

# Chris P Barnes

## List of Publications by Year in descending order

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Version: 2024-02-01

63  
papers

9,389  
citations

117625

34  
h-index

133252

59  
g-index

84  
all docs

84  
docs citations

84  
times ranked

18611  
citing authors

#	ARTICLE	IF	CITATIONS
1	Fundamental Building Blocks of Whole-Cell Biosensor Design. , 2022, , 383-405.		0
2	Fluctuating methylation clocks for cell lineage tracing at high temporal resolution in human tissues. Nature Biotechnology, 2022, 40, 720-730.	17.5	22
3	Automated design of synthetic microbial communities. Nature Communications, 2021, 12, 672.	12.8	58
4	Single strain control of microbial consortia. Nature Communications, 2021, 12, 1977.	12.8	37
5	Reconstructing single-cell karyotype alterations in colorectal cancer identifies punctuated and gradual diversification patterns. Nature Genetics, 2021, 53, 1187-1195.	21.4	37
6	Engineered acetoacetate-inducible whole-cell biosensors based on the AtoSC two-component system. Biotechnology and Bioengineering, 2021, 118, 4278-4289.	3.3	10
7	From Microbial Communities to Distributed Computing Systems. Frontiers in Bioengineering and Biotechnology, 2020, 8, 834.	4.1	19
8	Subclonal reconstruction of tumors by using machine learning and population genetics. Nature Genetics, 2020, 52, 898-907.	21.4	77
9	FlopR: An Open Source Software Package for Calibration and Normalization of Plate Reader and Flow Cytometry Data. ACS Synthetic Biology, 2020, 9, 2258-2266.	3.8	17
10	Evolutionary dynamics of neoantigens in growing tumors. Nature Genetics, 2020, 52, 1057-1066.	21.4	68
11	Measuring single cell divisions in human tissues from multi-region sequencing data. Nature Communications, 2020, 11, 1035.	12.8	41
12	Deep reinforcement learning for the control of microbial co-cultures in bioreactors. PLoS Computational Biology, 2020, 16, e1007783.	3.2	69
13	Fundamental Building Blocks of Whole-Cell Biosensor Design. , 2020, , 1-23.		4
14	Measuring the distribution of fitness effects in somatic evolution by combining clonal dynamics with dN/dS ratios. ELife, 2020, 9, .	6.0	32
15	Detecting Changes in the <i>Caenorhabditis elegans</i> Intestinal Environment Using an Engineered Bacterial Biosensor. ACS Synthetic Biology, 2019, 8, 2620-2628.	3.8	21
16	A Bayesian framework for the analysis of systems biology models of the brain. PLoS Computational Biology, 2019, 15, e1006631.	3.2	11
17	Modelling microbiome recovery after antibiotics using a stability landscape framework. ISME Journal, 2019, 13, 1845-1856.	9.8	98
18	Crypt fusion as a homeostatic mechanism in the human colon. Gut, 2019, 68, 1986-1993.	12.1	28

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19	Two New Plasmid Post-segregational Killing Mechanisms for the Implementation of Synthetic Gene Networks in Escherichia coli. <i>IScience</i> , 2019, 14, 323-334.	4.1	41
20	Combining a Toggle Switch and a Repressilator within the AC-DC Circuit Generates Distinct Dynamical Behaviors. <i>Cell Systems</i> , 2018, 6, 521-530.e3.	6.2	96
21	Reply to "Revisiting signatures of neutral tumor evolution in the light of complexity of cancer genomic data". <i>Nature Genetics</i> , 2018, 50, 1628-1630.	21.4	5
22	Towards an Aspect-Oriented Design and Modelling Framework for Synthetic Biology. <i>Processes</i> , 2018, 6, 167.	2.8	6
23	Reply to "Currently available bulk sequencing data do not necessarily support a model of neutral tumor evolution". <i>Nature Genetics</i> , 2018, 50, 1624-1626.	21.4	11
24	Reply to "Neutral tumor evolution?". <i>Nature Genetics</i> , 2018, 50, 1633-1637.	21.4	27
25	Quantification of subclonal selection in cancer from bulk sequencing data. <i>Nature Genetics</i> , 2018, 50, 895-903.	21.4	222
26	Synthetic Biology and Engineered Live Biotherapeutics: Toward Increasing System Complexity. <i>Cell Systems</i> , 2018, 7, 5-16.	6.2	107
27	Reply: Is the evolution of tumors Darwinian or non-Darwinian?. <i>National Science Review</i> , 2018, 5, 17-19.	9.5	3
28	Computing with biological switches and clocks. <i>Natural Computing</i> , 2018, 17, 761-779.	3.0	45
29	Catch my drift? Making sense of genomic intra-tumour heterogeneity. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2017, 1867, 95-100.	7.4	23
30	Reply: Uncertainties in tumor allele frequencies limit power to infer evolutionary pressures. <i>Nature Genetics</i> , 2017, 49, 1289-1291.	21.4	7
31	A computational method for the investigation of multistable systems and its application to genetic switches. <i>BMC Systems Biology</i> , 2016, 10, 130.	3.0	33
32	Robustness of MEK-ERK Dynamics and Origins of Cell-to-Cell Variability in MAPK Signaling. <i>Cell Reports</i> , 2016, 15, 2524-2535.	6.4	57
33	Identification of neutral tumor evolution across cancer types. <i>Nature Genetics</i> , 2016, 48, 238-244.	21.4	525
34	A Statistical Approach Reveals Designs for the Most Robust Stochastic Gene Oscillators. <i>ACS Synthetic Biology</i> , 2016, 5, 459-470.	3.8	61
35	Mechanistic Modelling and Bayesian Inference Elucidates the Variable Dynamics of Double-Strand Break Repair. <i>PLoS Computational Biology</i> , 2016, 12, e1005131.	3.2	8
36	Ptch1 and Gli regulate Shh signalling dynamics via multiple mechanisms. <i>Nature Communications</i> , 2015, 6, 6709.	12.8	123

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37	Directional Collective Cell Migration Emerges as a Property of Cell Interactions. <i>PLoS ONE</i> , 2014, 9, e104969.	2.5	68
38	Model Selection in Systems Biology Depends on Experimental Design. <i>PLoS Computational Biology</i> , 2014, 10, e1003650.	3.2	54
39	A theoretical framework for the regulation of Shh morphogen-controlled gene expression. <i>Development (Cambridge)</i> , 2014, 141, 3868-3878.	2.5	70
40	Reciprocal Duplication of the Williams-Beuren Syndrome Deletion on Chromosome 7q11.23 Is Associated with Schizophrenia. <i>Biological Psychiatry</i> , 2014, 75, 371-377.	1.3	66
41	A framework for parameter estimation and model selection from experimental data in systems biology using approximate Bayesian computation. <i>Nature Protocols</i> , 2014, 9, 439-456.	12.0	185
42	On Industrial Strength Bio-design Automation. <i>Communications in Computer and Information Science</i> , 2014, , 277-299.	0.5	0
43	Clinical drug resistance linked to interconvertible phenotypic and functional states of tumor-propagating cells in multiple myeloma. <i>Blood</i> , 2013, 121, 318-328.	1.4	112
44	On optimality of kernels for approximate Bayesian computation using sequential Monte Carlo. <i>Statistical Applications in Genetics and Molecular Biology</i> , 2013, 12, 87-107.	0.6	86
45	Reduced burden of very large and rare CNVs in bipolar affective disorder. <i>Bipolar Disorders</i> , 2013, 15, 893-898.	1.9	28
46	Considerate approaches to constructing summary statistics for ABC model selection. <i>Statistics and Computing</i> , 2012, 22, 1181-1197.	1.5	45
47	Calibrating spatio-temporal models of leukocyte dynamics against in vivo live-imaging data using approximate Bayesian computation. <i>Integrative Biology (United Kingdom)</i> , 2012, 4, 335.	1.3	31
48	Independent estimation of the frequency of rare CNVs in the UK population confirms their role in schizophrenia. <i>Schizophrenia Research</i> , 2012, 135, 1-7.	2.0	73
49	Genome-Wide Screen for Metabolic Syndrome Susceptibility Loci Reveals Strong Lipid Gene Contribution But No Evidence for Common Genetic Basis for Clustering of Metabolic Syndrome Traits. <i>Circulation: Cardiovascular Genetics</i> , 2012, 5, 242-249.	5.1	182
50	Clinical Drug Resistance Linked to Inter-Convertible Phenotypic and Functional States of Tumor-Propagating Cells in Multiple Myeloma.. <i>Blood</i> , 2012, 120, 2909-2909.	1.4	6
51	Computational design approaches and tools for synthetic biology. <i>Integrative Biology (United)</i> Tj ETQq1 1 0.784314 rgBT /Overlock 10	1.3	74
52	Bayesian design strategies for synthetic biology. <i>Interface Focus</i> , 2011, 1, 895-908.	3.0	29
53	GPU accelerated biochemical network simulation. <i>Bioinformatics</i> , 2011, 27, 874-876.	4.1	81
54	Designing attractive models via automated identification of chaotic and oscillatory dynamical regimes. <i>Nature Communications</i> , 2011, 2, 489.	12.8	62

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55	Bayesian design of synthetic biological systems. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 15190-15195.	7.1	82
56	Origins and functional impact of copy number variation in the human genome. Nature, 2010, 464, 704-712.	27.8	1,721
57	Genome-wide association study of CNVs in 16,000 cases of eight common diseases and 3,000 shared controls. Nature, 2010, 464, 713-720.	27.8	737
58	Integrating common and rare genetic variation in diverse human populations. Nature, 2010, 467, 52-58.	27.8	2,625
59	Founder population-specific HapMap panel increases power in GWA studies through improved imputation accuracy and CNV tagging. Genome Research, 2010, 20, 1344-1351.	5.5	52
60	ABC-SysBio—approximate Bayesian computation in Python with GPU support. Bioinformatics, 2010, 26, 1797-1799.	4.1	124
61	A Genome-Wide Association Study Confirms VKORC1, CYP2C9, and CYP4F2 as Principal Genetic Determinants of Warfarin Dose. PLoS Genetics, 2009, 5, e1000433.	3.5	554
62	A robust statistical method for case-control association testing with copy number variation. Nature Genetics, 2008, 40, 1245-1252.	21.4	151
63	Reply: Neutral tumor evolution?. , 0, , .		1