Michael Snyder

List of Publications by Year in descending order

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204 papers 72,859 citations

102 h-index 202 g-index

206 all docs

206 docs citations

206 times ranked 79002 citing authors

#	Article	IF	CITATIONS
1	RNA-Seq: a revolutionary tool for transcriptomics. Nature Reviews Genetics, 2009, 10, 57-63.	7.7	10,529
2	Identification and analysis of functional elements in 1% of the human genome by the ENCODE pilot project. Nature, 2007, 447, 799-816.	13.7	4,709
3	Functional profiling of the Saccharomyces cerevisiae genome. Nature, 2002, 418, 387-391.	13.7	3,938
4	Functional Characterization of the S. cerevisiae Genome by Gene Deletion and Parallel Analysis. Science, 1999, 285, 901-906.	6.0	3,761
5	Annotation of functional variation in personal genomes using RegulomeDB. Genome Research, 2012, 22, 1790-1797.	2.4	2,335
6	The Transcriptional Landscape of the Yeast Genome Defined by RNA Sequencing. Science, 2008, 320, 1344-1349.	6.0	2,180
7	Global Analysis of Protein Activities Using Proteome Chips. Science, 2001, 293, 2101-2105.	6.0	2,082
8	ChIP-seq guidelines and practices of the ENCODE and modENCODE consortia. Genome Research, 2012, 22, 1813-1831.	2.4	1,708
9	CNVnator: An approach to discover, genotype, and characterize typical and atypical CNVs from family and population genome sequencing. Genome Research, 2011, 21, 974-984.	2.4	1,387
10	Architecture of the human regulatory network derived from ENCODE data. Nature, 2012, 489, 91-100.	13.7	1,384
11	High-Quality Binary Protein Interaction Map of the Yeast Interactome Network. Science, 2008, 322, 104-110.	6.0	1,297
12	Genome-wide profiles of STAT1 DNA association using chromatin immunoprecipitation and massively parallel sequencing. Nature Methods, 2007, 4, 651-657.	9.0	1,254
13	A Bayesian Networks Approach for Predicting Protein-Protein Interactions from Genomic Data. Science, 2003, 302, 449-453.	6.0	1,183
14	Personal Omics Profiling Reveals Dynamic Molecular and Medical Phenotypes. Cell, 2012, 148, 1293-1307.	13.5	1,134
15	Genomic binding sites of the yeast cell-cycle transcription factors SBF and MBF. Nature, 2001, 409, 533-538.	13.7	1,030
16	Paired-End Mapping Reveals Extensive Structural Variation in the Human Genome. Science, 2007, 318, 420-426.	6.0	1,003
17	Mapping copy number variation by population-scale genome sequencing. Nature, 2011, 470, 59-65.	13.7	991
18	Global Identification of Human Transcribed Sequences with Genome Tiling Arrays. Science, 2004, 306, 2242-2246.	6.0	983

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19	Genomic analysis of regulatory network dynamics reveals large topological changes. Nature, 2004, 431, 308-312.	13.7	921
20	Global analysis of protein phosphorylation in yeast. Nature, 2005, 438, 679-684.	13.7	915
21	Integrative Analysis of the <i>Caenorhabditis elegans</i> Genome by the modENCODE Project. Science, 2010, 330, 1775-1787.	6.0	912
22	Protein chip technology. Current Opinion in Chemical Biology, 2003, 7, 55-63.	2.8	861
23	Analysis of yeast protein kinases using protein chips. Nature Genetics, 2000, 26, 283-289.	9.4	810
24	Sequence features and chromatin structure around the genomic regions bound by 119 human transcription factors. Genome Research, 2012, 22, 1798-1812.	2.4	762
25	Unlocking the secrets of the genome. Nature, 2009, 459, 927-930.	13.7	744
26	Subcellular localization of the yeast proteome. Genes and Development, 2002, 16, 707-719.	2.7	667
27	Linking disease associations with regulatory information in the human genome. Genome Research, 2012, 22, 1748-1759.	2.4	657
28	Protein analysis on a proteomic scale. Nature, 2003, 422, 208-215.	13.7	610
29	A single-molecule long-read survey of the human transcriptome. Nature Biotechnology, 2013, 31, 1009-1014.	9.4	600
30	What is a gene, post-ENCODE? History and updated definition. Genome Research, 2007, 17, 669-681.	2.4	530
31	Large-scale analysis of the yeast genome by transposon tagging and gene disruption. Nature, 1999, 402, 413-418.	13.7	521
32	Variation in Transcription Factor Binding Among Humans. Science, 2010, 328, 232-235.	6.0	521
33	PeakSeq enables systematic scoring of ChIP-seq experiments relative to controls. Nature Biotechnology, 2009, 27, 66-75.	9.4	514
34	Getting connected: analysis and principles of biological networks. Genes and Development, 2007, 21, 1010-1024.	2.7	477
35	Biochemical and genetic analysis of the yeast proteome with a movable ORF collection. Genes and Development, 2005, 19, 2816-2826.	2.7	443
36	An encyclopedia of mouse DNA elements (Mouse ENCODE). Genome Biology, 2012, 13, 418.	13.9	410

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37	Annotating non-coding regions of the genome. Nature Reviews Genetics, 2010, 11, 559-571.	7.7	398
38	Protein arrays and microarrays. Current Opinion in Chemical Biology, 2001, 5, 40-45.	2.8	376
39	The genetic architecture of Down syndrome phenotypes revealed by high-resolution analysis of human segmental trisomies. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 12031-12036.	3.3	342
40	Rapid analysis of the DNA-binding specificities of transcription factors with DNA microarrays. Nature Genetics, 2004, 36, 1331-1339.	9.4	341
41	Deciphering Protein Kinase Specificity Through Large-Scale Analysis of Yeast Phosphorylation Site Motifs. Science Signaling, 2010, 3, ra12.	1.6	341
42	Extensive Variation in Chromatin States Across Humans. Science, 2013, 342, 750-752.	6.0	338
43	Proteomics. Annual Review of Biochemistry, 2003, 72, 783-812.	5.0	332
44	A myelopoiesis-associated regulatory intergenic noncoding RNA transcript within the human HOXA cluster. Blood, 2009, 113, 2526-2534.	0.6	330
45	Divergence of Transcription Factor Binding Sites Across Related Yeast Species. Science, 2007, 317, 815-819.	6.0	320
46	Distribution of NF-ÂB-binding sites across human chromosome 22. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 12247-12252.	3.3	298
47	AlleleSeq: analysis of alleleâ€specific expression and binding in a network framework. Molecular Systems Biology, 2011, 7, 522.	3.2	284
48	Performance comparison of whole-genome sequencing platforms. Nature Biotechnology, 2012, 30, 78-82.	9.4	281
49	Analyzing antibody specificity with whole proteome microarrays. Nature Biotechnology, 2003, 21, 1509-1512.	9.4	270
50	Protein microarray technology. Mechanisms of Ageing and Development, 2007, 128, 161-167.	2.2	268
51	A Genomic Study of the Bipolar Bud Site Selection Pattern in <i>Saccharomyces cerevisiae</i> Molecular Biology of the Cell, 2001, 12, 2147-2170.	0.9	266
52	Lineage-specific dynamic and pre-established enhancer–promoter contacts cooperate in terminal differentiation. Nature Genetics, 2017, 49, 1522-1528.	9.4	255
53	Promise of personalized omics to precision medicine. Wiley Interdisciplinary Reviews: Systems Biology and Medicine, 2013, 5, 73-82.	6.6	245
54	Complex transcriptional circuitry at the G1/S transition in Saccharomyces cerevisiae. Genes and Development, 2002, 16, 3017-3033.	2.7	236

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55	Classification of human genomic regions based on experimentally determined binding sites of more than 100 transcription-related factors. Genome Biology, 2012, 13, R48.	13.9	233
56	Finding new components of the target of rapamycin (TOR) signaling network through chemical genetics and proteome chips. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 16594-16599.	3.3	225
57	Recurrent somatic mutations in regulatory regions of human cancer genomes. Nature Genetics, 2015, 47, 710-716.	9.4	225
58	Regulation of Gene Expression by a Metabolic Enzyme. Science, 2004, 306, 482-484.	6.0	223
59	PEMer: a computational framework with simulation-based error models for inferring genomic structural variants from massive paired-end sequencing data. Genome Biology, 2009, 10, R23.	13.9	223
60	RNAâ€Seq: A Method for Comprehensive Transcriptome Analysis. Current Protocols in Molecular Biology, 2010, 89, Unit 4.11.1-13.	2.9	218
61	TOS9 Regulates White-Opaque Switching in Candida albicans. Eukaryotic Cell, 2006, 5, 1674-1687.	3.4	207
62	Rnnotator: an automated de novo transcriptome assembly pipeline from stranded RNA-Seq reads. BMC Genomics, 2010, 11, 663.	1.2	201
63	Extensive In Vivo Metabolite-Protein Interactions Revealed by Large-Scale Systematic Analyses. Cell, 2010, 143, 639-650.	13.5	200
64	Pseudogenes in the ENCODE regions: Consensus annotation, analysis of transcription, and evolution. Genome Research, 2007, 17, 839-851.	2.4	191
65	Genome-Wide Occupancy of SREBP1 and Its Partners NFY and SP1 Reveals Novel Functional Roles and Combinatorial Regulation of Distinct Classes of Genes. PLoS Genetics, 2008, 4, e1000133.	1.5	191
66	Comprehensive annotation of the transcriptome of the human fungal pathogen <i>Candida albicans</i> using RNA-seq. Genome Research, 2010, 20, 1451-1458.	2.4	191
67	Genome-wide Association Analysis of Blood-Pressure Traits in African-Ancestry Individuals Reveals Common Associated Genes in African and Non-African Populations. American Journal of Human Genetics, 2013, 93, 545-554.	2.6	189
68	Mapping accessible chromatin regions using Sono-Seq. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 14926-14931.	3.3	186
69	Comparative analysis of regulatory information and circuits across distant species. Nature, 2014, 512, 453-456.	13.7	184
70	Mapping of transcription factor binding regions in mammalian cells by ChIP: Comparison of array- and sequencing-based technologies. Genome Research, 2007, 17, 898-909.	2.4	181
71	A question of size: the eukaryotic proteome and the problems in defining it. Nucleic Acids Research, 2002, 30, 1083-1090.	6.5	172
72	Molecular Dissection of a Yeast Septin: Distinct Domains Are Required for Septin Interaction, Localization, and Function. Molecular and Cellular Biology, 2003, 23, 2762-2777.	1.1	170

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73	Dynamic transcriptomes during neural differentiation of human embryonic stem cells revealed by short, long, and paired-end sequencing. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 5254-5259.	3.3	168
74	Ubiquitous heterogeneity and asymmetry of the chromatin environment at regulatory elements. Genome Research, 2012, 22, 1735-1747.	2.4	168
75	Close association of RNA polymerase II and many transcription factors with Pol III genes. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 3639-3644.	3.3	167
76	Understanding transcriptional regulation by integrative analysis of transcription factor binding data. Genome Research, 2012, 22, 1658-1667.	2.4	166
77	Genome-Wide Identification of Binding Sites Defines Distinct Functions for Caenorhabditis elegans PHA-4/FOXA in Development and Environmental Response. PLoS Genetics, 2010, 6, e1000848.	1.5	165
78	Genome-wide relationship between histone H3 lysine 4 mono- and tri-methylation and transcription factor binding. Genome Research, 2008, 18, 1906-1917.	2.4	163
79	Genetic analysis of variation in transcription factor binding in yeast. Nature, 2010, 464, 1187-1191.	13.7	162
80	Protein complexes take the bait. Nature, 2002, 415, 123-124.	13.7	161
81	Advances in functional protein microarray technology. FEBS Journal, 2005, 272, 5400-5411.	2.2	160
82	Nucleotide-resolution analysis of structural variants using BreakSeq and a breakpoint library. Nature Biotechnology, 2010, 28, 47-55.	9.4	158
83	Gene-centric Meta-analysis in 87,736 Individuals of European Ancestry Identifies Multiple Blood-Pressure-Related Loci. American Journal of Human Genetics, 2014, 94, 349-360.	2.6	158
84	Microarrays to characterize protein interactions on a whole-proteome scale. Proteomics, 2003, 3, 2190-2199.	1.3	155
85	Target hub proteins serve as master regulators of development in yeast. Genes and Development, 2006, 20, 435-448.	2.7	153
86	ChIP-chip: A genomic approach for identifying transcription factor binding sites. Methods in Enzymology, 2002, 350, 469-483.	0.4	151
87	Structured RNAs in the ENCODE selected regions of the human genome. Genome Research, 2007, 17, 852-864.	2.4	150
88	GATA-1 binding sites mapped in the Â-globin locus by using mammalian chlp-chip analysis. Proceedings of the National Academy of Sciences of the United States of America, 2002, 99, 2924-2929.	3.3	149
89	Dynamic trans-Acting Factor Colocalization in Human Cells. Cell, 2013, 155, 713-724.	13.5	142
90	Integrated systems analysis reveals a molecular network underlying autism spectrum disorders. Molecular Systems Biology, 2014, 10, 774.	3.2	138

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91	Efficient yeast ChIP-Seq using multiplex short-read DNA sequencing. BMC Genomics, 2009, 10, 37.	1.2	137
92	Landscape of cohesin-mediated chromatin loops in the human genome. Nature, 2020, 583, 737-743.	13.7	134
93	Integrated Network Analysis Reveals an Association between Plasma Mannose Levels and Insulin Resistance. Cell Metabolism, 2016, 24, 172-184.	7.2	133
94	Severe acute respiratory syndrome diagnostics using a coronavirus protein microarray. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 4011-4016.	3.3	131
95	Analysis of copy number variants and segmental duplications in the human genome: Evidence for a change in the process of formation in recent evolutionary history. Genome Research, 2008, 18, 1865-1874.	2.4	126
96	High-resolution mapping of DNA copy alterations in human chromosome 22 using high-density tiling oligonucleotide arrays. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 4534-4539.	3.3	125
97	Charging it up: global analysis of protein phosphorylation. Trends in Genetics, 2006, 22, 545-554.	2.9	123
98	DNA replication-timing analysis of human chromosome 22 at high resolution and different developmental states. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 17771-17776.	3.3	121
99	Personal genome sequencing: current approaches and challenges. Genes and Development, 2010, 24, 423-431.	2.7	119
100	Systematic evaluation of variability in ChIP-chip experiments using predefined DNA targets. Genome Research, 2008, 18, 393-403.	2.4	117
101	Regulatory analysis of the C. elegans genome with spatiotemporal resolution. Nature, 2014, 512, 400-405.	13.7	115
102	Impact of Chromatin Structures on DNA Processing for Genomic Analyses. PLoS ONE, 2009, 4, e6700.	1.1	115
103	GENOMICS: Defining Genes in the Genomics Era. Science, 2003, 300, 258-260.	6.0	114
104	An integrated approach for finding overlooked genes in yeast. Nature Biotechnology, 2002, 20, 58-63.	9.4	112
105	Predicting essential genes in fungal genomes. Genome Research, 2006, 16, 1126-1135.	2.4	109
106	Detecting and annotating genetic variations using the HugeSeq pipeline. Nature Biotechnology, 2012, 30, 226-229.	9.4	104
107	High-Resolution Copy-Number Variation Map Reflects Human Olfactory Receptor Diversity and Evolution. PLoS Genetics, 2008, 4, e1000249.	1.5	99
108	Long-Read Isoform Sequencing Reveals a Hidden Complexity of the Transcriptional Landscape of Herpes Simplex Virus Type 1. Frontiers in Microbiology, 2017, 8, 1079.	1.5	97

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109	Emerging technologies in yeast genomics. Nature Reviews Genetics, 2001, 2, 302-312.	7.7	96
110	Global changes in STAT target selection and transcription regulation upon interferon treatments. Genes and Development, 2005, 19, 2953-2968.	2.7	95
111	RSEQtools: a modular framework to analyze RNA-Seq data using compact, anonymized data summaries. Bioinformatics, 2011, 27, 281-283.	1.8	93
112	Full-Length Isoform Sequencing Reveals Novel Transcripts and Substantial Transcriptional Overlaps in a Herpesvirus. PLoS ONE, 2016, 11, e0162868.	1.1	93
113	The current excitement about copy-number variation: how it relates to gene duplications and protein families. Current Opinion in Structural Biology, 2008, 18, 366-374.	2.6	92
114	Construction and Analysis of an Integrated Regulatory Network Derived from High-Throughput Sequencing Data. PLoS Computational Biology, 2011, 7, e1002190.	1.5	92
115	Characterization of Enhancer Function from Genome-Wide Analyses. Annual Review of Genomics and Human Genetics, 2012, 13, 29-57.	2.5	86
116	Omics Profiling in Precision Oncology. Molecular and Cellular Proteomics, 2016, 15, 2525-2536.	2.5	84
117	Global analysis of the glycoproteome in <i>Saccharomyces cerevisiae</i> reveals new roles for protein glycosylation in eukaryotes. Molecular Systems Biology, 2009, 5, 308.	3.2	79
118	Systematic prediction and validation of breakpoints associated with copy-number variants in the human genome. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 10110-10115.	3.3	78
119	The Development of Protein Microarrays and Their Applications in DNA–Protein and Protein–Protein Interaction Analyses of Arabidopsis Transcription Factors. Molecular Plant, 2008, 1, 27-41.	3.9	78
120	A highly integrated and complex PPARGC1A transcription factor binding network in HepG2 cells. Genome Research, 2012, 22, 1668-1679.	2.4	75
121	Global analysis of gene expression in yeast. Functional and Integrative Genomics, 2002, 2, 171-180.	1.4	74
122	Negative regulation of calcineurin signaling by Hrr25p, a yeast homolog of casein kinase I. Genes and Development, 2003, 17, 2698-2708.	2.7	74
123	â€ ⁻ Omic' approaches for unraveling signaling networks. Current Opinion in Cell Biology, 2002, 14, 173-179.	2.6	73
124	AGAPE (Automated Genome Analysis PipelinE) for Pan-Genome Analysis of Saccharomyces cerevisiae. PLoS ONE, 2015, 10, e0120671.	1.1	73
125	Applications of DNA tiling arrays to experimental genome annotation and regulatory pathway discovery. Chromosome Research, 2005, 13, 259-274.	1.0	72
126	Modeling ChIP Sequencing In Silico with Applications. PLoS Computational Biology, 2008, 4, e1000158.	1.5	70

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127	Regulatory Variation Within and Between Species. Annual Review of Genomics and Human Genetics, 2011, 12, 327-346.	2.5	70
128	Proteome chips for whole-organism assays. Nature Reviews Molecular Cell Biology, 2006, 7, 617-622.	16.1	69
129	A novel mitochondrial protein, Tar1p, is encoded on the antisense strand of the nuclear 25S rDNA. Genes and Development, 2002, 16, 2755-2760.	2.7	67
130	iPOP Goes the World: Integrated Personalized Omics Profiling and the Road toward Improved Health Care. Chemistry and Biology, 2013, 20, 660-666.	6.2	67
131	Microfluidic isoform sequencing shows widespread splicing coordination in the human transcriptome. Genome Research, 2018, 28, 231-242.	2.4	64
132	Multi-Platform Sequencing Approach Reveals a Novel Transcriptome Profile in Pseudorabies Virus. Frontiers in Microbiology, 2018, 8, 2708.	1.5	64
133	Linking DNA-binding proteins to their recognition sequences by using protein microarrays. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 9940-9945.	3.3	63
134	Mapping the chromosomal targets of STAT1 by Sequence Tag Analysis of Genomic Enrichment (STAGE). Genome Research, 2007, 17, 910-916.	2.4	61
135	Proteome-wide survey of the autoimmune target repertoire in autoimmune polyendocrine syndrome type 1. Scientific Reports, 2016, 6, 20104.	1.6	61
136	Diverse protein kinase interactions identified by protein microarrays reveal novel connections between cellular processes. Genes and Development, 2011, 25, 767-778.	2.7	60
137	Accurate Identification and Analysis of Human mRNA Isoforms Using Deep Long Read Sequencing. G3: Genes, Genomes, Genetics, 2013, 3, 387-397.	0.8	59
138	Extensive Transcript Diversity and Novel Upstream Open Reading Frame Regulation in Yeast. G: Genes, Genemes, Genetics, 2013, 3, 343-352.	0.8	59
139	Genome-Wide Mapping of Copy Number Variation in Humans: Comparative Analysis of High Resolution Array Platforms. PLoS ONE, 2011, 6, e27859.	1.1	59
140	Identification of genomic indels and structural variations using split reads. BMC Genomics, 2011, 12, 375.	1.2	57
141	Statistical analysis of the genomic distribution and correlation of regulatory elements in the ENCODE regions. Genome Research, 2007, 17, 787-797.	2.4	56
142	Systematic analysis of transcribed loci in ENCODE regions using RACE sequencing reveals extensive transcription in the human genome. Genome Biology, 2008, 9, R3.	13.9	53
143	Cooperative transcription factor associations discovered using regulatory variation. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 13353-13358.	3.3	53
144	A genomic analysis of RNA polymerase II modification and chromatin architecture related to 3′ end RNA polyadenylation. Genome Research, 2008, 18, 1224-1237.	2.4	50

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145	Comparative annotation of functional regions in the human genome using epigenomic data. Nucleic Acids Research, 2013, 41, 4423-4432.	6.5	50
146	ChIPâ€Seq: A Method for Global Identification of Regulatory Elements in the Genome. Current Protocols in Molecular Biology, 2010, 91, Unit 21.19.1-14.	2.9	49
147	Design optimization methods for genomic DNA tiling arrays. Genome Research, 2005, 16, 271-281.	2.4	46
148	ProCAT: a data analysis approach for protein microarrays. Genome Biology, 2006, 7, R110.	13.9	46
149	Systems biology from a yeast omics perspective. FEBS Letters, 2009, 583, 3895-3899.	1.3	46
150	X chromosome-wide analyses of genomic DNA methylation states and gene expression in male and female neutrophils. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 3704-3709.	3.3	44
151	Multiple Long-Read Sequencing Survey of Herpes Simplex Virus Dynamic Transcriptome. Frontiers in Genetics, 2019, 10, 834.	1.1	44
152	Personal phenotypes to go with personal genomes. Molecular Systems Biology, 2009, 5, 273.	3.2	41
153	Proteomic Approaches for the Global Analysis of Proteins. BioTechniques, 2002, 33, 1308-1316.	0.8	40
154	Investigating metabolite–protein interactions: An overview of available techniques. Methods, 2012, 57, 459-466.	1.9	40
155	ExpressYourself: a modular platform for processing and visualizing microarray data. Nucleic Acids Research, 2003, 31, 3477-3482.	6.5	38
156	RNA polymerase II stalling: loading at the start prepares genes for a sprint. Genome Biology, 2008, 9, 220.	13.9	38
157	Yeast as a Model for Human Disease. Current Protocols in Human Genetics, 2006, 48, Unit 15.6.	3.5	36
158	Metabolites as global regulators: A new view of protein regulation. BioEssays, 2011, 33, 485-489.	1.2	36
159	Global analysis of protein function using protein microarrays. Mechanisms of Ageing and Development, 2005, 126, 171-175.	2.2	35
160	[33] High-throughput methods for the large-scale analysis of gene function by transposon tagging. Methods in Enzymology, 2000, 328, 550-574.	0.4	34
161	Large-scale mutagenesis: yeast genetics in the genome era. Current Opinion in Biotechnology, 2001, 12, 28-34.	3.3	34
162	MSB: A mean-shift-based approach for the analysis of structural variation in the genome. Genome Research, 2009, 19, 106-117.	2.4	33

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163	Systematic investigation of protein–small molecule interactions. IUBMB Life, 2013, 65, 2-8.	1.5	33
164	A supervised hidden markov model framework for efficiently segmenting tiling array data in transcriptional and chIP-chip experiments: systematically incorporating validated biological knowledge. Bioinformatics, 2006, 22, 3016-3024.	1.8	32
165	Large-scale identification of genes important for apical growth in Saccharomyces cerevisiae by directed allele replacement technology (DART) screening. Functional and Integrative Genomics, 2002, 1, 345-356.	1.4	31
166	Integrated analysis of experimental data sets reveals many novel promoters in 1% of the human genome. Genome Research, 2007, 17, 720-731.	2.4	31
167	Yeast proteomics and protein microarrays. Journal of Proteomics, 2010, 73, 2147-2157.	1.2	31
168	Proteogenomic Analysis of Human Colon Carcinoma Cell Lines LIM1215, LIM1899, and LIM2405. Journal of Proteome Research, 2013, 12, 1732-1742.	1.8	30
169	Personal genomes, quantitative dynamic omics and personalized medicine. Quantitative Biology, 2013, 1, 71-90.	0.3	29
170	Genome-wide profiling of human cap-independent translation-enhancing elements. Nature Methods, 2013, 10, 747-750.	9.0	29
171	Coherent Functional Modules Improve Transcription Factor Target Identification, Cooperativity Prediction, and Disease Association. PLoS Genetics, 2014, 10, e1004122.	1.5	29
172	MOTIPS: Automated Motif Analysis for Predicting Targets of Modular Protein Domains. BMC Bioinformatics, 2010, 11, 243.	1.2	28
173	Assessing the performance of different high-density tiling microarray strategies for mapping transcribed regions of the human genome. Genome Research, 2007, 17, 886-897.	2.4	25
174	The DART classification of unannotated transcription within the ENCODE regions: Associating transcription with known and novel loci. Genome Research, 2007, 17, 732-745.	2.4	25
175	Identification of Genes Critical for Resistance to Infection by West Nile Virus Using RNA-Seq Analysis. Viruses, 2013, 5, 1664-1681.	1.5	25
176	[15] Extrapolating Traditional DNA Microarray Statistics to Tiling and Protein Microarray Technologies. Methods in Enzymology, 2006, 411, 282-311.	0.4	23
177	ChIP-Seq. Methods in Enzymology, 2010, 470, 77-104.	0.4	23
178	Transcription factor binding site identification in yeast: a comparison of high-density oligonucleotide and PCR-based microarray platforms. Functional and Integrative Genomics, 2007, 7, 335-345.	1.4	22
179	Multiplatform next-generation sequencing identifies novel RNA molecules and transcript isoforms of the endogenous retrovirus isolated from cultured cells. FEMS Microbiology Letters, 2018, 365, .	0.7	21
180	Prospects and Challenges in Proteomics. Plant Physiology, 2005, 138, 560-562.	2.3	19

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181	Discovery of Novel Human Gene Regulatory Modules from Gene Co-expression and Promoter Motif Analysis. Scientific Reports, 2017, 7, 5557.	1.6	19
182	Systems Biology Approaches to Disease Marker Discovery. Disease Markers, 2010, 28, 209-224.	0.6	18
183	Protein Microarrays. Methods in Molecular Biology, 2009, 548, 209-222.	0.4	18
184	A Pilot Study of Transcription Unit Analysis in Rice Using Oligonucleotide Tiling-path Microarray. Plant Molecular Biology, 2005, 59, 137-149.	2.0	17
185	A plethora of sites. Nature Genetics, 2004, 36, 325-326.	9.4	16
186	Dissecting phosphorylation networks: lessons learned from yeast. Expert Review of Proteomics, 2011, 8, 775-786.	1.3	15
187	Integrating Sequencing Technologies in Personal Genomics: Optimal Low Cost Reconstruction of Structural Variants. PLoS Computational Biology, 2009, 5, e1000432.	1.5	14
188	Dual Platform Long-Read RNA-Sequencing Dataset of the Human Cytomegalovirus Lytic Transcriptome. Frontiers in Genetics, 2018, 9, 432.	1.1	14
189	RNA Sequencing. Methods in Molecular Biology, 2011, 759, 125-132.	0.4	14
190	Positional artifacts in microarrays: experimental verification and construction of COP, an automated detection tool. Nucleic Acids Research, 2006, 35, e8-e8.	6.5	12
191	Carbohydrate Analysis Prepares to Enter the "Omics―Era. Chemistry and Biology, 2002, 9, 400-401.	6.2	11
192	Fast Optimal Genome Tiling with Applications to Microarray Design and Homology Search. Journal of Computational Biology, 2004, 11, 766-785.	0.8	11
193	Yeast genomics: past, present, and future promise. Functional and Integrative Genomics, 2002, 2, 135-137.	1.4	7
194	Chromatin Immunoprecipitation and Multiplex Sequencing (ChIP-Seq) to Identify Global Transcription Factor Binding Sites in the Nematode Caenorhabditis Elegans. Methods in Enzymology, 2014, 539, 89-111.	0.4	7
195	Analyzing In Vivo Metaboliteâ€Protein Interactions by Largeâ€Scale Systematic Analyses. Current Protocols in Chemical Biology, 2011, 3, 181-196.	1.7	5
196	Preparation of Recombinant Protein Spotted Arrays for Proteomeâ€Wide Identification of Kinase Targets. Current Protocols in Protein Science, 2013, 72, Unit 27.4.	2.8	4
197	Global Analysis of Transcription Factor-Binding Sites in Yeast Using ChIP-Seq. Methods in Molecular Biology, 2014, 1205, 231-255.	0.4	4
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