

Xiao Liu

List of Publications by Year in descending order

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Version: 2024-02-01

99
papers

35,277
citations

57631

44
h-index

34900

98
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120
all docs

120
docs citations

120
times ranked

63945
citing authors

#	ARTICLE	IF	CITATIONS
1	Life History Recorded in the Vagino-cervical Microbiome Along with Multi-omes. Genomics, Proteomics and Bioinformatics, 2022, 20, 304-321.	3.0	18
2	Heterogeneous origin of IgE in atopic dermatitis and psoriasis revealed by B cell receptor repertoire analysis. Allergy: European Journal of Allergy and Clinical Immunology, 2022, 77, 559-568.	2.7	10
3	T Cell Repertoire Abnormality in Immunodeficiency Patients with DNA Repair and Methylation Defects. Journal of Clinical Immunology, 2022, 42, 375-393.	2.0	10
4	A genome-wide association study for gut metagenome in Chinese adults illuminates complex diseases. Cell Discovery, 2021, 7, 9.	3.1	49
5	Characterization of the human skin resistome and identification of two microbiota cutotypes. Microbiome, 2021, 9, 47.	4.9	42
6	Integrated genetic analyses revealed novel human longevity loci and reduced risks of multiple diseases in a cohort study of 15,651 Chinese individuals. Aging Cell, 2021, 20, e13323.	3.0	27
7	Diversity in immunogenomics: the value and the challenge. Nature Methods, 2021, 18, 588-591.	9.0	40
8	Sevoflurane inhibits ferroptosis: A new mechanism to explain its protective role against lipopolysaccharide-induced acute lung injury. Life Sciences, 2021, 275, 119391.	2.0	31
9	A transomic cohort as a reference point for promoting a healthy human gut microbiome. Medicine in Microecology, 2021, 8, 100039.	0.7	24
10	New genetic variants associated with major adverse cardiovascular events in patients with acute coronary syndromes and treated with clopidogrel and aspirin. Pharmacogenomics Journal, 2021, 21, 664-672.	0.9	5
11	A Positively Selected MAGEE2 LoF Allele Is Associated with Sexual Dimorphism in Human Brain Size and Shows Similar Phenotypes in Magee2 Null Mice. Molecular Biology and Evolution, 2021, 38, 5655-5663.	3.5	1
12	Distinct human Langerhans cell subsets orchestrate reciprocal functions and require different developmental regulation. Immunity, 2021, 54, 2305-2320.e11.	6.6	38
13	Characteristics of serum metabolites in sporadic amyotrophic lateral sclerosis patients based on gas chromatography-mass spectrometry. Scientific Reports, 2021, 11, 20786.	1.6	10
14	Dissecting the Landscape of Activated CMV-Stimulated CD4+ T Cells in Humans by Linking Single-Cell RNA-Seq With T-Cell Receptor Sequencing. Frontiers in Immunology, 2021, 12, 779961.	2.2	12
15	PIRD: Pan Immune Repertoire Database. Bioinformatics, 2020, 36, 897-903.	1.8	79
16	Tâ€cell receptor repertoire data provides new evidence for hygiene hypothesis of allergic diseases. Allergy: European Journal of Allergy and Clinical Immunology, 2020, 75, 681-683.	2.7	9
17	Developing an Unbiased Multiplex PCR System to Enrich the TRB Repertoire Toward Accurate Detection in Leukemia. Frontiers in Immunology, 2020, 11, 1631.	2.2	4
18	Comprehensive TCR repertoire analysis of CD4+ T-cell subsets in rheumatoid arthritis. Journal of Autoimmunity, 2020, 109, 102432.	3.0	29

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19	Single-cell RNA-seq unveils critical regulators of human FOXP3+ regulatory T cell stability. <i>Science Bulletin</i> , 2020, 65, 1114-1124.	4.3	10
20	Clinical implications of systematic phenotyping and exome sequencing in patients with primary antibody deficiency. <i>Genetics in Medicine</i> , 2019, 21, 243-251.	1.1	86
21	Alterations in the human gut microbiome associated with <i>Helicobacter pylori</i> infection. <i>FEBS Open Bio</i> , 2019, 9, 1552-1560.	1.0	30
22	Selection of potential cytokeratin-18 monoclonal antibodies following IGH repertoire evaluation in mice. <i>Journal of Immunological Methods</i> , 2019, 474, 112647.	0.6	2
23	Germline-Encoded TCR-MHC Contacts Promote TCR V Gene Bias in Umbilical Cord Blood T Cell Repertoire. <i>Frontiers in Immunology</i> , 2019, 10, 2064.	2.2	15
24	Identification of Variable and Joining Germline Genes and Alleles for Rhesus Macaque from B Cell Receptor Repertoires. <i>Journal of Immunology</i> , 2019, 202, 1612-1622.	0.4	9
25	T cell receptor \hat{I}^2 repertoires as novel diagnostic markers for systemic lupus erythematosus and rheumatoid arthritis. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, 1070-1078.	0.5	99
26	Single-cell transcriptomic landscape of nucleated cells in umbilical cord blood. <i>GigaScience</i> , 2019, 8, .	3.3	24
27	Myopia disease mouse models: a missense point mutation (S673G) and a protein-truncating mutation of the Zfp644 mimic human disease phenotype. <i>Cell and Bioscience</i> , 2019, 9, 21.	2.1	5
28	The bZIP transcription factor Afap1 mediates the oxidative stress response and aflatoxin biosynthesis in <i>Aspergillus flavus</i> . <i>Revista Argentina De Microbiologia</i> , 2019, 51, 292-301.	0.4	20
29	Efficient and unique cobarcoding of second-generation sequencing reads from long DNA molecules enabling cost-effective and accurate sequencing, haplotyping, and de novo assembly. <i>Genome Research</i> , 2019, 29, 798-808.	2.4	176
30	Compound Heterozygous Mutations of IL2-Inducible T cell Kinase in a Swedish Patient: the Importance of Early Genetic Diagnosis. <i>Journal of Clinical Immunology</i> , 2019, 39, 131-134.	2.0	9
31	The landscape and diagnostic potential of T and B cell repertoire in Immunoglobulin A Nephropathy. <i>Journal of Autoimmunity</i> , 2019, 97, 100-107.	3.0	25
32	History, applications, and challenges of immune repertoire research. <i>Cell Biology and Toxicology</i> , 2018, 34, 441-457.	2.4	81
33	Novel genetic loci associated HLA-B*08:01 positive myasthenia gravis. <i>Journal of Autoimmunity</i> , 2018, 88, 43-49.	3.0	20
34	A Comprehensive Analysis of the T and B Lymphocytes Repertoire Shaped by HIV Vaccines. <i>Frontiers in Immunology</i> , 2018, 9, 2194.	2.2	23
35	Sex Differences in Genetic Associations With Longevity. <i>JAMA Network Open</i> , 2018, 1, e181670.	2.8	60
36	A regulatory mutant on <i>TRIM26</i> conferring the risk of nasopharyngeal carcinoma by inducing low immune response. <i>Cancer Medicine</i> , 2018, 7, 3848-3861.	1.3	14

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37	Effect of Cinnamaldehyde on Morphological Alterations of <i>Aspergillus ochraceus</i> and Expression of Key Genes Involved in Ochratoxin A Biosynthesis. <i>Toxins</i> , 2018, 10, 340.	1.5	38
38	The correlation of copy number variations with longevity in a genome-wide association study of Han Chinese. <i>Aging</i> , 2018, 10, 1206-1222.	1.4	25
39	Aflatoxin B 1 inhibition in <i>Aspergillus flavus</i> by <i>Aspergillus niger</i> through down-regulating expression of major biosynthetic genes and AFB 1 degradation by atoxigenic <i>A. flavus</i> . <i>International Journal of Food Microbiology</i> , 2017, 256, 1-10.	2.1	54
40	Characterization of the B Cell Receptor Repertoire in the Intestinal Mucosa and of Tumor-Infiltrating Lymphocytes in Colorectal Adenoma and Carcinoma. <i>Journal of Immunology</i> , 2017, 198, 3719-3728.	0.4	39
41	The Different T-cell Receptor Repertoires in Breast Cancer Tumors, Draining Lymph Nodes, and Adjacent Tissues. <i>Cancer Immunology Research</i> , 2017, 5, 148-156.	1.6	87
42	An Unexpectedly Complex Architecture for Skin Pigmentation in Africans. <i>Cell</i> , 2017, 171, 1340-1353.e14.	13.5	134
43	A comprehensive profiling of T- and B-lymphocyte receptor repertoires from a Chinese-origin rhesus macaque by high-throughput sequencing. <i>PLoS ONE</i> , 2017, 12, e0182733.	1.1	18
44	Minimal Residual Disease Detection and Evolved IGH Clones Analysis in Acute B Lymphoblastic Leukemia Using IGH Deep Sequencing. <i>Frontiers in Immunology</i> , 2016, 7, 403.	2.2	37
45	IMPre: An Accurate and Efficient Software for Prediction of T- and B-Cell Receptor Germline Genes and Alleles from Rearranged Repertoire Data. <i>Frontiers in Immunology</i> , 2016, 7, 457.	2.2	47
46	Systematic Comparative Evaluation of Methods for Investigating the TCR β Repertoire. <i>PLoS ONE</i> , 2016, 11, e0152464.	1.1	58
47	Comparative Analysis of Immune Repertoires between Bactrian Camel's Conventional and Heavy-Chain Antibodies. <i>PLoS ONE</i> , 2016, 11, e0161801.	1.1	49
48	Species specific exome probes reveal new insights in positively selected genes in nonhuman primates. <i>Scientific Reports</i> , 2016, 6, 33876.	1.6	3
49	Novel Y-chromosomal microdeletions associated with non-obstructive azoospermia uncovered by high throughput sequencing of sequence-tagged sites (STSs). <i>Scientific Reports</i> , 2016, 6, 21831.	1.6	11
50	Novel loci and pathways significantly associated with longevity. <i>Scientific Reports</i> , 2016, 6, 21243.	1.6	145
51	Variation in fungal microbiome (mycobiome) and aflatoxins during simulated storage of in-shell peanuts and peanut kernels. <i>Scientific Reports</i> , 2016, 6, 25930.	1.6	33
52	The complete mitochondrial genome of the white-tailed tropicbird, <i>Phaethon lepturus</i> . <i>Mitochondrial DNA Part A: DNA Mapping, Sequencing, and Analysis</i> , 2016, 27, 4259-4260.	0.7	1
53	Deep sequencing of the MHC region in the Chinese population contributes to studies of complex disease. <i>Nature Genetics</i> , 2016, 48, 740-746.	9.4	188
54	Comprehensive Characterization of Oncogenic Drivers in Asian Lung Adenocarcinoma. <i>Journal of Thoracic Oncology</i> , 2016, 11, 2129-2140.	0.5	70

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55	A Cytocompatible Robust Hybrid Conducting Polymer Hydrogel for Use in a Magnesium Battery. <i>Advanced Materials</i> , 2016, 28, 9349-9355.	11.1	67
56	Genome-wide characteristics of de novo mutations in autism. <i>Npj Genomic Medicine</i> , 2016, 1, 160271-1602710.	1.7	200
57	Genomic and oncogenic preference of HBV integration in hepatocellular carcinoma. <i>Nature Communications</i> , 2016, 7, 12992.	5.8	228
58	Case report of a Liâ€“Fraumeni syndrome-like phenotype with a de novo mutation in CHEK2. <i>Medicine (United States)</i> , 2016, 95, e4251.	0.4	8
59	Inhibitory Effect of Cinnamaldehyde, Citral, and Eugenol on Aflatoxin Biosynthetic Gene Expression and Aflatoxin B₁ Biosynthesis in <i>Aspergillus flavus</i>. <i>Journal of Food Science</i> , 2015, 80, M2917-24.	1.5	79
60	Variation in fungal microbiome (mycobiome) and aflatoxin in stored in-shell peanuts at four different areas of China. <i>Frontiers in Microbiology</i> , 2015, 6, 1055.	1.5	37
61	Lung cancer in never-smoker Asian females is driven by oncogenic mutations, most often involving <i>EGFR</i>. <i>Oncotarget</i> , 2015, 6, 5465-5474.	0.8	116
62	De novo assembly of a haplotype-resolved human genome. <i>Nature Biotechnology</i> , 2015, 33, 617-622.	9.4	73
63	Genome-wide profiling of HPV integration in cervical cancer identifies clustered genomic hot spots and a potential microhomology-mediated integration mechanism. <i>Nature Genetics</i> , 2015, 47, 158-163.	9.4	393
64	Excess of Rare Variants in Genes that are Key Epigenetic Regulators of Spermatogenesis in the Patients with Non-Obstructive Azoospermia. <i>Scientific Reports</i> , 2015, 5, 8785.	1.6	39
65	Molecular analysis of gastric cancer identifies subtypes associated with distinct clinical outcomes. <i>Nature Medicine</i> , 2015, 21, 449-456.	15.2	1,592
66	A global reference for human genetic variation. <i>Nature</i> , 2015, 526, 68-74.	13.7	13,998
67	IMonitor: A Robust Pipeline for TCR and BCR Repertoire Analysis. <i>Genetics</i> , 2015, 201, 459-472.	1.2	119
68	Detection and Analysis of Human Papillomavirus (HPV) DNA in Breast Cancer Patients by an Effective Method of HPV Capture. <i>PLoS ONE</i> , 2014, 9, e90343.	1.1	15
69	Inhibitory Effect of Essential Oils on <i>Aspergillus ochraceus</i> Growth and Ochratoxin A Production. <i>PLoS ONE</i> , 2014, 9, e108285.	1.1	110
70	A Novel Excitation Assistance Switched Reluctance Wind Power Generator. <i>IEEE Transactions on Magnetics</i> , 2014, 50, 1-4.	1.2	21
71	Genomic landscape and genetic heterogeneity in gastric adenocarcinoma revealed by whole-genome sequencing. <i>Nature Communications</i> , 2014, 5, 5477.	5.8	166
72	Exome capture from saliva produces high quality genomic and metagenomic data. <i>BMC Genomics</i> , 2014, 15, 262.	1.2	34

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73	Decoding complex patterns of genomic rearrangement in hepatocellular carcinoma. <i>Genomics</i> , 2014, 103, 189-203.	1.3	49
74	Rapid detection of structural variation in a human genome using nanochannel-based genome mapping technology. <i>GigaScience</i> , 2014, 3, 34.	3.3	153
75	Association Between Variants of PRDM1 and NDP52 and Crohn's Disease, Based on Exome Sequencing and Functional Studies. <i>Gastroenterology</i> , 2013, 145, 339-347.	0.6	149
76	Deep sequencing identifies regulated small RNAs in <i>Dugesia japonica</i> . <i>Molecular Biology Reports</i> , 2013, 40, 4075-4081.	1.0	7
77	HIVID: An efficient method to detect HBV integration using low coverage sequencing. <i>Genomics</i> , 2013, 102, 338-344.	1.3	94
78	The Wilms Tumor Gene, <i>Wt1</i> , Is Critical for Mouse Spermatogenesis via Regulation of Sertoli Cell Polarity and Is Associated with Non-Obstructive Azoospermia in Humans. <i>PLoS Genetics</i> , 2013, 9, e1003645.	1.5	109
79	A population model for genotyping indels from next-generation sequence data. <i>Nucleic Acids Research</i> , 2013, 41, e46-e46.	6.5	12
80	Whole-genome sequencing identifies recurrent mutations in hepatocellular carcinoma. <i>Genome Research</i> , 2013, 23, 1422-1433.	2.4	457
81	An Integrated Tool to Study MHC Region: Accurate SNV Detection and HLA Genes Typing in Human MHC Region Using Targeted High-Throughput Sequencing. <i>PLoS ONE</i> , 2013, 8, e69388.	1.1	63
82	Genetic Aberrations in Imatinib-Resistant Dermatofibrosarcoma Protuberans Revealed by Whole Genome Sequencing. <i>PLoS ONE</i> , 2013, 8, e69752.	1.1	25
83	Abstract LB-229: Whole genome sequencing reveals genetic landscape of hepatocellular carcinoma.. , 2013, , .		0
84	Abstract LB-231: Decoding complex patterns of structural variations in hepatocellular carcinoma.. , 2013, , .		0
85	Genome-wide survey of recurrent HBV integration in hepatocellular carcinoma. <i>Nature Genetics</i> , 2012, 44, 765-769.	9.4	785
86	Single-Cell Exome Sequencing Reveals Single-Nucleotide Mutation Characteristics of a Kidney Tumor. <i>Cell</i> , 2012, 148, 886-895.	13.5	622
87	Single-cell sequencing analysis characterizes common and cell-lineage-specific mutations in a muscle-invasive bladder cancer. <i>GigaScience</i> , 2012, 1, 12.	3.3	99
88	An integrated map of genetic variation from 1,092 human genomes. <i>Nature</i> , 2012, 491, 56-65.	13.7	7,199
89	The oyster genome reveals stress adaptation and complexity of shell formation. <i>Nature</i> , 2012, 490, 49-54.	13.7	1,966
90	Frequent mutations of genes encoding ubiquitin-mediated proteolysis pathway components in clear cell renal cell carcinoma. <i>Nature Genetics</i> , 2012, 44, 17-19.	9.4	295

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91	An Abundance of Rare Functional Variants in 202 Drug Target Genes Sequenced in 14,002 People. <i>Science</i> , 2012, 337, 100-104.	6.0	626
92	Comprehensive comparison of three commercial human whole-exome capture platforms. <i>Genome Biology</i> , 2011, 12, R95.	13.9	145
93	BIPES, a cost-effective high-throughput method for assessing microbial diversity. <i>ISME Journal</i> , 2011, 5, 741-749.	4.4	160
94	Sequencing of 50 Human Exomes Reveals Adaptation to High Altitude. <i>Science</i> , 2010, 329, 75-78.	6.0	1,339
95	The sequence and de novo assembly of the giant panda genome. <i>Nature</i> , 2010, 463, 311-317.	13.7	1,058
96	Resequencing of 200 human exomes identifies an excess of low-frequency non-synonymous coding variants. <i>Nature Genetics</i> , 2010, 42, 969-972.	9.4	297
97	Archaeology Augments Tibet's Genetic History—Response. <i>Science</i> , 2010, 329, 1467-1468.	6.0	3
98	The effects of CYP1A1 gene polymorphism and p16 gene methylation on the risk of lung cancer in a Chinese population. <i>Chinese-German Journal of Clinical Oncology</i> , 2007, 6, 350-356.	0.1	2
99	M-GWAS for the Gut Microbiome in Chinese Adults Illuminates on Complex Diseases. <i>SSRN Electronic Journal</i> , 0, , .	0.4	4