

Melanie Ehrlich

List of Publications by Year in descending order

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109
papers

11,764
citations

53939

47
h-index

36203

101
g-index

115
all docs

115
docs citations

115
times ranked

13013
citing authors

#	ARTICLE	IF	CITATIONS
1	Epigenomic and Transcriptomic Prioritization of Candidate Obesity-Risk Regulatory GWAS SNPs. <i>International Journal of Molecular Sciences</i> , 2022, 23, 1271.	1.8	5
2	Risks and rewards of big-data in epigenomics research: an interview with Melanie Ehrlich. <i>Epigenomics</i> , 2022, 14, 351-358.	1.0	3
3	Epigenetics of Mitochondria-Associated Genes in Striated Muscle. <i>Epigenomes</i> , 2022, 6, 1.	0.8	3
4	Epigenetics and expression of key genes associated with cardiac fibrosis: <i>NLRP3</i> , <i>MMP2</i> , <i>MMP9</i> , <i>CCN2/CTGF</i> and <i>AGT</i> . <i>Epigenomics</i> , 2021, 13, 219-234.	1.0	16
5	Prioritization of Osteoporosis-Associated Genome-wide Association Study (GWAS) Single-Nucleotide Polymorphisms (SNPs) Using Epigenomics and Transcriptomics. <i>JBMR Plus</i> , 2021, 5, e10481.	1.3	14
6	Epigenetics of Muscle- and Brain-Specific Expression of <i>KLHL</i> Family Genes. <i>International Journal of Molecular Sciences</i> , 2020, 21, 8394.	1.8	14
7	Epigenetics of Skeletal Muscle-Associated Genes in the <i>ASB</i> , <i>LRRC</i> , <i>TMEM</i> , and <i>OSBPL</i> Gene Families. <i>Epigenomes</i> , 2020, 4, 1.	0.8	23
8	Osteoporosis- and obesity-risk interrelationships: an epigenetic analysis of GWAS-derived SNPs at the developmental gene <i>TBX15</i> . <i>Epigenetics</i> , 2020, 15, 728-749.	1.3	11
9	DNA hypermethylation in disease: mechanisms and clinical relevance. <i>Epigenetics</i> , 2019, 14, 1141-1163.	1.3	221
10	Tissue-specific epigenetics of atherosclerosis-related <i>ANGPT</i> and <i>ANGPTL</i> genes. <i>Epigenomics</i> , 2019, 11, 169-186.	1.0	30
11	Data showing atherosclerosis-associated differentially methylated regions are often at enhancers. <i>Data in Brief</i> , 2019, 23, 103812.	0.5	9
12	Atherosclerosis-associated differentially methylated regions can reflect the disease phenotype and are often at enhancers. <i>Atherosclerosis</i> , 2019, 280, 183-191.	0.4	29
13	Developmentally linked human DNA hypermethylation is associated with down-modulation, repression, and upregulation of transcription. <i>Epigenetics</i> , 2018, 13, 275-289.	1.3	31
14	The Sequence of Two Bacteriophages with Hypermodified Bases Reveals Novel Phage-Host Interactions. <i>Viruses</i> , 2018, 10, 217.	1.5	5
15	Association of 5-hydroxymethylation and 5-methylation of DNA cytosine with tissue-specific gene expression. <i>Epigenetics</i> , 2017, 12, 123-138.	1.3	61
16	Incomplete MyoD-induced transdifferentiation is associated with chromatin remodeling deficiencies. <i>Nucleic Acids Research</i> , 2017, 45, 11684-11699.	6.5	27
17	Epigenetics of the myotonic dystrophy-associated <i>DMPK</i> gene neighborhood. <i>Epigenomics</i> , 2016, 8, 13-31.	1.0	22
18	DNA Hypomethylation in Intragenic and Intergenic Enhancer Chromatin of Muscle-Specific Genes Usually Correlates with their Expression. <i>Yale Journal of Biology and Medicine</i> , 2016, 89, 441-455.	0.2	22

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19	Tissue-specific epigenetics in gene neighborhoods: myogenic transcription factor genes. <i>Human Molecular Genetics</i> , 2015, 24, 4660-4673.	1.4	21
20	Exploring DNA methylation changes in promoter, intragenic, and intergenic regions as early and late events in breast cancer formation. <i>BMC Cancer</i> , 2015, 15, 816.	1.1	86
21	Myogenic Differential Methylation: Diverse Associations with Chromatin Structure. <i>Biology</i> , 2014, 3, 426-451.	1.3	15
22	Notch signaling genes. <i>Epigenetics</i> , 2014, 9, 842-850.	1.3	49
23	DNA cytosine methylation and hydroxymethylation at the borders. <i>Epigenomics</i> , 2014, 6, 563-566.	1.0	38
24	DNA methylation and differentiation: HOX genes in muscle cells. <i>Epigenetics and Chromatin</i> , 2013, 6, 25.	1.8	49
25	DNA methylation and differentiation: silencing, upregulation and modulation of gene expression. <i>Epigenomics</i> , 2013, 5, 553-568.	1.0	176
26	DNA Hypomethylation and Hemimethylation in Cancer. <i>Advances in Experimental Medicine and Biology</i> , 2013, 754, 31-56.	0.8	126
27	Early de novo DNA methylation and prolonged demethylation in the muscle lineage. <i>Epigenetics</i> , 2013, 8, 317-332.	1.3	85
28	Modeling, simulation and analysis of methylation profiles from reduced representation bisulfite sequencing experiments. <i>Statistical Applications in Genetics and Molecular Biology</i> , 2013, 12, 723-42.	0.2	22
29	Deciphering transcription dysregulation in FSH muscular dystrophy. <i>Journal of Human Genetics</i> , 2012, 57, 477-484.	1.1	5
30	Diagnosis by sequencing: correction of misdiagnosis from FSHD2 to LGMD2A by whole-exome analysis. <i>European Journal of Human Genetics</i> , 2012, 20, 999-1003.	1.4	32
31	DNA ploidy, nuclear size, proliferation index and DNA-hypomethylation in ovarian cancer. <i>Gynecologic Oncology</i> , 2011, 121, 24-31.	0.6	49
32	Gene expression during normal and FSHD myogenesis. <i>BMC Medical Genomics</i> , 2011, 4, 67.	0.7	81
33	DNA Replication Timing Is Maintained Genome-Wide in Primary Human Myoblasts Independent of D4Z4 Contraction in FSH Muscular Dystrophy. <i>PLoS ONE</i> , 2011, 6, e27413.	1.1	21
34	Analysis of allele-specific RNA transcription in FSHD by RNA-DNA FISH in single myonuclei. <i>European Journal of Human Genetics</i> , 2010, 18, 448-456.	1.4	34
35	Hemimethylation footprints of DNA demethylation in cancer. <i>Epigenetics</i> , 2009, 4, 165-175.	1.3	32
36	Modeling Dependence in Methylation Patterns with Application to Ovarian Carcinomas. <i>Statistical Applications in Genetics and Molecular Biology</i> , 2009, 8, 1-27.	0.2	12

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37	DNaseI hypersensitivity at gene-poor, FSH dystrophy-linked 4q35.2. <i>Nucleic Acids Research</i> , 2009, 37, 7381-7393.	6.5	15
38	DNA hypomethylation in cancer cells. <i>Epigenomics</i> , 2009, 1, 239-259.	1.0	837
39	High-resolution analysis of DNaseI hypersensitive sites in a muscular dystrophy-linked subtelomeric region of 4q. <i>FASEB Journal</i> , 2009, 23, 487.1.	0.2	0
40	ICF, an immunodeficiency syndrome: DNA methyltransferase 3B involvement, chromosome anomalies, and gene dysregulation. <i>Autoimmunity</i> , 2008, 41, 253-271.	1.2	130
41	Epigenetics of a tandem DNA repeat: chromatin DNaseI sensitivity and opposite methylation changes in cancers. <i>Nucleic Acids Research</i> , 2008, 36, 2196-2207.	6.5	51
42	Cancer-Linked DNA Hypermethylation and Hypomethylation. , 2008, , 235-252.		0
43	RNAPol-ChIP analysis of transcription from FSHD-linked tandem repeats and satellite DNA. <i>Biochimica Et Biophysica Acta Gene Regulatory Mechanisms</i> , 2007, 1769, 29-40.	2.4	62
44	Hybridization analysis of D4Z4 repeat arrays linked to FSHD. <i>Chromosoma</i> , 2007, 116, 107-116.	1.0	25
45	Immunodeficiency, centromeric region instability, facial anomalies syndrome (ICF). <i>Orphanet Journal of Rare Diseases</i> , 2006, 1, 2.	1.2	109
46	Quantitative analysis of associations between DNA hypermethylation, hypomethylation, and DNMT RNA levels in ovarian tumors. <i>Oncogene</i> , 2006, 25, 2636-2645.	2.6	129
47	Frequent DNA hypomethylation of human juxtacentromeric BAGE loci in cancer. <i>Genes Chromosomes and Cancer</i> , 2005, 43, 11-24.	1.5	44
48	Interphase chromosomal abnormalities and mitotic missegregation of hypomethylated sequences in ICF syndrome cells. <i>Chromosoma</i> , 2005, 114, 118-126.	1.0	49
49	The Controversial Denouement of Vertebrate DNA Methylation Research. <i>Biochemistry (Moscow)</i> , 2005, 70, 568-575.	0.7	26
50	DNA Hypo- vs. Hypermethylation in Cancer. , 2005, , 31-41.		3
51	Both Hypomethylation and Hypermethylation in a 0.2-kb Region of a DNA Repeat in Cancer. <i>Molecular Cancer Research</i> , 2005, 3, 617-626.	1.5	40
52	A DNA repeat, NBL2, is hypermethylated in some cancers but hypomethylated in others. <i>Cancer Biology and Therapy</i> , 2005, 4, 446-454.	1.5	40
53	Analysis of repetitive element DNA methylation by MethyLight. <i>Nucleic Acids Research</i> , 2005, 33, 6823-6836.	6.5	636
54	DNA Methylation and Cancer-associated Genetic Instability. , 2005, 570, 363-392.		23

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55	DNA hypomethylation is prevalent even in low-grade breast cancers. <i>Cancer Biology and Therapy</i> , 2004, 3, 1225-1231.	1.5	117
56	DNA Hypomethylation and Ovarian Cancer Biology. <i>Cancer Research</i> , 2004, 64, 4472-4480.	0.4	221
57	Cytogenetic and immuno-FISH analysis of the 4q subtelomeric region, which is associated with facioscapulohumeral muscular dystrophy. <i>Chromosoma</i> , 2004, 112, 350-359.	1.0	36
58	Histone modification in constitutive heterochromatin versus unexpressed euchromatin in human cells. <i>Journal of Cellular Biochemistry</i> , 2004, 93, 286-300.	1.2	22
59	Satellite DNA hypomethylation in karyotyped Wilms tumors. <i>Cancer Genetics and Cytogenetics</i> , 2003, 141, 97-105.	1.0	81
60	Expression of various genes is controlled by DNA methylation during mammalian development. <i>Journal of Cellular Biochemistry</i> , 2003, 88, 899-910.	1.2	204
61	The ICF syndrome, a DNA methyltransferase 3B deficiency and immunodeficiency disease. <i>Clinical Immunology</i> , 2003, 109, 17-28.	1.4	179
62	Testing the position-effect variegation hypothesis for facioscapulohumeral muscular dystrophy by analysis of histone modification and gene expression in subtelomeric 4q. <i>Human Molecular Genetics</i> , 2003, 12, 2909-2921.	1.4	138
63	DNA Hypomethylation, Cancer, the Immunodeficiency, Centromeric Region Instability, Facial Anomalies Syndrome and Chromosomal Rearrangements. <i>Journal of Nutrition</i> , 2002, 132, 2424S-2429S.	1.3	113
64	DNA methylation in cancer: too much, but also too little. <i>Oncogene</i> , 2002, 21, 5400-5413.	2.6	1,390
65	Hypomethylation and hypermethylation of DNA in Wilms tumors. <i>Oncogene</i> , 2002, 21, 6694-6702.	2.6	165
66	Methylation of the FSHD Syndrome-Linked Subtelomeric Repeat in Normal and FSHD Cell Cultures and Tissues. <i>Molecular Genetics and Metabolism</i> , 2001, 74, 322-331.	0.5	31
67	DNA methyltransferase 3B mutations linked to the ICF syndrome cause dysregulation of lymphogenesis genes. <i>Human Molecular Genetics</i> , 2001, 10, 2917-2931.	1.4	108
68	Hypersensitivity to radiation-induced non-apoptotic and apoptotic death in cell lines from patients with the ICF chromosome instability syndrome. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2000, 456, 1-15.	0.4	19
69	Whole-genome methylation scan in ICF syndrome: hypomethylation of non-satellite DNA repeats D4Z4 and NBL2. <i>Human Molecular Genetics</i> , 2000, 9, 597-604.	1.4	153
70	Demethylation and expression of methylated plasmid DNA stably transfected into HeLa cells. <i>Nucleic Acids Research</i> , 1999, 27, 2332-2338.	6.5	20
71	A Methylation-responsive MDBP/RFX Site Is in the First Exon of the Collagen $\alpha 2(I)$ Promoter. <i>Journal of Biological Chemistry</i> , 1999, 274, 36649-36655.	1.6	37
72	Frequent Hypomethylation in Wilms Tumors of Pericentromeric DNA in Chromosomes 1 and 16. <i>Cancer Genetics and Cytogenetics</i> , 1999, 109, 34-39.	1.0	173

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73	Satellite DNA hypomethylation vs. overall genomic hypomethylation in ovarian epithelial tumors of different malignant potential. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 1999, 423, 91-101.	0.4	136
74	Hypomethylation of pericentromeric DNA in breast adenocarcinomas. , 1998, 77, 833-838.		218
75	Hypomethylation of pericentromeric DNA in breast adenocarcinomas. , 1998, 77, 833.		3
76	DNA demethylation and pericentromeric rearrangements of chromosome 1. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 1997, 379, 33-41.	0.4	161
77	Frequency of BCL-2/JH translocations in peripheral blood of follicular lymphoma patients. , 1997, 55, 205-207.		1
78	Increasing binding of a transcription factor immediately downstream of the cap site of a cytomegalovirus gene represses expression. <i>Nucleic Acids Research</i> , 1995, 23, 3026-3033.	6.5	26
79	A mutation in the 5' untranslated region of the human β -galactosidase A gene in high-activity variants inhibits specific protein binding. <i>FEBS Letters</i> , 1995, 371, 181-184.	1.3	9
80	Effect of DNA methylation on the binding of vertebrate and plant proteins to DNA. , 1993, 64, 145-168.		38
81	A broad bean cDNA clone encoding a DNA-binding protein resembling mammalian CREB in its sequence specificity and DNA methylation sensitivity. <i>Gene</i> , 1992, 117, 169-178.	1.0	29
82	DNA methylation inhibits propagation of tomato golden mosaic virus DNA in transfected protoplasts. <i>Plant Molecular Biology</i> , 1992, 18, 703-712.	2.0	77
83	Transfection of heteroduplexes containing uracil $\frac{1}{2}$ guanine or thymine $\frac{1}{2}$ guanine mispairs into plant cells. <i>Plant Molecular Biology</i> , 1992, 20, 123-131.	2.0	13
84	CpG methylation inhibits binding of several sequence-specific DNA-binding proteins from pea, wheat, soybean and cauliflower. <i>Plant Molecular Biology</i> , 1991, 17, 111-123.	2.0	118
85	Spontaneous deamination of cytosine and 5-methylcytosine residues in DNA and replacement of 5-methylcytosine residues with cytosine residues. <i>Mutation Research - Reviews in Genetic Toxicology</i> , 1990, 238, 277-286.	3.0	110
86	Binding sites in mammalian genes and viral gene regulatory regions recognized by methylated DNA-binding protein. <i>Nucleic Acids Research</i> , 1990, 18, 6253-6260.	6.5	60
87	Related sites in human and herpesvirus DNA recognized by methylated DNA-binding protein from human placenta. <i>Nucleic Acids Research</i> , 1989, 17, 1459-1474.	6.5	37
88	Methylated DNA-binding protein is present in various mammalian cell types. <i>Nucleic Acids Research</i> , 1988, 16, 8029-8044.	6.5	43
89	Hypermethylation of human DNA sequences in embryonal carcinoma cells and somatic tissues but not in sperm. <i>Nucleic Acids Research</i> , 1987, 15, 9429-9449.	6.5	45
90	A human DNA-binding protein is methylation-specific and sequence-specific. <i>Nucleic Acids Research</i> , 1986, 14, 1599-1614.	6.5	52

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91	DNA cytosine methylation and heat-induced deamination. <i>Bioscience Reports</i> , 1986, 6, 387-393.	1.1	166
92	Methylated DNA-binding protein from human placenta recognizes specific methylated sites on several prokaryotic DNAs. <i>Nucleic Acids Research</i> , 1986, 14, 9843-9860.	6.5	50
93	Effect of site-specific DNA methylation and mutagenesis on recognition by methylated DNA-binding protein from human placenta. <i>Nucleic Acids Research</i> , 1986, 14, 8387-8397.	6.5	42
94	The accessibility of 5-methylcytosine to specific antibodies in double-stranded DNA of <i>Xanthomonas</i> phage XP12. <i>FEBS Journal</i> , 1985, 152, 115-121.	0.2	18
95	DNA methylation in thermophilic bacteria: N4-methylcytosine, 5-methylcytosine, and N5methyladenine. <i>Nucleic Acids Research</i> , 1985, 13, 1399-1412.	6.5	143
96	Isolation of in Vitro Synthesized Covalently Closed Circular Double-Stranded DNA by Selective Denaturation and Filtration. <i>Preparative Biochemistry and Biotechnology</i> , 1984, 14, 485-497.	0.4	6
97	Quantitative reversed-phase high-performance liquid chromatography of major and modified nucleosides in dna. <i>Journal of Chromatography A</i> , 1984, 301, 199-219.	1.8	187
98	A protein from human placental nuclei binds preferentially to 5-methylcytosine-rich DNA. <i>Nature</i> , 1984, 308, 293-295.	13.7	102
99	Human placental DNA methyltransferase: DNA substrate and DNA binding specificity. <i>Nucleic Acids Research</i> , 1984, 12, 3473-3490.	6.5	38
100	Tissue-specific differences in DNA methylation in various mammals. <i>Biochimica Et Biophysica Acta Gene Regulatory Mechanisms</i> , 1983, 740, 212-219.	2.4	185
101	The 5-methylcytosine content of DNA from human tumors. <i>Nucleic Acids Research</i> , 1983, 11, 6883-6894.	6.5	797
102	The 5-Methylcytosine content of highly repeated sequences in human DNA. <i>Nucleic Acids Research</i> , 1983, 11, 3087-3095.	6.5	133
103	Digestion of highly modified bacteriophage DNA by restriction endonucleases. <i>Nucleic Acids Research</i> , 1982, 10, 1579-1591.	6.5	125
104	Heat- and alkali-induced deamination of 5-methylcytosine and cytosine residues in DNA. <i>Biochimica Et Biophysica Acta Gene Regulatory Mechanisms</i> , 1982, 697, 371-377.	2.4	121
105	Amount and distribution of 5-methylcytosine in human DNA from different types of tissues or cells. <i>Nucleic Acids Research</i> , 1982, 10, 2709-2721.	6.5	917
106	Quantitative reversed-phase high performance liquid chromatographic determination of major and modified deoxyribonucleosides in DNA. <i>Nucleic Acids Research</i> , 1980, 8, 4763-4776.	6.5	303
107	Comparison of bisulfite modification of 5-methyldeoxycytidine and deoxycytidine residues. <i>Nucleic Acids Research</i> , 1980, 8, 4777-4790.	6.5	240
108	Unusual properties of the DNA from <i>Xanthomonas</i> phage XP-12 in which 5-methylcytosine completely replaces cytosine. <i>Nucleic Acids and Protein Synthesis</i> , 1975, 395, 109-119.	1.7	97

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109	EFFECT OF BASE SEQUENCE ON THE ULTRAVIOLET IRRADIATION PRODUCTS OF DOUBLE-STRANDED POLYNUCLEOTIDES CONTAINING BROMOURACIL AND ADENINE. <i>Photochemistry and Photobiology</i> , 1974, 20, 159-165.	1.3	9