

Liqing Tian

List of Publications by Year in descending order

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Version: 2024-02-01

26
papers

1,381
citations

623699

14
h-index

752679

20
g-index

27
all docs

27
docs citations

27
times ranked

2678
citing authors

#	ARTICLE	IF	CITATIONS
1	LncRNA profile study reveals a three-lncRNA signature associated with the survival of patients with oesophageal squamous cell carcinoma. <i>Gut</i> , 2014, 63, 1700-1710.	12.1	385
2	Analysis of error profiles in deep next-generation sequencing data. <i>Genome Biology</i> , 2019, 20, 50.	8.8	196
3	Therapy-induced mutations drive the genomic landscape of relapsed acute lymphoblastic leukemia. <i>Blood</i> , 2020, 135, 41-55.	1.4	171
4	St. Jude Cloud: A Pediatric Cancer Genomic Data-Sharing Ecosystem. <i>Cancer Discovery</i> , 2021, 11, 1082-1099.	9.4	109
5	Acute depletion of CTCF directly affects MYC regulation through loss of enhancer-promoter looping. <i>Nucleic Acids Research</i> , 2019, 47, 6699-6713.	14.5	98
6	CICERO: a versatile method for detecting complex and diverse driver fusions using cancer RNA sequencing data. <i>Genome Biology</i> , 2020, 21, 126.	8.8	74
7	Improved side-chain modeling by coupling clash-detection guided iterative search with rotamer relaxation. <i>Bioinformatics</i> , 2011, 27, 785-790.	4.1	52
8	Mir-23a in amplified 19p13.13 loci targets metallothionein 2A and promotes growth in gastric cancer cells. <i>Journal of Cellular Biochemistry</i> , 2013, 114, 2160-2169.	2.6	49
9	Discovery of regulatory noncoding variants in individual cancer genomes by using cis-X. <i>Nature Genetics</i> , 2020, 52, 811-818.	21.4	47
10	MiRNA expression profile reveals a prognostic signature for esophageal squamous cell carcinoma. <i>Cancer Letters</i> , 2014, 350, 34-42.	7.2	43
11	Transcriptome profiling of esophageal squamous cell carcinoma reveals a long noncoding RNA acting as a tumor suppressor. <i>Oncotarget</i> , 2015, 6, 17065-17080.	1.8	39
12	Integrated Genomic Analysis Identifies <i>UBTF</i> Tandem Duplications as a Recurrent Lesion in Pediatric Acute Myeloid Leukemia. <i>Blood Cancer Discovery</i> , 2022, 3, 194-207.	5.0	38
13	Exploration of Coding and Non-coding Variants in Cancer Using GenomePaint. <i>Cancer Cell</i> , 2021, 39, 83-95.e4.	16.8	18
14	Long-read sequencing unveils IGH-DUX4 translocation into the silenced IGH allele in B-cell acute lymphoblastic leukemia. <i>Nature Communications</i> , 2019, 10, 2789.	12.8	14
15	Incorporation of Local Structural Preference Potential Improves Fold Recognition. <i>PLoS ONE</i> , 2011, 6, e17215.	2.5	13
16	Integrative network analysis reveals USP7 haploinsufficiency inhibits E-protein activity in pediatric T-lineage acute lymphoblastic leukemia (T-ALL). <i>Scientific Reports</i> , 2021, 11, 5154.	3.3	10
17	The landscape of coding RNA editing events in pediatric cancer. <i>BMC Cancer</i> , 2021, 21, 1233.	2.6	7
18	NCACO-score: An effective main-chain dependent scoring function for structure modeling. <i>BMC Bioinformatics</i> , 2011, 12, 208.	2.6	4

#	ARTICLE	IF	CITATIONS
19	A point mutation in the DNA-binding domain of HPV-2 E2 protein increases its DNA-binding capacity and reverses its transcriptional regulatory activity on the viral early promoter. BMC Molecular Biology, 2012, 13, 5.	3.0	4
20	Editorial: Big Data and Machine Learning in Cancer Genomics. Frontiers in Genetics, 2021, 12, 749584.	2.3	2
21	Abstract 1485: Allelic specificity of immunoglobulin heavy chain (IGH) translocation in B-cell acute lymphoblastic leukemia (B-ALL) unveiled by long-read sequencing. , 2018, , .		2
22	Abstract 1543: Mining cancer-specific isoforms as CAR T-cell therapy targets for pediatric solid and brain tumors. , 2021, , .		1
23	Mutational Landscape and Temporal Evolution during Treatment of Relapsed Acute Lymphoblastic Leukemia. Blood, 2018, 132, 917-917.	1.4	0
24	Abstract 2872: Acquisition of drug resistance mutations during chemotherapy treatment in pediatric acute lymphoblastic leukemia. , 2019, , .		0
25	Abstract 5478: CICERO: An accurate method for detecting complex and diverse driver fusions using cancer transcriptome sequencing (RNA-seq) data. , 2020, , .		0
26	Integrated Genomic Analysis Identifies UBTF Tandem Duplications As a Subtype-Defining Lesion in Pediatric Acute Myeloid Leukemia. Blood, 2021, 138, LBA-4-LBA-4.	1.4	0