Jim R Hughes

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6680978/publications.pdf

Version: 2024-02-01

71061 85498 6,676 73 41 71 h-index citations g-index papers 97 97 97 9171 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Making connections: enhancers in cellular differentiation. Trends in Genetics, 2022, 38, 395-408.	2.9	6
2	Capture-C: a modular and flexible approach for high-resolution chromosome conformation capture. Nature Protocols, 2022, 17, 445-475.	5 . 5	24
3	Dynamic Runx1 chromatin boundaries affect gene expression in hematopoietic development. Nature Communications, 2022, 13, 773.	5.8	10
4	Natural and Experimental Rewiring of Gene Regulatory Regions. Annual Review of Genomics and Human Genetics, 2022, 23, .	2.5	1
5	The chromatin remodeller ATRX facilitates diverse nuclear processes, in a stochastic manner, in both heterochromatin and euchromatin. Nature Communications, 2022, 13, .	5.8	20
6	Chromatin interaction maps identify Wnt responsive cis-regulatory elements coordinating Paupar-Pax6 expression in neuronal cells. PLoS Genetics, 2022, 18, e1010230.	1.5	6
7	Genetic and functional insights into CDA-I prevalence and pathogenesis. Journal of Medical Genetics, 2021, 58, 185-195.	1.5	9
8	High-resolution targeted 3C interrogation of cis-regulatory element organization at genome-wide scale. Nature Communications, 2021, 12, 531.	5.8	32
9	Fra-1 regulates its target genes via binding to remote enhancers without exerting major control on chromatin architecture in triple negative breast cancers. Nucleic Acids Research, 2021, 49, 2488-2508.	6.5	15
10	Enhancers predominantly regulate gene expression during differentiation via transcription initiation. Molecular Cell, 2021, 81, 983-997.e7.	4.5	27
11	Multi Locus View: an extensible web-based tool for the analysis of genomic data Communications Biology, 2021, 4, 623.	2.0	4
12	Defining genome architecture at base-pair resolution. Nature, 2021, 595, 125-129.	13.7	107
13	A gain-of-function single nucleotide variant creates a new promoter which acts as an orientation-dependent enhancer-blocker. Nature Communications, 2021, 12, 3806.	5.8	18
14	Reactivation of a developmentally silenced embryonic globin gene. Nature Communications, 2021, 12, 4439.	5.8	19
15	Recapitulation of erythropoiesis in congenital dyserythropoietic anemia type I (CDA-I) identifies defects in differentiation and nucleolar abnormalities. Haematologica, 2021, 106, 2960-2970.	1.7	10
16	Identification of LZTFL1 as a candidate effector gene at a COVID-19 risk locus. Nature Genetics, 2021, 53, 1606-1615.	9.4	93
17	Systematic integration of GATA transcription factors and epigenomes via IDEAS paints the regulatory landscape of hematopoietic cells. IUBMB Life, 2020, 72, 27-38.	1.5	8
18	Loss of Extreme Long-Range Enhancers in Human Neural Crest Drives a Craniofacial Disorder. Cell Stem Cell, 2020, 27, 765-783.e14.	5.2	101

#	Article	IF	CITATIONS
19	DeepC: predicting 3D genome folding using megabase-scale transfer learning. Nature Methods, 2020, 17, 1118-1124.	9.0	109
20	Dynamics of the 4D genome during in vivo lineage specification and differentiation. Nature Communications, 2020, 11, 2722.	5.8	79
21	An integrative view of the regulatory and transcriptional landscapes in mouse hematopoiesis. Genome Research, 2020, 30, 472-484.	2.4	38
22	A Dynamic Folded Hairpin Conformation Is Associated with $\hat{l}\pm$ -Globin Activation in Erythroid Cells. Cell Reports, 2020, 30, 2125-2135.e5.	2.9	38
23	Absolute Quantification of Transcription Factors Reveals Principles of Gene Regulation in Erythropoiesis. Molecular Cell, 2020, 78, 960-974.e11.	4.5	83
24	DOT1L inhibition reveals a distinct subset of enhancers dependent on H3K79 methylation. Nature Communications, 2019, 10, 2803.	5.8	99
25	Reconstruction of the Global Neural Crest Gene Regulatory Network InÂVivo. Developmental Cell, 2019, 51, 255-276.e7.	3.1	108
26	The bipartite TAD organization of the X-inactivation center ensures opposing developmental regulation of Tsix and Xist. Nature Genetics, 2019, 51, 1024-1034.	9.4	60
27	Nrf2 controls iron homoeostasis in haemochromatosis and thalassaemia via Bmp6 and hepcidin. Nature Metabolism, 2019, 1, 519-531.	5.1	88
28	A revised model for promoter competition based on multi-way chromatin interactions at the \hat{l}_{\pm} -globin locus. Nature Communications, 2019, 10, 5412.	5.8	60
29	HoxC5 and miR-615-3p target newly evolved genomic regions to repress hTERT and inhibit tumorigenesis. Nature Communications, 2018, 9, 100.	5.8	38
30	High-Throughput Genotyping of CRISPR/Cas Edited Cells in 96-Well Plates. Methods and Protocols, 2018, 1, 29.	0.9	6
31	A tissue-specific self-interacting chromatin domain forms independently of enhancer-promoter interactions. Nature Communications, 2018, 9, 3849.	5.8	62
32	Single-allele chromatin interactions identify regulatory hubs in dynamic compartmentalized domains. Nature Genetics, 2018, 50, 1744-1751.	9.4	150
33	MLL-AF4 binds directly to a BCL-2 specific enhancer and modulates H3K27Âacetylation. Experimental Hematology, 2017, 47, 64-75.	0.2	25
34	How best to identify chromosomal interactions: a comparison of approaches. Nature Methods, 2017, 14, 125-134.	9.0	124
35	Functional characterisation of cis-regulatory elements governing dynamic <i>Eomes</i> expression in the early mouse embryo. Development (Cambridge), 2017, 144, 1249-1260.	1.2	32
36	DNA methylation of intragenic CpG islands depends on their transcriptional activity during differentiation and disease. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E7526-E7535.	3.3	125

#	Article	IF	CITATIONS
37	Comparative analysis of three-dimensional chromosomal architecture identifies a novel fetal hemoglobin regulatory element. Genes and Development, 2017, 31, 1704-1713.	2.7	113
38	Sasquatch: predicting the impact of regulatory SNPs on transcription factor binding from cell- and tissue-specific DNase footprints. Genome Research, 2017, 27, 1730-1742.	2.4	33
39	<i>VHL</i> Deficiency Drives Enhancer Activation of Oncogenes in Clear Cell Renal Cell Carcinoma. Cancer Discovery, 2017, 7, 1284-1305.	7.7	111
40	Hepcidin is regulated by promoter-associated histone acetylation and HDAC3. Nature Communications, 2017, 8, 403.	5.8	45
41	Editing an \hat{l} ±-globin enhancer in primary human hematopoietic stem cells as a treatment for \hat{l}^2 -thalassemia. Nature Communications, 2017, 8, 424.	5 . 8	85
42	Tissue-specific CTCF–cohesin-mediated chromatin architecture delimits enhancer interactions and function in vivo. Nature Cell Biology, 2017, 19, 952-961.	4.6	179
43	Between form and function: the complexity of genome folding. Human Molecular Genetics, 2017, 26, R208-R215.	1.4	20
44	Robust detection of chromosomal interactions from small numbers of cells using low-input Capture-C. Nucleic Acids Research, 2017, 45, e184-e184.	6.5	27
45	Low-input Capture-C: A Chromosome Conformation Capture Assay to Analyze Chromatin Architecture in Small Numbers of Cells. Bio-protocol, 2017, 7, .	0.2	15
46	Comparison of Fetal and Adult Erythroid Chromosomal Architectures Identifies a Novel Fetal Hemoglobin Regulatory Region. Blood, 2017, 130, 774-774.	0.6	0
47	Genetic dissection of the α-globin super-enhancer in vivo. Nature Genetics, 2016, 48, 895-903.	9.4	308
48	Capture reveals preformed chromatin interactions between <scp>HIF</scp> â€binding sites and distant promoters. EMBO Reports, 2016, 17, 1410-1421.	2.0	63
49	Unlinking an IncRNA from Its Associated cis Element. Molecular Cell, 2016, 62, 104-110.	4.5	216
50	A genome-editing strategy to treat \hat{l}^2 -hemoglobinopathies that recapitulates a mutation associated with a benign genetic condition. Nature Medicine, 2016, 22, 987-990.	15.2	279
51	Epigenomic profiling of primary gastric adenocarcinoma reveals super-enhancer heterogeneity. Nature Communications, 2016, 7, 12983.	5.8	123
52	Predicting the three-dimensional folding of cis-regulatory regions in mammalian genomes using bioinformatic data and polymer models. Genome Biology, 2016, 17, 59.	3.8	97
53	Multiplexed analysis of chromosome conformation at vastly improved sensitivity. Nature Methods, 2016, 13, 74-80.	9.0	225
54	ATRX Plays a Key Role in Maintaining Silencing at Interstitial Heterochromatic Loci and Imprinted Genes. Cell Reports, 2015, 11, 405-418.	2.9	152

#	Article	IF	Citations
55	Analysis of hundreds of cis-regulatory landscapes at high resolution in a single, high-throughput experiment. Nature Genetics, 2014, 46, 205-212.	9.4	417
56	Chromatin signatures at transcriptional start sites separate two equally populated yet distinct classes of intergenic long noncoding RNAs. Genome Biology, 2013, 14, R131.	13.9	183
57	Analysis of Sequence Variation Underlying Tissue-specific Transcription Factor Binding and Gene Expression. Human Mutation, 2013, 34, 1140-1148.	1.1	10
58	Causes and Consequences of Chromatin Variation between Inbred Mice. PLoS Genetics, 2013, 9, e1003570.	1.5	18
59	MIG: Multi-Image Genome viewer. Bioinformatics, 2013, 29, 2477-2478.	1.8	15
60	High-resolution analysis of $\langle i \rangle$ cis $\langle i \rangle$ -acting regulatory networks at the α-globin locus. Philosophical Transactions of the Royal Society B: Biological Sciences, 2013, 368, 20120361.	1.8	12
61	Intragenic Enhancers Act as Alternative Promoters. Molecular Cell, 2012, 45, 447-458.	4.5	237
62	Nprl3 is required for normal development of the cardiovascular system. Mammalian Genome, 2012, 23, 404-415.	1.0	38
63	Genome-wide identification of TAL1's functional targets: Insights into its mechanisms of action in primary erythroid cells. Genome Research, 2010, 20, 1064-1083.	2.4	154
64	Adventitious changes in long-range gene expression caused by polymorphic structural variation and promoter competition. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 21771-21776.	3.3	77
65	Alternative Runx1 promoter usage in mouse developmental hematopoiesis. Blood Cells, Molecules, and Diseases, 2009, 43, 35-42.	0.6	52
66	Manipulating the Mouse Genome to Engineer Precise Functional Syntenic ReplacementsÂwith Human Sequence. Cell, 2007, 128, 197-209.	13.5	150
67	Tissue-specific histone modification and transcription factor binding in $\hat{l}\pm$ globin gene expression. Blood, 2007, 110, 4503-4510.	0.6	69
68	A Regulatory SNP Causes a Human Genetic Disease by Creating a New Transcriptional Promoter. Science, 2006, 312, 1215-1217.	6.0	254
69	Annotation of cis-regulatory elements by identification, subclassification, and functional assessment of multispecies conserved sequences. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 9830-9835.	3.3	133
70	Globin gene activation during haemopoiesis is driven by protein complexes nucleated by GATA-1 and GATA-2. EMBO Journal, 2004, 23, 2841-2852.	3.5	193
71	Comparative analysis of the polycystic kidney disease 1 (PKD1) gene reveals an integral membrane glycoprotein with multiple evolutionary conserved domains. Human Molecular Genetics, 1997, 6, 1483-1489.	1.4	141
72	Deletion of the TSC2 and PKD1 genes associated with severe infantile polycystic kidney disease — a contiguous gene syndrome. Nature Genetics, 1994, 8, 328-332.	9.4	466

#	Article	IF	CITATIONS
73	Tri-C. Protocol Exchange, 0, , .	0.3	3