

# Shimin Shuai

## List of Publications by Citations

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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	Analyses of non-coding somatic drivers in 2,658 cancer whole genomes. <i>Nature</i> , <b>2020</b> , 578, 102-111	50.4	220
14	The U1 spliceosomal RNA is recurrently mutated in multiple cancers. <i>Nature</i> , <b>2019</b> , 574, 712-716	50.4	79
13	MEDU-39. HIGHLY RECURRENT U1 SMALL NUCLEAR RNA HOTSPOT MUTATIONS DRIVE ALTERNATIVE SPLICING IN SONIC HEDGEHOG MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , <b>2019</b> , 21, ii111-ii111		78
12	Recurrent noncoding U1 snRNA mutations drive cryptic splicing in SHH medulloblastoma. <i>Nature</i> , <b>2019</b> , 574, 707-711	50.4	78
11	Cancer LncRNA Census reveals evidence for deep functional conservation of long noncoding RNAs in tumorigenesis. <i>Communications Biology</i> , <b>2020</b> , 3, 56	6.7	77
10	Integrative pathway enrichment analysis of multivariate omics data. <i>Nature Communications</i> , <b>2020</b> , 11, 735	17.4	53
9	Recurrent noncoding regulatory mutations in pancreatic ductal adenocarcinoma. <i>Nature Genetics</i> , <b>2017</b> , 49, 825-833	36.3	41
8	Pathway and network analysis of more than 2500 whole cancer genomes. <i>Nature Communications</i> , <b>2020</b> , 11, 729	17.4	38
7	Candidate Cancer Driver Mutations in Distal Regulatory Elements and Long-Range Chromatin Interaction Networks. <i>Molecular Cell</i> , <b>2020</b> , 77, 1307-1321.e10	17.6	20
6	Combined burden and functional impact tests for cancer driver discovery using DriverPower. <i>Nature Communications</i> , <b>2020</b> , 11, 734	17.4	16
5	Discovery and characterization of coding and non-coding driver mutations in more than 2,500 whole cancer genomes		12
4	Candidate cancer driver mutations in superenhancers and long-range chromatin interaction networks		5
3	DriverPower: Combined burden and functional impact tests for cancer driver discovery		4
2	PanCancer analysis of somatic mutations in repetitive regions reveals recurrent mutations in snRNA U2.. <i>Npj Genomic Medicine</i> , <b>2022</b> , 7, 19	6.2	1
1	Interruption of Klf5 acetylation in basal progenitor cells promotes luminal commitment by activating Notch signaling.. <i>Journal of Genetics and Genomics</i> , <b>2021</b> ,	4	0