## Ni Huang

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Accessible Region Conformation Capture (ARC-C) gives high-resolution insights into genome architecture and regulation. Genome Research, 2022, 32, 357-366.	5.5	6
2	Cross-tissue immune cell analysis reveals tissue-specific features in humans. Science, 2022, 376, eabl5197.	12.6	265
3	Developmental cell programs are co-opted in inflammatory skin disease. Science, 2021, 371, .	12.6	264
4	User-friendly, scalable tools and workflows for single-cell RNA-seq analysis. Nature Methods, 2021, 18, 327-328.	19.0	26
5	Transcriptional characterization of human megakaryocyte polyploidization and lineage commitment. Journal of Thrombosis and Haemostasis, 2021, 19, 1236-1249.	3.8	15
6	DREAM represses distinct targets by cooperating with different THAP domain proteins. Cell Reports, 2021, 37, 109835.	6.4	6
7	Expression Atlas update: from tissues to single cells. Nucleic Acids Research, 2020, 48, D77-D83.	14.5	363
8	A single-cell RNA-sequencing training and analysis suite using the Galaxy framework. GigaScience, 2020, 9, .	6.4	14
9	Putative cell type discovery from single-cell gene expression data. Nature Methods, 2020, 17, 621-628.	19.0	91
10	Physical and functional interaction between SET1/COMPASS complex component CFP-1 and a Sin3S HDAC complex in C. elegans. Nucleic Acids Research, 2019, 47, 11164-11180.	14.5	54
11	Rare mutations in the complement regulatory gene CSMD1 are associated with male and female infertility. Nature Communications, 2019, 10, 4626.	12.8	24
12	Chromatin accessibility dynamics across C. elegans development and ageing. ELife, 2018, 7, .	6.0	76
13	Dynamic landscape and regulation of RNA editing in mammals. Nature, 2017, 550, 249-254.	27.8	495
14	A team of heterochromatin factors collaborates with small RNA pathways to combat repetitive elements and germline stress. ELife, 2017, 6, .	6.0	87
15	Stable <i>Caenorhabditis elegans</i> chromatin domains separate broadly expressed and developmentally regulated genes. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, E7020-E7029.	7.1	73
16	A Screen for Genomic Disorders of Infertility Identifies MAST2 Duplications Associated with Nonobstructive Azoospermia in Humans1. Biology of Reproduction, 2015, 93, 61.	2.7	30
17	Human Spermatogenic Failure Purges Deleterious Mutation Load from the Autosomes and Both Sex Chromosomes, including the Gene DMRT1. PLoS Genetics, 2013, 9, e1003349.	3.5	118
18	Large, rare chromosomal deletions associated with severe early-onset obesity. Nature, 2010, 463, 666-670.	27.8	487