Cheryl A Keller

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	A comparative encyclopedia of DNA elements in the mouse genome. Nature, 2014, 515, 355-364.	27.8	1,444
2	Expanded encyclopaedias of DNA elements in the human and mouse genomes. Nature, 2020, 583, 699-710.	27.8	1,252
3	An encyclopedia of mouse DNA elements (Mouse ENCODE). Genome Biology, 2012, 13, 418.	9.6	410
4	Principles of regulatory information conservation between mouse and human. Nature, 2014, 515, 371-375.	27.8	259
5	The Â2 Subunit of GABAA Receptors Is a Substrate for Palmitoylation by GODZ. Journal of Neuroscience, 2004, 24, 5881-5891.	3.6	225
6	Chromatin structure dynamics during the mitosis-to-G1 phase transition. Nature, 2019, 576, 158-162.	27.8	167
7	GODZ-Mediated Palmitoylation of GABAA Receptors Is Required for Normal Assembly and Function of GABAergic Inhibitory Synapses. Journal of Neuroscience, 2006, 26, 12758-12768.	3.6	148
8	Perspectives on ENCODE. Nature, 2020, 583, 693-698.	27.8	123
9	Domain-focused CRISPR screen identifies HRI as a fetal hemoglobin regulator in human erythroid cells. Science, 2018, 361, 285-290.	12.6	119
10	Transcriptional Burst Initiation and Polymerase Pause Release Are Key Control Points of Transcriptional Regulation. Molecular Cell, 2019, 73, 519-532.e4.	9.7	118
11	The BET Protein BRD2 Cooperates with CTCF to Enforce Transcriptional and Architectural Boundaries. Molecular Cell, 2017, 66, 102-116.e7.	9.7	114
12	Comparative analysis of three-dimensional chromosomal architecture identifies a novel fetal hemoglobin regulatory element. Genes and Development, 2017, 31, 1704-1713.	5.9	113
13	Dynamics of the epigenetic landscape during erythroid differentiation after GATA1 restoration. Genome Research, 2011, 21, 1659-1671.	5.5	110
14	Genome accessibility is widely preserved and locally modulated during mitosis. Genome Research, 2015, 25, 213-225.	5.5	103
15	Occupancy by key transcription factors is a more accurate predictor of enhancer activity than histone modifications or chromatin accessibility. Epigenetics and Chromatin, 2015, 8, 16.	3.9	100
16	Functions of BET proteins in erythroid gene expression. Blood, 2015, 125, 2825-2834.	1.4	93
17	A hyperactive transcriptional state marks genome reactivation at the mitosis–G1 transition. Genes and Development, 2016, 30, 1423-1439.	5.9	92
18	Divergent functions of hematopoietic transcription factors in lineage priming and differentiation during erythro-megakaryopoiesis. Genome Research, 2014, 24, 1932-1944.	5.5	88

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19	Dynamic shifts in occupancy by TAL1 are guided by GATA factors and drive large-scale reprogramming of gene expression during hematopoiesis. Genome Research, 2014, 24, 1945-1962.	5.5	71
20	Distinct properties and functions of CTCF revealed by a rapidly inducible degron system. Cell Reports, 2021, 34, 108783.	6.4	53
21	Interrogating Histone Acetylation and BRD4 as Mitotic Bookmarks of Transcription. Cell Reports, 2019, 27, 400-415.e5.	6.4	52
22	Establishment of regulatory elements during erythro-megakaryopoiesis identifies hematopoietic lineage-commitment points. Epigenetics and Chromatin, 2018, 11, 22.	3.9	49
23	ZNF410ÂUniquely Activates the NuRD Component CHD4 to Silence Fetal Hemoglobin Expression. Molecular Cell, 2021, 81, 239-254.e8.	9.7	48
24	Loss-of-Function Mutations Reveal That the Drosophila nautilus Gene Is Not Essential for Embryonic Myogenesis or Viability. Developmental Biology, 2001, 231, 374-382.	2.0	44
25	The HRI-regulated transcription factor ATF4 activates BCL11A transcription to silence fetal hemoglobin expression. Blood, 2020, 135, 2121-2132.	1.4	42
26	Misexpression ofnautilusInduces Myogenesis in Cardioblasts and Alters the Pattern of Somatic Muscle Fibers. Developmental Biology, 1997, 181, 197-212.	2.0	40
27	An integrative view of the regulatory and transcriptional landscapes in mouse hematopoiesis. Genome Research, 2020, 30, 472-484.	5.5	38
28	Exploiting genetic variation to uncover rules of transcription factor binding and chromatin accessibility. Nature Communications, 2018, 9, 782.	12.8	36
29	Alteration of genome folding via contact domain boundary insertion. Nature Genetics, 2020, 52, 1076-1087.	21.4	35
30	Genome-Wide Organization of GATA1 and TAL1 Determined at High Resolution. Molecular and Cellular Biology, 2016, 36, 157-172.	2.3	32
31	CTCF and transcription influence chromatin structure re-configuration after mitosis. Nature Communications, 2021, 12, 5157.	12.8	32
32	S3norm: simultaneous normalization of sequencing depth and signal-to-noise ratio in epigenomic data. Nucleic Acids Research, 2020, 48, e43-e43.	14.5	31
33	Understanding heterogeneity of fetal hemoglobin induction through comparative analysis of F and A erythroblasts. Blood, 2020, 135, 1957-1968.	1.4	30
34	Frequent somatic <i>TET2</i> mutations in chronic NK-LGL leukemia with distinct patterns of cytopenias. Blood, 2021, 138, 662-673.	1.4	30
35	The E3 ligase adaptor molecule SPOP regulates fetal hemoglobin levels in adult erythroid cells. Blood Advances, 2019, 3, 1586-1597.	5.2	25
36	Protein Phosphatase 6C (PPP6C) Loss Significantly Raises Fetal Hemoglobin Levels and Reduces Cell Sickling. Blood, 2021, 138, 2031-2031.	1.4	18

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37	HRI depletion cooperates with pharmacologic inducers to elevate fetal hemoglobin and reduce sickle cell formation. Blood Advances, 2020, 4, 4560-4572.	5.2	15
38	Ubiquitination, proteasomes and GABAA receptors. Nature Cell Biology, 2001, 3, E232-E233.	10.3	14
39	Dual function NFI factors control fetal hemoglobin silencing in adult erythroid cells. Nature Genetics, 2022, 54, 874-884.	21.4	13
40	Dynamics of GATA1 binding and expression response in a GATA1-induced erythroid differentiation system. Genomics Data, 2015, 4, 1-7.	1.3	10
41	Systematic integration of GATA transcription factors and epigenomes via IDEAS paints the regulatory landscape of hematopoietic cells. IUBMB Life, 2020, 72, 27-38.	3.4	8
42	Comparison of Expression and Epigenetic Profiles in Human and Mouse Erythropoiesis and Megakaryopoiesis Using a Systems Biology Model. Blood, 2015, 126, 2383-2383.	1.4	5
43	Identification and characterization of RBM12 as a novel regulator ofÂfetal hemoglobin expression. Blood Advances, 2022, 6, 5956-5968.	5.2	5
44	SBR-Blood: systems biology repository for hematopoietic cells. Nucleic Acids Research, 2016, 44, D925-D931.	14.5	4
45	Effects of sheared chromatin length on ChIP-seq quality and sensitivity. G3: Genes, Genomes, Genetics, 2021, 11, .	1.8	3
46	HIC2 Controls Developmental Hemoglobin Switching By Repressing BCL11A Transcription. Blood, 2021, 138, 571-571.	1.4	2
47	Control of Fetal Hemoglobin Levels By NFI Transcription Factors. Blood, 2020, 136, 54-54.	1.4	2
48	De Novo DNA Methylation Is Associated with Granulopoiesis and Megakaryopoiesis but Not Erythropoiesis. Blood, 2016, 128, 3868-3868.	1.4	1
49	Epigenetics of Cellular Memory: Insights from the Chromatin Accessibility Landscape of the Mitotic Genome. Blood, 2014, 124, 4342-4342.	1.4	1
50	Isolated Changes in Chromatin Accessibility and Enhancer-Promoter Contacts at the β-Globin Locus Distinguish Fetal Hemoglobin Producing F-Cells from a-Cells. Blood, 2021, 138, 855-855.	1.4	1
51	Systems Biology in Heterogenous Tissues: Integrating Multiple *Omics Datasets to Understand Hematopoietic Differentiation. , 2019, , .		0
52	Enhancer Accessibility during Erythropoiesis and Megakaryopoiesis Correlates with Lineage-Specific Gene Expression. Blood, 2015, 126, 3576-3576.	1.4	0
53	Establishment of Enhancer Elements during Erythro-Megakaryopoiesis. Blood, 2016, 128, 1486-1486.	1.4	0
54	Mouse Erythroid Cells Originate from a Megakaryocyte Precursor in Common Myeloid Progenitors. Blood, 2019, 134, 337-337.	1.4	0

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55	Interrogating Post-Transcriptional Mechanisms of Fetal Hemoglobin Regulation. Blood, 2021, 138, 3079-3079.	1.4	0
56	A β-Globin Locus-Intrinsic Epigenetic Mechanism Underlies Fetal Globin Production in F-Cells. Blood, 2020, 136, 16-17.	1.4	0
57	Systematic Integration of Epigenomic Landscapes in Human and Mouse Blood Cells to Predict Activity and Targets of Regulatory Elements. FASEB Journal, 2022, 36, .	0.5	0