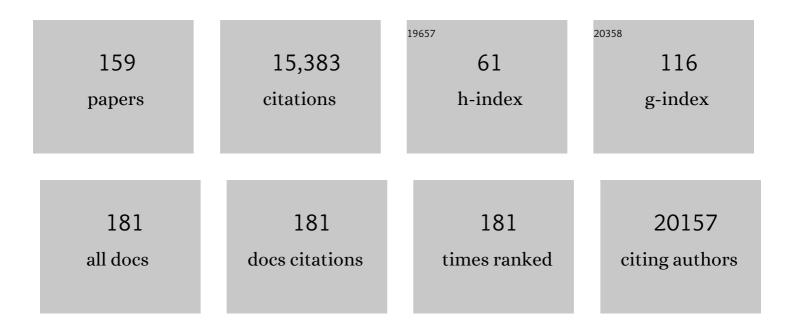
## Leonard Cornelis Schalkwyk

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	The Collaborative Cross, a community resource for the genetic analysis of complex traits. Nature Genetics, 2004, 36, 1133-1137.	21.4	1,034
2	A data-driven approach to preprocessing Illumina 450K methylation array data. BMC Genomics, 2013, 14, 293.	2.8	850
3	Alzheimer's disease: early alterations in brain DNA methylation at ANK1, BIN1, RHBDF2 and other loci. Nature Neuroscience, 2014, 17, 1156-1163.	14.8	800
4	Functional annotation of the human brain methylome identifies tissue-specific epigenetic variation across brain and blood. Genome Biology, 2012, 13, R43.	9.6	585
5	DNA methylation aging clocks: challenges and recommendations. Genome Biology, 2019, 20, 249.	8.8	552
6	Methylomic profiling implicates cortical deregulation of ANK1 in Alzheimer's disease. Nature Neuroscience, 2014, 17, 1164-1170.	14.8	488
7	Disease-associated epigenetic changes in monozygotic twins discordant for schizophrenia and bipolar disorder. Human Molecular Genetics, 2011, 20, 4786-4796.	2.9	407
8	Interindividual methylomic variation across blood, cortex, and cerebellum: implications for epigenetic studies of neurological and neuropsychiatric phenotypes. Epigenetics, 2015, 10, 1024-1032.	2.7	393
9	The nature and identification of quantitative trait loci: a community's view. Nature Reviews Genetics, 2003, 4, 911-916.	16.3	390
10	Moderation of breastfeeding effects on the IQ by genetic variation in fatty acid metabolism. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 18860-18865.	7.1	324
11	Methylation QTLs in the developing brain and their enrichment in schizophrenia risk loci. Nature Neuroscience, 2016, 19, 48-54.	14.8	306
12	Genetic analysis of the mouse brain proteome. Nature Genetics, 2002, 30, 385-393.	21.4	293
13	Transformation methods for halophilic archaebacteria. Canadian Journal of Microbiology, 1989, 35, 148-152.	1.7	290
14	An integrated genetic-epigenetic analysis of schizophrenia: evidence for co-localization of genetic associations and differential DNA methylation. Genome Biology, 2016, 17, 176.	8.8	287
15	Methylomic analysis of monozygotic twins discordant for autism spectrum disorder and related behavioural traits. Molecular Psychiatry, 2014, 19, 495-503.	7.9	280
16	High resolution cosmid and P1 maps spanning the 14 Mb genome of the fission yeast S. pombe. Cell, 1993, 73, 109-120.	28.9	271
17	Methylomic trajectories across human fetal brain development. Genome Research, 2015, 25, 338-352.	5.5	250
18	Allelic Skewing of DNA Methylation Is Widespread across the Genome. American Journal of Human Genetics, 2010, 86, 196-212.	6.2	228

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19	Genomic and phenotypic insights from an atlas of genetic effects on DNA methylation. Nature Genetics, 2021, 53, 1311-1321.	21.4	218
20	Longitudinal changes of telomere length and epigenetic age related to traumatic stress and post-traumatic stress disorder. Psychoneuroendocrinology, 2015, 51, 506-512.	2.7	186
21	Towards high resolution maps of the mouse and human genomes—a facility for ordering markers to 0.1 cM resolution. Human Molecular Genetics, 1994, 3, 621-627.	2.9	185
22	Genome-wide DNA methylation levels and altered cortisol stress reactivity following childhood trauma in humans. Nature Communications, 2016, 7, 10967.	12.8	175
23	Loss of adenylyl cyclase I activity disrupts patterning of mouse somatosensory cortex. Nature Genetics, 1998, 19, 289-291.	21.4	156
24	Assessing Reliability, Heritability and General Cognitive Ability in a Battery of Cognitive Tasks for Laboratory Mice. Behavior Genetics, 2005, 35, 675-692.	2.1	146
25	Methylomic profiling of human brain tissue supports a neurodevelopmental origin for schizophrenia. Genome Biology, 2014, 15, 483.	8.8	141
26	Elevated DNA methylation across a 48â€kb region spanning the <i>HOXA</i> gene cluster is associated with Alzheimer's disease neuropathology. Alzheimer's and Dementia, 2018, 14, 1580-1588.	0.8	138
27	A histone acetylome-wide association study of Alzheimer's disease identifies disease-associated H3K27ac differences in the entorhinal cortex. Nature Neuroscience, 2018, 21, 1618-1627.	14.8	138
28	Differential methylation of the TRPA1 promoter in pain sensitivity. Nature Communications, 2014, 5, 2978.	12.8	132
29	Mitochondrial genes are altered in blood early in Alzheimer's disease. Neurobiology of Aging, 2017, 53, 36-47.	3.1	132
30	Allele-specific methylation in the human genome. Epigenetics, 2010, 5, 578-582.	2.7	129
31	Expression quantitative trait loci in the developing human brain and their enrichment in neuropsychiatric disorders. Genome Biology, 2018, 19, 194.	8.8	126
32	Leveraging DNA-Methylation Quantitative-Trait Loci to Characterize the Relationship between Methylomic Variation, Gene Expression, and Complex Traits. American Journal of Human Genetics, 2018, 103, 654-665.	6.2	126
33	Maternal separation is associated with strainâ€specific responses to stress and epigenetic alterations to <i>Nr3c1</i> , <i> Avp</i> , and <i>Nr4a1</i> in mouse. Brain and Behavior, 2012, 2, 455-467.	2.2	123
34	Genetics of behavioural domains across the neuropsychiatric spectrum; of mice and men. Molecular Psychiatry, 2007, 12, 324-330.	7.9	117
35	Cross-region reduction in 5-hydroxymethylcytosine in Alzheimer's disease brain. Neurobiology of Aging, 2014, 35, 1850-1854.	3.1	114
36	Quantitative trait locus association scan of early reading disability and ability using pooled DNA and 100K SNP microarrays in a sample of 5760 children. Molecular Psychiatry, 2008, 13, 729-740.	7.9	101

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37	A genomeâ€wide association study identifies multiple loci associated with mathematics ability and disability. Genes, Brain and Behavior, 2010, 9, 234-247.	2.2	100
38	Recalibrating the epigenetic clock: implications for assessing biological age in the human cortex. Brain, 2020, 143, 3763-3775.	7.6	100
39	Systematic underestimation of the epigenetic clock and age acceleration in older subjects. Genome Biology, 2019, 20, 283.	8.8	97
40	Advanced paternal age effects in neurodevelopmental disorders—review of potential underlying mechanisms. Translational Psychiatry, 2017, 7, e1019-e1019.	4.8	94
41	Performance deficit of α7 nicotinic receptor knockout mice in a delayed matching-to-place task suggests a mild impairment of working/episodic-like memory. Genes, Brain and Behavior, 2006, 5, 433-440.	2.2	92
42	Genotyping pooled DNA using 100K SNP microarrays: a step towards genomewide association scans. Nucleic Acids Research, 2006, 34, e27-e27.	14.5	92
43	Behavioural battery testing: Evaluation and behavioural outcomes in 8 inbred mouse strains. Physiology and Behavior, 2010, 99, 301-316.	2.1	92
44	SNPs, microarrays and pooled DNA: identification of four loci associated with mild mental impairment in a sample of 6000 children. Human Molecular Genetics, 2005, 14, 1315-1325.	2.9	91
45	Intermediate DNA methylation is a conserved signature of genome regulation. Nature Communications, 2015, 6, 6363.	12.8	91
46	Detailed physical map and set of overlapping clones covering the genome of the archaebacterium Haloferax volcanii DS2. Journal of Molecular Biology, 1991, 222, 509-524.	4.2	90
47	Genotyping Pooled DNA on Microarrays: A Systematic Genome Screen of Thousands of SNPs in Large Samples to Detect QTLs for Complex Traits. Behavior Genetics, 2004, 34, 549-555.	2.1	89
48	The mouse homeobox gene <i>Not</i> is required for caudal notochord development and affected by the truncate mutation. Genes and Development, 2004, 18, 1725-1736.	5.9	84
49	Genetics of the hippocampal transcriptome in mouse: a systematic survey and online neurogenomics resource. Frontiers in Neuroscience, 2009, 3, 55.	2.8	84
50	Variation in 5-hydroxymethylcytosine across human cortex and cerebellum. Genome Biology, 2016, 17, 27.	8.8	83
51	Depressionâ€Related Behavioral Tests. Current Protocols in Mouse Biology, 2012, 2, 119-127.	1.2	82
52	Increased DNA methylation near TREM2 is consistently seen in the superior temporal gyrus in Alzheimer's disease brain. Neurobiology of Aging, 2016, 47, 35-40.	3.1	79
53	Full-length transcript sequencing of human and mouse cerebral cortex identifies widespread isoform diversity and alternative splicing. Cell Reports, 2021, 37, 110022.	6.4	79
54	Hippocampal gene expression profiling across eight mouse inbred strains: towards understanding the molecular basis for behaviour. European Journal of Neuroscience, 2004, 19, 2576-2582.	2.6	77

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55	Advancing Paternal Age Is Associated with Deficits in Social and Exploratory Behaviors in the Offspring: A Mouse Model. PLoS ONE, 2009, 4, e8456.	2.5	77
56	Molecular and biochemical characterisation of DNA-dependent protein kinase-defective rodent mutant irs-20. Nucleic Acids Research, 1998, 26, 1965-1973.	14.5	74
57	A Three-Stage Genome-Wide Association Study of General Cognitive Ability: Hunting the Small Effects. Behavior Genetics, 2010, 40, 759-767.	2.1	74
58	Schizophrenia-associated methylomic variation: molecular signatures of disease and polygenic risk burden across multiple brain regions. Human Molecular Genetics, 2017, 26, ddw373.	2.9	74
59	The correlation between reading and mathematics ability at age twelve has a substantial genetic component. Nature Communications, 2014, 5, 4204.	12.8	72
60	A meta-analysis of epigenome-wide association studies in Alzheimer's disease highlights novel differentially methylated loci across cortex. Nature Communications, 2021, 12, 3517.	12.8	72
61	DNA methylation in interleukin-11 predicts clinical response to antidepressants in GENDEP. Translational Psychiatry, 2013, 3, e300-e300.	4.8	71
62	Genome-wide DNA methylation profiling identifies convergent molecular signatures associated with idiopathic and syndromic autism in post-mortem human brain tissue. Human Molecular Genetics, 2019, 28, 2201-2211.	2.9	70
63	Tumor necrosis factor and its targets in the inflammatory cytokine pathway are identified as putative transcriptomic biomarkers for escitalopram response. European Neuropsychopharmacology, 2013, 23, 1105-1114.	0.7	68
64	Tetraodon fluviatilis,a New Puffer Fish Model for Genome Studies. Genomics, 1997, 41, 177-184.	2.9	66
65	5-hydroxymethylcytosine is highly dynamic across human fetal brain development. BMC Genomics, 2017, 18, 738.	2.8	63
66	The differential transcriptome and ontology profiles of floating and cumulus granulosa cells in stimulated human antral follicles. Molecular Human Reproduction, 2010, 16, 229-240.	2.8	61
67	Genotyping DNA pools on microarrays: Tackling the QTL problem of large samples and large numbers of SNPs. BMC Genomics, 2005, 6, 52.	2.8	60
68	The role of nicotinic receptor α7 subunits in nicotine discrimination. Neuropharmacology, 2004, 46, 363-371.	4.1	56
69	Integrative mouse and human mRNA studies using WGCNA nominates novel candidate genes involved in the pathogenesis of major depressive disorder. Pharmacogenomics, 2013, 14, 1979-1990.	1.3	55
70	Reduced Anxiety and Depression-Like Behaviours in the Circadian Period Mutant Mouse Afterhours. PLoS ONE, 2012, 7, e38263.	2.5	54
71	A 5× genome coverage bovine BAC library: production, characterization, and distribution. Mammalian Genome, 1999, 10, 706-709.	2.2	53
72	Stochastic Choice of Allelic Expression in Human Neural Stem Cells. Stem Cells, 2012, 30, 1938-1947.	3.2	53

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73	Epigenomic and transcriptomic signatures of a Klinefelter syndrome (47,XXY) karyotype in the brain. Epigenetics, 2014, 9, 587-599.	2.7	53
74	The endogenous and reactive depression subtypes revisited: integrative animal and human studies implicate multiple distinct molecular mechanisms underlying major depressive disorder. BMC Medicine, 2014, 12, 73.	5.5	52
75	Microarrays. Developmental Science, 2007, 10, 19-23.	2.4	51
76	Wfs1 gene deletion causes growth retardation in mice and interferes with the growth hormone pathway. Physiological Genomics, 2009, 37, 249-259.	2.3	49
77	A Genome-Wide Association Study of Social and Non-Social Autistic-Like Traits in the General Population Using Pooled DNA, 500ÂK SNP Microarrays and Both Community and Diagnosed Autism Replication Samples. Behavior Genetics, 2010, 40, 31-45.	2.1	49
78	Bigmelon: tools for analysing large DNA methylation datasets. Bioinformatics, 2019, 35, 981-986.	4.1	49
79	Genome mapping in halobacteria. Canadian Journal of Microbiology, 1989, 35, 21-29.	1.7	48
80	Association analysis of mild mental impairment using DNA pooling to screen 432 brain-expressed single-nucleotide polymorphisms. Molecular Psychiatry, 2005, 10, 384-392.	7.9	46
81	<i>TCF4</i> ( <i>e2â€2; ITF2</i> ): A schizophreniaâ€associated gene with pleiotropic effects on human disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 1-16.	1.7	46
82	Interpretation of knockout experiments: the congenic footprint. Genes, Brain and Behavior, 2007, 6, 299-303.	2.2	45
83	Applicability of DNA pools on 500 K SNP microarrays for cost-effective initial screens in genomewide association studies. BMC Genomics, 2007, 8, 214.	2.8	43
84	Identification of genes and gene pathways associated with major depressive disorder by integrative brain analysis of rat and human prefrontal cortex transcriptomes. Translational Psychiatry, 2015, 5, e519-e519.	4.8	43
85	ISH51: a large, degenerate family of insertion sequence-like elements in the genome of the archaehacterium,Halobacterium volcanii. Nucleic Acids Research, 1986, 14, 6983-7000.	14.5	42
86	Construction and Characterization of a 10-Genome Equivalent Yeast Artificial Chromosome Library for the Laboratory Rat,Rattus norvegicus. Genomics, 1997, 39, 385-392.	2.9	42
87	ViPAR: a software platform for the Virtual Pooling and Analysis of Research Data. International Journal of Epidemiology, 2016, 45, 408-416.	1.9	42
88	A central resource for accurate allele frequency estimation from pooled DNA genotyped on DNA microarrays. Nucleic Acids Research, 2005, 33, e25-e25.	14.5	39
89	Putative Transcriptomic Biomarkers in the Inflammatory Cytokine Pathway Differentiate Major Depressive Disorder Patients from Control Subjects and Bipolar Disorder Patients. PLoS ONE, 2014, 9, e91076.	2.5	39
90	Using hippocampal microRNA expression differences between mouse inbred strains to characterise miRNA function. Mammalian Genome, 2008, 19, 552-60.	2.2	38

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91	Peripheral blood RNA gene expression profiling in patients with bacterial meningitis. Frontiers in Neuroscience, 2013, 7, 33.	2.8	38
92	Efficient high-resolution genetic mapping of mouse interspersed repetitive sequence PCR products, toward integrated genetic and physical mapping of the mouse genome Proceedings of the National Academy of Sciences of the United States of America, 1995, 92, 5302-5306.	7.1	36
93	Application of microarrays to the analysis of the inactivation status of human X-linked genes expressed in lymphocytes. European Journal of Human Genetics, 2004, 12, 639-646.	2.8	35
94	Transcriptome analysis of genes and gene networks involved in aggressive behavior in mouse and zebrafish. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2016, 171, 827-838.	1.7	35
95	Regional differences in mitochondrial DNA methylation in human post-mortem brain tissue. Clinical Epigenetics, 2017, 9, 47.	4.1	34
96	Characterising sex differences of autosomal DNA methylation in whole blood using the Illumina EPIC array. Clinical Epigenetics, 2022, 14, 62.	4.1	34
97	Tissue-specific patterns of allelically-skewed DNA methylation. Epigenetics, 2016, 11, 24-35.	2.7	32
98	Advanced paternal age is associated with altered DNA methylation at brain-expressed imprinted loci in inbred mice: implications for neuropsychiatric disease. Molecular Psychiatry, 2013, 18, 635-636.	7.9	31
99	The inflammatory cytokines: molecular biomarkers for major depressive disorder?. Biomarkers in Medicine, 2015, 9, 169-180.	1.4	31
100	Panel of Microsatellite Markers for Whole-Genome Scans and Radiation Hybrid Mapping and a Mouse Family Tree. Genome Research, 1999, 9, 878-887.	5.5	30
101	Convergent Animal and Human Evidence Suggests a Role of PPM1A Gene in Response to Antidepressants. Biological Psychiatry, 2011, 69, 360-365.	1.3	30
102	Genes and Gene Networks Implicated in Aggression Related Behaviour. Neurogenetics, 2014, 15, 255-266.	1.4	30
103	Pervasive and opposing effects of Unpredictable Chronic Mild Stress (UCMS) on hippocampal gene expression in BALB/cJ and C57BL/6J mouse strains. BMC Genomics, 2015, 16, 262.	2.8	30
104	Gene expression profiling reveals upregulation of Tlr4 receptors in Cckb receptor deficient mice. Behavioural Brain Research, 2008, 188, 62-70.	2.2	29
105	Genetic and environmental (inter)actions in male mouse lines selected for aggressive and nonaggressive behavior. Genes, Brain and Behavior, 2004, 3, 101-109.	2.2	28
106	Antidepressants and the resilience to early-life stress in inbred mouse strains. Pharmacogenetics and Genomics, 2011, 21, 779-789.	1.5	28
107	Antidepressant-dependent mRNA changes in mouse associated with hippocampal neurogenesis in a mouse model of depression. Pharmacogenetics and Genomics, 2012, 22, 765-776.	1.5	28
108	Quantitative traits for the tail suspension test: automation, optimization, and BXD RI mapping. Mammalian Genome, 2007, 18, 482-491.	2.2	27

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109	The SNPMaP package for R: a framework for genome-wide association using DNA pooling on microarrays. Bioinformatics, 2009, 25, 281-283.	4.1	27
110	A bovine YAC library containing four- to five-fold genome equivalents. Mammalian Genome, 1999, 10, 837-838.	2.2	25
111	Behavioral Characterization of Wild Derived Male Mice (Mus musculus musculus) of the PWD/Ph Inbred Strain: High Exploration Compared to C57BL/6J. Behavior Genetics, 2004, 34, 621-630.	2.1	25
112	A B2 SINE insertion in the <i>Comt1</i> gene ( <i>Comt1</i> <sup><i>B2i</i></sup> ) results in an overexpressing, behavior modifying allele present in classical inbred mouse strains. Genes, Brain and Behavior, 2010, 9, 925-932.	2.2	24
113	Interspecies comparisons of functional genetic variations and their implications in neuropsychiatry. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2009, 150B, 309-317.	1.7	22
114	Effects of advanced paternal age on trajectories of social behavior in offspring. Genes, Brain and Behavior, 2015, 14, 443-453.	2.2	22
115	Silencing of the <i>WFS1</i> gene in HEK cells induces pathways related to neurodegeneration and mitochondrial damage. Physiological Genomics, 2013, 45, 182-190.	2.3	21
116	Genetic variation in hippocampal microRNA expression differences in C57BL/6 J X DBA/2 J (BXD) recombinant inbred mouse strains. BMC Genomics, 2012, 13, 476.	2.8	20
117	ATP-binding cassette sub-family F member 1 (ABCF1) is identified as a putative therapeutic target of escitalopram in the inflammatory cytokine pathway. Journal of Psychopharmacology, 2013, 27, 609-615.	4.0	20
118	Paternal Age Alters Social Development in Offspring. Journal of the American Academy of Child and Adolescent Psychiatry, 2017, 56, 383-390.	0.5	20
119	Techniques in mammalian genome mapping. Current Opinion in Biotechnology, 1995, 6, 37-43.	6.6	19
120	Toward the construction of integrated physical and genetic maps of the mouse genome using interspersed repetitive sequence PCR (IRS-PCR) genomics Genome Research, 1996, 6, 290-299.	5.5	18
121	Construction and characterisation of a gridded chicken cosmid library with four-fold genomic coverage. Animal Genetics, 1998, 29, 295-301.	1.7	18
122	Pharmacoproteomic investigation into antidepressant response in two mouse inbred strains. Proteomics, 2012, 12, 2355-2365.	2.2	18
123	A yeast artificial chromosome (YAC) library containing 10 haploid chicken genome equivalents. Mammalian Genome, 1997, 8, 274-276.	2.2	16
124	High-Throughput Scanning of the Rat Genome Using Interspersed Repetitive Sequence-PCR Markers. Genomics, 2000, 69, 287-294.	2.9	16
125	IRS-PCR-based genetic mapping of the huntingtin interacting protein gene (HIP1) on mouse Chromosome 5. Mammalian Genome, 1998, 9, 26-31.	2.2	15
126	To what extent is blood a reasonable surrogate for brain in gene expression studies: estimation from mouse hippocampus and spleen. Frontiers in Neuroscience, 2009, 3, 54.	2.8	15

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127	DNA methylation at the lgf2/H19 imprinting control region is associated with cerebellum mass in outbred mice. Molecular Brain, 2012, 5, 42.	2.6	15
128	Home-cage activity in heterogeneous stock (HS) mice as a model of baseline activity. Genes, Brain and Behavior, 2002, 1, 166-173.	2.2	14
129	Moodâ€stabilizers differentially affect housekeeping gene expression in human cells. International Journal of Methods in Psychiatric Research, 2014, 23, 279-288.	2.1	14
130	RNA sequencing of identical twins discordant for autism reveals blood-based signatures implicating immune and transcriptional dysregulation. Molecular Autism, 2019, 10, 38.	4.9	14
131	The DNA methylome of human sperm is distinct from blood with little evidence for tissue-consistent obesity associations. PLoS Genetics, 2020, 16, e1009035.	3.5	13
132	Transcriptomic changes in the frontal cortex associated with paternal age. Molecular Autism, 2014, 5, 24.	4.9	11
133	ETn insertion in the mouse Adcy1 gene: transcriptional and phylogenetic analyses. Mammalian Genome, 2000, 11, 97-103.	2.2	10
134	Highly polygenic architecture of antidepressant treatment response: Comparative analysis of SSRI and NRI treatment in an animal model of depression. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2017, 174, 235-250.	1.7	10
135	Gene set enrichment; a problem of pathways. Briefings in Functional Genomics, 2010, 9, 385-390.	2.7	9
136	Comparative mRNA analysis of behavioral and genetic mouse models of aggression. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2016, 171, 427-436.	1.7	9
137	DNA methylation-based sex classifier to predict sex and identify sex chromosome aneuploidy. BMC Genomics, 2021, 22, 484.	2.8	9
138	Chapter 15 Halobacterial genes and genomes. New Comprehensive Biochemistry, 1993, , 467-496.	0.1	8
139	Technology development at the interface of proteome research and genomics: Mapping nonpolymorphic proteins on the physical map of mouse chromosomes. Electrophoresis, 1999, 20, 1027-1032.	2.4	8
140	Effect of Chronic Valproic Acid Treatment on Hepatic Gene Expression Profile inWfs1Knockout Mouse. PPAR Research, 2014, 2014, 1-11.	2.4	8
141	Advanced Integrated Mouse YAC Map Including BAC Framework. Genome Research, 2001, 11, 2142-2150.	5.5	7
142	Hypothalamic gene expression profile indicates a reduction in G protein signaling in the <i>Wfs1</i> mutant mice. Physiological Genomics, 2011, 43, 1351-1358.	2.3	7
143	Refined radiation hybrid map of mouse Chromosome 17. Mammalian Genome, 1998, 9, 807-811.	2.2	6
144	Long-Term Effects of Gestational Nicotine Exposure and Food-Restriction on Gene Expression in the Striatum of Adolescent Rats. PLoS ONE, 2014, 9, e88896.	2.5	5

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145	InterpolatedXY: a two-step strategy to normalize DNA methylation microarray data avoiding sex bias. Bioinformatics, 2022, 38, 3950-3957.	4.1	5
146	Insertion sequences on plasmid pHV1 of <i>Haloferax volcanii</i> . Canadian Journal of Microbiology, 1993, 39, 201-206.	1.7	4
147	Complex probes for high-throughput parallel genetic mapping of genomic mouse BAC clones. Mammalian Genome, 1998, 9, 611-616.	2.2	4
148	Assessing Individual Differences in Genome-Wide Gene Expression in Human Whole Blood: Reliability Over Four Hours and Stability Over 10 Months. Twin Research and Human Genetics, 2009, 12, 372-380.	0.6	4
149	Utility of the pooling approach as applied to whole genome association scans with high-density Affymetrix microarrays. BMC Research Notes, 2010, 3, 274.	1.4	3
150	Peripheral blood RNA expression profiling in illicit methcathinone users reveals effect on immune system. Frontiers in Genetics, 2011, 2, 42.	2.3	3
151	Genetic polymorphisms and their association with brain and behavioural measures in heterogeneous stock mice. Scientific Reports, 2017, 7, 41204.	3.3	2
152	New tools for the high throughput characterization of rat genomic DNA samples. Journal of Experimental Animal Science, 2000, 41, 35-37.	0.5	1
153	Introduction to mammalian genome special issue: genetics of behavior. Mammalian Genome, 2014, 25, 1-2.	2.2	1
154	Introduction to the Special Issue: Aggression Across Species. Behavior Genetics, 2003, 33, 457-460.	2.1	0
155	S.16.02 Depression: the GENDEP study. European Neuropsychopharmacology, 2006, 16, S187.	0.7	0
156	P.1.31 Gene expression analyses of mouse fibroblast cell line L929 after antidepressant treatment. European Neuropsychopharmacology, 2007, 17, S27-S28.	0.7	0
157	Response to comment by Stuart Macgregor. Behavior Genetics, 2010, 40, 48-48.	2.1	0
158	O3-04-03: CROSS-TISSUE METHYLOMIC PROFILING IN ALZHEIMER'S DISEASE. , 2014, 10, P215-P215.		0
159	Technology development at the interface of proteome research and genomics: Mapping nonpolymorphic proteins on the physical map of mouse chromosomes. Electrophoresis, 1999, 20,	2.4	0

1027-1032