

# David R Liu

## List of Publications by Year in descending order

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127  
papers

36,177  
citations

11235

73  
h-index

12940

136  
g-index

149  
all docs

149  
docs citations

149  
times ranked

26220  
citing authors

#	ARTICLE	IF	CITATIONS
1	Engineered pegRNAs improve prime editing efficiency. <i>Nature Biotechnology</i> , 2022, 40, 402-410.	9.4	293
2	Disruption of HIV-1 co-receptors CCR5 and CXCR4 in primary human T cells and hematopoietic stem and progenitor cells using base editing. <i>Molecular Therapy</i> , 2022, 30, 130-144.	3.7	23
3	Engineered virus-like particles for efficient in vivo delivery of therapeutic proteins. <i>Cell</i> , 2022, 185, 250-265.e16.	13.5	251
4	Programmable deletion, replacement, integration and inversion of large DNA sequences with twin prime editing. <i>Nature Biotechnology</i> , 2022, 40, 731-740.	9.4	230
5	CRISPR-free base editors with enhanced activity and expanded targeting scope in mitochondrial and nuclear DNA. <i>Nature Biotechnology</i> , 2022, 40, 1378-1387.	9.4	81
6	In vivo base editing rescues cone photoreceptors in a mouse model of early-onset inherited retinal degeneration. <i>Nature Communications</i> , 2022, 13, 1830.	5.8	42
7	Prioritization of autoimmune disease-associated genetic variants that perturb regulatory element activity in T cells. <i>Nature Genetics</i> , 2022, 54, 603-612.	9.4	15
8	Therapeutic in vivo delivery of gene editing agents. <i>Cell</i> , 2022, 185, 2806-2827.	13.5	131
9	Restoration of visual function in adult mice with an inherited retinal disease via adenine base editing. <i>Nature Biomedical Engineering</i> , 2021, 5, 169-178.	11.6	90
10	Precision genome editing using cytosine and adenine base editors in mammalian cells. <i>Nature Protocols</i> , 2021, 16, 1089-1128.	5.5	90
11	Laboratory evolution of a sortase enzyme that modifies amyloid- $\beta$ protein. <i>Nature Chemical Biology</i> , 2021, 17, 317-325.	3.9	34
12	In vivo base editing rescues Hutchinson Gilford progeria syndrome in mice. <i>Nature</i> , 2021, 589, 608-614.	13.7	275
13	Phage-assisted evolution of botulinum neurotoxin proteases with reprogrammed specificity. <i>Science</i> , 2021, 371, 803-810.	6.0	46
14	Massively parallel assessment of human variants with base editor screens. <i>Cell</i> , 2021, 184, 1064-1080.e20.	13.5	175
15	Prime editing in mice reveals the essentiality of a single base in driving tissue-specific gene expression. <i>Genome Biology</i> , 2021, 22, 83.	3.8	62
16	The NIH Somatic Cell Genome Editing program. <i>Nature</i> , 2021, 592, 195-204.	13.7	84
17	Mechanisms of angiogenic incompetence in Hutchinson Gilford progeria via downregulation of endothelial NOS. <i>Aging Cell</i> , 2021, 20, e13388.	3.0	11
18	Base editor treats progeria in mice. <i>Nature</i> , 2021, , .	13.7	4

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19	Base editing of haematopoietic stem cells rescues sickle cell disease in mice. <i>Nature</i> , 2021, 595, 295-302.	13.7	175
20	Efficient C-to-G base editors developed using CRISPRi screens, target-library analysis, and machine learning. <i>Nature Biotechnology</i> , 2021, 39, 1414-1425.	9.4	118
21	A rechargeable anti-thrombotic coating for blood-contacting devices. <i>Biomaterials</i> , 2021, 276, 121011.	5.7	8
22	In vivo somatic cell base editing and prime editing. <i>Molecular Therapy</i> , 2021, 29, 3107-3124.	3.7	87
23	Functional correction of CFTR mutations in human airway epithelial cells using adenine base editors. <i>Nucleic Acids Research</i> , 2021, 49, 10558-10572.	6.5	25
24	Enhanced prime editing systems by manipulating cellular determinants of editing outcomes. <i>Cell</i> , 2021, 184, 5635-5652.e29.	13.5	332
25	Reconstruction of evolving gene variants and fitness from short sequencing reads. <i>Nature Chemical Biology</i> , 2021, 17, 1188-1198.	3.9	8
26	Disulfide-compatible phage-assisted continuous evolution in the periplasmic space. <i>Nature Communications</i> , 2021, 12, 5959.	5.8	13
27	Adenine base editing in an adult mouse model of tyrosinaemia. <i>Nature Biomedical Engineering</i> , 2020, 4, 125-130.	11.6	136
28	Base Editor Correction of COL7A1 in Recessive Dystrophic Epidermolysis Bullosa Patient-Derived Fibroblasts and iPSCs. <i>Journal of Investigative Dermatology</i> , 2020, 140, 338-347.e5.	0.3	69
29	Phage-assisted continuous and non-continuous evolution. <i>Nature Protocols</i> , 2020, 15, 4101-4127.	5.5	42
30	DNA capture by a CRISPR-Cas9-guided adenine base editor. <i>Science</i> , 2020, 369, 566-571.	6.0	114
31	Multimodal small-molecule screening for human prion protein binders. <i>Journal of Biological Chemistry</i> , 2020, 295, 13516-13531.	1.6	14
32	Glucose Response by Stem Cell-Derived $\beta^2$ Cells In Vitro Is Inhibited by a Bottleneck in Glycolysis. <i>Cell Reports</i> , 2020, 31, 107623.	2.9	72
33	Genome editing with CRISPR-Cas nucleases, base editors, transposases and prime editors. <i>Nature Biotechnology</i> , 2020, 38, 824-844.	9.4	1,277
34	In vivo base editing restores sensory transduction and transiently improves auditory function in a mouse model of recessive deafness. <i>Science Translational Medicine</i> , 2020, 12, .	5.8	114
35	Determinants of Base Editing Outcomes from Target Library Analysis and Machine Learning. <i>Cell</i> , 2020, 182, 463-480.e30.	13.5	166
36	Phage-assisted evolution of an adenine base editor with improved Cas domain compatibility and activity. <i>Nature Biotechnology</i> , 2020, 38, 883-891.	9.4	502

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37	Prime genome editing in rice and wheat. <i>Nature Biotechnology</i> , 2020, 38, 582-585.	9.4	544
38	Programmable m6A modification of cellular RNAs with a Cas13-directed methyltransferase. <i>Nature Biotechnology</i> , 2020, 38, 1431-1440.	9.4	173
39	A bacterial cytidine deaminase toxin enables CRISPR-free mitochondrial base editing. <i>Nature</i> , 2020, 583, 631-637.	13.7	409
40	Continuous evolution of SpCas9 variants compatible with non-G PAMs. <i>Nature Biotechnology</i> , 2020, 38, 471-481.	9.4	234
41	Evaluation and minimization of Cas9-independent off-target DNA editing by cytosine base editors. <i>Nature Biotechnology</i> , 2020, 38, 620-628.	9.4	272
42	High-throughput analysis of the activities of xCas9, SpCas9-NG and SpCas9 at matched and mismatched target sequences in human cells. <i>Nature Biomedical Engineering</i> , 2020, 4, 111-124.	11.6	98
43	Cytosine and adenine base editing of the brain, liver, retina, heart and skeletal muscle of mice via adeno-associated viruses. <i>Nature Biomedical Engineering</i> , 2020, 4, 97-110.	11.6	293
44	Chemical modifications of adenine base editor mRNA and guide RNA expand its application scope. <i>Nature Communications</i> , 2020, 11, 1979.	5.8	66
45	The developing toolkit of continuous directed evolution. <i>Nature Chemical Biology</i> , 2020, 16, 610-619.	3.9	80
46	Adenosine Base Editing of $\beta$ -Globin Promoters Induces Fetal Hemoglobin and Inhibit Erythroid Sickling. <i>Blood</i> , 2020, 136, 21-22.	0.6	8
47	Continuous evolution of base editors with expanded target compatibility and improved activity. <i>Nature Biotechnology</i> , 2019, 37, 1070-1079.	9.4	215
48	An anionic human protein mediates cationic liposome delivery of genome editing proteins into mammalian cells. <i>Nature Communications</i> , 2019, 10, 2905.	5.8	20
49	Search-and-replace genome editing without double-strand breaks or donor DNA. <i>Nature</i> , 2019, 576, 149-157.	13.7	2,662
50	Analysis and minimization of cellular RNA editing by DNA adenine base editors. <i>Science Advances</i> , 2019, 5, eaax5717.	4.7	206
51	Development of hRad51-Cas9 nickase fusions that mediate HDR without double-stranded breaks. <i>Nature Communications</i> , 2019, 10, 2212.	5.8	76
52	Circularly permuted and PAM-modified Cas9 variants broaden the targeting scope of base editors. <i>Nature Biotechnology</i> , 2019, 37, 626-631.	9.4	207
53	Substrate-selective inhibitors that reprogram the activity of insulin-degrading enzyme. <i>Nature Chemical Biology</i> , 2019, 15, 565-574.	3.9	36
54	A High-Throughput Platform to Identify Small-Molecule Inhibitors of CRISPR-Cas9. <i>Cell</i> , 2019, 177, 1067-1079.e19.	13.5	133

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55	High-resolution specificity profiling and off-target prediction for site-specific DNA recombinases. <i>Nature Communications</i> , 2019, 10, 1937.	5.8	22
56	Phage-Assisted Evolution of <i>Bacillus methanolicus</i> Methanol Dehydrogenase 2. <i>ACS Synthetic Biology</i> , 2019, 8, 796-806.	1.9	61
57	Simultaneous targeting of linked loci in mouse embryos using base editing. <i>Scientific Reports</i> , 2019, 9, 1662.	1.6	12
58	Side chain determinants of biopolymer function during selection and replication. <i>Nature Chemical Biology</i> , 2019, 15, 419-426.	3.9	17
59	CRISPResso2 provides accurate and rapid genome editing sequence analysis. <i>Nature Biotechnology</i> , 2019, 37, 224-226.	9.4	891
60	CREB5 Promotes Resistance to Androgen-Receptor Antagonists and Androgen Deprivation in Prostate Cancer. <i>Cell Reports</i> , 2019, 29, 2355-2370.e6.	2.9	45
61	Evolved Cas9 variants with broad PAM compatibility and high DNA specificity. <i>Nature</i> , 2018, 556, 57-63.	13.7	1,195
62	Evolution of sequence-defined highly functionalized nucleic acid polymers. <i>Nature Chemistry</i> , 2018, 10, 420-427.	6.6	83
63	Rewritable multi-event analog recording in bacterial and mammalian cells. <i>Science</i> , 2018, 360, .	6.0	193
64	Treatment of autosomal dominant hearing loss by in vivo delivery of genome editing agents. <i>Nature</i> , 2018, 553, 217-221.	13.7	412
65	Editing the Genome Without Double-Stranded DNA Breaks. <i>ACS Chemical Biology</i> , 2018, 13, 383-388.	1.6	89
66	Development of a formaldehyde biosensor with application to synthetic methylotrophy. <i>Biotechnology and Bioengineering</i> , 2018, 115, 206-215.	1.7	44
67	Targeting fidelity of adenine and cytosine base editors in mouse embryos. <i>Nature Communications</i> , 2018, 9, 4804.	5.8	72
68	Predictable and precise template-free CRISPR editing of pathogenic variants. <i>Nature</i> , 2018, 563, 646-651.	13.7	414
69	Base editing: precision chemistry on the genome and transcriptome of living cells. <i>Nature Reviews Genetics</i> , 2018, 19, 770-788.	7.7	1,072
70	One-Pot Dual Labeling of IgG 1 and Preparation of C-to-C Fusion Proteins Through a Combination of Sortase A and Butelase 1. <i>Bioconjugate Chemistry</i> , 2018, 29, 3245-3249.	1.8	72
71	Improving cytidine and adenine base editors by expression optimization and ancestral reconstruction. <i>Nature Biotechnology</i> , 2018, 36, 843-846.	9.4	644
72	Ensemble cryoEM elucidates the mechanism of insulin capture and degradation by human insulin degrading enzyme. <i>ELife</i> , 2018, 7, .	2.8	45

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73	Continuous directed evolution of proteins with improved soluble expression. <i>Nature Chemical Biology</i> , 2018, 14, 972-980.	3.9	71
74	Green fluorescent proteins engineered for cartilage-targeted drug delivery: Insights for transport into highly charged avascular tissues. <i>Biomaterials</i> , 2018, 183, 218-233.	5.7	50
75	In vivo base editing of post-mitotic sensory cells. <i>Nature Communications</i> , 2018, 9, 2184.	5.8	166
76	Increasing the genome-targeting scope and precision of base editing with engineered Cas9-cytidine deaminase fusions. <i>Nature Biotechnology</i> , 2017, 35, 371-376.	9.4	609
77	Improving the DNA specificity and applicability of base editing through protein engineering and protein delivery. <i>Nature Communications</i> , 2017, 8, 15790.	5.8	343
78	Programmable base editing of Aâ€¢T to Gâ€¢C in genomic DNA without DNA cleavage. <i>Nature</i> , 2017, 551, 464-471.	13.7	2,807
79	Crystal structures reveal an elusive functional domain of pyrrolysyl-tRNA synthetase. <i>Nature Chemical Biology</i> , 2017, 13, 1261-1266.	3.9	73
80	Phage-assisted continuous evolution of proteases with altered substrate specificity. <i>Nature Communications</i> , 2017, 8, 956.	5.8	85
81	Continuous directed evolution of aminoacyl-tRNA synthetases. <i>Nature Chemical Biology</i> , 2017, 13, 1253-1260.	3.9	185
82	Improved base excision repair inhibition and bacteriophage Mu Gam protein yields C:G-to-T:A base editors with higher efficiency and product purity. <i>Science Advances</i> , 2017, 3, eaao4774.	4.7	582
83	Discovery of a Covalent Kinase Inhibitor from a DNA-Encoded Small-Molecule Library Ã— Protein Library Selection. <i>Journal of the American Chemical Society</i> , 2017, 139, 10192-10195.	6.6	67
84	Aptazyme-embedded guide RNAs enable ligand-responsive genome editing and transcriptional activation. <i>Nature Communications</i> , 2017, 8, 15939.	5.8	169
85	CRISPR-Based Technologies for the Manipulation of Eukaryotic Genomes. <i>Cell</i> , 2017, 168, 20-36.	13.5	783
86	Sequence Determinants of Intracellular Phase Separation by Complex Coacervation of a Disordered Protein. <i>Molecular Cell</i> , 2016, 63, 72-85.	4.5	622
87	Efficient delivery of genome-editing proteins using bioreducible lipid nanoparticles. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, 2868-2873.	3.3	495
88	Programmable editing of a target base in genomic DNA without double-stranded DNA cleavage. <i>Nature</i> , 2016, 533, 420-424.	13.7	3,662
89	Continuous evolution of <i>Bacillus thuringiensis</i> toxins overcomes insect resistance. <i>Nature</i> , 2016, 533, 58-63.	13.7	159
90	Structural and Biochemical Basis for Intracellular Kinase Inhibition by Src-specific Peptidic Macrocycles. <i>Cell Chemical Biology</i> , 2016, 23, 1103-1112.	2.5	12

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91	A programmable Cas9-serine recombinase fusion protein that operates on DNA sequences in mammalian cells. <i>Nucleic Acids Research</i> , 2016, 44, gkw707.	6.5	46
92	In situ regeneration of bioactive coatings enabled by an evolved <i>Staphylococcus aureus</i> sortase A. <i>Nature Communications</i> , 2016, 7, 11140.	5.8	33
93	Chemical Biology Approaches to Genome Editing: Understanding, Controlling, and Delivering Programmable Nucleases. <i>Cell Chemical Biology</i> , 2016, 23, 57-73.	2.5	42
94	Analytical Devices Based on Direct Synthesis of DNA on Paper. <i>Analytical Chemistry</i> , 2016, 88, 725-731.	3.2	38
95	Methods for the directed evolution of proteins. <i>Nature Reviews Genetics</i> , 2015, 16, 379-394.	7.7	699
96	Novel selection methods for DNA-encoded chemical libraries. <i>Current Opinion in Chemical Biology</i> , 2015, 26, 55-61.	2.8	54
97	Small molecule-triggered Cas9 protein with improved genome-editing specificity. <i>Nature Chemical Biology</i> , 2015, 11, 316-318.	3.9	364
98	Continuous directed evolution of DNA-binding proteins to improve TALEN specificity. <i>Nature Methods</i> , 2015, 12, 939-942.	9.0	88
99	Development of potent in vivo mutagenesis plasmids with broad mutational spectra. <i>Nature Communications</i> , 2015, 6, 8425.	5.8	138
100	Discovery and Characterization of a Peptide That Enhances Endosomal Escape of Delivered Proteins in Vitro and in Vivo. <i>Journal of the American Chemical Society</i> , 2015, 137, 14084-14093.	6.6	109
101	In vivo continuous directed evolution. <i>Current Opinion in Chemical Biology</i> , 2015, 24, 1-10.	2.8	65
102	Cationic lipid-mediated delivery of proteins enables efficient protein-based genome editing in vitro and in vivo. <i>Nature Biotechnology</i> , 2015, 33, 73-80.	9.4	1,180
103	A naturally occurring, noncanonical GTP aptamer made of simple tandem repeats. <i>RNA Biology</i> , 2014, 11, 682-692.	1.5	9
104	Immobilization of Actively Thromboresistant Assemblies on Sterile Blood-Contacting Surfaces. <i>Advanced Healthcare Materials</i> , 2014, 3, 30-35.	3.9	28
105	Determining the Specificities of TALENs, Cas9, and Other Genome-Editing Enzymes. <i>Methods in Enzymology</i> , 2014, 546, 47-78.	0.4	59
106	Broad specificity profiling of TALENs results in engineered nucleases with improved DNA-cleavage specificity. <i>Nature Methods</i> , 2014, 11, 429-435.	9.0	182
107	Fusion of catalytically inactive Cas9 to FokI nuclease improves the specificity of genome modification. <i>Nature Biotechnology</i> , 2014, 32, 577-582.	9.4	740
108	Anti-diabetic activity of insulin-degrading enzyme inhibitors mediated by multiple hormones. <i>Nature</i> , 2014, 511, 94-98.	13.7	207

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109	Negative selection and stringency modulation in phage-assisted continuous evolution. <i>Nature Chemical Biology</i> , 2014, 10, 216-222.	3.9	129
110	Electrophilic activity-based RNA probes reveal a self-alkylating RNA for RNA labeling. <i>Nature Chemical Biology</i> , 2014, 10, 1049-1054.	3.9	30
111	A system for the continuous directed evolution of proteases rapidly reveals drug-resistance mutations. <i>Nature Communications</i> , 2014, 5, 5352.	5.8	82
112	A DNA-based molecular probe for optically reporting cellular traction forces. <i>Nature Methods</i> , 2014, 11, 1229-1232.	9.0	171
113	Reprogramming the specificity of sortase enzymes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 13343-13348.	3.3	151
114	Identification of Ligand-Target Pairs from Combined Libraries of Small Molecules and Unpurified Protein Targets in Cell Lysates. <i>Journal of the American Chemical Society</i> , 2014, 136, 3264-3270.	6.6	74
115	High-throughput profiling of off-target DNA cleavage reveals RNA-programmed Cas9 nuclease specificity. <i>Nature Biotechnology</i> , 2013, 31, 839-843.	9.4	1,303
116	Experimental interrogation of the path dependence and stochasticity of protein evolution using phage-assisted continuous evolution. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 9007-9012.	3.3	92
117	DNA Ligase-Mediated Translation of DNA Into Densely Functionalized Nucleic Acid Polymers. <i>Journal of the American Chemical Society</i> , 2013, 135, 98-101.	6.6	65
118	A Population-Based Experimental Model for Protein Evolution: Effects of Mutation Rate and Selection Stringency on Evolutionary Outcomes. <i>Biochemistry</i> , 2013, 52, 1490-1499.	1.2	37
119	Cellular Uptake Mechanisms and Endosomal Trafficking of Supercharged Proteins. <i>Chemistry and Biology</i> , 2012, 19, 831-843.	6.2	80
120	Revealing off-target cleavage specificities of zinc-finger nucleases by in vitro selection. <i>Nature Methods</i> , 2011, 8, 765-770.	9.0	404
121	A system for the continuous directed evolution of biomolecules. <i>Nature</i> , 2011, 472, 499-503.	13.7	518
122	A Class of Human Proteins that Deliver Functional Proteins into Mammalian Cells In Vitro and In Vivo. <i>Chemistry and Biology</i> , 2011, 18, 833-838.	6.2	98
123	A general strategy for the evolution of bond-forming enzymes using yeast display. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 11399-11404.	3.3	479
124	Enhanced Functional Potential of Nucleic Acid Aptamer Libraries Patterned to Increase Secondary Structure. <i>Journal of the American Chemical Society</i> , 2010, 132, 9453-9464.	6.6	70
125	Potent Delivery of Functional Proteins into Mammalian Cells <i>in Vitro</i> and <i>in Vivo</i> Using a Supercharged Protein. <i>ACS Chemical Biology</i> , 2010, 5, 747-752.	1.6	185
126	Supercharging Proteins Can Impart Unusual Resilience. <i>Journal of the American Chemical Society</i> , 2007, 129, 10110-10112.	6.6	438



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127	Analysis of Active Site Residues in <i>Escherichia coli</i> Chorismate Mutase by Site-Directed Mutagenesis. <i>Journal of the American Chemical Society</i> , 1996, 118, 1789-1790.	6.6	65