

Richard P Woychik

List of Publications by Year in descending order

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78
papers

8,850
citations

61945

43
h-index

66879

78
g-index

84
all docs

84
docs citations

84
times ranked

7555
citing authors

#	ARTICLE	IF	CITATIONS
1	A brain-specific <i>pgc1</i> fusion transcript affects gene expression and behavioural outcomes in mice. <i>Life Science Alliance</i> , 2021, 4, e202101122.	1.3	2
2	Rapid Scaling Up of Covid-19 Diagnostic Testing in the United States – The NIH RADx Initiative. <i>New England Journal of Medicine</i> , 2020, 383, 1071-1077.	13.9	182
3	Single Nucleotide Resolution Analysis Reveals Pervasive, Long-Lasting DNA Methylation Changes by Developmental Exposure to a Mitochondrial Toxicant. <i>Cell Reports</i> , 2020, 32, 108131.	2.9	22
4	Mitochondrial acetyl-CoA reversibly regulates locus-specific histone acetylation and gene expression. <i>Life Science Alliance</i> , 2019, 2, e201800228.	1.3	35
5	A Leveraged Signal-to-Noise Ratio (LSTNR) Method to Extract Differentially Expressed Genes and Multivariate Patterns of Expression From Noisy and Low-Replication RNAseq Data. <i>Frontiers in Genetics</i> , 2018, 9, 176.	1.1	13
6	Mitochondrial nicotinamide adenine dinucleotide reduced (NADH) oxidation links the tricarboxylic acid (TCA) cycle with methionine metabolism and nuclear DNA methylation. <i>PLoS Biology</i> , 2018, 16, e2005707.	2.6	77
7	A Novel Analytical Strategy to Identify Fusion Transcripts between Repetitive Elements and Protein Coding-Exons Using RNA-Seq. <i>PLoS ONE</i> , 2016, 11, e0159028.	1.1	11
8	TCA Cycle and Mitochondrial Membrane Potential Are Necessary for Diverse Biological Functions. <i>Molecular Cell</i> , 2016, 61, 199-209.	4.5	396
9	Unraveling the Health Effects of Environmental Mixtures: An NIEHS Priority. <i>Environmental Health Perspectives</i> , 2013, 121, A6-8.	2.8	147
10	Laser surgery for mouse geneticists. <i>Nature Biotechnology</i> , 2007, 25, 59-60.	9.4	10
11	The Knockout Mouse Project. <i>Nature Genetics</i> , 2004, 36, 921-924.	9.4	556
12	Liver-specific expression of the agouti gene in transgenic mice promotes liver carcinogenesis in the absence of obesity and diabetes. <i>Molecular Cancer</i> , 2004, 3, 17.	7.9	11
13	Loss of the Tg737 protein results in skeletal patterning defects. <i>Developmental Dynamics</i> , 2003, 227, 78-90.	0.8	121
14	Our small relative. <i>Nature Genetics</i> , 2003, 33, 3-4.	9.4	12
15	Massively parallel signature sequencing (MPSS) as a tool for in-depth quantitative gene expression profiling in all organisms. <i>Briefings in Functional Genomics</i> , 2002, 1, 95-104.	1.3	134
16	Molecular and Phenotypic Analysis of 25 Recessive, Homozygous-Viable Alleles at the Mouse <i>agouti</i> Locus. <i>Genetics</i> , 2002, 160, 659-674.	1.2	37
17	Expression of <i>Pcdh15</i> in the inner ear, nervous system and various epithelia of the developing embryo. <i>Mechanisms of Development</i> , 2001, 105, 163-166.	1.7	34
18	Phenotypic variations of <i>orpk</i> mutation and chromosomal localization of modifiers influencing kidney phenotype. <i>Physiological Genomics</i> , 2001, 7, 127-134.	1.0	16

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19	The mouse Ames waltzer hearing-loss mutant is caused by mutation of Pcdh15, a novel protocadherin gene. <i>Nature Genetics</i> , 2001, 27, 99-102.	9.4	276
20	Mutations in the novel protocadherin PCDH15 cause Usher syndrome type 1F. <i>Human Molecular Genetics</i> , 2001, 10, 1709-1718.	1.4	257
21	Genotype-based screen for ENU-induced mutations in mouse embryonic stem cells. <i>Nature Genetics</i> , 2000, 24, 314-317.	9.4	156
22	Persistent Hyperplastic Tunica Vasculosa Lentis and Persistent Hyperplastic Primary Vitreous in Transgenic Line TgN3261Rpw. <i>Veterinary Pathology</i> , 2000, 37, 422-427.	0.8	13
23	Neuroepithelial defects of the inner ear in a new allele of the mouse mutation Ames waltzer. <i>Hearing Research</i> , 2000, 148, 181-191.	0.9	43
24	An agouti mutation lacking the basic domain induces yellow pigmentation but not obesity in transgenic mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1999, 96, 8579-8584.	3.3	30
25	Effective chemical mutagenesis in FVB/N mice requires low doses of ethylnitrosourea. <i>Mammalian Genome</i> , 1999, 10, 308-310.	1.0	24
26	Alternative processing of the human and mouse Raly genes. <i>Biochimica Et Biophysica Acta Gene Regulatory Mechanisms</i> , 1999, 1447, 107-112.	2.4	8
27	Functional genomics in the post-genome era. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 1998, 400, 3-14.	0.4	45
28	Utilization of microhomologous recombination in yeast to generate targeting constructs for mammalian genes. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 1998, 401, 11-25.	0.4	6
29	The molecular biology of polycystic kidney disease. <i>Pediatric Nephrology</i> , 1998, 12, 721-726.	0.9	21
30	Characterization of Growth Factor Responsiveness and Alterations in Growth Factor Homeostasis Involved in the Tumorigenic Conversion of Mouse Oval Cells. <i>Growth Factors</i> , 1998, 15, 81-94.	0.5	16
31	Epidermal growth factor receptor activity mediates renal cyst formation in polycystic kidney disease.. <i>Journal of Clinical Investigation</i> , 1998, 101, 935-939.	3.9	178
32	The Role of the agouti Gene in the Yellow Obese Syndrome ., <i>Journal of Nutrition</i> , 1997, 127, 1902S-1907S.	1.3	143
33	Using Targeted Large Deletions and High-EfficiencyN-Ethyl-N-nitrosourea Mutagenesis for Functional Analyses of the Mammalian Genome. <i>Methods</i> , 1997, 13, 423-436.	1.9	76
34	Combined effects of insulin treatment and adipose tissue-specific agouti expression on the development of obesity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1997, 94, 919-922.	3.3	72
35	Agouti regulation of intracellular calcium: role of melanocortin receptors. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , 1997, 272, E379-E384.	1.8	41
36	The tetratricopeptide repeat containing Tg737 gene is a liver neoplasia tumor suppressor gene. <i>Oncogene</i> , 1997, 15, 1797-1803.	2.6	33

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37	Efficacy of taxol in the orpk mouse model of polycystic kidney disease. <i>Pediatric Nephrology</i> , 1997, 11, 728-733.	0.9	24
38	Role of the agouti gene in obesity. <i>Journal of Endocrinology</i> , 1997, 155, 207-209.	1.2	51
39	Upregulation of adipocyte metabolism by agouti protein: possible paracrine actions in yellow mouse obesity. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , 1996, 270, E192-E196.	1.8	67
40	The effects of calcium channel blockade on agouti-induced obesity. <i>FASEB Journal</i> , 1996, 10, 1646-1652.	0.2	85
41	Functional correction of renal defects in a mouse model for ARPKD through expression of the cloned wild-type Tg737 cDNA. <i>Kidney International</i> , 1996, 50, 1240-1248.	2.6	34
42	Ectopic expression of the agouti gene in transgenic mice causes obesity, features of type II diabetes, and yellow fur.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1995, 92, 4728-4732.	3.3	273
43	Agouti regulation of intracellular calcium: role in the insulin resistance of viable yellow mice.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1995, 92, 4733-4737.	3.3	141
44	Analysis of Polymerase Chain Reaction-Amplified DNA Products by Mass Spectrometry Using Matrix-Assisted Laser Desorption and Electrospray: Current Status. <i>Analytical Biochemistry</i> , 1995, 230, 205-214.	1.1	67
45	Sequence analysis of the human hTg737 gene and its polymorphic sites in patients with autosomal recessive polycystic kidney disease. <i>Mammalian Genome</i> , 1995, 6, 805-808.	1.0	15
46	Forefronts in Nephrology: The molecular basis of renal cystic disease. <i>Kidney International</i> , 1995, 47, 732.	2.6	0
47	Deficiency of the $\beta 3$ subunit of the type A α -aminobutyric acid receptor causes cleft palate in mice. <i>Nature Genetics</i> , 1995, 11, 344-346.	9.4	118
48	Characterization of the human homologue of the mouse Tg737 candidate polycystic kidney disease gene. <i>Human Molecular Genetics</i> , 1995, 4, 559-567.	1.4	43
49	Molecular analysis of reverse mutations from nonagouti (a) to black-and-tan (a(t)) and white-bellied agouti (Aw) reveals alternative forms of agouti transcripts.. <i>Genes and Development</i> , 1994, 8, 481-490.	2.7	114
50	Candidate gene associated with a mutation causing recessive polycystic kidney disease in mice. <i>Science</i> , 1994, 264, 1329-1333.	6.0	336
51	Agouti protein is an antagonist of the melanocyte-stimulating-hormone receptor. <i>Nature</i> , 1994, 371, 799-802.	13.7	999
52	Differential expression of a new dominant agouti allele (Aiapy) is correlated with methylation state and is influenced by parental lineage.. <i>Genes and Development</i> , 1994, 8, 1463-1472.	2.7	262
53	Molecular structure and chromosomal mapping of the human homolog of the agouti gene.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1994, 91, 9760-9764.	3.3	165
54	A molecular model for the genetic and phenotypic characteristics of the mouse lethal yellow (Ay) mutation.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1994, 91, 2562-2566.	3.3	188

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55	TBASE: a computerized database for transgenic animals and targeted mutations. <i>Nature</i> , 1993, 363, 375-376.	13.7	23
56	The embryonic lethality of homozygous lethal yellow mice (Ay/Ay) is associated with the disruption of a novel RNA-binding protein.. <i>Genes and Development</i> , 1993, 7, 1203-1213.	2.7	184
57	Scanning tunneling microscopy of DNA: The chemical modification of gold surfaces for immobilization of DNA. <i>Journal of Vacuum Science and Technology A: Vacuum, Surfaces and Films</i> , 1992, 10, 591-595.	0.9	33
58	Immobilization of DNA for scanning probe microscopy.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1992, 89, 10129-10133.	3.3	53
59	Molecular characterization of the mouse agouti locus. <i>Cell</i> , 1992, 71, 1195-1204.	13.5	802
60	Scanning tunneling microscopy of DNA: a novel technique using radiolabeled DNA to evaluate chemically mediated attachment of DNA to surfaces. <i>Ultramicroscopy</i> , 1992, 42-44, 1088-1094.	0.8	7
61	An approach to the use of stable isotopes for DNA sequencing. <i>Genomics</i> , 1991, 9, 51-59.	1.3	29
62	Potential application of sputter-initiated resonance ionization spectroscopy for DNA sequencing. <i>Analytical Chemistry</i> , 1991, 63, 402-407.	3.2	28
63	Molecular characterization of a region of DNA associated with mutations at the agouti locus in the mouse.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1991, 88, 8062-8066.	3.3	34
64	Electrochemically induced adsorption of radio-labeled DNA on gold and HOPG substrates for STM investigations. <i>Ultramicroscopy</i> , 1991, 38, 253-264.	0.8	14
65	Resonance ionization spectroscopy for multiplex sequencing of Tin-labeled DNA. <i>Genetic Analysis, Techniques and Applications</i> , 1991, 8, 167-170.	1.5	5
66	Molecular and genetic characterization of a radiation-induced structural rearrangement in mouse chromosome 2 causing mutations at the limb deformity and agouti loci.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1990, 87, 2588-2592.	3.3	58
67	'Formins': proteins deduced from the alternative transcripts of the limb deformity gene. <i>Nature</i> , 1990, 346, 850-853.	13.7	195
68	Disruption of formin-encoding transcripts in two mutant limb deformity alleles. <i>Nature</i> , 1990, 346, 853-855.	13.7	101
69	Location of the gene involving the Small eye mutation on mouse chromosome 2 suggests homology with human aniridia 2 (AN2). <i>Genomics</i> , 1990, 7, 270-275.	1.3	48
70	Structure and Regulated Expression of Bovine Prolactin and Bovine Growth Hormone Genes. <i>Advances in Experimental Medicine and Biology</i> , 1986, 205, 281-299.	0.8	3
71	Differential Effects of Polyadenylation Regions on Gene Expression in Mammalian Cells. <i>DNA and Cell Biology</i> , 1986, 5, 115-122.	5.1	79
72	Synthesis of bovine growth hormone in primates by using a herpesvirus vector.. <i>Molecular and Cellular Biology</i> , 1985, 5, 2796-2803.	1.1	44

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73	An inherited limb deformity created by insertional mutagenesis in a transgenic mouse. <i>Nature</i> , 1985, 318, 36-40.	13.7	300
74	Characterization of the Bovine Prolactin Gene. <i>DNA and Cell Biology</i> , 1984, 3, 237-249.	5.1	66
75	Requirement for the 3' flanking region of the bovine growth hormone gene for accurate polyadenylation.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1984, 81, 3944-3948.	3.3	132
76	Variation in the polyadenylation site of bovine prolactin mRNA.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1982, 79, 223-227.	3.3	76
77	Cloning and nucleotide sequencing of the bovine growth hormone gene. <i>Nucleic Acids Research</i> , 1982, 10, 7197-7210.	6.5	214
78	Quantitation of the interaction of Escherichia coli RNA polymerase holoenzyme with double-helical DNA using a thermodynamically rigorous centrifugation method. <i>Biochemistry</i> , 1981, 20, 250-256.	1.2	30