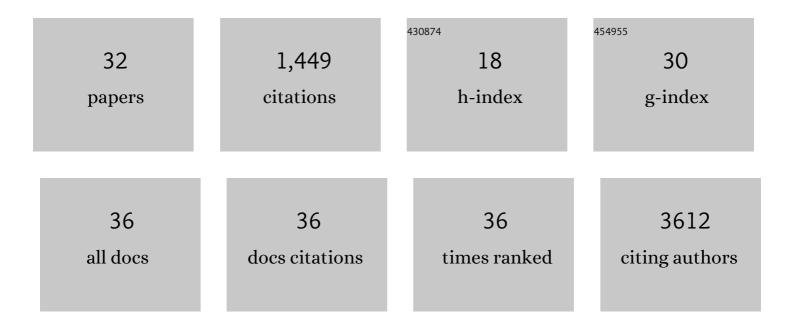
Zhongyang Zhang

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Massive parallel sequencing uncovers actionable FGFR2–PPHLN1 fusion and ARAF mutations in in in intrahepatic cholangiocarcinoma. Nature Communications, 2015, 6, 6087.	12.8	240
2	Molecular predictors of prevention of recurrence in HCC with sorafenib as adjuvant treatment and prognostic factors in the phase 3 STORM trial. Gut, 2019, 68, 1065-1075.	12.1	195
3	Trunk mutational events present minimal intra- and inter-tumoral heterogeneity in hepatocellular carcinoma. Journal of Hepatology, 2017, 67, 1222-1231.	3.7	121
4	Mixed hepatocellular cholangiocarcinoma tumors: Cholangiolocellular carcinoma is a distinct molecular entity. Journal of Hepatology, 2017, 66, 952-961.	3.7	120
5	Unique Genomic Profile of Fibrolamellar Hepatocellular Carcinoma. Gastroenterology, 2015, 148, 806-818.e10.	1.3	109
6	NPHP1 (Nephrocystin-1) Gene Deletions Cause Adult-Onset ESRD. Journal of the American Society of Nephrology: JASN, 2018, 29, 1772-1779.	6.1	74
7	Development and clinical application of an integrative genomic approach to personalized cancer therapy. Genome Medicine, 2016, 8, 62.	8.2	71
8	Genome-Wide Association Study of the Modified Stumvoll Insulin Sensitivity Index Identifies <i>BCL2</i> and <i>FAM19A2</i> as Novel Insulin Sensitivity Loci. Diabetes, 2016, 65, 3200-3211.	0.6	67
9	Understanding the Hidden Complexity of Latin American Population Isolates. American Journal of Human Genetics, 2018, 103, 707-726.	6.2	48
10	Airway microbiome is associated with respiratory functions and responses to ambient particulate matter exposure. Ecotoxicology and Environmental Safety, 2019, 167, 269-277.	6.0	48
11	SAAS-CNV: A Joint Segmentation Approach on Aggregated and Allele Specific Signals for the Identification of Somatic Copy Number Alterations with Next-Generation Sequencing Data. PLoS Computational Biology, 2015, 11, e1004618.	3.2	40
12	Genetic regulation of the placental transcriptome underlies birth weight and risk of childhood obesity. PLoS Genetics, 2018, 14, e1007799.	3.5	38
13	Recipient APOL1 risk alleles associate with death-censored renal allograft survival and rejection episodes. Journal of Clinical Investigation, 2021, 131, .	8.2	33
14	Genome-wide approach identifies a novel gene-maternal pre-pregnancy BMI interaction on preterm birth. Nature Communications, 2017, 8, 15608.	12.8	31
15	Collagenous Colitis Is Associated With HLA Signature and Shares Genetic Risks With Other Immune-Mediated Diseases. Gastroenterology, 2020, 159, 549-561.e8.	1.3	31
16	Contribution of common and rare variants to bipolar disorder susceptibility in extended pedigrees from population isolates. Translational Psychiatry, 2020, 10, 74.	4.8	25
17	Genome-wide non-HLA donor-recipient genetic differences influence renal allograft survival via early allograft fibrosis. Kidney International, 2020, 98, 758-768.	5.2	25
18	Reconstructing DNA copy number by penalized estimation and imputation. Annals of Applied Statistics, 2010. 4. 1749-1773.	1.1	22

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#	Article	IF	CITATIONS
19	Prenatal exposure to ambient air multi-pollutants significantly impairs intrauterine fetal development trajectory. Ecotoxicology and Environmental Safety, 2020, 201, 110726.	6.0	20
20	Identification of human age-associated gene co-expressions in functional modules using liquid association. Oncotarget, 2018, 9, 1063-1074.	1.8	16
21	Reconstructing DNA copy number by joint segmentation of multiple sequences. BMC Bioinformatics, 2012, 13, 205.	2.6	15
22	EnsembleCNV: an ensemble machine learning algorithm to identify and genotype copy number variation using SNP array data. Nucleic Acids Research, 2019, 47, e39-e39.	14.5	15
23	Non-HLA donor–recipient mismatches in kidney transplantation—A stone left unturned. American Journal of Transplantation, 2020, 20, 19-24.	4.7	13
24	Intestinal Dysbiosis in Young Cystic Fibrosis Rabbits. Journal of Personalized Medicine, 2021, 11, 132.	2.5	6
25	Ambient Air Pollutants and Traffic Factors Were Associated with Blood and Urine Biomarkers and Asthma Risk. Environmental Science & Technology, 2022, 56, 7298-7307.	10.0	6
26	Using SAAS-CNV to Detect and Characterize Somatic Copy Number Alterations in Cancer Genomes from Next Generation Sequencing and SNP Array Data. Methods in Molecular Biology, 2018, 1833, 29-47.	0.9	5
27	A Novel Targeted Learning Method for Quantitative Trait Loci Mapping. Genetics, 2014, 198, 1369-1376.	2.9	4
28	APOL1 G2 risk allele—clarifying nomenclature. Kidney International, 2017, 92, 518-519.	5.2	4
29	Bio3Air, an integrative system for monitoring individual-level air pollutant exposure with high time and spatial resolution. Ecotoxicology and Environmental Safety, 2019, 169, 756-763.	6.0	3
30	Polygenic risk score for alcohol drinking behavior improves prediction of inflammatory bowel disease risk. Human Molecular Genetics, 2021, 30, 514-523.	2.9	2
31	A Dynamic Pooling Approach to Extract Complete Allele Signal Information in Somatic Copy Number Alternations Detection. , 2018, , .		0
32	Enabling Clinical Trials for AMR in the Era of Precision Medicine. Transplantation, 2021, 105, 482-483.	1.0	0