

# Douglas M Fowler

## List of Publications by Year in descending order

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67  
papers

8,567  
citations

117625

34  
h-index

102487

66  
g-index

92  
all docs

92  
docs citations

92  
times ranked

9468  
citing authors

#	ARTICLE	IF	CITATIONS
1	Functional amyloid “ from bacteria to humans. Trends in Biochemical Sciences, 2007, 32, 217-224.	7.5	953
2	Deep mutational scanning: a new style of protein science. Nature Methods, 2014, 11, 801-807.	19.0	885
3	Functional Amyloid Formation within Mammalian Tissue. PLoS Biology, 2005, 4, e6.	5.6	672
4	Semen-Derived Amyloid Fibrils Drastically Enhance HIV Infection. Cell, 2007, 131, 1059-1071.	28.9	510
5	High-resolution mapping of protein sequence-function relationships. Nature Methods, 2010, 7, 741-746.	19.0	482
6	Multiplex assessment of protein variant abundance by massively parallel sequencing. Nature Genetics, 2018, 50, 874-882.	21.4	323
7	Elevated levels of oxidized cholesterol metabolites in Lewy body disease brains accelerate $\alpha$ -synuclein fibrilization. Nature Chemical Biology, 2006, 2, 249-253.	8.0	312
8	Structure of the Sec13/31 COPII coat cage. Nature, 2006, 439, 234-238.	27.8	286
9	Variant Interpretation: Functional Assays to the Rescue. American Journal of Human Genetics, 2017, 101, 315-325.	6.2	275
10	Massively Parallel Functional Analysis of BRCA1 RING Domain Variants. Genetics, 2015, 200, 413-422.	2.9	272
11	A fundamental protein property, thermodynamic stability, revealed solely from large-scale measurements of protein function. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 16858-16863.	7.1	226
12	Quantitative Missense Variant Effect Prediction Using Large-Scale Mutagenesis Data. Cell Systems, 2018, 6, 116-124.e3.	6.2	176
13	Deep mutational scanning: assessing protein function on a massive scale. Trends in Biotechnology, 2011, 29, 435-442.	9.3	174
14	The dynamin middle domain is critical for tetramerization and higher-order self-assembly. EMBO Journal, 2007, 26, 559-566.	7.8	164
15	Activity-enhancing mutations in an E3 ubiquitin ligase identified by high-throughput mutagenesis. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, E1263-72.	7.1	158
16	A statistical framework for analyzing deep mutational scanning data. Genome Biology, 2017, 18, 150.	8.8	155
17	Measuring the activity of protein variants on a large scale using deep mutational scanning. Nature Protocols, 2014, 9, 2267-2284.	12.0	147
18	A framework for exhaustively mapping functional missense variants. Molecular Systems Biology, 2017, 13, 957.	7.2	146

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19	Biophysical and Mechanistic Models for Disease-Causing Protein Variants. <i>Trends in Biochemical Sciences</i> , 2019, 44, 575-588.	7.5	143
20	Pharmacogenomics of CYP2C9: Functional and Clinical Considerations. <i>Journal of Personalized Medicine</i> , 2018, 8, 1.	2.5	136
21	MaveDB: an open-source platform to distribute and interpret data from multiplexed assays of variant effect. <i>Genome Biology</i> , 2019, 20, 223.	8.8	130
22	Enrich: software for analysis of protein function by enrichment and depletion of variants. <i>Bioinformatics</i> , 2011, 27, 3430-3431.	4.1	112
23	Partial Restoration of Mutant Enzyme Homeostasis in Three Distinct Lysosomal Storage Disease Cell Lines by Altering Calcium Homeostasis. <i>PLoS Biology</i> , 2008, 6, e26.	5.6	110
24	Bedside Back to Bench: Building Bridges between Basic and Clinical Genomic Research. <i>Cell</i> , 2017, 169, 6-12.	28.9	103
25	N-terminal Domains Elicit Formation of Functional Pmel17 Amyloid Fibrils. <i>Journal of Biological Chemistry</i> , 2009, 284, 35543-35555.	3.4	101
26	Analysis of Large-Scale Mutagenesis Data To Assess the Impact of Single Amino Acid Substitutions. <i>Genetics</i> , 2017, 207, 53-61.	2.9	101
27	Massively parallel variant characterization identifies <i>NUDT15</i> alleles associated with thiopurine toxicity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 5394-5401.	7.1	95
28	Rapidly inducible Cas9 and DSB-ddPCR to probe editing kinetics. <i>Nature Methods</i> , 2017, 14, 891-896.	19.0	88
29	A platform for functional assessment of large variant libraries in mammalian cells. <i>Nucleic Acids Research</i> , 2017, 45, e102-e102.	14.5	80
30	Understanding the Origins of Loss of Protein Function by Analyzing the Effects of Thousands of Variants on Activity and Abundance. <i>Molecular Biology and Evolution</i> , 2021, 38, 3235-3246.	8.9	65
31	Multiplexed measurement of variant abundance and activity reveals VKOR topology, active site and human variant impact. <i>ELife</i> , 2020, 9, .	6.0	58
32	Massively parallel characterization of CYP2C9 variant enzyme activity and abundance. <i>American Journal of Human Genetics</i> , 2021, 108, 1735-1751.	6.2	53
33	An improved platform for functional assessment of large protein libraries in mammalian cells. <i>Nucleic Acids Research</i> , 2020, 48, e1.	14.5	51
34	Recommendations for the collection and use of multiplexed functional data for clinical variant interpretation. <i>Genome Medicine</i> , 2019, 11, 85.	8.2	47
35	High-throughput, microscope-based sorting to dissect cellular heterogeneity. <i>Molecular Systems Biology</i> , 2020, 16, e9442.	7.2	46
36	A Combined Approach Reveals a Regulatory Mechanism Coupling Src's Kinase Activity, Localization, and Phosphotransferase-Independent Functions. <i>Molecular Cell</i> , 2019, 74, 393-408.e20.	9.7	45

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37	Suppression of unwanted CRISPR-Cas9 editing by co-administration of catalytically inactivating truncated guide RNAs. <i>Nature Communications</i> , 2020, 11, 2697.	12.8	42
38	High-throughput discovery of trafficking-deficient variants in the cardiac potassium channel KV11.1. <i>Heart Rhythm</i> , 2020, 17, 2180-2189.	0.7	42
39	Closing the gap: Systematic integration of multiplexed functional data resolves variants of uncertain significance in BRCA1, TP53, and PTEN. <i>American Journal of Human Genetics</i> , 2021, 108, 2248-2258.	6.2	42
40	Expression and Functional Characterization of Breast Cancer-Associated Cytochrome P450 4Z1 in <i>Saccharomyces cerevisiae</i> . <i>Drug Metabolism and Disposition</i> , 2017, 45, 1364-1371.	3.3	35
41	Contemporary, yeast-based approaches to understanding human genetic variation. <i>Current Opinion in Genetics and Development</i> , 2013, 23, 658-664.	3.3	34
42	Engineering A-kinase Anchoring Protein (AKAP)-selective Regulatory Subunits of Protein Kinase A (PKA) through Structure-based Phage Selection. <i>Journal of Biological Chemistry</i> , 2013, 288, 17111-17121.	3.4	34
43	New Pharmacogenomics Research Network: An Open Community Catalyzing Research and Translation in Precision Medicine. <i>Clinical Pharmacology and Therapeutics</i> , 2017, 102, 897-902.	4.7	34
44	Deep Mutational Scan of an <i>SCN5A</i> Voltage Sensor. <i>Circulation Genomic and Precision Medicine</i> , 2020, 13, e002786.	3.6	33
45	Elucidating the Molecular Determinants of A $\beta$ 2 Aggregation with Deep Mutational Scanning. <i>G3: Genes, Genomes, Genetics</i> , 2019, 9, 3683-3689.	1.8	32
46	Systematic misclassification of missense variants in BRCA1 and BRCA2 "coldspots". <i>Genetics in Medicine</i> , 2020, 22, 825-830.	2.4	32
47	Comprehensive exploration of the translocation, stability and substrate recognition requirements in VIM-2 lactamase. <i>ELife</i> , 2020, 9, .	6.0	26
48	Suppression of statin effectiveness by copper and zinc in yeast and human cells. <i>Molecular BioSystems</i> , 2011, 7, 533-544.	2.9	23
49	Classifying disease-associated variants using measures of protein activity and stability. , 2020, , 91-107.		21
50	Probing ion channel functional architecture and domain recombination compatibility by massively parallel domain insertion profiling. <i>Nature Communications</i> , 2021, 12, 7114.	12.8	19
51	Applying Multiplex Assays to Understand Variation in Pharmacogenes. <i>Clinical Pharmacology and Therapeutics</i> , 2019, 106, 290-294.	4.7	15
52	Functional Amyloidogenesis and Cytotoxicity"Insights into Biology and Pathology. <i>PLoS Biology</i> , 2012, 10, e1001459.	5.6	14
53	Integrating thousands of PTEN variant activity and abundance measurements reveals variant subgroups and new dominant negatives in cancers. <i>Genome Medicine</i> , 2021, 13, 165.	8.2	14
54	Novel Rhizosphere Soil Alleles for the Enzyme 1-Aminocyclopropane-1-Carboxylate Deaminase Queried for Function with an <i>In Vivo</i> Competition Assay. <i>Applied and Environmental Microbiology</i> , 2016, 82, 1050-1059.	3.1	13

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55	Rheostatic Control of Cas9-Mediated DNA Double Strand Break (DSB) Generation and Genome Editing. ACS Chemical Biology, 2018, 13, 438-442.	3.4	13
56	Keeping up with the genomes: scaling genomic variant interpretation. Genome Medicine, 2020, 12, 5.	8.2	13
57	A Premalignant Cell-Based Model for Functionalization and Classification of <i>PTEN</i> Variants. Cancer Research, 2020, 80, 2775-2789.	0.9	11
58	Aggregating Knowledge about Prions and Amyloid. Cell, 2009, 137, 20-22.	28.9	10
59	Environmental selection and epistasis in an empirical phenotypeâ€“environmentâ€“fitness landscape. Nature Ecology and Evolution, 2022, 6, 427-438.	7.8	10
60	The Impact of Genetic Variants on PTEN Molecular Functions and Cellular Phenotypes. Cold Spring Harbor Perspectives in Medicine, 2019, 9, a036228.	6.2	9
61	MaveRegistry: a collaboration platform for multiplexed assays of variant effect. Bioinformatics, 2021, 37, 3382-3383.	4.1	9
62	Measuring Pharmacogene Variant Function at Scale Using Multiplexed Assays. Annual Review of Pharmacology and Toxicology, 2022, 62, 531-550.	9.4	9
63	Early emergence of negative regulation of the tyrosine kinase Src by the C-terminal Src kinase. Journal of Biological Chemistry, 2017, 292, 18518-18529.	3.4	7
64	Parallel Chemoselective Profiling for Mapping Protein Structure. Cell Chemical Biology, 2020, 27, 1084-1096.e4.	5.2	6
65	Mutagenesis-based protein structure determination. Nature Genetics, 2019, 51, 1072-1073.	21.4	5
66	Temporal and rheostatic control of genome editing with a chemically-inducible Cas9. Methods in Enzymology, 2020, 633, 119-141.	1.0	3
67	Protocol for rapidly inducible Cas9 and DSB-ddPCR. Protocol Exchange, 0, , .	0.3	2