

Naomi R Wray

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

399 papers	42,488 citations	93 h-index	201 g-index
472 ext. papers	55,930 ext. citations	10.6 avg, IF	7.65 L-index

#	Paper	IF	Citations
399	Impact of CYP2C19 metaboliser status on SSRI response: a retrospective study of 9500 participants of the Australian Genetics of Depression Study.. <i>Pharmacogenomics Journal</i> , 2022 ,	3.5	1
398	Genome-wide identification of the genetic basis of amyotrophic lateral sclerosis.. <i>Neuron</i> , 2022 ,	13.9	8
397	Functional characterisation of the amyotrophic lateral sclerosis risk locus GPX3/TNIP1.. <i>Genome Medicine</i> , 2022 , 14, 7	14.4	0
396	Investigating the phenotypic and genetic associations between personality traits and suicidal behavior across major mental health diagnoses.. <i>European Archives of Psychiatry and Clinical Neuroscience</i> , 2022 , 1	5.1	0
395	Genome-wide study of DNA methylation shows alterations in metabolic, inflammatory, and cholesterol pathways in ALS.. <i>Science Translational Medicine</i> , 2022 , 14, eabj0264	17.5	4
394	Mapping genomic loci implicates genes and synaptic biology in schizophrenia.. <i>Nature</i> , 2022 ,	50.4	35
393	Comprehensive genetic analysis of the human lipidome identifies loci associated with lipid homeostasis with links to coronary artery disease. <i>Nature Communications</i> , 2022 , 13,	17.4	5
392	Common and rare variant association analyses in amyotrophic lateral sclerosis identify 15 risk loci with distinct genetic architectures and neuron-specific biology. <i>Nature Genetics</i> , 2021 , 53, 1636-1648	36.3	19
391	Understanding genetic risk factors for common side effects of antidepressant medications. <i>Communications Medicine</i> , 2021 , 1,		2
390	Autism-related dietary preferences mediate autism-gut microbiome associations. <i>Cell</i> , 2021 , 184, 5916-5931.e17	59.1	17
389	Polygenic burden could explain high rates of affective disorders in a community with restricted founder population. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2021 , 186, 367-375	3.5	
388	The Australian Genetics of Depression Study: New Risk Loci and Dissecting Heterogeneity Between Subtypes.. <i>Biological Psychiatry</i> , 2021 ,	7.9	2
387	Conditional GWAS analysis to identify disorder-specific SNPs for psychiatric disorders. <i>Molecular Psychiatry</i> , 2021 , 26, 2070-2081	15.1	19
386	Meta-analysis of genome-wide DNA methylation identifies shared associations across neurodegenerative disorders. <i>Genome Biology</i> , 2021 , 22, 90	18.3	6
385	Genome-wide gene expression changes in postpartum depression point towards an altered immune landscape. <i>Translational Psychiatry</i> , 2021 , 11, 155	8.6	5
384	Polygenic risk score analysis for amyotrophic lateral sclerosis leveraging cognitive performance, educational attainment and schizophrenia. <i>European Journal of Human Genetics</i> , 2021 ,	5.3	7
383	Comorbid Chronic Pain and Depression: Shared Risk Factors and Differential Antidepressant Effectiveness. <i>Frontiers in Psychiatry</i> , 2021 , 12, 643609	5	13

382	Risk of Early-Onset Depression Associated With Polygenic Liability, Parental Psychiatric History, and Socioeconomic Status. <i>JAMA Psychiatry</i> , 2021 , 78, 387-397	14.5	11
381	A Comparison of Ten Polygenic Score Methods for Psychiatric Disorders Applied Across Multiple Cohorts. <i>Biological Psychiatry</i> , 2021 , 90, 611-620	7.9	17
380	Genome-wide association study of more than 40,000 bipolar disorder cases provides new insights into the underlying biology. <i>Nature Genetics</i> , 2021 , 53, 817-829	36.3	83
379	Estimation of non-additive genetic variance in human complex traits from a large sample of unrelated individuals. <i>American Journal of Human Genetics</i> , 2021 , 108, 786-798	11	19
378	Gene action, genetic variation, and GWAS: A user-friendly web tool. <i>PLoS Genetics</i> , 2021 , 17, e1009548	6	0
377	Leveraging both individual-level genetic data and GWAS summary statistics increases polygenic prediction. <i>American Journal of Human Genetics</i> , 2021 , 108, 1001-1011	11	2
376	Examining Sex-Differentiated Genetic Effects Across Neuropsychiatric and Behavioral Traits. <i>Biological Psychiatry</i> , 2021 , 89, 1127-1137	7.9	12
375	Association of Antihypertensive Drug Target Genes With Psychiatric Disorders: A Mendelian Randomization Study. <i>JAMA Psychiatry</i> , 2021 , 78, 623-631	14.5	0
374	Genetic risk for chronic pain is associated with lower antidepressant effectiveness: Converging evidence for a depression subtype. <i>Australian and New Zealand Journal of Psychiatry</i> , 2021 , 48, 674-681	26.1	0
373	Genetic association study of childhood aggression across raters, instruments, and age. <i>Translational Psychiatry</i> , 2021 , 11, 413	8.6	7
372	Risk in Relatives, Heritability, SNP-Based Heritability, and Genetic Correlations in Psychiatric Disorders: A Review. <i>Biological Psychiatry</i> , 2021 , 89, 11-19	7.9	20
371	Cardiovascular disease, psychiatric diagnosis and sex differences in the multistep hypothesis of amyotrophic lateral sclerosis. <i>European Journal of Neurology</i> , 2021 , 28, 421-429	6	7
370	From Basic Science to Clinical Application of Polygenic Risk Scores: A Primer. <i>JAMA Psychiatry</i> , 2021 , 78, 101-109	14.5	49
369	Could Polygenic Risk Scores Be Useful in Psychiatry?: A Review. <i>JAMA Psychiatry</i> , 2021 , 78, 210-219	14.5	53
368	Genome-wide analyses of behavioural traits are subject to bias by misreports and longitudinal changes. <i>Nature Communications</i> , 2021 , 12, 20211	17.4	16
367	GWAS of peptic ulcer disease implicates <i>Helicobacter pylori</i> infection, other gastrointestinal disorders and depression. <i>Nature Communications</i> , 2021 , 12, 1146	17.4	20
366	Widespread signatures of natural selection across human complex traits and functional genomic categories. <i>Nature Communications</i> , 2021 , 12, 1164	17.4	12
365	Analysis of common genetic variation and rare CNVs in the Australian Autism Biobank. <i>Molecular Autism</i> , 2021 , 12, 12	6.5	4

364	Schizophrenia polygenic risk scores in youth mental health: preliminary associations with diagnosis, clinical stage and functioning. <i>BJPsych Open</i> , 2021 , 7, e58	5	1
363	Phenotypic covariance across the entire spectrum of relatedness for 86 billion pairs of individuals. <i>Nature Communications</i> , 2021 , 12, 1050	17.4	7
362	Genomic partitioning of inbreeding depression in humans. <i>American Journal of Human Genetics</i> , 2021 , 108, 1488-1501	11	3
361	Polygenic Risk Scores Derived From Varying Definitions of Depression and Risk of Depression. <i>JAMA Psychiatry</i> , 2021 , 78, 1152-1160	14.5	3
360	Continuity of Genetic Risk for Aggressive Behavior Across the Life-Course. <i>Behavior Genetics</i> , 2021 , 51, 592-606	3.2	2
359	Investigating Shared Genetic Basis Across Tourette Syndrome and Comorbid Neurodevelopmental Disorders Along the Impulsivity-Compulsivity Spectrum. <i>Biological Psychiatry</i> , 2021 , 90, 317-327	7.9	12
358	MiND AUS partnership: a roadmap for the cure and management of motor Neurone disease. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2021 , 1-8	3.6	0
357	Discovery and implications of polygenicity of common diseases. <i>Science</i> , 2021 , 373, 1468-1473	33.3	13
356	Genomic and phenotypic insights from an atlas of genetic effects on DNA methylation. <i>Nature Genetics</i> , 2021 , 53, 1311-1321	36.3	27
355	The Genetic Architecture of Depression in Individuals of East Asian Ancestry: A Genome-Wide Association Study. <i>JAMA Psychiatry</i> , 2021 , 78, 1258-1269	14.5	7
354	Dissecting the Shared Genetic Architecture of Suicide Attempt, Psychiatric Disorders, and Known Risk Factors. <i>Biological Psychiatry</i> , 2021 ,	7.9	11
353	Refining Attention-Deficit/Hyperactivity Disorder and Autism Spectrum Disorder Genetic Loci by Integrating Summary Data From Genome-wide Association, Gene Expression, and DNA Methylation Studies. <i>Biological Psychiatry</i> , 2020 , 88, 470-479	7.9	6
352	Nick Martin and the Genetics of Depression: Sample Size, Sample Size, Sample Size. <i>Twin Research and Human Genetics</i> , 2020 , 23, 109-111	2.2	
351	ALS in Danish Registries: Heritability and links to psychiatric and cardiovascular disorders. <i>Neurology: Genetics</i> , 2020 , 6, e398	3.8	15
350	Cohort profile: the Australian genetics of depression study. <i>BMJ Open</i> , 2020 , 10, e032580	3	13
349	Mutations in heat shock protein beta-1 (HSPB1) are associated with a range of clinical phenotypes related to different patterns of motor neuron dysfunction: A case series. <i>Journal of the Neurological Sciences</i> , 2020 , 413, 116809	3.2	3
348	Genetic stratification of depression in UK Biobank. <i>Translational Psychiatry</i> , 2020 , 10, 163	8.6	8
347	Bayesian reassessment of the epigenetic architecture of complex traits. <i>Nature Communications</i> , 2020 , 11, 2865	17.4	18

346	What do we know about the variability in survival of patients with amyotrophic lateral sclerosis?. <i>Expert Review of Neurotherapeutics</i> , 2020 , 20, 921-941	4.3	4
345	Analysis of DNA methylation associates the cystine-glutamate antiporter SLC7A11 with risk of Parkinson's disease. <i>Nature Communications</i> , 2020 , 11, 1238	17.4	25
344	Significant out-of-sample classification from methylation profile scoring for amyotrophic lateral sclerosis. <i>Npj Genomic Medicine</i> , 2020 , 5, 10	6.2	11
343	Association of Mental Disorder in Childhood and Adolescence With Subsequent Educational Achievement. <i>JAMA Psychiatry</i> , 2020 , 77, 797-805	14.5	31
342	Multi-method genome- and epigenome-wide studies of inflammatory protein levels in healthy older adults. <i>Genome Medicine</i> , 2020 , 12, 60	14.4	9
341	Progression and survival of patients with motor neuron disease relative to their fecal microbiota. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2020 , 21, 549-562	3.6	11
340	Genome-wide association study of dietary intake in the UK biobank study and its associations with schizophrenia and other traits. <i>Translational Psychiatry</i> , 2020 , 10, 51	8.6	10
339	Genome-wide gene-environment analyses of major depressive disorder and reported lifetime traumatic experiences in UK Biobank. <i>Molecular Psychiatry</i> , 2020 , 25, 1430-1446	15.1	47
338	Promoter-anchored chromatin interactions predicted from genetic analysis of epigenomic data. <i>Nature Communications</i> , 2020 , 11, 2061	17.4	1
337	Genome-wide association study identifies 143 loci associated with 25 hydroxyvitamin D concentration. <i>Nature Communications</i> , 2020 , 11, 1647	17.4	58
336	Blood DNA methylation sites predict death risk in a longitudinal study of 12, 300 individuals. <i>Aging</i> , 2020 , 12, 14092-14124	5.6	6
335	repeat expansions confer risk for amyotrophic lateral sclerosis and contribute to TDP-43 mislocalization. <i>Brain Communications</i> , 2020 , 2, fcaa064	4.5	12
334	A unified framework for association and prediction from vertex-wise grey-matter structure. <i>Human Brain Mapping</i> , 2020 , 41, 4062-4076	5.9	3
333	Genetic comorbidity between major depression and cardio-metabolic traits, stratified by age at onset of major depression. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2020 , 183, 309-330	3.5	8
332	Rare Variant Burden Analysis within Enhancers Identifies CAV1 as an ALS Risk Gene. <i>Cell Reports</i> , 2020 , 33, 108456	10.6	6
331	Genome-wide Meta-analysis Finds the ACSL5-ZDHHC6 Locus Is Associated with ALS and Links Weight Loss to the Disease Genetics. <i>Cell Reports</i> , 2020 , 33, 108323	10.6	18
330	Risk prediction of late-onset Alzheimer's disease implies an oligogenic architecture. <i>Nature Communications</i> , 2020 , 11, 4799	17.4	41
329	Genetic control of temperament traits across species: association of autism spectrum disorder risk genes with cattle temperament. <i>Genetics Selection Evolution</i> , 2020 , 52, 51	4.9	9

328	RICOPILI: Rapid Imputation for CONsortias PIPELine. <i>Bioinformatics</i> , 2020 , 36, 930-933	7.2	72
327	Evaluating the Impact of Nonrandom Mating: Psychiatric Outcomes Among the Offspring of Pairs Diagnosed With Schizophrenia and Bipolar Disorder. <i>Biological Psychiatry</i> , 2020 , 87, 253-262	7.9	4
326	Classical Human Leukocyte Antigen Alleles and C4 Haplotypes Are Not Significantly Associated With Depression. <i>Biological Psychiatry</i> , 2020 , 87, 419-430	7.9	9
325	Improved precision of epigenetic clock estimates across tissues and its implication for biological ageing. <i>Genome Medicine</i> , 2019 , 11, 54	14.4	81
324	Extreme inbreeding in a European ancestry sample from the contemporary UK population. <i>Nature Communications</i> , 2019 , 10, 3719	17.4	14
323	A genome-wide association study of shared risk across psychiatric disorders implicates gene regulation during fetal neurodevelopment. <i>Nature Neuroscience</i> , 2019 , 22, 353-361	25.5	93
322	Quantifying between-cohort and between-sex genetic heterogeneity in major depressive disorder. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2019 , 180, 439-447	3.5	16
321	Sleep Disorders and Risk of Incident Depression: A Population Case-Control Study. <i>Twin Research and Human Genetics</i> , 2019 , 22, 140-146	2.2	12
320	Genetic correlations of polygenic disease traits: from theory to practice. <i>Nature Reviews Genetics</i> , 2019 , 20, 567-581	30.1	98
319	OSCA: a tool for omic-data-based complex trait analysis. <i>Genome Biology</i> , 2019 , 20, 107	18.3	40
318	GWAS of Suicide Attempt in Psychiatric Disorders and Association With Major Depression Polygenic Risk Scores. <i>American Journal of Psychiatry</i> , 2019 , 176, 651-660	11.9	103
317	Gut microbiota in ALS: possible role in pathogenesis?. <i>Expert Review of Neurotherapeutics</i> , 2019 , 19, 785-805	19.5	21
316	Examining the Impact of Imputation Errors on Fine-Mapping Using DNA Methylation QTL as a Model Trait. <i>Genetics</i> , 2019 , 212, 577-586	4	1
315	Genotype-covariate correlation and interaction disentangled by a whole-genome multivariate reaction norm model. <i>Nature Communications</i> , 2019 , 10, 2239	17.4	23
314	Genome-wide association study of medication-use and associated disease in the UK Biobank. <i>Nature Communications</i> , 2019 , 10, 1891	17.4	48
313	Genome-wide association study identifies 30 loci associated with bipolar disorder. <i>Nature Genetics</i> , 2019 , 51, 793-803	36.3	662
312	Complex Trait Prediction from Genome Data: Contrasting EBV in Livestock to PRS in Humans: Genomic Prediction. <i>Genetics</i> , 2019 , 211, 1131-1141	4	47
311	A DIRECT TEST OF THE DIATHESIS-STRESS MODEL FOR DEPRESSION. <i>European Neuropsychopharmacology</i> , 2019 , 29, S805-S806	1.2	2

310	Evidence of causal effect of major depression on alcohol dependence: findings from the psychiatric genomics consortium. <i>Psychological Medicine</i> , 2019 , 49, 1218-1226	6.9	33
309	Attention deficit hyperactivity disorder symptoms as antecedents of later psychotic outcomes in 22q11.2 deletion syndrome. <i>Schizophrenia Research</i> , 2019 , 204, 320-325	3.6	11
308	Genotype-by-environment interactions inferred from genetic effects on phenotypic variability in the UK Biobank. <i>Science Advances</i> , 2019 , 5, eaaw3538	14.3	59
307	Assortative Mating in Autism Spectrum Disorder: Toward an Evidence Base From DNA Data, but Not There Yet. <i>Biological Psychiatry</i> , 2019 , 86, 250-252	7.9	1
306	Genome and epigenome wide studies of neurological protein biomarkers in the Lothian Birth Cohort 1936. <i>Nature Communications</i> , 2019 , 10, 3160	17.4	21
305	"Arte et Labore"-A Blackburn Rovers fan's legacy in human complex trait genetics. <i>Journal of Animal Breeding and Genetics</i> , 2019 , 136, 273-278	2.9	0
304	Association of Schizophrenia Risk With Disordered Niacin Metabolism in an Indian Genome-wide Association Study. <i>JAMA Psychiatry</i> , 2019 , 76, 1026-1034	14.5	24
303	Genetic risk scores for major psychiatric disorders and the risk of postpartum psychiatric disorders. <i>Translational Psychiatry</i> , 2019 , 9, 288	8.6	10
302	Genetic correlates of social stratification in Great Britain. <i>Nature Human Behaviour</i> , 2019 , 3, 1332-1342	12.8	83
301	Genome-wide meta-analysis of depression identifies 102 independent variants and highlights the importance of the prefrontal brain regions. <i>Nature Neuroscience</i> , 2019 , 22, 343-352	25.5	639
300	Improved polygenic prediction by Bayesian multiple regression on summary statistics. <i>Nature Communications</i> , 2019 , 10, 5086	17.4	114
299	A resource-efficient tool for mixed model association analysis of large-scale data. <i>Nature Genetics</i> , 2019 , 51, 1749-1755	36.3	102
298	Association of Whole-Genome and NETRIN1 Signaling Pathway-Derived Polygenic Risk Scores for Major Depressive Disorder and White Matter Microstructure in the UK Biobank. <i>Biological Psychiatry: Cognitive Neuroscience and Neuroimaging</i> , 2019 , 4, 91-100	3.4	12
297	Cumulative influence of parity-related genomic changes in multiple sclerosis. <i>Journal of Neuroimmunology</i> , 2019 , 328, 38-49	3.5	4
296	Is Schizophrenia a Risk Factor for Breast Cancer?-Evidence From Genetic Data. <i>Schizophrenia Bulletin</i> , 2019 , 45, 1251-1256	1.3	11
295	Improving genetic prediction by leveraging genetic correlations among human diseases and traits. <i>Nature Communications</i> , 2018 , 9, 989	17.4	76
294	Reply to Kardos et al.: Estimation of inbreeding depression from SNP data. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018 , 115, E2494-E2495	11.5	4
293	Common schizophrenia alleles are enriched in mutation-intolerant genes and in regions under strong background selection. <i>Nature Genetics</i> , 2018 , 50, 381-389	36.3	787

292	Integrative analysis of omics summary data reveals putative mechanisms underlying complex traits. <i>Nature Communications</i> , 2018 , 9, 918	17.4	110
291	Sizing up whole-genome sequencing studies of common diseases. <i>Nature Genetics</i> , 2018 , 50, 635-637	36.3	10
290	Signatures of negative selection in the genetic architecture of human complex traits. <i>Nature Genetics</i> , 2018 , 50, 746-753	36.3	178
289	GWAS of epigenetic aging rates in blood reveals a critical role for TERT. <i>Nature Communications</i> , 2018 , 9, 387	17.4	106
288	Causal associations between risk factors and common diseases inferred from GWAS summary data. <i>Nature Communications</i> , 2018 , 9, 224	17.4	346
287	Hypermetabolism in ALS is associated with greater functional decline and shorter survival. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018 , 89, 1016-1023	5.5	96
286	Genome-wide association analyses identify 44 risk variants and refine the genetic architecture of major depression. <i>Nature Genetics</i> , 2018 , 50, 668-681	36.3	1301
285	A DNA methylation biomarker of alcohol consumption. <i>Molecular Psychiatry</i> , 2018 , 23, 422-433	15.1	164
284	Brain age predicts mortality. <i>Molecular Psychiatry</i> , 2018 , 23, 1385-1392	15.1	260
283	A direct test of the diathesis-stress model for depression. <i>Molecular Psychiatry</i> , 2018 , 23, 1590-1596	15.1	114
282	Does Childhood Trauma Moderate Polygenic Risk for Depression? A Meta-analysis of 5765 Subjects From the Psychiatric Genomics Consortium. <i>Biological Psychiatry</i> , 2018 , 84, 138-147	7.9	48
281	Embracing polygenicity: a review of methods and tools for psychiatric genetics research. <i>Psychological Medicine</i> , 2018 , 48, 1055-1067	6.9	48
280	The epigenetic clock and telomere length are independently associated with chronological age and mortality. <i>International Journal of Epidemiology</i> , 2018 , 45, 424-432	7.8	153
279	Accuracy of Inferred APOE Genotypes for a Range of Genotyping Arrays and Imputation Reference Panels. <i>Journal of Alzheimer's Disease</i> , 2018 , 64, 49-54	4.3	5
278	Analysis of the Influence of microRNAs in Lithium Response in Bipolar Disorder. <i>Frontiers in Psychiatry</i> , 2018 , 9, 207	5	15
277	Analysis of shared heritability in common disorders of the brain. <i>Science</i> , 2018 , 360,	33.3	666
276	DNA Methylation Signatures of Depressive Symptoms in Middle-aged and Elderly Persons: Meta-analysis of Multiethnic Epigenome-wide Studies. <i>JAMA Psychiatry</i> , 2018 , 75, 949-959	14.5	51
275	Familiality of Psychiatric Disorders and Risk of Postpartum Psychiatric Episodes: A Population-Based Cohort Study. <i>American Journal of Psychiatry</i> , 2018 , 175, 783-791	11.9	14

274	Comparison of Genotypic and Phenotypic Correlations: Cheverud's Conjecture in Humans. <i>Genetics</i> , 2018 , 209, 941-948	4	48
273	Common Disease Is More Complex Than Implied by the Core Gene Omnigenic Model. <i>Cell</i> , 2018 , 173, 1573-1580	56.2	151
272	Identifying gene targets for brain-related traits using transcriptomic and methylomic data from blood. <i>Nature Communications</i> , 2018 , 9, 2282	17.4	147
271	Genome-wide gene-environment interaction in depression: A systematic evaluation of candidate genes: The childhood trauma working-group of PGC-MDD. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2018 , 177, 40-49	3.5	43
270	A Genetic Investigation of Sex Bias in the Prevalence of Attention-Deficit/Hyperactivity Disorder. <i>Biological Psychiatry</i> , 2018 , 83, 1044-1053	7.9	93
269	Identification of 55,000 Replicated DNA Methylation QTL. <i>Scientific Reports</i> , 2018 , 8, 17605	4.9	78
268	Imprint of assortative mating on the human genome. <i>Nature Human Behaviour</i> , 2018 , 2, 948-954	12.8	45
267	Trajectories of inflammatory biomarkers over the eighth decade and their associations with immune cell profiles and epigenetic ageing. <i>Clinical Epigenetics</i> , 2018 , 10, 159	7.7	17
266	Dissection of genetic variation and evidence for pleiotropy in male pattern baldness. <i>Nature Communications</i> , 2018 , 9, 5407	17.4	37
265	The association between neonatal vitamin D status and risk of schizophrenia. <i>Scientific Reports</i> , 2018 , 8, 17692	4.9	49
264	PPD ACT: an app-based genetic study of postpartum depression. <i>Translational Psychiatry</i> , 2018 , 8, 260	8.6	10
263	Epigenetic prediction of complex traits and death. <i>Genome Biology</i> , 2018 , 19, 136	18.3	77
262	Genotype effects contribute to variation in longitudinal methylome patterns in older people. <i>Genome Medicine</i> , 2018 , 10, 75	14.4	21
261	Study protocol for the Australian autism biobank: an international resource to advance autism discovery research. <i>BMC Pediatrics</i> , 2018 , 18, 284	2.6	9
260	GWAS on family history of Alzheimer's disease. <i>Translational Psychiatry</i> , 2018 , 8, 99	8.6	238
259	Estimation of Genetic Correlation via Linkage Disequilibrium Score Regression and Genomic Restricted Maximum Likelihood. <i>American Journal of Human Genetics</i> , 2018 , 102, 1185-1194	11	55
258	Age at first birth in women is genetically associated with increased risk of schizophrenia. <i>Scientific Reports</i> , 2018 , 8, 10168	4.9	11
257	Trans-eQTLs identified in whole blood have limited influence on complex disease biology. <i>European Journal of Human Genetics</i> , 2018 , 26, 1361-1368	5.3	1

256	Misestimation of heritability and prediction accuracy of male-pattern baldness. <i>Nature Communications</i> , 2018 , 9, 2537	17.4	14
255	A Combined Pathway and Regional Heritability Analysis Indicates NETRIN1 Pathway Is Associated With Major Depressive Disorder. <i>Biological Psychiatry</i> , 2017 , 81, 336-346	7.9	25
254	Genome-wide Association for Major Depression Through Age at Onset Stratification: Major Depressive Disorder Working Group of the Psychiatric Genomics Consortium. <i>Biological Psychiatry</i> , 2017 , 81, 325-335	7.9	129
253	Using information of relatives in genomic prediction to apply effective stratified medicine. <i>Scientific Reports</i> , 2017 , 7, 42091	4.9	31
252	Genome-wide association study of borderline personality disorder reveals genetic overlap with bipolar disorder, major depression and schizophrenia. <i>Translational Psychiatry</i> , 2017 , 7, e1155	8.6	100
251	Genetic signatures of high-altitude adaptation in Tibetans. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, 4189-4194	11.5	93
250	Genetic effects influencing risk for major depressive disorder in China and Europe. <i>Translational Psychiatry</i> , 2017 , 7, e1074	8.6	48
249	Extra-motor abnormalities in amyotrophic lateral sclerosis: another layer of heterogeneity. <i>Expert Review of Neurotherapeutics</i> , 2017 , 17, 561-577	4.3	14
248	Genome-wide Regional Heritability Mapping Identifies a Locus Within the TOX2 Gene Associated With Major Depressive Disorder. <i>Biological Psychiatry</i> , 2017 , 82, 312-321	7.9	17
247	CNV-association meta-analysis in 191,161 European adults reveals new loci associated with anthropometric traits. <i>Nature Communications</i> , 2017 , 8, 744	17.4	37
246	Inference in Psychiatry via 2-Sample Mendelian Randomization-From Association to Causal Pathway?. <i>JAMA Psychiatry</i> , 2017 , 74, 1191-1192	14.5	11
245	Genetic Association of Major Depression With Atypical Features and Obesity-Related Immunometabolic Dysregulations. <i>JAMA Psychiatry</i> , 2017 , 74, 1214-1225	14.5	109
244	Association of Body Mass Index with DNA Methylation and Gene Expression in Blood Cells and Relations to Cardiometabolic Disease: A Mendelian Randomization Approach. <i>PLoS Medicine</i> , 2017 , 14, e1002215	11.6	162
243	Performance of risk prediction for inflammatory bowel disease based on genotyping platform and genomic risk score method. <i>BMC Medical Genetics</i> , 2017 , 18, 94	2.1	20
242	Whole-exome sequencing in amyotrophic lateral sclerosis suggests NEK1 is a risk gene in Chinese. <i>Genome Medicine</i> , 2017 , 9, 97	14.4	17
241	Concepts, estimation and interpretation of SNP-based heritability. <i>Nature Genetics</i> , 2017 , 49, 1304-1310	36.3	217
240	Comparison of faecal microbe diversity between motor neurone disease (mnd) and control participants. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017 , 88, e1.83-e1	5.5	
239	Cross-ethnic meta-analysis identifies association of the GPX3-TNIP1 locus with amyotrophic lateral sclerosis. <i>Nature Communications</i> , 2017 , 8, 611	17.4	45

238	Whole exome sequencing and DNA methylation analysis in a clinical amyotrophic lateral sclerosis cohort. <i>Molecular Genetics & Genomic Medicine</i> , 2017 , 5, 418-428	2.3	8
237	Detection and quantification of inbreeding depression for complex traits from SNP data. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, 8602-8607	11.5	20
236	Large-scale GWAS identifies multiple loci for hand grip strength providing biological insights into muscular fitness. <i>Nature Communications</i> , 2017 , 8, 16015	17.4	80
235	10 Years of GWAS Discovery: Biology, Function, and Translation. <i>American Journal of Human Genetics</i> , 2017 , 101, 5-22	11	1651
234	Investigating the relationship between iron and depression. <i>Journal of Psychiatric Research</i> , 2017 , 94, 148-155	5.2	3
233	Rare DNA variants in the brain-derived neurotrophic factor gene increase risk for attention-deficit hyperactivity disorder: a next-generation sequencing study. <i>Molecular Psychiatry</i> , 2017 , 22, 580-584	15.1	25
232	Contribution of copy number variants to schizophrenia from a genome-wide study of 41,321 subjects. <i>Nature Genetics</i> , 2017 , 49, 27-35	36.3	530
231	High loading of polygenic risk in cases with chronic schizophrenia. <i>Molecular Psychiatry</i> , 2016 , 21, 969-74	15.1	44
230	Evidence of CNH3 involvement in opioid dependence. <i>Molecular Psychiatry</i> , 2016 , 21, 608-14	15.1	74
229	GCTA-GREML accounts for linkage disequilibrium when estimating genetic variance from genome-wide SNPs. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016 , 113, E4579-80	11.5	25
228	Epigenetic Signatures of Cigarette Smoking. <i>Circulation: Cardiovascular Genetics</i> , 2016 , 9, 436-447		442
227	Genome-wide association analyses identify new risk variants and the genetic architecture of amyotrophic lateral sclerosis. <i>Nature Genetics</i> , 2016 , 48, 1043-8	36.3	328
226	Exploring Boundaries for the Genetic Consequences of Assortative Mating for Psychiatric Traits. <i>JAMA Psychiatry</i> , 2016 , 73, 1189-1195	14.5	36
225	Genome-wide association study reveals greater polygenic loading for schizophrenia in cases with a family history of illness. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2016 , 171B, 276-89	3.5	23
224	Meta-analysis of genome-wide association studies of anxiety disorders. <i>Molecular Psychiatry</i> , 2016 , 21, 1391-9	15.1	213
223	Genetic variants associated with response to lithium treatment in bipolar disorder: a genome-wide association study. <i>Lancet, The</i> , 2016 , 387, 1085-1093	40	216
222	Evidence for Genetic Overlap Between Schizophrenia and Age at First Birth in Women. <i>JAMA Psychiatry</i> , 2016 , 73, 497-505	14.5	40
221	Integration of summary data from GWAS and eQTL studies predicts complex trait gene targets. <i>Nature Genetics</i> , 2016 , 48, 481-7	36.3	929

220	Disease and Polygenic Architecture: Avoid Trio Design and Appropriately Account for Unscreened Control Subjects for Common Disease. <i>American Journal of Human Genetics</i> , 2016 , 98, 382-91	11	27
219	Across-cohort QC analyses of GWAS summary statistics from complex traits. <i>European Journal of Human Genetics</i> , 2016 , 25, 137-146	5.3	13
218	DNA methylation-based measures of biological age: meta-analysis predicting time to death. <i>Aging</i> , 2016 , 8, 1844-1865	5.6	531
217	Predicting gene targets from integrative analyses of summary data from GWAS and eQTL studies for 28 human complex traits. <i>Genome Medicine</i> , 2016 , 8, 84	14.4	59
216	A method to decipher pleiotropy by detecting underlying heterogeneity driven by hidden subgroups applied to autoimmune and neuropsychiatric diseases. <i>Nature Genetics</i> , 2016 , 48, 803-10	36.3	45
215	Risk of psychiatric illness from advanced paternal age is not predominantly from de novo mutations. <i>Nature Genetics</i> , 2016 , 48, 718-24	36.3	74
214	Genetic overlap between diagnostic subtypes of ischemic stroke. <i>Stroke</i> , 2015 , 46, 615-9	6.7	33
213	Improving Phenotypic Prediction by Combining Genetic and Epigenetic Associations. <i>American Journal of Human Genetics</i> , 2015 , 97, 75-85	11	85
212	Simultaneous discovery, estimation and prediction analysis of complex traits using a bayesian mixture model. <i>PLoS Genetics</i> , 2015 , 11, e1004969	6	206
211	Cohort Profile Update: The Mater-University of Queensland Study of Pregnancy (MUSP). <i>International Journal of Epidemiology</i> , 2015 , 44, 78-78f	7.8	82
210	Polygenic Risk Score, Parental Socioeconomic Status, Family History of Psychiatric Disorders, and the Risk for Schizophrenia: A Danish Population-Based Study and Meta-analysis. <i>JAMA Psychiatry</i> , 2015 , 72, 635-41	14.5	177
209	DNA methylation age of blood predicts all-cause mortality in later life. <i>Genome Biology</i> , 2015 , 16, 25	18.3	670
208	Heritability of Transforming Growth Factor- β and Tumor Necrosis Factor-Receptor Type 1 Expression and Vitamin D Levels in Healthy Adolescent Twins. <i>Twin Research and Human Genetics</i> , 2015 , 18, 28-35	2.2	16
207	Contrasting genetic architectures of schizophrenia and other complex diseases using fast variance-components analysis. <i>Nature Genetics</i> , 2015 , 47, 1385-92	36.3	299
206	Novel directions for G \times E analysis in psychiatry. <i>Epidemiology and Psychiatric Sciences</i> , 2015 , 24, 12-9	5.1	10
205	The epigenetic clock is correlated with physical and cognitive fitness in the Lothian Birth Cohort 1936. <i>International Journal of Epidemiology</i> , 2015 , 44, 1388-96	7.8	357
204	Quantitative genetics of disease traits. <i>Journal of Animal Breeding and Genetics</i> , 2015 , 132, 198-203	2.9	9
203	C9orf72 hexanucleotide repeat expansions in Chinese sporadic amyotrophic lateral sclerosis. <i>Neurobiology of Aging</i> , 2015 , 36, 2660.e1-8	5.6	40

202	Genetic variance estimation with imputed variants finds negligible missing heritability for human height and body mass index. <i>Nature Genetics</i> , 2015 , 47, 1114-20	36.3	522
201	New data and an old puzzle: the negative association between schizophrenia and rheumatoid arthritis. <i>International Journal of Epidemiology</i> , 2015 , 44, 1706-21	7.8	43
200	Concepts and Misconceptions about the Polygenic Additive Model Applied to Disease. <i>Human Heredity</i> , 2015 , 80, 165-70	1.1	15
199	The association between lower educational attainment and depression owing to shared genetic effects? Results in ~25,000 subjects. <i>Molecular Psychiatry</i> , 2015 , 20, 735-43	15.1	39
198	Purification of neural precursor cells reveals the presence of distinct, stimulus-specific subpopulations of quiescent precursors in the adult mouse hippocampus. <i>Journal of Neuroscience</i> , 2015 , 35, 8132-44	6.6	38
197	Lifetime stress accelerates epigenetic aging in an urban, African American cohort: relevance of glucocorticoid signaling. <i>Genome Biology</i> , 2015 , 16, 266	18.3	234
196	EMX1 regulates NRP1-mediated wiring of the mouse anterior cingulate cortex. <i>Development (Cambridge)</i> , 2015 , 142, 3746-57	6.6	17
195	DNA modification study of major depressive disorder: beyond locus-by-locus comparisons. <i>Biological Psychiatry</i> , 2015 , 77, 246-255	7.9	49
194	Joint analysis of psychiatric disorders increases accuracy of risk prediction for schizophrenia, bipolar disorder, and major depressive disorder. <i>American Journal of Human Genetics</i> , 2015 , 96, 283-94	11	161
193	Seasonality shows evidence for polygenic architecture and genetic correlation with schizophrenia and bipolar disorder. <i>Journal of Clinical Psychiatry</i> , 2015 , 76, 128-34	4.6	18
192	NFIB-mediated repression of the epigenetic factor Ezh2 regulates cortical development. <i>Journal of Neuroscience</i> , 2014 , 34, 2921-30	6.6	53
191	Testing the role of circadian genes in conferring risk for psychiatric disorders. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2014 , 165B, 254-60	3.5	32
190	Large-scale genomics unveils the genetic architecture of psychiatric disorders. <i>Nature Neuroscience</i> , 2014 , 17, 782-90	25.5	269
189	Partitioning heritability of regulatory and cell-type-specific variants across 11 common diseases. <i>American Journal of Human Genetics</i> , 2014 , 95, 535-52	11	411
188	Overlap of expression quantitative trait loci (eQTL) in human brain and blood. <i>BMC Medical Genomics</i> , 2014 , 7, 31	3.7	41
187	Genetic studies of major depressive disorder: why are there no genome-wide association study findings and what can we do about it?. <i>Biological Psychiatry</i> , 2014 , 76, 510-2	7.9	125
186	The contribution of genetic variants to disease depends on the ruler. <i>Nature Reviews Genetics</i> , 2014 , 15, 765-76	30.1	105
185	Genetic and environmental exposures constrain epigenetic drift over the human life course. <i>Genome Research</i> , 2014 , 24, 1725-33	9.7	123

184	Research review: Polygenic methods and their application to psychiatric traits. <i>Journal of Child Psychology and Psychiatry and Allied Disciplines</i> , 2014 , 55, 1068-87	7.9	410
183	Genetic predisposition to schizophrenia associated with increased use of cannabis. <i>Molecular Psychiatry</i> , 2014 , 19, 1201-4	15.1	136
182	A comparative study of techniques for differential expression analysis on RNA-Seq data. <i>PLoS ONE</i> , 2014 , 9, e103207	3.7	152
181	Estimation and partitioning of (co)heritability of inflammatory bowel disease from GWAS and immunoChip data. <i>Human Molecular Genetics</i> , 2014 , 23, 4710-20	5.6	73
180	Genetic Basis of Complex Genetic Disease: The Contribution of Disease Heterogeneity to Missing Heritability. <i>Current Epidemiology Reports</i> , 2014 , 1, 220-227	2.9	43
179	Response to 'Predicting the diagnosis of autism spectrum disorder using gene pathway analysis'. <i>Molecular Psychiatry</i> , 2014 , 19, 859-61	15.1	14
178	The association between family history of mental disorders and general cognitive ability. <i>Translational Psychiatry</i> , 2014 , 4, e412	8.6	16
177	Statistical power to detect genetic (co)variance of complex traits using SNP data in unrelated samples. <i>PLoS Genetics</i> , 2014 , 10, e1004269	6	236
176	Applying polygenic risk scores to postpartum depression. <i>Archives of Women's Mental Health</i> , 2014 , 17, 519-28	5	49
175	A recessive genetic model and runs of homozygosity in major depressive disorder. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2014 , 165B, 157-66	3.5	19
174	Hypermethylation in the ZBTB20 gene is associated with major depressive disorder. <i>Genome Biology</i> , 2014 , 15, R56	18.3	73
173	Association of OPRD1 polymorphisms with heroin dependence in a large case-control series. <i>Addiction Biology</i> , 2014 , 19, 111-21	4.6	59
172	Explaining additional genetic variation in complex traits. <i>Trends in Genetics</i> , 2014 , 30, 124-32	8.5	110
171	Future Directions in Genetics of Psychiatric Disorders 2014 , 311-337		0
170	The neuroprogressive nature of major depressive disorder: pathways to disease evolution and resistance, and therapeutic implications. <i>Molecular Psychiatry</i> , 2013 , 18, 595-606	15.1	338
169	Genome-wide association analysis identifies 13 new risk loci for schizophrenia. <i>Nature Genetics</i> , 2013 , 45, 1150-9	36.3	1153
168	Author reply to A commentary on Pitfalls of predicting complex traits from SNPs. <i>Nature Reviews Genetics</i> , 2013 , 14, 894	30.1	4
167	A genome-wide association study of sleep habits and insomnia. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2013 , 162B, 439-51	3.5	81

166	Where GWAS and epidemiology meet: opportunities for the simultaneous study of genetic and environmental risk factors in schizophrenia. <i>Schizophrenia Bulletin</i> , 2013 , 39, 955-9	1.3	59
165	Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. <i>Nature Genetics</i> , 2013 , 45, 984-94	36.3	1628
164	Protocol for a collaborative meta-analysis of 5-HTTLPR, stress, and depression. <i>BMC Psychiatry</i> , 2013 , 13, 304	4.2	33
163	Estimation and partition of heritability in human populations using whole-genome analysis methods. <i>Annual Review of Genetics</i> , 2013 , 47, 75-95	14.5	110
162	A genome wide survey supports the involvement of large copy number variants in schizophrenia with and without intellectual disability. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2013 , 162B, 847-54	3.5	13
161	Are surgical trials with negative results being interpreted correctly?. <i>Journal of the American College of Surgeons</i> , 2013 , 216, 158-66	4.4	12
160	Estimation and partitioning of polygenic variation captured by common SNPs for Alzheimer's disease, multiple sclerosis and endometriosis. <i>Human Molecular Genetics</i> , 2013 , 22, 832-41	5.6	147
159	Estimation of SNP heritability from dense genotype data. <i>American Journal of Human Genetics</i> , 2013 , 93, 1151-5	11	85
158	Additive genetic variation in schizophrenia risk is shared by populations of African and European descent. <i>American Journal of Human Genetics</i> , 2013 , 93, 463-70	11	55
157	A mega-analysis of genome-wide association studies for major depressive disorder. <i>Molecular Psychiatry</i> , 2013 , 18, 497-511	15.1	853
156	The heritability of delusional-like experiences. <i>Acta Psychiatrica Scandinavica</i> , 2013 , 127, 48-52	6.5	3
155	Interpreting the role of de novo protein-coding mutations in neuropsychiatric disease. <i>Nature Genetics</i> , 2013 , 45, 234-8	36.3	64
154	Identification of risk loci with shared effects on five major psychiatric disorders: a genome-wide analysis. <i>Lancet, The</i> , 2013 , 381, 1371-1379	40	2112
153	Pitfalls of predicting complex traits from SNPs. <i>Nature Reviews Genetics</i> , 2013 , 14, 507-15	30.1	457
152	Polygenic transmission and complex neuro developmental network for attention deficit hyperactivity disorder: genome-wide association study of both common and rare variants. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2013 , 162B, 419-430	3.5	125
151	Research review: the role of cytokines in depression in adolescents: a systematic review. <i>Journal of Child Psychology and Psychiatry and Allied Disciplines</i> , 2013 , 54, 816-35	7.9	65
150	ANKK1, TTC12, and NCAM1 polymorphisms and heroin dependence: importance of considering drug exposure. <i>JAMA Psychiatry</i> , 2013 , 70, 325-33	14.5	45
149	Partitioning the heritability of Tourette syndrome and obsessive compulsive disorder reveals differences in genetic architecture. <i>PLoS Genetics</i> , 2013 , 9, e1003864	6	189

148	Monozygotic twins affected with major depressive disorder have greater variance in methylation than their unaffected co-twin. <i>Translational Psychiatry</i> , 2013 , 3, e269	8.6	78
147	Assumptions and properties of limiting pathway models for analysis of epistasis in complex traits. <i>PLoS ONE</i> , 2013 , 8, e68913	3.7	11
146	Assessment of Response to Lithium Maintenance Treatment in Bipolar Disorder: A Consortium on Lithium Genetics (ConLiGen) Report. <i>PLoS ONE</i> , 2013 , 8, e65636	3.7	113
145	Novel genetic analysis for case-control genome-wide association studies: quantification of power and genomic prediction accuracy. <i>PLoS ONE</i> , 2013 , 8, e71494	3.7	30
144	A genome-wide meta-analysis of association studies of Cloninger's Temperament Scales. <i>Translational Psychiatry</i> , 2012 , 2, e116	8.6	85
143	Estimating the proportion of variation in susceptibility to schizophrenia captured by common SNPs. <i>Nature Genetics</i> , 2012 , 44, 247-50	36.3	471
142	Evidence-based psychiatric genetics, AKA the false dichotomy between common and rare variant hypotheses. <i>Molecular Psychiatry</i> , 2012 , 17, 474-85	15.1	108
141	Don't give up on GWAS. <i>Molecular Psychiatry</i> , 2012 , 17, 2-3	15.1	46
140	Using summary data from the danish national registers to estimate heritabilities for schizophrenia, bipolar disorder, and major depressive disorder. <i>Frontiers in Genetics</i> , 2012 , 3, 118	4.5	140
139	Meta-analyses of genome-wide linkage scans of anxiety-related phenotypes. <i>European Journal of Human Genetics</i> , 2012 , 20, 1078-84	5.3	22
138	A better coefficient of determination for genetic profile analysis. <i>Genetic Epidemiology</i> , 2012 , 36, 214-242.6		158
137	Appraisals of Stressful Life Events as a Genetically-Linked Mechanism in the Stress-Depression Relationship. <i>Cognitive Therapy and Research</i> , 2012 , 36, 338-347	2.7	18
136	Genome-wide association study of major depressive disorder: new results, meta-analysis, and lessons learned. <i>Molecular Psychiatry</i> , 2012 , 17, 36-48	15.1	335
135	Common SNPs explain some of the variation in the personality dimensions of neuroticism and extraversion. <i>Translational Psychiatry</i> , 2012 , 2, e102	8.6	137
134	Impact of diagnostic misclassification on estimation of genetic correlations using genome-wide genotypes. <i>European Journal of Human Genetics</i> , 2012 , 20, 668-74	5.3	51
133	Multi-locus genome-wide association analysis supports the role of glutamatergic synaptic transmission in the etiology of major depressive disorder. <i>Translational Psychiatry</i> , 2012 , 2, e184	8.6	62
132	Genome-wide association analysis of coffee drinking suggests association with CYP1A1/CYP1A2 and NRCAM. <i>Molecular Psychiatry</i> , 2012 , 17, 1116-29	15.1	93
131	Estimation of pleiotropy between complex diseases using single-nucleotide polymorphism-derived genomic relationships and restricted maximum likelihood. <i>Bioinformatics</i> , 2012 , 28, 2540-2	7.2	414

130	Genetic co-morbidity between neuroticism, anxiety/depression and somatic distress in a population sample of adolescent and young adult twins. <i>Psychological Medicine</i> , 2012 , 42, 1249-60	6.9	64
129	Unraveling the genetic etiology of adult antisocial behavior: a genome-wide association study. <i>PLoS ONE</i> , 2012 , 7, e45086	3.7	63
128	Identification of tag haplotypes for 5HTTLPR for different genome-wide SNP platforms. <i>Molecular Psychiatry</i> , 2011 , 16, 1073-5	15.1	17
127	Underestimated effect sizes in GWAS: fundamental limitations of single SNP analysis for dichotomous phenotypes. <i>PLoS ONE</i> , 2011 , 6, e27964	3.7	37
126	Association between ORMDL3, IL1RL1 and a deletion on chromosome 17q21 with asthma risk in Australia. <i>European Journal of Human Genetics</i> , 2011 , 19, 458-64	5.3	92
125	Genetic risk profiles for depression and anxiety in adult and elderly cohorts. <i>Molecular Psychiatry</i> , 2011 , 16, 773-83	15.1	116
124	Glutamate cysteine ligase (GCL) and self reported depression: an association study from the HUNT. <i>Journal of Affective Disorders</i> , 2011 , 131, 207-13	6.6	12
123	Estimating missing heritability for disease from genome-wide association studies. <i>American Journal of Human Genetics</i> , 2011 , 88, 294-305	11	737
122	A 3p26-3p25 genetic linkage finding for DSM-IV major depression in heavy smoking families. <i>American Journal of Psychiatry</i> , 2011 , 168, 848-52	11.9	33
121	The genetic association between personality and major depression or bipolar disorder. A polygenic score analysis using genome-wide association data. <i>Translational Psychiatry</i> , 2011 , 1, e50	8.6	83
120	Synthetic associations created by rare variants do not explain most GWAS results. <i>PLoS Biology</i> , 2011 , 9, e1000579	9.7	129
119	Sporadic cases are the norm for complex disease. <i>European Journal of Human Genetics</i> , 2010 , 18, 1039-43	3.3	78
118	The genetic interpretation of area under the ROC curve in genomic profiling. <i>PLoS Genetics</i> , 2010 , 6, e1000864	6	239
117	Choosing the best tools for comparative analyses of texts. <i>International Journal of Corpus Linguistics</i> , 2010 , 15, 429-473	0.8	5
116	Genetic differences between five European populations. <i>Human Heredity</i> , 2010 , 70, 141-9	1.1	24
115	A genome-wide association study of Cloninger's temperament scales: implications for the evolutionary genetics of personality. <i>Biological Psychology</i> , 2010 , 85, 306-17	3.2	128
114	Narrowing the boundaries of the genetic architecture of schizophrenia. <i>Schizophrenia Bulletin</i> , 2010 , 36, 14-23	1.3	86
113	Multi-locus models of genetic risk of disease. <i>Genome Medicine</i> , 2010 , 2, 10	14.4	54

112	Do 5HTTLPR and stress interact in risk for depression and suicidality? Item response analyses of a large sample. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2010 , 153B, 757-65	3.5	21
111	Comparing apples and oranges: equating the power of case-control and quantitative trait association studies. <i>Genetic Epidemiology</i> , 2010 , 34, 254-7	2.6	51
110	A versatile gene-based test for genome-wide association studies. <i>American Journal of Human Genetics</i> , 2010 , 87, 139-45	11	648
109	Phenotypic and discordant-monozygotic analyses of stress and perceived social support as antecedents to or sequelae of risk for depression. <i>Twin Research and Human Genetics</i> , 2009 , 12, 469-88	2.2	6
108	Twins in the World: The Legends They Inspire and the Lives That They Lead Alessandra Piontelli (2008). New York: Palgrave Macmillan, 272 pp, US\$26.95, ISBN-13: 978-0-230-60597-8.. <i>Twin Research and Human Genetics</i> , 2009 , 12, 407-407	2.2	
107	Harnessing the information contained within genome-wide association studies to improve individual prediction of complex disease risk. <i>Human Molecular Genetics</i> , 2009 , 18, 3525-31	5.6	237
106	Suggestive linkage on chromosome 2, 8, and 17 for lifetime major depression. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2009 , 150B, 352-8	3.5	17
105	Genetic and environmental influences on the co-morbidity between depression, panic disorder, agoraphobia, and social phobia: a twin study. <i>Depression and Anxiety</i> , 2009 , 26, 1004-11	8.4	74
104	Genome-wide association for major depressive disorder: a possible role for the presynaptic protein piccolo. <i>Molecular Psychiatry</i> , 2009 , 14, 359-75	15.1	322
103	Common polygenic variation contributes to risk of schizophrenia and bipolar disorder. <i>Nature</i> , 2009 , 460, 748-52	50.4	3568
102	Accurate, Large-Scale Genotyping of 5HTTLPR and Flanking Single Nucleotide Polymorphisms in an Association Study of Depression, Anxiety, and Personality Measures. <i>Biological Psychiatry</i> , 2009 , 66, 468-78	7.9	93
101	Estimating Effects and Making Predictions from Genome-Wide Marker Data. <i>Statistical Science</i> , 2009 , 24,	2.4	105
100	Heritability in the genomics era--concepts and misconceptions. <i>Nature Reviews Genetics</i> , 2008 , 9, 255-66	30.1	1155
99	A whole genome association study of neuroticism using DNA pooling. <i>Molecular Psychiatry</i> , 2008 , 13, 302-12	15.1	127
98	Prediction of individual genetic risk of complex disease. <i>Current Opinion in Genetics and Development</i> , 2008 , 18, 257-63	4.9	129
97	Emotionally Healthy Twins: A New Philosophy for Parenting Two Unique Individuals Joan A. Friedman (2008). Da Capo Press, Life Long Books, 224 pp., US\$15.95, ISBN 13 978 0 7382 1087 2. <i>Twin Research and Human Genetics</i> , 2008 , 11, 241-242	2.2	
96	COPD education and cognitive behavioral therapy group treatment for clinically significant symptoms of depression and anxiety in COPD patients: a randomized controlled trial. <i>Psychological Medicine</i> , 2008 , 38, 385-96	6.9	145
95	Shared temperament risk factors for anorexia nervosa: a twin study. <i>Psychosomatic Medicine</i> , 2008 , 70, 239-44	3.7	70

94	Association study of candidate variants from brain-derived neurotrophic factor and dystrobrevin-binding protein 1 with neuroticism, anxiety, and depression. <i>Psychiatric Genetics</i> , 2008 , 18, 219-25	2.9	18
93	Genome-wide linkage analysis of multiple measures of neuroticism of 2 large cohorts from Australia and the Netherlands. <i>Archives of General Psychiatry</i> , 2008 , 65, 649-58		31
92	Association study of candidate variants of COMT with neuroticism, anxiety and depression. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2008 , 147B, 1314-8	3.5	44
91	Use of monozygotic twins to investigate the relationship between 5HTTLPR genotype, depression and stressful life events: an application of Item Response Theory. <i>Novartis Foundation Symposium</i> , 2008 , 293, 48-59; discussion 59-70		18
90	Prediction of individual genetic risk to disease from genome-wide association studies. <i>Genome Research</i> , 2007 , 17, 1520-8	9.7	436
89	Haplotype analysis and a novel allele-sharing method refines a chromosome 4p locus linked to bipolar affective disorder. <i>Biological Psychiatry</i> , 2007 , 61, 797-805	7.9	22
88	Association analysis of the chromosome 4p15-p16 candidate region for bipolar disorder and schizophrenia. <i>Molecular Psychiatry</i> , 2007 , 12, 1011-25	15.1	31
87	Expression of eEF1A2 is associated with clear cell histology in ovarian carcinomas: overexpression of the gene is not dependent on modifications at the EEF1A2 locus. <i>British Journal of Cancer</i> , 2007 , 96, 1613-20	8.7	32
86	Anxiety and comorbid measures associated with PLXNA2. <i>Archives of General Psychiatry</i> , 2007 , 64, 318-26		45
85	Genetic and phenotypic stability of measures of neuroticism over 22 years. <i>Twin Research and Human Genetics</i> , 2007 , 10, 695-702	2.2	61
84	Genes, Environment and Psychopathology: Understanding the Causes of Psychiatric and Substance Use Disorders Kenneth S. Kendler and Carol A. Prescott (2006). New York: The Guildford Press, 412 pp, US\$45.00, ISBN-10: 1-59385-316-5.. <i>Twin Research and Human Genetics</i> , 2007 , 10, 231-233	2.2	
83	Sex Differences in Symptoms of Depression in Unrelated Individuals and Opposite-Sex Twin and Sibling Pairs. <i>Twin Research and Human Genetics</i> , 2006 , 9, 632-636	2.2	20
82	Empirical Evaluation of the Genetic Similarity of Samples From Twin Registries in Australia and the Netherlands Using 359 STRP Markers. <i>Twin Research and Human Genetics</i> , 2006 , 9, 600-602	2.2	10
81	Association analysis of the chromosome 4p-located G protein-coupled receptor 78 (GPR78) gene in bipolar affective disorder and schizophrenia. <i>Molecular Psychiatry</i> , 2006 , 11, 384-94	15.1	19
80	Sex differences in symptoms of depression in unrelated individuals and opposite-sex twin and sibling pairs. <i>Twin Research and Human Genetics</i> , 2006 , 9, 632-6	2.2	8
79	Empirical evaluation of the genetic similarity of samples from twin registries in Australia and the Netherlands using 359 STRP markers. <i>Twin Research and Human Genetics</i> , 2006 , 9, 600-2	2.2	6
78	Allele Frequencies and the r2 Measure of Linkage Disequilibrium: Impact on Design and Interpretation of Association Studies. <i>Twin Research and Human Genetics</i> , 2005 , 8, 87-94	2.2	276
77	Indivisible by Two: Lives of Extraordinary Twins Nancy L. Segal (2005). Harvard University Press. 280pp, \$US24.95, ISBN 0-674-01933-4.. <i>Twin Research and Human Genetics</i> , 2005 , 8, 666-668	2.2	1

76	Sex-specific association between bipolar affective disorder in women and GPR50, an X-linked orphan G protein-coupled receptor. <i>Molecular Psychiatry</i> , 2005 , 10, 470-8	15.1	84
75	Association between the TRAX/DISC locus and both bipolar disorder and schizophrenia in the Scottish population. <i>Molecular Psychiatry</i> , 2005 , 10, 657-68, 616	15.1	149
74	Association between the TRAX/DISC locus and both bipolar disorder and schizophrenia in the Scottish population. <i>Molecular Psychiatry</i> , 2005 , 10, 616-616	15.1	14
73	Translation elongation factor eEF1A2 is a potential oncoprotein that is overexpressed in two-thirds of breast tumours. <i>BMC Cancer</i> , 2005 , 5, 113	4.8	110
72	Allele frequencies and the r^2 measure of linkage disequilibrium: impact on design and interpretation of association studies. <i>Twin Research and Human Genetics</i> , 2005 , 8, 87-94	2.2	46
71	Indivisible by Two: Lives of Extraordinary Twins Nancy L. Segal (2005). Harvard University Press. 280pp, \$US24.95, ISBN 0-674-01933-4.. <i>Twin Research and Human Genetics</i> , 2005 , 8, 666-668	2.2	
70	Conventional multipoint nonparametric linkage analysis is not necessarily inherently biased. <i>American Journal of Human Genetics</i> , 2004 , 75, 718-20; author reply 723-7	11	4
69	Response to Amar J. Klar: The chromosome 1;11 translocation provides the best evidence supporting genetic etiology for schizophrenia and bipolar affective disorders. <i>Genetics</i> , 2003 , 163, 833-5; author reply 837-8	4	9
68	Genetics of schizophrenia and bipolar affective disorder: strategies to identify candidate genes. <i>Cold Spring Harbor Symposia on Quantitative Biology</i> , 2003 , 68, 383-94	3.9	5
67	A comparison of some simple methods to identify geographical areas with excess incidence of a rare disease such as childhood leukaemia. <i>Statistics in Medicine</i> , 1999 , 18, 1501-16	2.3	21
66	Population density and childhood leukaemia: results of the EUROCLUS Study. <i>European Journal of Cancer</i> , 1999 , 35, 439-44	7.5	38
65	Spatial clustering of childhood leukaemia: summary results from the EUROCLUS project. <i>British Journal of Cancer</i> , 1998 , 77, 818-24	8.7	53
64	Assigning pedigree beef performance records to contemporary groups taking account of within-herd calving patterns. <i>Animal Science</i> , 1997 , 65, 193-198		34
63	Aggregation of childhood leukemia in geographic areas of Greece. <i>Cancer Causes and Control</i> , 1997 , 8, 239-45	2.8	18
62	Use of MOET in Merino breeding programmes: a practical and economic appraisal. <i>Animal Science</i> , 1996 , 62, 241-254		4
61	MOET breeding schemes for wool sheep 1. Design alternatives. <i>Animal Science</i> , 1994 , 59, 71-86		11
60	MOET breeding schemes for wool sheep 2. Selection for adult fleece traits. <i>Animal Science</i> , 1994 , 59, 87-98		
59	Prediction of rates of inbreeding in populations undergoing index selection. <i>Theoretical and Applied Genetics</i> , 1994 , 87, 878-92	6	29

58	Calculation of prediction error variances using sparse matrix methods. <i>Journal of Animal Breeding and Genetics</i> , 1994 , 111, 102-9	2.9	9
57	Prediction of long-term contributions and inbreeding in populations undergoing mass selection. <i>Genetical Research</i> , 1993 , 62, 231-242	1.1	39
56	Prediction of asymptotic rates of response from selection on multiple traits using univariate and multivariate best linear unbiased predictors. <i>Animal Science</i> , 1993 , 57, 1-13		14
55	Accounting for Mutation Effects in the Additive Genetic Variance-Covariance Matrix and Its Inverse. <i>Biometrics</i> , 1990 , 46, 177	1.8	25
54	Prediction of rates of inbreeding in selected populations. <i>Genetical Research</i> , 1990 , 55, 41-54	1.1	133
53	Methods for predicting rates of inbreeding in selected populations. <i>Theoretical and Applied Genetics</i> , 1990 , 80, 503-12	6	32
52	Electronics in animal breeding. <i>Proceedings of the British Society of Animal Production</i> (1972), 1990 , 1990, 107-107		
51	Asymptotic rates of response from index selection. <i>Animal Science</i> , 1989 , 49, 217-227		71
50	Breeding Value Estimation for Pigs in Closed Nucleus Herds. <i>Proceedings of the British Society of Animal Production</i> (1972), 1988 , 1988, 12-12		1
49	Analysis of gestation length in American Simmental cattle. <i>Journal of Animal Science</i> , 1987 , 65, 970-4	0.7	11
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- 1 Genome-wide study of DNA methylation in Amyotrophic Lateral Sclerosis identifies differentially methylated loci and implicates metabolic, inflammatory and cholesterol pathways 1