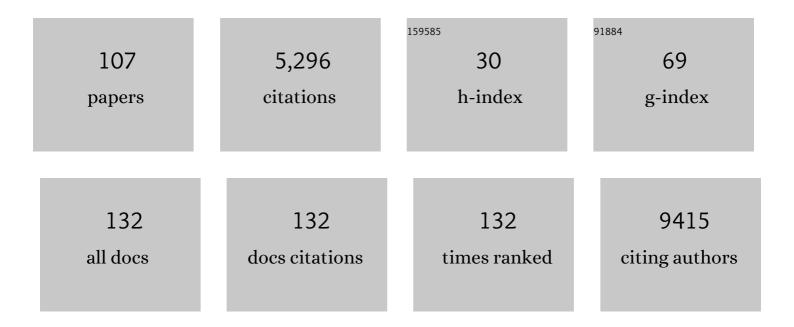


List of Publications by Year in descending order

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HENC XU

#	Article	IF	CITATIONS
1	Genomic evolution and diverse models of systemic metastases in colorectal cancer. Gut, 2022, 71, 322-332.	12.1	51
2	Prenatal diagnosis of ductal origin of distal pulmonary artery: presentation of three cases and literature review. Ultrasound in Obstetrics and Gynecology, 2022, 60, 284-290.	1.7	1
3	Deep learningâ€based AI model for signetâ€ring cell carcinoma diagnosis and chemotherapy response prediction in gastric cancer. Medical Physics, 2022, 49, 1535-1546.	3.0	17
4	Association of recurrent APOBEC3B alterations with the prognosis of gastric-type cervical adenocarcinoma. Gynecologic Oncology, 2022, 165, 105-113.	1.4	7
5	PHLDB2 Mediates Cetuximab Resistance via Interacting With EGFR in Latent Metastasis of Colorectal Cancer. Cellular and Molecular Gastroenterology and Hepatology, 2022, 13, 1223-1242.	4.5	16
6	Thrombin induces ACSL4-dependent ferroptosis during cerebral ischemia/reperfusion. Signal Transduction and Targeted Therapy, 2022, 7, 59.	17.1	88
7	ANI analysis of poxvirus genomes reveals its potential application to viral species rank demarcation. Virus Evolution, 2022, 8, .	4.9	7
8	Case Report: Brachydactyly Type A1 Induced by a Novel Variant of in-Frame Insertion in the IHH Gene. Frontiers in Genetics, 2022, 13, .	2.3	1
9	Fas-Associated Factor 1 Promotes Hepatic Insulin Resistance via JNK Signaling Pathway. Oxidative Medicine and Cellular Longevity, 2021, 2021, 1-10.	4.0	2
10	Novel Recurrent Altered Genes in Chinese Patients With Anaplastic Thyroid Cancer. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e988-e998.	3.6	33
11	<i>ORAOV1-B</i> Promotes OSCC Metastasis via the NF-κB-TNFα Loop. Journal of Dental Research, 2021, 100, 002203452199633.	5.2	15
12	Multi-omics approaches identify SF3B3 and SIRT3 as candidate autophagic regulators and druggable targets in invasive breast carcinoma. Acta Pharmaceutica Sinica B, 2021, 11, 1227-1245.	12.0	26
13	LINC-PINT impedes DNA repair and enhances radiotherapeutic response by targeting DNA-PKcs in nasopharyngeal cancer. Cell Death and Disease, 2021, 12, 454.	6.3	14
14	Improvement of Neoantigen Identification Through Convolution Neural Network. Frontiers in Immunology, 2021, 12, 682103.	4.8	4
15	Urinary complement proteins and risk of end-stage renal disease: quantitative urinary proteomics in patients with type 2 diabetes and biopsy-proven diabetic nephropathy. Journal of Endocrinological Investigation, 2021, 44, 2709-2723.	3.3	16
16	Longitudinal Genomic Evolution of Conventional Papillary Thyroid Cancer With Brain Metastasis. Frontiers in Oncology, 2021, 11, 620924.	2.8	4
17	Characterizing dedifferentiation of thyroid cancer by integrated analysis. Science Advances, 2021, 7, .	10.3	76
18	Alpha defensin-1 attenuates surgically induced osteoarthritis in association with promoting M1 to M2 macrophage polarization. Osteoarthritis and Cartilage, 2021, 29, 1048-1059.	1.3	21

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19	Chrysophanol Relieves Cisplatin-Induced Nephrotoxicity via Concomitant Inhibition of Oxidative Stress, Apoptosis, and Inflammation. Frontiers in Physiology, 2021, 12, 706359.	2.8	3
20	Age-Associated Proteomic Signatures and Potential Clinically Actionable Targets of Colorectal Cancer. Molecular and Cellular Proteomics, 2021, 20, 100115.	3.8	29
21	Personalized neoantigen pulsed dendritic cell vaccine for advanced lung cancer. Signal Transduction and Targeted Therapy, 2021, 6, 26.	17.1	112
22	LncRNA linc00312 suppresses radiotherapy resistance by targeting DNA-PKcs and impairing DNA damage repair in nasopharyngeal carcinoma. Cell Death and Disease, 2021, 12, 69.	6.3	56
23	Genome-wide analysis identify novel germline genetic variations in ADCY1 influencing platinum-based chemotherapy response in non-small cell lung cancer. Acta Pharmaceutica Sinica B, 2021, 12, 1514-1522.	12.0	2
24	Long Non-coding RNA Expression Profiling in Biopsy to Identify Renal Allograft at Risk of Chronic Damage and Future Graft Loss. Applied Biochemistry and Biotechnology, 2020, 190, 660-673.	2.9	11
25	Prognostic and predictive role of DNA mismatch repair status in stage IIâ€III colorectal cancer: A systematic review and metaâ€analysis. Clinical Genetics, 2020, 97, 25-38.	2.0	19
26	<i>ARID5B</i> Influences Antimetabolite Drug Sensitivity and Prognosis of Acute Lymphoblastic Leukemia. Clinical Cancer Research, 2020, 26, 256-264.	7.0	25
27	A functional variant in CHK1 contributes to increased risk of nasopharyngeal carcinoma in a Han Chinese population. Journal of Cellular Biochemistry, 2020, 121, 3248-3255.	2.6	1
28	Personalized neoantigen-pulsed dendritic cell vaccines show superior immunogenicity to neoantigen-adjuvant vaccines in mouse tumor models. Cancer Immunology, Immunotherapy, 2020, 69, 135-145.	4.2	42
29	Structural and Functional Insights into an Archaeal Lipid Synthase. Cell Reports, 2020, 33, 108294.	6.4	11
30	DP7-C-modified liposomes enhance immune responses and the antitumor effect of a neoantigen-based mRNA vaccine. Journal of Controlled Release, 2020, 328, 210-221.	9.9	54
31	The Significance of the CLDN18-ARHGAP Fusion Gene in Gastric Cancer: A Systematic Review and Meta-Analysis. Frontiers in Oncology, 2020, 10, 1214.	2.8	23
32	The association of hematuria on kidney clinicopathologic features and renal outcome in patients with diabetic nephropathy: a biopsy-based study. Journal of Endocrinological Investigation, 2020, 43, 1213-1220.	3.3	13
33	The Role of ARID5B in Acute Lymphoblastic Leukemia and Beyond. Frontiers in Genetics, 2020, 11, 598.	2.3	21
34	GPR39 Overexpression in OSCC Promotes YAP-Sustained Malignant Progression. Journal of Dental Research, 2020, 99, 949-958.	5.2	22
35	Rational Discovery of Response Biomarkers: Candidate Prognostic Factors and Biomarkers for Checkpoint Inhibitor-Based Immunotherapy. Advances in Experimental Medicine and Biology, 2020, 1248, 143-166.	1.6	3
36	Cause-specific risk of major adverse cardiovascular outcomes and hypoglycemic in patients with type 2 diabetes: a multicenter prospective cohort study. Endocrine, 2019, 63, 44-51.	2.3	8

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37	Polytropic Influence of TRIB3 rs2295490 Genetic Polymorphism on Response to Antihypertensive Agents in Patients With Essential Hypertension. Frontiers in Pharmacology, 2019, 10, 236.	3.5	5
38	Long-Term Correction of Copper Metabolism in Wilson's Disease Mice with AAV8 Vector Delivering Truncated ATP7B. Human Gene Therapy, 2019, 30, 1494-1504.	2.7	15
39	Prevotella copri is associated with carboplatin-induced gut toxicity. Cell Death and Disease, 2019, 10, 714.	6.3	32
40	Regulatory Network and Prognostic Effect Investigation of PIP4K2A in Leukemia and Solid Cancers. Frontiers in Genetics, 2019, 9, 721.	2.3	12
41	A Functional 5'-UTR Polymorphism of MYC Contributes to Nasopharyngeal Carcinoma Susceptibility and Chemoradiotherapy Induced Toxicities. Journal of Cancer, 2019, 10, 147-155.	2.5	12
42	ARL15 overexpression attenuates high glucose-induced impairment of insulin signaling and oxidative stress in human umbilical vein endothelial cells. Life Sciences, 2019, 220, 127-135.	4.3	12
43	Topical calcineurin inhibitors in the treatment of oral lichen planus: a systematic review and metaâ€analysis. British Journal of Dermatology, 2019, 181, 1166-1176.	1.5	38
44	NUDT15 Polymorphism Confer Increased Susceptibility to Thiopurine-Induced Leukopenia in Patients With Autoimmune Hepatitis and Related Cirrhosis. Frontiers in Pharmacology, 2019, 10, 346.	3.5	22
45	Metabolismâ€induced tumor activator 1 (MITA1), an Energy Stress–Inducible Long Noncoding RNA, Promotes Hepatocellular Carcinoma Metastasis. Hepatology, 2019, 70, 215-230.	7.3	65
46	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. Endocrine, 2019, 64, 536-543.	2.3	19
47	Reduced cortical thickness related to single nucleotide polymorphisms in the major histocompatibility complex region in antipsychoticâ€naive schizophrenia. Brain and Behavior, 2019, 9, e01253.	2.2	2
48	Genome-Wide Association Study of Susceptibility Loci for T-Cell Acute Lymphoblastic Leukemia in Children. Journal of the National Cancer Institute, 2019, 111, 1350-1357.	6.3	32
49	Proteomic Maps of Human Gastrointestinal Stromal Tumor Subgroups*. Molecular and Cellular Proteomics, 2019, 18, 923a-935.	3.8	10
50	Genetic polymorphisms near IL-21 gene associated with Th17 cytokines confer risk for systemic lupus erythematosus in Chinese Han population. Lupus, 2019, 28, 406-413.	1.6	2
51	Subtypeâ€specific inherited predisposition to pemphigus in the Chinese population. British Journal of Dermatology, 2019, 180, 828-835.	1.5	25
52	Novel susceptibility variants at the ERG locus for childhood acute lymphoblastic leukemia in Hispanics. Blood, 2019, 133, 724-729.	1.4	44
53	The influence of telmisartan on metformin pharmacokinetics and pharmacodynamics. Journal of Pharmacological Sciences, 2019, 139, 37-41.	2.5	4
54	Significant benefits of osimertinib in treating acquired resistance to first-generation EGFR-TKIs in lung squamous cell cancer: A case report. World Journal of Clinical Cases, 2019, 7, 1221-1229.	0.8	3

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55	Novel GLI3 Mutations in Chinese Patients with Non-syndromic Post-axial Polydactyly. Current Molecular Medicine, 2019, 19, 228-235.	1.3	5
56	Characteristics of genomic alterations of lung adenocarcinoma in young neverâ€smokers. International Journal of Cancer, 2018, 143, 1696-1705.	5.1	45
57	Insights of Acute Lymphoblastic Leukemia with Development of Genomic Investigation. Methods in Molecular Biology, 2018, 1754, 387-413.	0.9	1
58	Combination of common and novel rare <i>NUDT15</i> variants improves predictive sensitivity of thiopurine-induced leukopenia in children with acute lymphoblastic leukemia. Haematologica, 2018, 103, e293-e295.	3.5	24
59	Triterpenoids with antioxidant activities from <i>Myricaria squamosa</i> . Journal of Asian Natural Products Research, 2018, 20, 292-298.	1.4	6
60	Evidence of genetic factors involved in oral lichen planus pathogenesis. Oral Diseases, 2018, 24, 864-865.	3.0	3
61	<i>TP53</i> Germline Variations Influence the Predisposition and Prognosis of B-Cell Acute Lymphoblastic Leukemia in Children. Journal of Clinical Oncology, 2018, 36, 591-599.	1.6	121
62	Prognostic significance of frequent CLDN18-ARHGAP26/6 fusion in gastric signet-ring cell cancer. Nature Communications, 2018, 9, 2447.	12.8	100
63	Presence and diagnostic value of circulating tsncRNA for ovarian tumor. Molecular Cancer, 2018, 17, 163.	19.2	19
64	Intensive Glucose Control Reduces the Risk Effect of TRIB3, SMARCD3, and ATF6 Genetic Variation on Diabetic Vascular Complications. Frontiers in Pharmacology, 2018, 9, 1422.	3.5	10
65	Prognostic Factors for Checkpoint Inhibitor Based Immunotherapy: An Update With New Evidences. Frontiers in Pharmacology, 2018, 9, 1050.	3.5	48
66	CSF glutamate level decreases in heavy smokers and negatively correlates with BDI scores. Psychiatry Research, 2018, 270, 627-630.	3.3	4
67	Structural and functional insights into the regulation of the lysis–lysogeny decision in viral communities. Nature Microbiology, 2018, 3, 1285-1294.	13.3	49
68	Precise control of maxillary multidirectional movement in Le Fort I osteotomy using a surgical guiding device. British Journal of Oral and Maxillofacial Surgery, 2018, 56, 797-804.	0.8	11
69	Induction and Amelioration of Methotrexate-Induced Gastrointestinal Toxicity are Related to Immune Response and Gut Microbiota. EBioMedicine, 2018, 33, 122-133.	6.1	80
70	Association of the independent polymorphisms in CDKN2A with susceptibility of acute lymphoblastic leukemia. Bioscience Reports, 2018, 38, .	2.4	10
71	MicroRNA-214 promotes hepatic stellate cell activation and liver fibrosis by suppressing Sufu expression. Cell Death and Disease, 2018, 9, 718.	6.3	72
72	Leukemia Risk Gene ARID5B is a Crucial Regulator of B-Cell Development. Blood, 2018, 132, 385-385.	1.4	2

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73	The genetic landscape of benign thyroid nodules revealed by whole exome and transcriptome sequencing. Nature Communications, 2017, 8, 15533.	12.8	53
74	Rare gene variants in a patient with azathioprine-induced lethal myelosuppression. Annals of Hematology, 2017, 96, 2131-2133.	1.8	6
75	Novel Gene and Network Associations Found for Acute Lymphoblastic Leukemia Using Case–Control and Family-Based Studies in Multiethnic Populations. Cancer Epidemiology Biomarkers and Prevention, 2017, 26, 1531-1539.	2.5	7
76	Impact of <i>NUDT15</i> polymorphisms on thiopurines-induced myelotoxicity and thiopurines tolerance dose. Oncotarget, 2017, 8, 13575-13585.	1.8	40
77	Regulatory network of <i>GATA3</i> in pediatric acute lymphoblastic leukemia. Oncotarget, 2017, 8, 36040-36053.	1.8	30
78	The novel complex combination of alum, CpG ODN and HH2 as adjuvant in cancer vaccine effectively suppresses tumor growth <i>in vivo</i> . Oncotarget, 2017, 8, 45951-45964.	1.8	18
79	Prognostic effect of high-flux hemodialysis in patients with chronic kidney disease. Brazilian Journal of Medical and Biological Research, 2016, 49, e4708.	1.5	7
80	Efficacy evaluation of clonazepam for symptom remission in burning mouth syndrome: a metaâ€analysis. Oral Diseases, 2016, 22, 503-511.	3.0	52
81	Association Between PIP4K2A Polymorphisms and Acute Lymphoblastic Leukemia Susceptibility. Medicine (United States), 2016, 95, e3542.	1.0	27
82	Quantum yield and lifetime data analysis for the UV curable quantum dot nanocomposites. Data in Brief, 2016, 6, 614-618.	1.0	2
83	CRISPR/Cas9 facilitates investigation of neural circuit disease using human iPSCs: mechanism of epilepsy caused by an SCN1A loss-of-function mutation. Translational Psychiatry, 2016, 6, e703-e703.	4.8	82
84	Genome-Wide Association Studies of HIV-1 Host Control in Ethnically Diverse Chinese Populations. Scientific Reports, 2015, 5, 10879.	3.3	23
85	A genome-wide association study of susceptibility to acute lymphoblastic leukemia in adolescents and young adults. Blood, 2015, 125, 680-686.	1.4	110
86	Common variants in ACYP2 influence susceptibility to cisplatin-induced hearing loss. Nature Genetics, 2015, 47, 263-266.	21.4	109
87	RGD-modifided oncolytic adenovirus exhibited potent cytotoxic effect on CAR-negative bladder cancer-initiating cells. Cell Death and Disease, 2015, 6, e1760-e1760.	6.3	18
88	Inherited coding variants at the CDKN2A locus influence susceptibility to acute lymphoblastic leukaemia in children. Nature Communications, 2015, 6, 7553.	12.8	72
89	Bioconcentration Factors and Potential Human Health Risks of Heavy Metals in Cultivated Lentinus edodes in Chengdu, People's Republic of China. Journal of Food Protection, 2015, 78, 390-395.	1.7	9
90	Germline genetic variation in ETV6 and risk of childhood acute lymphoblastic leukaemia: a systematic genetic study. Lancet Oncology, The, 2015, 16, 1659-1666.	10.7	161

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91	Uncovering the Rare Variants of DLC1 Isoform 1 and Their Functional Effects in a Chinese Sporadic Congenital Heart Disease Cohort. PLoS ONE, 2014, 9, e90215.	2.5	14
92	A New System Identification Approach to Identify Genetic Variants in Sequencing Studies for a Binary Phenotype. Human Heredity, 2014, 78, 104-116.	0.8	19
93	ARID5B Regulates Leukemia Sensitivity to Antimetabolites in Children with Acute Lymphoblastic Leukemia Via Effects on Cell Cycle Progression. Blood, 2014, 124, 791-791.	1.4	28
94	Inherited GATA3 variants are associated with Ph-like childhood acute lymphoblastic leukemia and risk of relapse. Nature Genetics, 2013, 45, 1494-1498.	21.4	264
95	Response. Journal of the National Cancer Institute, 2013, 105, 1512-1513.	6.3	6
96	Novel Susceptibility Variants at 10p12.31-12.2 for Childhood Acute Lymphoblastic Leukemia in Ethnically Diverse Populations. Journal of the National Cancer Institute, 2013, 105, 733-742.	6.3	208
97	CEBPE promoter SNPs, caught red handed? A commentary on identification of functional nucleotide and haplotype variants in the promoter of the CEBPE gene. Journal of Human Genetics, 2013, 58, 571-572.	2.3	0
98	Colocalization of Inflammatory Response with B7-H1 Expression in Human Melanocytic Lesions Supports an Adaptive Resistance Mechanism of Immune Escape. Science Translational Medicine, 2012, 4, 127ra37.	12.4	1,837
99	<i>ARID5B</i> Genetic Polymorphisms Contribute to Racial Disparities in the Incidence and Treatment Outcome of Childhood Acute Lymphoblastic Leukemia. Journal of Clinical Oncology, 2012, 30, 751-757.	1.6	165
100	PDE4B Modulates Glucocorticoid Sensitivity in Childhood Acute Lymphoblastic Leukemia. Blood, 2012, 120, 530-530.	1.4	3
101	Genome-Wide Association Study Identifies a Novel Susceptibility Locus At 10p12.31-12.2 for Childhood Acute Lymphoblastic Leukemia in Ethinically Diverse Populations. Blood, 2012, 120, 877-877.	1.4	2
102	Length of the ORF, position of the first AUG and the Kozak motif are important factors in potential dual-coding transcripts. Cell Research, 2010, 20, 445-457.	12.0	35
103	Screening of Kozak-motif-located SNPs and analysis of their association with human diseases. Biochemical and Biophysical Research Communications, 2010, 392, 89-94.	2.1	15
104	ARID5B Genetic Polymorphisms Contribute to Racial Disparities In Childhood Acute Lymphoblastic Leukemia: A Children's Oncology Group Study. Blood, 2010, 116, 8-8.	1.4	1
105	Assessment of Glomerular Filtration Rate in Renal Transplant Patients Using Serum Cystatin C. Transplantation Proceedings, 2006, 38, 2006-2008.	0.6	29
106	Mitomycin C reduces haze formation in rabbits after excimer laser photorefractive keratectomy. Journal of Refractive Surgery, 2001, 17, 342-9.	2.3	9
107	Characteristics of an established retinoblastoma cell line HXO-Rb44. Yan Ke Xue Bao = Eye Science, 1995, 11, 16-21.	0.1	4