

Robert K Bradley

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.

66

papers

7,369

citations

31

h-index

75

g-index

75

ext. papers

9,159

ext. citations

15.8

avg, IF

5.6

L-index

#	Paper	IF	Citations
66	Integrative clinical genomics of advanced prostate cancer. <i>Cell</i> , 2015 , 161, 1215-1228	56.2	1765
65	Evolution of genes and genomes on the Drosophila phylogeny. <i>Nature</i> , 2007 , 450, 203-18	50.4	1586
64	Braveheart, a long noncoding RNA required for cardiovascular lineage commitment. <i>Cell</i> , 2013 , 152, 570-82	38.2	701
63	RNA splicing factors as oncoproteins and tumour suppressors. <i>Nature Reviews Cancer</i> , 2016 , 16, 413-30	31.3	383
62	SRSF2 Mutations Contribute to Myelodysplasia by Mutant-Specific Effects on Exon Recognition. <i>Cancer Cell</i> , 2015 , 27, 617-30	24.3	337
61	Fast statistical alignment. <i>PLoS Computational Biology</i> , 2009 , 5, e1000392	5	252
60	Modulation of splicing catalysis for therapeutic targeting of leukemia with mutations in genes encoding spliceosomal proteins. <i>Nature Medicine</i> , 2016 , 22, 672-8	50.5	227
59	Widespread intron retention diversifies most cancer transcriptomes. <i>Genome Medicine</i> , 2015 , 7, 45	14.4	188
58	U2AF1 mutations alter splice site recognition in hematological malignancies. <i>Genome Research</i> , 2015 , 25, 14-26	9.7	174
57	Binding site turnover produces pervasive quantitative changes in transcription factor binding between closely related Drosophila species. <i>PLoS Biology</i> , 2010 , 8, e1000343	9.7	154
56	Synthetic Lethal and Convergent Biological Effects of Cancer-Associated Spliceosomal Gene Mutations. <i>Cancer Cell</i> , 2018 , 34, 225-241.e8	24.3	96
55	Genome-wide RNAi screens in human brain tumor isolates reveal a novel viability requirement for PHF5A. <i>Genes and Development</i> , 2013 , 27, 1032-45	12.6	88
54	Massive mitochondrial gene transfer in a parasitic flowering plant clade. <i>PLoS Genetics</i> , 2013 , 9, e1003265	12.6	86
53	Spliceosomal disruption of the non-canonical BAF complex in cancer. <i>Nature</i> , 2019 , 574, 432-436	50.4	80
52	Coordinated alterations in RNA splicing and epigenetic regulation drive leukaemogenesis. <i>Nature</i> , 2019 , 574, 273-277	50.4	76
51	Alternative splicing of RNA triplets is often regulated and accelerates proteome evolution. <i>PLoS Biology</i> , 2012 , 10, e1001229	9.7	74
50	A feedback loop between nonsense-mediated decay and the retrogene DUX4 in facioscapulohumeral muscular dystrophy. <i>ELife</i> , 2015 , 4,	8.9	73

49	Horizontal transfer of expressed genes in a parasitic flowering plant. <i>BMC Genomics</i> , 2012 , 13, 227	4.5	73
48	Spliceosomal gene mutations in myelodysplasia: molecular links to clonal abnormalities of hematopoiesis. <i>Genes and Development</i> , 2016 , 30, 989-1001	12.6	73
47	Wild-Type U2AF1 Antagonizes the Splicing Program Characteristic of U2AF1-Mutant Tumors and Is Required for Cell Survival. <i>PLoS Genetics</i> , 2016 , 12, e1006384	6	57
46	RNA components of the spliceosome regulate tissue- and cancer-specific alternative splicing. <i>Genome Research</i> , 2019 , 29, 1591-1604	9.7	53
45	Sample processing obscures cancer-specific alterations in leukemic transcriptomes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 16802-7	11.5	50
44	Most human introns are recognized via multiple and tissue-specific branchpoints. <i>Genes and Development</i> , 2018 , 32, 577-591	12.6	49
43	Model systems of DUX4 expression recapitulate the transcriptional profile of FSHD cells. <i>Human Molecular Genetics</i> , 2016 , 25, 4419-4431	5.6	49
42	Exon Junction Complex Shapes the Transcriptome by Repressing Recursive Splicing. <i>Molecular Cell</i> , 2018 , 72, 496-509.e9	17.6	43
41	XRate: a fast prototyping, training and annotation tool for phylo-grammars. <i>BMC Bioinformatics</i> , 2006 , 7, 428	3.6	40
40	Altered RNA Splicing by Mutant p53 Activates Oncogenic RAS Signaling in Pancreatic Cancer. <i>Cancer Cell</i> , 2020 , 38, 198-211.e8	24.3	38
39	The TAGteam motif facilitates binding of 21 sequence-specific transcription factors in the Drosophila embryo. <i>Genome Research</i> , 2012 , 22, 656-65	9.7	38
38	Impaired hematopoiesis and leukemia development in mice with a conditional knock-in allele of a mutant splicing factor gene. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018 , 115, E10437-E10446	11.5	38
37	DUX4 Suppresses MHC Class I to Promote Cancer Immune Evasion and Resistance to Checkpoint Blockade. <i>Developmental Cell</i> , 2019 , 50, 658-671.e7	10.2	37
36	The RNA Surveillance Factor UPF1 Represses Myogenesis via Its E3 Ubiquitin Ligase Activity. <i>Molecular Cell</i> , 2017 , 67, 239-251.e6	17.6	33
35	Specific alignment of structured RNA: stochastic grammars and sequence annealing. <i>Bioinformatics</i> , 2008 , 24, 2677-83	7.2	31
34	Dissecting the Contributions of Cooperating Gene Mutations to Cancer Phenotypes and Drug Responses with Patient-Derived iPSCs. <i>Stem Cell Reports</i> , 2018 , 10, 1610-1624	8	30
33	RNA isoform screens uncover the essentiality and tumor-suppressor activity of ultraconserved poison exons. <i>Nature Genetics</i> , 2020 , 52, 84-94	36.3	30
32	Transducers: an emerging probabilistic framework for modeling indels on trees. <i>Bioinformatics</i> , 2007 , 23, 3258-62	7.2	29

31	Pharmacologic modulation of RNA splicing enhances anti-tumor immunity. <i>Cell</i> , 2021 , 184, 4032-4047.e316.2	36.2	25
30	Translational plasticity facilitates the accumulation of nonsense genetic variants in the human population. <i>Genome Research</i> , 2016 , 26, 1639-1650	9.7	24
29	Single-cell genomics reveals the genetic and molecular bases for escape from mutational epistasis in myeloid neoplasms. <i>Blood</i> , 2020 , 136, 1477-1486	2.2	21
28	Recurrent SRSF2 mutations in MDS affect both splicing and NMD. <i>Genes and Development</i> , 2020 , 34, 413-427	12.7	19
27	Tools for simulating evolution of aligned genomic regions with integrated parameter estimation. <i>Genome Biology</i> , 2008 , 9, R147	18.3	18
26	Transgenic mice expressing tunable levels of DUX4 develop characteristic facioscapulohumeral muscular dystrophy-like pathophysiology ranging in severity. <i>Skeletal Muscle</i> , 2020 , 10, 8	5.1	18
25	Minor intron retention drives clonal hematopoietic disorders and diverse cancer predisposition. <i>Nature Genetics</i> , 2021 , 53, 707-718	36.3	13
24	Quantitative proteomics reveals key roles for post-transcriptional gene regulation in the molecular pathology of facioscapulohumeral muscular dystrophy. <i>ELife</i> , 2019 , 8,	8.9	12
23	Convergent organization of aberrant MYB complex controls oncogenic gene expression in acute myeloid leukemia. <i>ELife</i> , 2021 , 10,	8.9	12
22	Short H2A histone variants are expressed in cancer. <i>Nature Communications</i> , 2021 , 12, 490	17.4	12
21	Evolutionary modeling and prediction of non-coding RNAs in Drosophila. <i>PLoS ONE</i> , 2009 , 4, e6478	3.7	10
20	Rare and private spliceosomal gene mutations drive partial, complete, and dual phenocopies of hotspot alterations. <i>Blood</i> , 2020 , 135, 1032-1043	2.2	8
19	Characterization of neoantigen-specific T cells in cancer resistant to immune checkpoint therapies. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021 , 118,	11.5	8
18	Discovery of synthetic lethal and tumor suppressor paralog pairs in the human genome. <i>Cell Reports</i> , 2021 , 36, 109597	10.6	8
17	Evolutionary triplet models of structured RNA. <i>PLoS Computational Biology</i> , 2009 , 5, e1000483	5	4
16	Degenerate eigenvalues for Hamiltonians with no obvious symmetries. <i>Advances in Theoretical and Mathematical Physics</i> , 2005 , 9, 593-602	1.3	4
15	Integrative oncogene-dependency mapping identifies RIT1 vulnerabilities and synergies in lung cancer. <i>Nature Communications</i> , 2021 , 12, 4789	17.4	4
14	RNA components of the spliceosome regulate tissue- and cancer-specific alternative splicing		3

13	Wild-type U2AF1 antagonizes the splicing program characteristic of U2AF1-mutant tumors and is required for cell survival		3
12	Coordinated mis-splicing of TMEM14C and ABCB7 causes ring sideroblast formation in SF3B1-mutant myelodysplastic syndrome. <i>Blood</i> , 2021 ,	2.2	2
11	The origins and consequences of variants in pancreatic adenosquamous carcinoma. <i>ELife</i> , 2021 , 10,	8.9	2
10	Synthetic introns enable splicing factor mutation-dependent targeting of cancer cells.. <i>Nature Biotechnology</i> , 2022 ,	44.5	2
9	Congenital myotonic dystrophy-an RNA-mediated disease across a developmental continuum. <i>Genes and Development</i> , 2017 , 31, 1067-1068	12.6	1
8	Nonsense-mediated mRNA decay uses complementary mechanisms to suppress mRNA and protein accumulation. <i>Life Science Alliance</i> , 2022 , 5,	5.8	1
7	Coordinated Mis-Splicing of Multiple Mitochondrial Iron Metabolism Genes Causes Ring Sideroblast Formation in SF3B1-Mutant MDS. <i>Blood</i> , 2020 , 136, 4-4	2.2	1
6	Discovery of synthetic lethal and tumor suppressive paralog pairs in the human genome		1
5	Transgenic mice expressing tunable levels of DUX4 develop characteristic facioscapulohumeral muscular dystrophy-like pathophysiology ranging in severity		1
4	Nonsense-mediated mRNA decay utilizes complementary mechanisms to suppress mRNA and protein accumulation		1
3	ZRSR2 Mutation Induced Minor Intron Retention Drives MDS and Diverse Cancer Predisposition Via Aberrant Splicing of LZTR1. <i>Blood</i> , 2020 , 136, 10-11	2.2	0
2	is mutated in clonal hematopoiesis and myelodysplastic syndromes and impacts RNA splicing. <i>Blood Cancer Discovery</i> , 2021 , 2, 500-517	7	0
1	Probing Aberrant Splicing in a Novel Model of SF3B1-Mutant Myelodysplastic Syndromes. <i>Blood</i> , 2019 , 134, 1706-1706	2.2	