Matthew C Canver

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.

48 47 2,797 20 g-index h-index citations papers 48 11.7 3,707 4.92 avg, IF L-index ext. citations ext. papers

#	Paper	IF	Citations
47	Technologies and Computational Analysis Strategies for CRISPR Applications. <i>Molecular Cell</i> , 2020 , 79, 11-29	17.6	7
46	Accelerated thrombin times are associated with thrombotic risk. <i>American Journal of Hematology</i> , 2020 , 95, E113-E114	7.1	
45	A saturating mutagenesis CRISPR-Cas9-mediated functional genomic screen identifies and regulatory elements of in murine ESCs. <i>Journal of Biological Chemistry</i> , 2020 , 295, 15797-15809	5.4	2
44	CRISPRitz: rapid, high-throughput and variant-aware in silico off-target site identification for CRISPR genome editing. <i>Bioinformatics</i> , 2020 , 36, 2001-2008	7.2	15
43	Thawed solvent/detergent-treated plasma demonstrates comparable clinical efficacy to thawed plasma. <i>Transfusion</i> , 2020 , 60, 1940-1949	2.9	O
42	Visual evidence of a hemolytic transfusion reaction identified by blood bank testing after emergency blood transfusion. <i>Transfusion</i> , 2019 , 59, 3301-3302	2.9	
41	Performance of Five Commercial Identification Platforms for Identification of Staphylococcus delphini. <i>Journal of Clinical Microbiology</i> , 2019 , 57,	9.7	4
40	Improved Performance of a Rapid Immunochromatographic Assay for Detection of PBP2a in Non-Staphylococcus aureus Staphylococcal Species. <i>Journal of Clinical Microbiology</i> , 2019 , 57,	9.7	7
39	TAF5L and TAF6L Maintain Self-Renewal of Embryonic Stem Cells via the MYC Regulatory Network. <i>Molecular Cell</i> , 2019 , 74, 1148-1163.e7	17.6	19
38	CRISPResso2 provides accurate and rapid genome editing sequence analysis. <i>Nature Biotechnology</i> , 2019 , 37, 224-226	44.5	326
37	Rational targeting of a NuRD subcomplex guided by comprehensive in situ mutagenesis. <i>Nature Genetics</i> , 2019 , 51, 1149-1159	36.3	44
36	DrugThatGene: integrative analysis to streamline the identification of druggable genes, pathways and protein complexes from CRISPR screens. <i>Bioinformatics</i> , 2019 , 35, 1981-1984	7.2	3
35	Genome-wide CRISPR-Cas9 Screen Identifies Leukemia-Specific Dependence on a Pre-mRNA Metabolic Pathway Regulated by DCPS. <i>Cancer Cell</i> , 2018 , 33, 386-400.e5	24.3	57
34	Impact of Genetic Variation on CRISPR-Cas Targeting. CRISPR Journal, 2018, 1, 159-170	2.5	16
33	Integrated design, execution, and analysis of arrayed and pooled CRISPR genome-editing experiments. <i>Nature Protocols</i> , 2018 , 13, 946-986	18.8	42
32	Rational Targeting of a NuRD Sub-Complex for Fetal Hemoglobin Induction Following Comprehensive in Situ Mutagenesis. <i>Blood</i> , 2018 , 132, 2342-2342	2.2	
31	CRISPR-SURF: discovering regulatory elements by deconvolution of CRISPR tiling screen data. Nature Methods, 2018, 15, 992-993	21.6	17

(2014-2018)

30	CRISPRO: identification of functional protein coding sequences based on genome editing dense mutagenesis. <i>Genome Biology</i> , 2018 , 19, 169	18.3	20
29	Downregulation of Endothelin Receptor B Contributes to Defective B Cell Lymphopoiesis in Trisomy 21 Pluripotent Stem Cells. <i>Scientific Reports</i> , 2018 , 8, 8001	4.9	10
28	Variant-aware saturating mutagenesis using multiple Cas9 nucleases identifies regulatory elements at trait-associated loci. <i>Nature Genetics</i> , 2017 , 49, 625-634	36.3	73
27	Functional interrogation of non-coding DNA through CRISPR genome editing. <i>Methods</i> , 2017 , 121-122, 118-129	4.6	19
26	High-Throughput Approaches to Pinpoint Function within the Noncoding Genome. <i>Molecular Cell</i> , 2017 , 68, 44-59	17.6	37
25	Human genetic variation alters CRISPR-Cas9 on- and off-targeting specificity at therapeutically implicated loci. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, E11257-E11266	11.5	66
24	Genome-Wide CRISPR/Cas9 Screen Reveals That the Dcps Scavenger Decapping Enzyme Is Essential for AML Cell Survival. <i>Blood</i> , 2017 , 130, 782-782	2.2	
23	Interferon-Bignaling promotes embryonic HSC maturation. <i>Blood</i> , 2016 , 128, 204-16	2.2	28
22	Transcription factors LRF and BCL11A independently repress expression of fetal hemoglobin. <i>Science</i> , 2016 , 351, 285-9	33.3	187
21	Analyzing CRISPR genome-editing experiments with CRISPResso. <i>Nature Biotechnology</i> , 2016 , 34, 695-7	44.5	286
21		44·5 2.2	286
	Customizing the genome as therapy for the Ehemoglobinopathies. <i>Blood</i> , 2016 , 127, 2536-45 miRNA-embedded shRNAs for Lineage-specific BCL11A Knockdown and Hemoglobin F Induction.		38
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20 19 18	Customizing the genome as therapy for the Ehemoglobinopathies. <i>Blood</i> , 2016 , 127, 2536-45 miRNA-embedded shRNAs for Lineage-specific BCL11A Knockdown and Hemoglobin F Induction. <i>Molecular Therapy</i> , 2015 , 23, 1465-74 BCL11A enhancer dissection by Cas9-mediated in situ saturating mutagenesis. <i>Nature</i> , 2015 , 527, 192-7 PRC2 Is Required to Maintain Expression of the Maternal Gtl2-Rian-Mirg Locus by Preventing De Novo DNA Methylation in Mouse Embryonic Stem Cells. <i>Cell Reports</i> , 2015 , 12, 1456-70	2.2 11.7 50.4	38 82 528
20 19 18	Customizing the genome as therapy for the Ehemoglobinopathies. <i>Blood</i> , 2016 , 127, 2536-45 miRNA-embedded shRNAs for Lineage-specific BCL11A Knockdown and Hemoglobin F Induction. <i>Molecular Therapy</i> , 2015 , 23, 1465-74 BCL11A enhancer dissection by Cas9-mediated in situ saturating mutagenesis. <i>Nature</i> , 2015 , 527, 192-7 PRC2 Is Required to Maintain Expression of the Maternal Gtl2-Rian-Mirg Locus by Preventing De Novo DNA Methylation in Mouse Embryonic Stem Cells. <i>Cell Reports</i> , 2015 , 12, 1456-70 EHMT1 and EHMT2 inhibition induces fetal hemoglobin expression. <i>Blood</i> , 2015 , 126, 1930-9	2.2 11.7 50.4	38 82 528 46
20 19 18 17	Customizing the genome as therapy for the Ehemoglobinopathies. <i>Blood</i> , 2016 , 127, 2536-45 miRNA-embedded shRNAs for Lineage-specific BCL11A Knockdown and Hemoglobin F Induction. <i>Molecular Therapy</i> , 2015 , 23, 1465-74 BCL11A enhancer dissection by Cas9-mediated in situ saturating mutagenesis. <i>Nature</i> , 2015 , 527, 192-7 PRC2 Is Required to Maintain Expression of the Maternal Gtl2-Rian-Mirg Locus by Preventing De Novo DNA Methylation in Mouse Embryonic Stem Cells. <i>Cell Reports</i> , 2015 , 12, 1456-70 EHMT1 and EHMT2 inhibition induces fetal hemoglobin expression. <i>Blood</i> , 2015 , 126, 1930-9 Generation of genomic deletions in mammalian cell lines via CRISPR/Cas9. <i>Journal of Visualized Experiments</i> , 2015 , e52118	2.2 11.7 50.4 10.6	38 82 528 46

12	Characterization of genomic deletion efficiency mediated by clustered regularly interspaced short palindromic repeats (CRISPR)/Cas9 nuclease system in mammalian cells. <i>Journal of Biological Chemistry</i> , 2014 , 289, 21312-24	5.4	236
11	Optimization of Bcl11a Knockdown By miRNA Scaffold Embedded Shrnas Leading to Enhanced Induction of Fetal Hemoglobin in Erythroid Cells for the Treatment of Beta-Hemoglobinopathies. <i>Blood</i> , 2014 , 124, 2150-2150	2.2	1
10	An erythroid enhancer of BCL11A subject to genetic variation determines fetal hemoglobin level. <i>Science</i> , 2013 , 342, 253-7	33.3	400
9	Looking Forward [Perspectives on Graduate Life]. <i>IEEE Pulse</i> , 2013 , 4, 11-65	0.7	
8	Independence Day [Perspectives on Graduate Life]. IEEE Pulse, 2013, 4, 12-12	0.7	
7	Making the big decision [Perspectives on Graduate Life]. <i>IEEE Pulse</i> , 2013 , 4, 9-10	0.7	
6	Another One in the Books [Perspectives on Graduate Life]. IEEE Pulse, 2013, 4, 8-62	0.7	
5	Fine-Mapping and Genome Editing Reveal An Essential Erythroid Enhancer At The HbF-Associated BCL11A Locus. <i>Blood</i> , 2013 , 122, 437-437	2.2	1
4	Graduate school: the problem of choice. <i>IEEE Pulse</i> , 2012 , 3, 8, 10	0.7	1
3	Bioengineering Training in Medical Education [Perspectives on Graduate Life]. <i>IEEE Pulse</i> , 2010 , 1, 9-11	0.7	
2	Histamine H1 and H2 receptor-mediated vasoreactivity of human internal thoracic and radial arteries. <i>Surgery</i> , 2004 , 136, 458-63	3.6	8
1	Analysis and comparison of genome editing using CRISPResso2		4