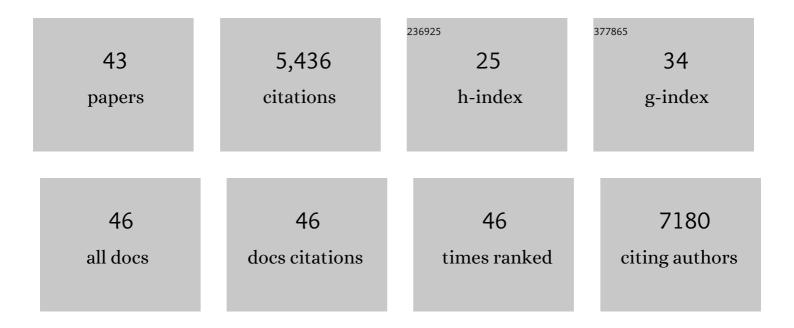
## David Garrick

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Distinct Factors Control Histone Variant H3.3 Localization at Specific Genomic Regions. Cell, 2010, 140, 678-691.	28.9	1,069
2	Repeat-induced gene silencing in mammals. Nature Genetics, 1998, 18, 56-59.	21.4	804
3	Transcription of antisense RNA leading to gene silencing and methylation as a novel cause of human genetic disease. Nature Genetics, 2003, 34, 157-165.	21.4	505
4	Mutations in ATRX, encoding a SWI/SNF-like protein, cause diverse changes in the pattern of DNA methylation. Nature Genetics, 2000, 24, 368-371.	21.4	476
5	ATR-X Syndrome Protein Targets Tandem Repeats and Influences Allele-Specific Expression in a Size-Dependent Manner. Cell, 2010, 143, 367-378.	28.9	365
6	Intragenic Enhancers Act as Alternative Promoters. Molecular Cell, 2012, 45, 447-458.	9.7	237
7	Combinatorial readout of histone H3 modifications specifies localization of ATRX to heterochromatin. Nature Structural and Molecular Biology, 2011, 18, 777-782.	8.2	187
8	An interspecies analysis reveals a key role for unmethylated CpG dinucleotides in vertebrate Polycomb complex recruitment. EMBO Journal, 2012, 31, 317-329.	7.8	173
9	The chromatin-remodeling protein ATRX is critical for neuronal survival during corticogenesis. Journal of Clinical Investigation, 2005, 115, 258-267.	8.2	169
10	Position-dependent variegation of globin transgene expression in mice Proceedings of the National Academy of Sciences of the United States of America, 1995, 92, 5371-5375.	7.1	164
11	Loss of Atrx Affects Trophoblast Development and the Pattern of X-Inactivation in Extraembryonic Tissues. PLoS Genetics, 2006, 2, e58.	3.5	140
12	ldentification of acquired somatic mutations in the gene encoding chromatin-remodeling factor ATRX in the α-thalassemia myelodysplasia syndrome (ATMDS). Nature Genetics, 2003, 34, 446-449.	21.4	132
13	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.	7.1	125
14	The chromatin-remodeling protein ATRX is critical for neuronal survival during corticogenesis. Journal of Clinical Investigation, 2005, 115, 258-267.	8.2	119
15	Age-Associated Decrease of the Histone Methyltransferase SUV39H1 in HSC Perturbs Heterochromatin and B Lymphoid Differentiation. Stem Cell Reports, 2016, 6, 970-984.	4.8	88
16	Age-dependent silencing of globin transgenes in the mouse. Nucleic Acids Research, 1996, 24, 1465-1471.	14.5	84
17	Polycomb eviction as a new distant enhancer function. Genes and Development, 2011, 25, 1583-1588.	5.9	78
18	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.	7.1	77

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19	Generation of bivalent chromatin domains during cell fate decisions. Epigenetics and Chromatin, 2011, 4, 9.	3.9	54
20	A conserved truncated isoform of the ATR-X syndrome protein lacking the SWI/SNF-homology domain. Gene, 2004, 326, 23-34.	2.2	53
21	Variegated Expression of a Globin Transgene Correlates with Chromatin Accessibility But Not Methylation Status. Nucleic Acids Research, 1996, 24, 4902-4909.	14.5	52
22	The role of the polycomb complex in silencing α-globin gene expression in nonerythroid cells. Blood, 2008, 112, 3889-3899.	1.4	51
23	Understanding α-Globin Gene Regulation: Aiming to Improve the Management of Thalassemia. Annals of the New York Academy of Sciences, 2005, 1054, 92-102.	3.8	47
24	Defining the Cause of Skewed X-Chromosome Inactivation in X-Linked Mental Retardation by Use of a Mouse Model. American Journal of Human Genetics, 2007, 80, 1138-1149.	6.2	32
25	Epigenetic Effects on Transgene Expression. , 2001, 158, 351-368.		31
26	A large deletion in the human Â-globin cluster caused by a replication error is associated with an unexpectedly mild phenotype. Human Molecular Genetics, 2008, 17, 3084-3093.	2.9	26
27	The long non-coding RNA CRNDE regulates growth of multiple myeloma cells via an effect on IL6 signalling. Leukemia, 2021, 35, 1710-1721.	7.2	26
28	CpG binding protein (CFP1) occupies open chromatin regions of active genes, including enhancers and non-CpG islands. Epigenetics and Chromatin, 2018, 11, 59.	3.9	19
29	Analysis of Sequence Variation Underlying Tissue-specific Transcription Factor Binding and Gene Expression. Human Mutation, 2013, 34, 1140-1148.	2.5	10
30	Identification of the transcription factor MAZ as a regulator of erythropoiesis. Blood Advances, 2021, 5, 3002-3015.	5.2	8
31	CARâ€₹ cells derived from multiple myeloma patients at diagnosis have improved cytotoxic functions compared to those produced at relapse or following daratumumab treatment. EJHaem, 2022, 3, 970-974.	1.0	8
32	Epigenetic mechanisms. , 0, , 62-74.		7
33	How transcriptional and epigenetic programmes are played out on an individual mammalian gene cluster during lineage commitment and differentiation. Biochemical Society Symposia, 2006, 73, 11-22.	2.7	7
34	Switching genes on and off in haemopoiesis. Biochemical Society Transactions, 2008, 36, 613-618.	3.4	6
35	Aging of Human Haematopoietic Stem Cells. , 2015, , 127-147.		2
36	Hematopoietic Stem Cell Aging and Malignant Hemopathies. , 2018, , 1-13.		2

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#	Article	IF	CITATIONS
37	CTCF, cohesin and higher-order chromatin structure. Epigenomics, 2009, 1, 232.	2.1	1
38	Research Highlights. Epigenomics, 2009, 1, 231-234.	2.1	0
39	Noncoding RNA and Epigenetic Change in Hematopoietic Stem Cell Aging. , 2018, , 1-29.		0
40	Noncoding RNA and Epigenetic Change in Hematopoietic Stem Cell Aging. , 2019, , 1011-1038.		0
41	Hematopoietic Stem Cell Aging and Malignant Hemopathies. , 2020, , 169-181.		0
42	The regulatory interplay of CpG islands and nucleosome remodeling at mammalian primary response genes. Epigenomics, 2009, 1, 233.	2.1	0
43	XNP/ATRX at sites of nucleosome replacement. Epigenomics, 2009, 1, 233-4.	2.1	0