Primary Biliary Cholangitis. Gastroenterology, 2021, 160, 2483-2495.e26.	0.6	27
38	Structural Analysis of the SARS-CoV-2 Omicron Variant Proteins. Research, 2021, 2021, 9769586.	2.8	27
39	Genetic association between $\langle i \rangle$ NRG1 $\langle i \rangle$ and schizophrenia, major depressive disorder, bipolar disorder in Han Chinese population. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2016, 171, 468-478.	1.1	26
40	Identifying the Genotypes of Hepatitis B Virus (HBV) with DNA Origami Label. Small, 2018, 14, 1701718.	5.2	23
41	Investigation of Variants in UCP2 in Chinese Type 2 Diabetes and Diabetic Retinopathy. PLoS ONE, 2014, 9, e112670.	1.1	22
42	Association between SREBF2 gene polymorphisms and metabolic syndrome in clozapine-treated patients with schizophrenia. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2015, 56, 136-141.	2.5	19
43	The schizophrenia genetics knowledgebase: a comprehensive update of findings from candidate gene studies. Translational Psychiatry, 2019, 9, 205.	2.4	19
44	Prediction of causal genes and gene expression analysis of attention-deficit hyperactivity disorder in the different brain region, a comprehensive integrative analysis of ADHD. Behavioural Brain Research, 2019, 364, 183-192.	1.2	18
45	ITIH family genes confer risk to schizophrenia and major depressive disorder in the Han Chinese population. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2014, 51, 34-38.	2.5	17
46	The NVL gene confers risk for both major depressive disorder and schizophrenia in the Han Chinese population. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2015, 62, 7-13.	2.5	17
47	Genome-Wide Association Study of Obstructive Sleep Apnea and Objective Sleep-related Traits Identifies Novel Risk Loci in Han Chinese Individuals. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 1534-1545.	2.5	17
48	Endothelial Nitric Oxide Synthase (eNOS) T-786C, 4a4b, and G894T Polymorphisms and Male Infertility: Study for Idiopathic Asthenozoospermia and Meta-Analysis1. Biology of Reproduction, 2015, 92, 38.	1.2	16
49	Renal hypouricemia caused by novel compound heterozygous mutations in the SLC22A12 gene: a case report with literature review. BMC Medical Genetics, 2018, 19, 142.	2.1	16
50	Identification of serum microRNAs as diagnostic biomarkers for schizophrenia. Hereditas, 2019, 156, 23.	0.5	16
51	Performance comparison of four types of target enrichment baits for exome DNA sequencing. Hereditas, 2021, 158, 10.	0.5	16
52	Structural Comparison and Drug Screening of Spike Proteins of Ten SARS-CoV-2 Variants. Research, 2022, 2022, 9781758.	2.8	15
53	The Potential Effect of Aberrant Testosterone Levels on Common Diseases: A Mendelian Randomization Study. Genes, 2020, 11, 721.	1.0	14
54	Association study of 15q14 and 15q25 with high myopia in the Han Chinese population. BMC Genetics, 2014, 15, 51.	2.7	12

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55	Association between <i>SCAP</i> and <i>SREBF1</i> gene polymorphisms and metabolic syndrome in schizophrenia patients treated with atypical antipsychotics. World Journal of Biological Psychiatry, 2016, 17, 467-474.	1.3	12
56	Association of SCN10A Polymorphisms with the Recurrence of Atrial Fibrillation after Catheter Ablation in a Chinese Han Population. Scientific Reports, 2017, 7, 44003.	1.6	11
57	A genome-wide assessment of rare copy number variants in colorectal cancer. Oncotarget, 2015, 6, 26411-26423.	0.8	11
58	Identification of a novel susceptibility locus at 16q23.1 associated with childhood acute lymphoblastic leukemia in Han Chinese. Human Molecular Genetics, 2016, 25, ddw112.	1.4	10
59	A new risk locus in the ZEB2 gene for schizophrenia in the Han Chinese population. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2016, 66, 97-103.	2.5	10
60	Polymorphisms in NRGN are associated with schizophrenia, major depressive disorder and bipolar disorder in the Han Chinese population. Journal of Affective Disorders, 2016, 194, 180-187.	2.0	10
61	Genomeâ€wide twoâ€locus interaction analysis identifies multiple epistatic SNP pairs that confer risk of prostate cancer: A crossâ€population study. International Journal of Cancer, 2017, 140, 2075-2084.	2.3	10
62	Association between the variability of the <i> ABCA13 < /i > gene and the risk of major depressive disorder and schizophrenia in the Han Chinese population. World Journal of Biological Psychiatry, 2017, 18, 550-556.</i>	1.3	9
63	PPARG Polymorphisms Are Associated with Unexplained Mild Vision Loss in Patients with Type 2 Diabetes Mellitus. Journal of Ophthalmology, 2019, 2019, 1-7.	0.6	9
64	Elevated levels of IL â€18 associated with schizophrenia and first episode psychosis: A systematic review and metaâ€analysis. Microbial Biotechnology, 2020, 15, 896-905.	0.9	9
65	The polymorphism rs671 at ALDH2 associated with serum uric acid levels in Chinese Han males: A genome-wide association study. Gene, 2018, 651, 62-69.	1.0	8
66	ACTN3 is associated with children's physical fitness in Han Chinese. Molecular Genetics and Genomics, 2019, 294, 47-56.	1.0	8
67	The amino acid variants in HLA II molecules explain the major association with adult-onset Still's disease in the Han Chinese population. Journal of Autoimmunity, 2021, 116, 102562.	3.0	8
68	Colchicine prophylaxis is associated with fewer gout flares after COVID-19 vaccination. Annals of the Rheumatic Diseases, 2022, 81, 1189-1193.	0.5	8
69	Diacylglycerol kinase κ (<scp>DGKK</scp>) variants and hypospadias in <scp>H</scp> an <scp>C</scp> hinese: association and metaâ€analysis. BJU International, 2015, 116, 634-640.	1.3	7
70	SLC39A8 is a risk factor for schizophrenia in Uygur Chinese: a case-control study. BMC Psychiatry, 2019, 19, 293.	1.1	7
71	Four Loci Are Associated with Cardiorespiratory Fitness and Endurance Performance in Young Chinese Females. Scientific Reports, 2020, 10, 10117.	1.6	7
72	Fineâ€mapping of <i>ZDHHC2</i> identifies risk variants for schizophrenia in the Han Chinese population. Molecular Genetics & Denomic Medicine, 2020, 8, e1190.	0.6	7

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73	Psychiatric genetics in China: achievements and challenges. Molecular Psychiatry, 2016, 21, 4-9.	4.1	6
74	Association study of <i>NDST3</i> gene for schizophrenia, bipolar disorder, major depressive disorder in the Han Chinese population. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2018, 177, 3-9.	1.1	6
75	Identification of rare and common variants in BNIP3L: a schizophrenia susceptibility gene. Human Genomics, 2020, 14, 16.	1.4	6
76	Genome-wide analysis of DNA methylation identifies \$100A13 as an epigenetic biomarker in individuals with chronic (≥ 30 years) type 2 diabetes without diabetic retinopathy. Clinical Epigenetics, 2020, 12, 77.	1.8	6
77	Analysis of association between common variants of uncoupling proteins genes and diabetic retinopathy in a Chinese population. BMC Medical Genetics, 2020, 21, 25.	2.1	6
78	Cigarette smoking and schizophrenia: Mendelian randomisation study. British Journal of Psychiatry, 2021, 218, 98-103.	1.7	6
79	Identification of SHANK2 Pathogenic Variants in a Chinese Uygur Population with Schizophrenia. Journal of Molecular Neuroscience, 2021, 71, 1-8.	1.1	6
80	eRFSVM: a hybrid classifier to predict enhancers-integrating random forests with support vector machines. Hereditas, 2016, 153, 6.	0.5	5
81	Common variants in the SLC28A2 gene are associated with serum uric acid level and hyperuricemia and gout in Han Chinese. Hereditas, 2019, 156, 4.	0.5	5
82	Common variants in FAN1, located in 15q13.3, confer risk for schizophrenia and bipolar disorder in Han Chinese. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2020, 103, 109973.	2.5	5
83	STRsearch: a new pipeline for targeted profiling of short tandem repeats in massively parallel sequencing data. Hereditas, 2020, 157, 8.	0.5	5
84	Systematic comparative study of computational methods for HLA typing from nextâ€generation sequencing. Hla, 2021, 97, 481-492.	0.4	5
85	Integrative omics analysis reveals effective stratification and potential prognosis markers of pan-gastrointestinal cancers. IScience, 2021, 24, 102824.	1.9	5
86	The Relationship between Alcohol Consumption and Gout: A Mendelian Randomization Study. Genes, 2022, 13, 557.	1.0	5
87	Analysis of association between common variants in the <i>SLCO6A1 </i> gene with schizophrenia, bipolar disorder and major depressive disorder in the Han Chinese population. World Journal of Biological Psychiatry, 2016, 17, 140-146.	1.3	4
88	Association of fat mass and obesity-associated and retinitis pigmentosa guanosine triphosphatase (GTPase) regulator-interacting protein-1 like polymorphisms with body mass index in Chinese women. Endocrine Journal, 2018, 65, 783-791.	0.7	4
89	Amplicon targeted resequencing for <i>SLC2A9</i> and <i>SLC22A12</i> identified novel mutations in hypouricemia subjects. Molecular Genetics & Genomic Medicine, 2019, 7, e00722.	0.6	4
90	The association between rs12807809 polymorphism in neurogranin gene and risk of schizophrenia. Medicine (United States), 2019, 98, e18518.	0.4	4

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91	Mechanistic Modeling of Gene Regulation and Metabolism Identifies Potential Targets for Hepatocellular Carcinoma. Frontiers in Genetics, 2020, 11, 595242.	1.1	4
92	Rare and common variants analysis of the EMB gene in patients with schizophrenia. BMC Psychiatry, 2020, 20, 135.	1.1	4
93	Functional annotation of regulatory single nucleotide polymorphisms associated with schizophrenia. Schizophrenia Research, 2020, 218, 326-328.	1.1	4
94	Common variants in <i>QPCT</i> gene confer risk of schizophrenia in the Han Chinese population. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2016, 171, 237-242.	1.1	3
95	Role played by the SP4 gene in schizophrenia and major depressive disorder in the Han Chinese population. British Journal of Psychiatry, 2016, 208, 441-445.	1.7	3
96	Polymorphism of the PPARD Gene and Dynamic Balance Performance in Han Chinese Children. Hereditas, 2019, 156, 15.	0.5	3
97	VariFAST: a variant filter by automated scoring based on tagged-signatures. BMC Bioinformatics, 2019, 20, 713.	1.2	3
98	A Polynesian-specific copy number variant encompassing the MICA gene associates with gout. Human Molecular Genetics, 2022, 31, 3757-3768.	1.4	3
99	Common variants in SATB2 are associated with schizophrenia in Uygur Chinese population. Psychiatric Genetics, 2019, 29, 120-126.	0.6	2
100	Polymorphisms and rare variants identified by next-generation sequencing confer risk for lung cancer in han Chinese population. Pathology Research and Practice, 2020, 216, 152873.	1.0	2
101	A natural marmoset model of genetic generalized epilepsy. Molecular Brain, 2022, 15, 16.	1.3	2
102	SHEsisPCA: A GPU-Based Software to Correct for Population Stratification that Efficiently Accelerates the Process for Handling Genome-Wide Datasets. Journal of Genetics and Genomics, 2015, 42, 445-453.	1.7	1
103	The TNF-α-308G/A Polymorphism is Not Associated with Ocular <i>Chlamydia trachomatis</i> Infection in Han Chinese Children. Ophthalmic Genetics, 2016, 37, 245-247.	0.5	1
104	Noninvasive fetal trisomy detection by multiplexed semiconductor sequencing: a barcoding analysis strategy. Journal of Human Genetics, 2016, 61, 247-252.	1.1	1
105	Biological data processing based on bio-processor unit (BPU), a new concept for next generation computational biology. Science China Life Sciences, 2018, 61, 597-598.	2.3	1
106	A meta-analysis of genome-wide association studies using Japanese and Taiwanese has revealed novel loci associated with gout susceptibility. Human Cell, 2022, 35, 767.	1.2	1
107	Rare variations in the SHANK3 gene confers susceptibility to schizophrenia in Uygur Chinese population. Schizophrenia Research, 2021, 228, 597-599.	1.1	0