

Shimin Shuai

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

15
papers

722
citations

11
h-index

16
g-index

16
ext. papers

1,075
ext. citations

21.9
avg, IF

3.87
L-index

#	Paper	IF	Citations
15	PanCancer analysis of somatic mutations in repetitive regions reveals recurrent mutations in snRNA U2.. <i>Npj Genomic Medicine</i> , 2022 , 7, 19	6.2	1
14	Interruption of KLF5 acetylation in basal progenitor cells promotes luminal commitment by activating Notch signaling.. <i>Journal of Genetics and Genomics</i> , 2021 ,	4	0
13	Candidate Cancer Driver Mutations in Distal Regulatory Elements and Long-Range Chromatin Interaction Networks. <i>Molecular Cell</i> , 2020 , 77, 1307-1321.e10	17.6	20
12	Combined burden and functional impact tests for cancer driver discovery using DriverPower. <i>Nature Communications</i> , 2020 , 11, 734	17.4	16
11	Integrative pathway enrichment analysis of multivariate omics data. <i>Nature Communications</i> , 2020 , 11, 735	17.4	53
10	Pathway and network analysis of more than 2500 whole cancer genomes. <i>Nature Communications</i> , 2020 , 11, 729	17.4	38
9	Analyses of non-coding somatic drivers in 2,658 cancer whole genomes. <i>Nature</i> , 2020 , 578, 102-111	50.4	220
8	Cancer LncRNA Census reveals evidence for deep functional conservation of long noncoding RNAs in tumorigenesis. <i>Communications Biology</i> , 2020 , 3, 56	6.7	77
7	The U1 spliceosomal RNA is recurrently mutated in multiple cancers. <i>Nature</i> , 2019 , 574, 712-716	50.4	79
6	MEDU-39. HIGHLY RECURRENT U1 SMALL NUCLEAR RNA HOTSPOT MUTATIONS DRIVE ALTERNATIVE SPLICING IN SONIC HEDGEHOG MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2019 , 21, ii111-ii111	11	78
5	Recurrent noncoding U1 snRNA mutations drive cryptic splicing in SHH medulloblastoma. <i>Nature</i> , 2019 , 574, 707-711	50.4	78
4	Recurrent noncoding regulatory mutations in pancreatic ductal adenocarcinoma. <i>Nature Genetics</i> , 2017 , 49, 825-833	36.3	41
3	DriverPower: Combined burden and functional impact tests for cancer driver discovery		4
2	Candidate cancer driver mutations in superenhancers and long-range chromatin interaction networks		5
1	Discovery and characterization of coding and non-coding driver mutations in more than 2,500 whole cancer genomes		12