

Yan Zhang

List of Publications by Year in descending order

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22
papers

1,263
citations

933447

10
h-index

677142

22
g-index

23
all docs

23
docs citations

23
times ranked

2121
citing authors

#	ARTICLE	IF	CITATIONS
1	Genome-Wide Association Study in Asian Populations Identifies Variants in ETS1 and WDFY4 Associated with Systemic Lupus Erythematosus. <i>PLoS Genetics</i> , 2010, 6, e1000841.	3.5	378
2	Genome-wide association meta-analysis in Chinese and European individuals identifies ten new loci associated with systemic lupus erythematosus. <i>Nature Genetics</i> , 2016, 48, 940-946.	21.4	283
3	Meta-analysis Followed by Replication Identifies Loci in or near CDKN1B, TET3, CD80, DRAM1, and ARID5B as Associated with Systemic Lupus Erythematosus in Asians. <i>American Journal of Human Genetics</i> , 2013, 92, 41-51.	6.2	184
4	Identification of 38 novel loci for systemic lupus erythematosus and genetic heterogeneity between ancestral groups. <i>Nature Communications</i> , 2021, 12, 772.	12.8	128
5	ELF1 is associated with systemic lupus erythematosus in Asian populations. <i>Human Molecular Genetics</i> , 2011, 20, 601-607.	2.9	78
6	Meta-analysis of GWAS on two Chinese populations followed by replication identifies novel genetic variants on the X chromosome associated with systemic lupus erythematosus. <i>Human Molecular Genetics</i> , 2015, 24, 274-284.	2.9	35
7	Identification of <i>ST3AGL4</i> , <i>MFHAS1</i> , <i>CSNK2A2</i> and <i>CD226</i> as loci associated with systemic lupus erythematosus (SLE) and evaluation of SLE genetics in drug repositioning. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, 1078-1084.	0.9	34
8	Meta-analysis of GWAS on both Chinese and European populations identifies GPR173 as a novel X chromosome susceptibility gene for SLE. <i>Arthritis Research and Therapy</i> , 2018, 20, 92.	3.5	19
9	Epistatic Interaction between Genetic Variants in Susceptibility Gene <i>ETS1</i> Correlates with IL-17 Levels in SLE Patients. <i>Annals of Human Genetics</i> , 2013, 77, 344-350.	0.8	16
10	Genome-wide association study on Northern Chinese identifies <i>KLF2</i> and <i>DOT1L</i> and <i>STAB2</i> associated with systemic lupus erythematosus. <i>Rheumatology</i> , 2021, 60, 4407-4417.	1.9	16
11	Association of polymorphism in the <i>VEGFA</i> gene 3'UTR +936T/C with susceptibility to biliary atresia in a Southern Chinese Han population. <i>Journal of Clinical Laboratory Analysis</i> , 2018, 32, e22342.	2.1	12
12	Epistatic Association of CD14 and NOTCH2 Genetic Polymorphisms with Biliary Atresia in a Southern Chinese Population. <i>Molecular Therapy - Nucleic Acids</i> , 2018, 13, 590-595.	5.1	10
13	Identification of Regulatory Modules That Stratify Lupus Disease Mechanism through Integrating Multi-Omics Data. <i>Molecular Therapy - Nucleic Acids</i> , 2020, 19, 318-329.	5.1	10
14	<i>PDGFA</i> gene rs9690350 polymorphism increases biliary atresia risk in Chinese children. <i>Bioscience Reports</i> , 2020, 40, .	2.4	10
15	Association of IL18 genetic polymorphisms with increased risk of Biliary atresia susceptibility in Southern Chinese children. <i>Gene</i> , 2018, 677, 228-231.	2.2	9
16	Independent Replication on Genome-Wide Association Study Signals Identifies IRF3 as a Novel Locus for Systemic Lupus Erythematosus. <i>Frontiers in Genetics</i> , 2020, 11, 600.	2.3	9
17	Associations of <i>SLC6A20</i> genetic polymorphisms with Hirschsprung's disease in a Southern Chinese population. <i>Bioscience Reports</i> , 2019, 39, .	2.4	9
18	Meta-analysis of two Chinese populations identifies an autoimmune disease risk allele in 22q11.21 as associated with systemic lupus erythematosus. <i>Arthritis Research and Therapy</i> , 2015, 17, 67.	3.5	6

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19	Associations between common genetic variants in microRNAs and Hirschsprung disease susceptibility in Southern Chinese children. <i>Journal of Gene Medicine</i> , 2021, 23, e3301.	2.8	6
20	Down-regulation of STAT3 enhanced chemokine expression and neutrophil recruitment in biliary atresia. <i>Clinical Science</i> , 2021, 135, 865-884.	4.3	5
21	Association between DSCAM polymorphisms and non-syndromic Hirschsprung disease in Chinese population. <i>BMC Medical Genetics</i> , 2018, 19, 116.	2.1	4
22	Significant Association of rs2147555 Genetic Polymorphism in the EDNRB Gene with Hirschsprung Disease in Southern Chinese Children. <i>BioMed Research International</i> , 2020, 2020, 1-6.	1.9	1