

# Britt Adamson

## 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.

20  
papers

4,942  
citations

16  
h-index

20  
g-index

20  
ext. papers

6,608  
ext. citations

34.3  
avg, IF

5.22  
L-index

#	Paper	IF	Citations
20	Enhanced prime editing systems by manipulating cellular determinants of editing outcomes. <i>Cell</i> , <b>2021</b> , 184, 5635-5652.e29	56.2	48
19	Mapping the genetic landscape of DNA double-strand break repair. <i>Cell</i> , <b>2021</b> , 184, 5653-5669.e25	56.2	12
18	Efficient C <sub>T</sub> -to-G <sub>T</sub> base editors developed using CRISPRi screens, target-library analysis, and machine learning. <i>Nature Biotechnology</i> , <b>2021</b> , 39, 1414-1425	44.5	32
17	Pervasive functional translation of noncanonical human open reading frames. <i>Science</i> , <b>2020</b> , 367, 1140-1146	34.6	168
16	Combinatorial single-cell CRISPR screens by direct guide RNA capture and targeted sequencing. <i>Nature Biotechnology</i> , <b>2020</b> , 38, 954-961	44.5	85
15	Prime Editing: Precision Genome Editing by Reverse Transcription. <i>Molecular Cell</i> , <b>2020</b> , 77, 210-212	17.6	11
14	Molecular recording of mammalian embryogenesis. <i>Nature</i> , <b>2019</b> , 570, 77-82	50.4	140
13	Mapping the Genetic Landscape of Human Cells. <i>Cell</i> , <b>2018</b> , 174, 953-967.e22	56.2	136
12	Compact and highly active next-generation libraries for CRISPR-mediated gene repression and activation. <i>ELife</i> , <b>2016</b> , 5,	8.9	343
11	Perturb-Seq: Dissecting Molecular Circuits with Scalable Single-Cell RNA Profiling of Pooled Genetic Screens. <i>Cell</i> , <b>2016</b> , 167, 1853-1866.e17	56.2	675
10	A Multiplexed Single-Cell CRISPR Screening Platform Enables Systematic Dissection of the Unfolded Protein Response. <i>Cell</i> , <b>2016</b> , 167, 1867-1882.e21	56.2	518
9	A Systematic Analysis of Factors Localized to Damaged Chromatin Reveals PARP-Dependent Recruitment of Transcription Factors. <i>Cell Reports</i> , <b>2015</b> , 11, 1486-500	10.6	100
8	Genome-Scale CRISPR-Mediated Control of Gene Repression and Activation. <i>Cell</i> , <b>2014</b> , 159, 647-61	56.2	1556
7	Polyubiquitinated PCNA recruits the ZRANB3 translocase to maintain genomic integrity after replication stress. <i>Molecular Cell</i> , <b>2012</b> , 47, 396-409	17.6	194
6	A genome-wide homologous recombination screen identifies the RNA-binding protein RBMX as a component of the DNA-damage response. <i>Nature Cell Biology</i> , <b>2012</b> , 14, 318-28	23.4	300
5	A bioinformatics method identifies prominent off-targeted transcripts in RNAi screens. <i>Nature Methods</i> , <b>2012</b> , 9, 363-6	21.6	119
4	A chromatin localization screen reveals poly (ADP ribose)-regulated recruitment of the repressive polycomb and NuRD complexes to sites of DNA damage. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2010</b> , 107, 18475-80	11.5	408

3	A genome-wide camptothecin sensitivity screen identifies a mammalian MMS22L-NFKBIL2 complex required for genomic stability. <i>Molecular Cell</i> , <b>2010</b> , 40, 645-57	17.6	81
2	Approaches to maximize sgRNA-barcode coupling in Perturb-seq screens		14
1	Mapping the Genetic Landscape of DNA Double-strand Break Repair		2