

Heng Xu

List of Publications by Year in descending order

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Version: 2024-02-01

107
papers

5,296
citations

159585

30
h-index

91884

69
g-index

132
all docs

132
docs citations

132
times ranked

9415
citing authors

#	ARTICLE	IF	CITATIONS
1	Colocalization of Inflammatory Response with B7-H1 Expression in Human Melanocytic Lesions Supports an Adaptive Resistance Mechanism of Immune Escape. <i>Science Translational Medicine</i> , 2012, 4, 127ra37.	12.4	1,837
2	Inherited GATA3 variants are associated with Ph-like childhood acute lymphoblastic leukemia and risk of relapse. <i>Nature Genetics</i> , 2013, 45, 1494-1498.	21.4	264
3	Novel Susceptibility Variants at 10p12.31-12.2 for Childhood Acute Lymphoblastic Leukemia in Ethnically Diverse Populations. <i>Journal of the National Cancer Institute</i> , 2013, 105, 733-742.	6.3	208
4	<i>ARID5B</i> Genetic Polymorphisms Contribute to Racial Disparities in the Incidence and Treatment Outcome of Childhood Acute Lymphoblastic Leukemia. <i>Journal of Clinical Oncology</i> , 2012, 30, 751-757.	1.6	165
5	Germline genetic variation in <i>ETV6</i> and risk of childhood acute lymphoblastic leukaemia: a systematic genetic study. <i>Lancet Oncology</i> , 2015, 16, 1659-1666.	10.7	161
6	<i>TP53</i> Germline Variations Influence the Predisposition and Prognosis of B-Cell Acute Lymphoblastic Leukemia in Children. <i>Journal of Clinical Oncology</i> , 2018, 36, 591-599.	1.6	121
7	Personalized neoantigen pulsed dendritic cell vaccine for advanced lung cancer. <i>Signal Transduction and Targeted Therapy</i> , 2021, 6, 26.	17.1	112
8	A genome-wide association study of susceptibility to acute lymphoblastic leukemia in adolescents and young adults. <i>Blood</i> , 2015, 125, 680-686.	1.4	110
9	Common variants in <i>ACYP2</i> influence susceptibility to cisplatin-induced hearing loss. <i>Nature Genetics</i> , 2015, 47, 263-266.	21.4	109
10	Prognostic significance of frequent <i>CLDN18-ARHGAP26/6</i> fusion in gastric signet-ring cell cancer. <i>Nature Communications</i> , 2018, 9, 2447.	12.8	100
11	Thrombin induces <i>ACSL4</i> -dependent ferroptosis during cerebral ischemia/reperfusion. <i>Signal Transduction and Targeted Therapy</i> , 2022, 7, 59.	17.1	88
12	CRISPR/Cas9 facilitates investigation of neural circuit disease using human iPSCs: mechanism of epilepsy caused by an <i>SCN1A</i> loss-of-function mutation. <i>Translational Psychiatry</i> , 2016, 6, e703-e703.	4.8	82
13	Induction and Amelioration of Methotrexate-Induced Gastrointestinal Toxicity are Related to Immune Response and Gut Microbiota. <i>EBioMedicine</i> , 2018, 33, 122-133.	6.1	80
14	Characterizing dedifferentiation of thyroid cancer by integrated analysis. <i>Science Advances</i> , 2021, 7, .	10.3	76
15	Inherited coding variants at the <i>CDKN2A</i> locus influence susceptibility to acute lymphoblastic leukaemia in children. <i>Nature Communications</i> , 2015, 6, 7553.	12.8	72
16	MicroRNA-214 promotes hepatic stellate cell activation and liver fibrosis by suppressing <i>Sufu</i> expression. <i>Cell Death and Disease</i> , 2018, 9, 718.	6.3	72
17	Metabolism-induced tumor activator 1 (MITA1), an Energy Stress-Inducible Long Noncoding RNA, Promotes Hepatocellular Carcinoma Metastasis. <i>Hepatology</i> , 2019, 70, 215-230.	7.3	65
18	LncRNA linc00312 suppresses radiotherapy resistance by targeting DNA-PKcs and impairing DNA damage repair in nasopharyngeal carcinoma. <i>Cell Death and Disease</i> , 2021, 12, 69.	6.3	56

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19	DP7-C-modified liposomes enhance immune responses and the antitumor effect of a neoantigen-based mRNA vaccine. <i>Journal of Controlled Release</i> , 2020, 328, 210-221.	9.9	54
20	The genetic landscape of benign thyroid nodules revealed by whole exome and transcriptome sequencing. <i>Nature Communications</i> , 2017, 8, 15533.	12.8	53
21	Efficacy evaluation of clonazepam for symptom remission in burning mouth syndrome: a meta-analysis. <i>Oral Diseases</i> , 2016, 22, 503-511.	3.0	52
22	Genomic evolution and diverse models of systemic metastases in colorectal cancer. <i>Gut</i> , 2022, 71, 322-332.	12.1	51
23	Structural and functional insights into the regulation of the lysis-lysogeny decision in viral communities. <i>Nature Microbiology</i> , 2018, 3, 1285-1294.	13.3	49
24	Prognostic Factors for Checkpoint Inhibitor Based Immunotherapy: An Update With New Evidences. <i>Frontiers in Pharmacology</i> , 2018, 9, 1050.	3.5	48
25	Characteristics of genomic alterations of lung adenocarcinoma in young never-smokers. <i>International Journal of Cancer</i> , 2018, 143, 1696-1705.	5.1	45
26	Novel susceptibility variants at the ERG locus for childhood acute lymphoblastic leukemia in Hispanics. <i>Blood</i> , 2019, 133, 724-729.	1.4	44
27	Personalized neoantigen-pulsed dendritic cell vaccines show superior immunogenicity to neoantigen-adjuvant vaccines in mouse tumor models. <i>Cancer Immunology, Immunotherapy</i> , 2020, 69, 135-145.	4.2	42
28	Impact of <i>NUDT15</i> polymorphisms on thiopurines-induced myelotoxicity and thiopurines tolerance dose. <i>Oncotarget</i> , 2017, 8, 13575-13585.	1.8	40
29	Topical calcineurin inhibitors in the treatment of oral lichen planus: a systematic review and meta-analysis. <i>British Journal of Dermatology</i> , 2019, 181, 1166-1176.	1.5	38
30	Length of the ORF, position of the first AUG and the Kozak motif are important factors in potential dual-coding transcripts. <i>Cell Research</i> , 2010, 20, 445-457.	12.0	35
31	Novel Recurrent Altered Genes in Chinese Patients With Anaplastic Thyroid Cancer. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021, 106, e988-e998.	3.6	33
32	<i>Prevotella copri</i> is associated with carboplatin-induced gut toxicity. <i>Cell Death and Disease</i> , 2019, 10, 714.	6.3	32
33	Genome-Wide Association Study of Susceptibility Loci for T-Cell Acute Lymphoblastic Leukemia in Children. <i>Journal of the National Cancer Institute</i> , 2019, 111, 1350-1357.	6.3	32
34	Regulatory network of <i>GATA3</i> in pediatric acute lymphoblastic leukemia. <i>Oncotarget</i> , 2017, 8, 36040-36053.	1.8	30
35	Assessment of Glomerular Filtration Rate in Renal Transplant Patients Using Serum Cystatin C. <i>Transplantation Proceedings</i> , 2006, 38, 2006-2008.	0.6	29
36	Age-Associated Proteomic Signatures and Potential Clinically Actionable Targets of Colorectal Cancer. <i>Molecular and Cellular Proteomics</i> , 2021, 20, 100115.	3.8	29

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37	ARID5B Regulates Leukemia Sensitivity to Antimetabolites in Children with Acute Lymphoblastic Leukemia Via Effects on Cell Cycle Progression. <i>Blood</i> , 2014, 124, 791-791.	1.4	28
38	Association Between PIP4K2A Polymorphisms and Acute Lymphoblastic Leukemia Susceptibility. <i>Medicine (United States)</i> , 2016, 95, e3542.	1.0	27
39	Multi-omics approaches identify SF3B3 and SIRT3 as candidate autophagic regulators and druggable targets in invasive breast carcinoma. <i>Acta Pharmaceutica Sinica B</i> , 2021, 11, 1227-1245.	12.0	26
40	Subtype-specific inherited predisposition to pemphigus in the Chinese population. <i>British Journal of Dermatology</i> , 2019, 180, 828-835.	1.5	25
41	ARID5B Influences Antimetabolite Drug Sensitivity and Prognosis of Acute Lymphoblastic Leukemia. <i>Clinical Cancer Research</i> , 2020, 26, 256-264.	7.0	25
42	Combination of common and novel rare NUDT15 variants improves predictive sensitivity of thiopurine-induced leukopenia in children with acute lymphoblastic leukemia. <i>Haematologica</i> , 2018, 103, e293-e295.	3.5	24
43	Genome-Wide Association Studies of HIV-1 Host Control in Ethnically Diverse Chinese Populations. <i>Scientific Reports</i> , 2015, 5, 10879.	3.3	23
44	The Significance of the CLDN18-ARHGAP Fusion Gene in Gastric Cancer: A Systematic Review and Meta-Analysis. <i>Frontiers in Oncology</i> , 2020, 10, 1214.	2.8	23
45	NUDT15 Polymorphism Confer Increased Susceptibility to Thiopurine-Induced Leukopenia in Patients With Autoimmune Hepatitis and Related Cirrhosis. <i>Frontiers in Pharmacology</i> , 2019, 10, 346.	3.5	22
46	GPR39 Overexpression in OSCC Promotes YAP-Sustained Malignant Progression. <i>Journal of Dental Research</i> , 2020, 99, 949-958.	5.2	22
47	The Role of ARID5B in Acute Lymphoblastic Leukemia and Beyond. <i>Frontiers in Genetics</i> , 2020, 11, 598.	2.3	21
48	Alpha defensin-1 attenuates surgically induced osteoarthritis in association with promoting M1 to M2 macrophage polarization. <i>Osteoarthritis and Cartilage</i> , 2021, 29, 1048-1059.	1.3	21
49	A New System Identification Approach to Identify Genetic Variants in Sequencing Studies for a Binary Phenotype. <i>Human Heredity</i> , 2014, 78, 104-116.	0.8	19
50	Presence and diagnostic value of circulating tsncRNA for ovarian tumor. <i>Molecular Cancer</i> , 2018, 17, 163.	19.2	19
51	Prognostic impact of visit-to-visit glycemic variability on the risks of major adverse cardiovascular outcomes and hypoglycemia in patients with different glycemic control and type 2 diabetes. <i>Endocrine</i> , 2019, 64, 536-543.	2.3	19
52	Prognostic and predictive role of DNA mismatch repair status in stage II colorectal cancer: A systematic review and meta-analysis. <i>Clinical Genetics</i> , 2020, 97, 25-38.	2.0	19
53	RGD-modified oncolytic adenovirus exhibited potent cytotoxic effect on CAR-negative bladder cancer-initiating cells. <i>Cell Death and Disease</i> , 2015, 6, e1760-e1760.	6.3	18
54	The novel complex combination of alum, CpG ODN and HH2 as adjuvant in cancer vaccine effectively suppresses tumor growth <i>in vivo</i> . <i>Oncotarget</i> , 2017, 8, 45951-45964.	1.8	18

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55	Deep learning-based AI model for signet-ring cell carcinoma diagnosis and chemotherapy response prediction in gastric cancer. <i>Medical Physics</i> , 2022, 49, 1535-1546.	3.0	17
56	Urinary complement proteins and risk of end-stage renal disease: quantitative urinary proteomics in patients with type 2 diabetes and biopsy-proven diabetic nephropathy. <i>Journal of Endocrinological Investigation</i> , 2021, 44, 2709-2723.	3.3	16
57	PHLDB2 Mediates Cetuximab Resistance via Interacting With EGFR in Latent Metastasis of Colorectal Cancer. <i>Cellular and Molecular Gastroenterology and Hepatology</i> , 2022, 13, 1223-1242.	4.5	16
58	Screening of Kozak-motif-located SNPs and analysis of their association with human diseases. <i>Biochemical and Biophysical Research Communications</i> , 2010, 392, 89-94.	2.1	15
59	Long-Term Correction of Copper Metabolism in Wilson's Disease Mice with AAV8 Vector Delivering Truncated ATP7B. <i>Human Gene Therapy</i> , 2019, 30, 1494-1504.	2.7	15
60	<i>O</i> R _{AOV1-B} Promotes OSCC Metastasis via the NF- κ B-TNF α Loop. <i>Journal of Dental Research</i> , 2021, 100, 002203452199633.	5.2	15
61	Uncovering the Rare Variants of DLC1 Isoform 1 and Their Functional Effects in a Chinese Sporadic Congenital Heart Disease Cohort. <i>PLoS ONE</i> , 2014, 9, e90215.	2.5	14
62	LINC-PINT impedes DNA repair and enhances radiotherapeutic response by targeting DNA-PKcs in nasopharyngeal cancer. <i>Cell Death and Disease</i> , 2021, 12, 454.	6.3	14
63	The association of hematuria on kidney clinicopathologic features and renal outcome in patients with diabetic nephropathy: a biopsy-based study. <i>Journal of Endocrinological Investigation</i> , 2020, 43, 1213-1220.	3.3	13
64	Regulatory Network and Prognostic Effect Investigation of PIP4K2A in Leukemia and Solid Cancers. <i>Frontiers in Genetics</i> , 2019, 9, 721.	2.3	12
65	A Functional 5'-UTR Polymorphism of MYC Contributes to Nasopharyngeal Carcinoma Susceptibility and Chemoradiotherapy Induced Toxicities. <i>Journal of Cancer</i> , 2019, 10, 147-155.	2.5	12
66	ARL15 overexpression attenuates high glucose-induced impairment of insulin signaling and oxidative stress in human umbilical vein endothelial cells. <i>Life Sciences</i> , 2019, 220, 127-135.	4.3	12
67	Precise control of maxillary multidirectional movement in Le Fort I osteotomy using a surgical guiding device. <i>British Journal of Oral and Maxillofacial Surgery</i> , 2018, 56, 797-804.	0.8	11
68	Long Non-coding RNA Expression Profiling in Biopsy to Identify Renal Allograft at Risk of Chronic Damage and Future Graft Loss. <i>Applied Biochemistry and Biotechnology</i> , 2020, 190, 660-673.	2.9	11
69	Structural and Functional Insights into an Archaeal Lipid Synthase. <i>Cell Reports</i> , 2020, 33, 108294.	6.4	11
70	Intensive Glucose Control Reduces the Risk Effect of TRIB3, SMARCD3, and ATF6 Genetic Variation on Diabetic Vascular Complications. <i>Frontiers in Pharmacology</i> , 2018, 9, 1422.	3.5	10
71	Association of the independent polymorphisms in CDKN2A with susceptibility of acute lymphoblastic leukemia. <i>Bioscience Reports</i> , 2018, 38, .	2.4	10
72	Proteomic Maps of Human Gastrointestinal Stromal Tumor Subgroups*. <i>Molecular and Cellular Proteomics</i> , 2019, 18, 923a-935.	3.8	10

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73	Bioconcentration Factors and Potential Human Health Risks of Heavy Metals in Cultivated <i>Lentinus edodes</i> in Chengdu, People's Republic of China. <i>Journal of Food Protection</i> , 2015, 78, 390-395.	1.7	9
74	Mitomycin C reduces haze formation in rabbits after excimer laser photorefractive keratectomy. <i>Journal of Refractive Surgery</i> , 2001, 17, 342-9.	2.3	9
75	Cause-specific risk of major adverse cardiovascular outcomes and hypoglycemic in patients with type 2 diabetes: a multicenter prospective cohort study. <i>Endocrine</i> , 2019, 63, 44-51.	2.3	8
76	Prognostic effect of high-flux hemodialysis in patients with chronic kidney disease. <i>Brazilian Journal of Medical and Biological Research</i> , 2016, 49, e4708.	1.5	7
77	Novel Gene and Network Associations Found for Acute Lymphoblastic Leukemia Using Case-€“Control and Family-Based Studies in Multiethnic Populations. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2017, 26, 1531-1539.	2.5	7
78	Association of recurrent APOBEC3B alterations with the prognosis of gastric-type cervical adenocarcinoma. <i>Gynecologic Oncology</i> , 2022, 165, 105-113.	1.4	7
79	ANI analysis of poxvirus genomes reveals its potential application to viral species rank demarcation. <i>Virus Evolution</i> , 2022, 8, .	4.9	7
80	Response. <i>Journal of the National Cancer Institute</i> , 2013, 105, 1512-1513.	6.3	6
81	Rare gene variants in a patient with azathioprine-induced lethal myelosuppression. <i>Annals of Hematology</i> , 2017, 96, 2131-2133.	1.8	6
82	Triterpenoids with antioxidant activities from <i>Myricaria squamosa</i> . <i>Journal of Asian Natural Products Research</i> , 2018, 20, 292-298.	1.4	6
83	Polytropic Influence of TRIB3 rs2295490 Genetic Polymorphism on Response to Antihypertensive Agents in Patients With Essential Hypertension. <i>Frontiers in Pharmacology</i> , 2019, 10, 236.	3.5	5
84	Novel GLI3 Mutations in Chinese Patients with Non-syndromic Post-axial Polydactyly. <i>Current Molecular Medicine</i> , 2019, 19, 228-235.	1.3	5
85	CSF glutamate level decreases in heavy smokers and negatively correlates with BDI scores. <i>Psychiatry Research</i> , 2018, 270, 627-630.	3.3	4
86	The influence of telmisartan on metformin pharmacokinetics and pharmacodynamics. <i>Journal of Pharmacological Sciences</i> , 2019, 139, 37-41.	2.5	4
87	Improvement of Neoantigen Identification Through Convolution Neural Network. <i>Frontiers in Immunology</i> , 2021, 12, 682103.	4.8	4
88	Longitudinal Genomic Evolution of Conventional Papillary Thyroid Cancer With Brain Metastasis. <i>Frontiers in Oncology</i> , 2021, 11, 620924.	2.8	4
89	Characteristics of an established retinoblastoma cell line HXO-Rb44. <i>Yan Ke Xue Bao = Eye Science</i> , 1995, 11, 16-21.	0.1	4
90	Evidence of genetic factors involved in oral lichen planus pathogenesis. <i>Oral Diseases</i> , 2018, 24, 864-865.	3.0	3

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91	Chrysophanol Relieves Cisplatin-Induced Nephrotoxicity via Concomitant Inhibition of Oxidative Stress, Apoptosis, and Inflammation. <i>Frontiers in Physiology</i> , 2021, 12, 706359.	2.8	3
92	Rational Discovery of Response Biomarkers: Candidate Prognostic Factors and Biomarkers for Checkpoint Inhibitor-Based Immunotherapy. <i>Advances in Experimental Medicine and Biology</i> , 2020, 1248, 143-166.	1.6	3
93	Significant benefits of osimertinib in treating acquired resistance to first-generation EGFR-TKIs in lung squamous cell cancer: A case report. <i>World Journal of Clinical Cases</i> , 2019, 7, 1221-1229.	0.8	3
94	PDE4B Modulates Glucocorticoid Sensitivity in Childhood Acute Lymphoblastic Leukemia. <i>Blood</i> , 2012, 120, 530-530.	1.4	3
95	Quantum yield and lifetime data analysis for the UV curable quantum dot nanocomposites. <i>Data in Brief</i> , 2016, 6, 614-618.	1.0	2
96	Reduced cortical thickness related to single nucleotide polymorphisms in the major histocompatibility complex region in antipsychotic-naïve schizophrenia. <i>Brain and Behavior</i> , 2019, 9, e01253.	2.2	2
97	Genetic polymorphisms near IL-21 gene associated with Th17 cytokines confer risk for systemic lupus erythematosus in Chinese Han population. <i>Lupus</i> , 2019, 28, 406-413.	1.6	2
98	Fas-Associated Factor 1 Promotes Hepatic Insulin Resistance via JNK Signaling Pathway. <i>Oxidative Medicine and Cellular Longevity</i> , 2021, 2021, 1-10.	4.0	2
99	Leukemia Risk Gene ARID5B is a Crucial Regulator of B-Cell Development. <i>Blood</i> , 2018, 132, 385-385.	1.4	2
100	Genome-wide analysis identify novel germline genetic variations in ADCY1 influencing platinum-based chemotherapy response in non-small cell lung cancer. <i>Acta Pharmaceutica Sinica B</i> , 2021, 12, 1514-1522.	12.0	2
101	Genome-Wide Association Study Identifies a Novel Susceptibility Locus At 10p12.31-12.2 for Childhood Acute Lymphoblastic Leukemia in Ethnically Diverse Populations. <i>Blood</i> , 2012, 120, 877-877.	1.4	2
102	Insights of Acute Lymphoblastic Leukemia with Development of Genomic Investigation. <i>Methods in Molecular Biology</i> , 2018, 1754, 387-413.	0.9	1
103	A functional variant in CHK1 contributes to increased risk of nasopharyngeal carcinoma in a Han Chinese population. <i>Journal of Cellular Biochemistry</i> , 2020, 121, 3248-3255.	2.6	1
104	ARID5B Genetic Polymorphisms Contribute to Racial Disparities In Childhood Acute Lymphoblastic Leukemia: A Children's Oncology Group Study. <i>Blood</i> , 2010, 116, 8-8.	1.4	1
105	Prenatal diagnosis of ductal origin of distal pulmonary artery: presentation of three cases and literature review. <i>Ultrasound in Obstetrics and Gynecology</i> , 2022, 60, 284-290.	1.7	1
106	Case Report: Brachydactyly Type A1 Induced by a Novel Variant of in-Frame Insertion in the IHH Gene. <i>Frontiers in Genetics</i> , 2022, 13, .	2.3	1
107	CEBPE promoter SNPs, caught red handed? A commentary on identification of functional nucleotide and haplotype variants in the promoter of the CEBPE gene. <i>Journal of Human Genetics</i> , 2013, 58, 571-572.	2.3	0