

# Masako Suzuki

## List of Publications by Year in descending order

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Version: 2024-02-01

90  
papers

3,935  
citations

101543

36  
h-index

128289

60  
g-index

105  
all docs

105  
docs citations

105  
times ranked

7547  
citing authors

#	ARTICLE	IF	CITATIONS
1	Cell type-specific chromatin accessibility analysis in the mouse and human brain. <i>Epigenetics</i> , 2022, 17, 202-219.	2.7	13
2	Tet-mediated DNA demethylation regulates specification of hematopoietic stem and progenitor cells during mammalian embryogenesis. <i>Science Advances</i> , 2022, 8, eabm3470.	10.3	13
3	Vitamin D Deficiency During Development Permanently Alters Liver Cell Composition and Function. <i>Frontiers in Endocrinology</i> , 2022, 13, .	3.5	2
4	Genetic Variations of Vitamin A-Absorption and Storage-Related Genes, and Their Potential Contribution to Vitamin A Deficiency Risks Among Different Ethnic Groups. <i>Frontiers in Nutrition</i> , 2022, 9, 861619.	3.7	6
5	Disproportionate Vitamin A Deficiency in Women of Specific Ethnicities Linked to Differences in Allele Frequencies of Vitamin A-Related Polymorphisms. <i>Nutrients</i> , 2021, 13, 1743.	4.1	8
6	Premature differentiation of nephron progenitor cell and dysregulation of gene pathways critical to kidney development in a model of preterm birth. <i>Scientific Reports</i> , 2021, 11, 21667.	3.3	4
7	The SEQC2 epigenomics quality control (EpiQC) study. <i>Genome Biology</i> , 2021, 22, 332.	8.8	20
8	The Role of Egr1 in Driving Ventral Hippocampal Gene Expression Patterns Underlying Sex Hormone-Dependent Changes in Anxiety-Like Behavior Within Females. <i>Biological Psychiatry</i> , 2020, 87, S453.	1.3	0
9	A Cellular Stress Response Induced by the CRISPR-dCas9 Activation System Is Not Heritable Through Cell Divisions. <i>CRISPR Journal</i> , 2020, 3, 188-197.	2.9	2
10	Functional Genomics of the Pediatric Obese Asthma Phenotype Reveal Enrichment of Rho-GTPase Pathways. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 202, 259-274.	5.6	17
11	The shape of gene expression distributions matter: how incorporating distribution shape improves the interpretation of cancer transcriptomic data. <i>BMC Bioinformatics</i> , 2020, 21, 562.	2.6	19
12	Functional genetic variants can mediate their regulatory effects through alteration of transcription factor binding. <i>Nature Communications</i> , 2019, 10, 3472.	12.8	39
13	High-efficiency genomic editing in Epstein-Barr virus-transformed lymphoblastoid B cells using a single-stranded donor oligonucleotide strategy. <i>Communications Biology</i> , 2019, 2, 312.	4.4	5
14	Chromatin organization in the female mouse brain fluctuates across the oestrous cycle. <i>Nature Communications</i> , 2019, 10, 2851.	12.8	68
15	2011 - ENGINEERING A HEMATOPOIETIC STEM CELL NICHE BY REVITALIZING MESENCHYMAL STEM CELLS. <i>Experimental Hematology</i> , 2019, 76, S45.	0.4	0
16	Insights from deconvolution of cell subtype proportions enhance the interpretation of functional genomic data. <i>PLoS ONE</i> , 2019, 14, e0215987.	2.5	21
17	Engineering a haematopoietic stem cell niche by revitalizing mesenchymal stromal cells. <i>Nature Cell Biology</i> , 2019, 21, 560-567.	10.3	74
18	Ascorbic acid-induced TET activation mitigates adverse hydroxymethylcytosine loss in renal cell carcinoma. <i>Journal of Clinical Investigation</i> , 2019, 129, 1612-1625.	8.2	64

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19	Lactate-mediated epigenetic reprogramming regulates formation of human pancreatic cancer-associated fibroblasts. <i>ELife</i> , 2019, 8, .	6.0	103
20	Engineering a Hematopoietic Stem Cell Niche By Revitalizing Mesenchymal Stem Cells with Five Transcription Factors. <i>Blood</i> , 2019, 134, 5004-5004.	1.4	0
21	Intrauterine Hyperglycemia Is Associated with an Impaired Postnatal Response to Oxidative Damage. <i>Stem Cells and Development</i> , 2018, 27, 683-691.	2.1	10
22	Interleukin 2 modulates thymic-derived regulatory T cell epigenetic landscape. <i>Nature Communications</i> , 2018, 9, 5368.	12.8	26
23	Fetalâ€™Not Maternalâ€™APOL1 Genotype Associated with Risk for Preeclampsia in Those with African Ancestry. <i>American Journal of Human Genetics</i> , 2018, 103, 367-376.	6.2	49
24	Mechanisms of establishment and functional significance of DNA demethylation during erythroid differentiation. <i>Blood Advances</i> , 2018, 2, 1833-1852.	5.2	15
25	Detecting, quantifying, and discriminating the mechanism of mosaic chromosomal aneuploidies using MAD-seq. <i>Genome Research</i> , 2018, 28, 1039-1052.	5.5	3
26	O10. Neuronal Chromatin Dynamics and Anxiety-Related Phenotypes Across the Estrous Cycle. <i>Biological Psychiatry</i> , 2018, 83, S112.	1.3	0
27	Selective modulation of local linkages between active transcription and oxidative demethylation activity shapes cardiomyocyte-specific gene-body epigenetic status in mice. <i>BMC Genomics</i> , 2018, 19, 349.	2.8	4
28	Whole-genome bisulfite sequencing with improved accuracy and cost. <i>Genome Research</i> , 2018, 28, 1364-1371.	5.5	64
29	Notch Pathway Is Activated via Genetic and Epigenetic Alterations and Is a Therapeutic Target in Clear Cell Renal Cancer. <i>Journal of Biological Chemistry</i> , 2017, 292, 837-846.	3.4	43
30	Geneticâ€™epigenetic interactions in cis: a major focus in the post-GWAS era. <i>Genome Biology</i> , 2017, 18, 120.	8.8	109
31	In Utero Exposure to a High-Fat Diet Programs Hepatic Hypermethylation and Gene Dysregulation and Development of Metabolic Syndrome in Male Mice. <i>Endocrinology</i> , 2017, 158, 2860-2872.	2.8	42
32	Altered hydroxymethylation is seen at regulatory regions in pancreatic cancer and regulates oncogenic pathways. <i>Genome Research</i> , 2017, 27, 1830-1842.	5.5	51
33	Abstract 1170: Notch pathway is overexpressed and is a therapeutic target in clear cell renal cancer. , 2017, , .		0
34	Loss of <i>MEN1</i> activates DNMT1 implicating DNA hypermethylation as a driver of MEN1 tumorigenesis. <i>Oncotarget</i> , 2016, 7, 12633-12650.	1.8	25
35	Amnion as a surrogate tissue reporter of the effects of maternal preeclampsia on the fetus. <i>Clinical Epigenetics</i> , 2016, 8, 67.	4.1	9
36	An integrative analysis sheds light on methylation profiles. <i>Science Translational Medicine</i> , 2016, 8, .	12.4	0

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37	Balance of nutrients: Is more better?. Science Translational Medicine, 2016, 8, .	12.4	0
38	Post-conversion targeted capture of modified cytosines in mammalian and plant genomes. Nucleic Acids Research, 2015, 43, e81-e81.	14.5	62
39	DNA demethylation by 5-aza-2â€²-deoxycytidine is imprinted, targeted to euchromatin, and has limited transcriptional consequences. Epigenetics and Chromatin, 2015, 8, 11.	3.9	38
40	DNA methylation: A link between genome variation and blood pressure. Science Translational Medicine, 2015, 7, .	12.4	1
41	Genome Wide Methylome Alterations in Lung Cancer. PLoS ONE, 2015, 10, e0143826.	2.5	30
42	Childhood memory. Science Translational Medicine, 2015, 7, .	12.4	0
43	Timing is everything. Science Translational Medicine, 2015, 7, .	12.4	0
44	Diverse diseases, diverse variants. Science Translational Medicine, 2015, 7, .	12.4	0
45	A brite idea for noncoding variants. Science Translational Medicine, 2015, 7, .	12.4	0
46	Above the landscape. Science Translational Medicine, 2015, 7, .	12.4	0
47	Kidney Cancer Is Characterized by Aberrant Methylation of Tissue-Specific Enhancers That Are Prognostic for Overall Survival. Clinical Cancer Research, 2014, 20, 4349-4360.	7.0	60
48	Mosaic Epigenetic Dysregulation of Ectodermal Cells in Autism Spectrum Disorder. PLoS Genetics, 2014, 10, e1004402.	3.5	93
49	DNA Methylation is Developmentally Regulated for Genes Essential for Cardiogenesis. Journal of the American Heart Association, 2014, 3, e000976.	3.7	71
50	Redistribution of H3K27me3 upon DNA hypomethylation results in de-repression of Polycomb target genes. Genome Biology, 2013, 14, R25.	9.6	200
51	Cytosine methylation changes in enhancer regions of core pro-fibrotic genes characterize kidney fibrosis development. Genome Biology, 2013, 14, R108.	9.6	187
52	Hypomethylation of Noncoding DNA Regions and Overexpression of the Long Noncoding RNA, AFAP1-AS1, in Barrett's Esophagus and Esophageal Adenocarcinoma. Gastroenterology, 2013, 144, 956-966.e4.	1.3	216
53	Myeloma Is Characterized by Stage-Specific Alterations in DNA Methylation That Occur Early during Myelomagenesis. Journal of Immunology, 2013, 190, 2966-2975.	0.8	90
54	Genome-wide DNA Methylation Analysis Using Massively Parallel Sequencing Technologies. Seminars in Hematology, 2013, 50, 70-77.	3.4	11

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55	Epigenetic Functions of Smchd1 Repress Gene Clusters on the Inactive X Chromosome and on Autosomes. <i>Molecular and Cellular Biology</i> , 2013, 33, 3150-3165.	2.3	99
56	Large, Male Germ Cell-Specific Hypomethylated DNA Domains With Unique Genomic and Epigenomic Features on the Mouse X Chromosome. <i>DNA Research</i> , 2013, 20, 549-565.	3.4	10
57	Methylome Profiling Reveals Distinct Alterations in Phenotypic and Mutational Subgroups of Myeloproliferative Neoplasms. <i>Cancer Research</i> , 2013, 73, 1076-1085.	0.9	50
58	High Resolution Methylome Analysis Reveals Widespread Functional Hypomethylation during Adult Human Erythropoiesis. <i>Journal of Biological Chemistry</i> , 2013, 288, 8805-8814.	3.4	37
59	Genome-wide hydroxymethylation tested using the HELP-GT assay shows redistribution in cancer. <i>Nucleic Acids Research</i> , 2013, 41, e157-e157.	14.5	69
60	Differential epigenome-wide DNA methylation patterns in childhood obesity-associated asthma. <i>Scientific Reports</i> , 2013, 3, 2164.	3.3	94
61	Lsh regulates LTR retrotransposon repression independently of Dnmt3b function. <i>Genome Biology</i> , 2013, 14, R146.	9.6	54
62	Abstract 4231: Genome wide hydroxymethylation tested using the HELP-GT assay shows redistribution in cancer.. , 2013, , .		0
63	The Wasp System: An open source environment for managing and analyzing genomic data. <i>Genomics</i> , 2012, 100, 345-351.	2.9	24
64	Maternal gametic transmission of translocations or inversions of human chromosome 11p15.5 results in regional DNA hypermethylation and downregulation of CDKN1C expression. <i>Genomics</i> , 2012, 99, 25-35.	2.9	18
65	Epigenetic Silencing of the Circadian Clock Gene CRY1 is Associated with an Indolent Clinical Course in Chronic Lymphocytic Leukemia. <i>PLoS ONE</i> , 2012, 7, e34347.	2.5	44
66	Whole Methylome Explorations Of Paired Lung Tumor And Non-Tumor Clinical Samples. , 2012, , .		0
67	Automated Computational Analysis of Genome-Wide DNA Methylation Profiling Data from HELP-Tagging Assays. <i>Methods in Molecular Biology</i> , 2012, 815, 79-87.	0.9	13
68	Abstract 4230: Senescence reversion contributes to acquired drug resistance. , 2012, , .		1
69	Abstract A38: Whole methylome explorations of paired lung tumor and non-tumor clinical samples. <i>Clinical Cancer Research</i> , 2012, 18, A38-A38.	7.0	0
70	DNA methylation changes in murine breast adenocarcinomas allow the identification of candidate genes for human breast carcinogenesis. <i>Mammalian Genome</i> , 2011, 22, 249-259.	2.2	11
71	Epigenetic changes in B lymphocytes associated with house dust mite allergic asthma. <i>Epigenetics</i> , 2011, 6, 1131-1137.	2.7	62
72	Late-replicating heterochromatin is characterized by decreased cytosine methylation in the human genome. <i>Genome Research</i> , 2011, 21, 1833-1840.	5.5	38

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73	Aberrant Epigenetic and Genetic Marks Are Seen in Myelodysplastic Leukocytes and Reveal Dock4 as a Candidate Pathogenic Gene on Chromosome 7q. <i>Journal of Biological Chemistry</i> , 2011, 286, 25211-25223.	3.4	41
74	Widespread Hypomethylation Occurs Early and Synergizes with Gene Amplification during Esophageal Carcinogenesis. <i>PLoS Genetics</i> , 2011, 7, e1001356.	3.5	112
75	Abstract 4820: Whole methylome explorations of paired tumor and non-tumor clinical samples using HELP. , 2011, , .		0
76	DNA methylation profiling using HpaII tiny fragment enrichment by ligation-mediated PCR (HELP). <i>Methods</i> , 2010, 52, 218-222.	3.8	26
77	Optimized design and data analysis of tag-based cytosine methylation assays. <i>Genome Biology</i> , 2010, 11, R36.	9.6	76
78	Epigenomic Profiling of Myeloproliferative Diseases Reveal Idiopathic Myelofibrosis as An Epigenetically Distinct Subgroup and Highlights the Epigenetic Effects of Jak2V617F Mutation. <i>Blood</i> , 2010, 116, 627-627.	1.4	2
79	A pipeline for the quantitative analysis of CG dinucleotide methylation using mass spectrometry. <i>Bioinformatics</i> , 2009, 25, 2164-2170.	4.1	69
80	High-resolution genome-wide cytosine methylation profiling with simultaneous copy number analysis and optimization for limited cell numbers. <i>Nucleic Acids Research</i> , 2009, 37, 3829-3839.	14.5	141
81	Institutional Profile: The Einstein Center for Epigenomics: studying the role of epigenomic dysregulation in human disease. <i>Epigenomics</i> , 2009, 1, 33-38.	2.1	1
82	Establishment of trophoblast stem cell lines from somatic cell nuclear-transferred embryos. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 16293-16297.	7.1	23
83	Genome-wide and locus-specific DNA hypomethylation in G9a deficient mouse embryonic stem cells. <i>Genes To Cells</i> , 2007, 12, 1-11.	1.2	79
84	Cell type-specific methylation profiles occurring disproportionately in CpG-less regions that delineate developmental similarity. <i>Genes To Cells</i> , 2007, 12, 1123-1132.	1.2	41
85	A new class of tissue-specifically methylated regions involving entire CpG islands in the mouse. <i>Genes To Cells</i> , 2007, 12, 1305-1314.	1.2	27
86	Comparative isoschizomer profiling of cytosine methylation: The HELP assay. <i>Genome Research</i> , 2006, 16, 1046-1055.	5.5	355
87	Equivalency of Nuclear Transfer-Derived Embryonic Stem Cells to Those Derived from Fertilized Mouse Blastocysts. <i>Stem Cells</i> , 2006, 24, 2023-2033.	3.2	156
88	DNA Methylation Profiles of Donor Nuclei Cells and Tissues of Cloned Bovine Fetuses. <i>Journal of Reproduction and Development</i> , 2006, 52, 259-266.	1.4	34
89	Regulation of conceptus adhesion by endometrial CXC chemokines during the implantation period in sheep. <i>Molecular Reproduction and Development</i> , 2006, 73, 850-858.	2.0	52
90	Preference of DNA Methyltransferases for CpG Islands in Mouse Embryonic Stem Cells. <i>Genome Research</i> , 2004, 14, 1733-1740.	5.5	70