## Cathryn M Lewis

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Genome-wide association study of 14,000 cases of seven common diseases and 3,000 shared controls. Nature, 2007, 447, 661-678.	13.7	8,895
2	A strong candidate for the breast and ovarian cancer susceptibility gene BRCA1. Science, 1994, 266, 66-71.	6.0	5,747
3	Genome-wide association analyses identify 44 risk variants and refine the genetic architecture of major depression. Nature Genetics, 2018, 50, 668-681.	9.4	2,224
4	Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. Nature Genetics, 2013, 45, 984-994.	9.4	2,067
5	Localization of a breast cancer susceptibility gene, BRCA2, to chromosome 13q12-13. Science, 1994, 265, 2088-2090.	6.0	1,725
6	Genome-wide meta-analysis of depression identifies 102 independent variants and highlights the importance of the prefrontal brain regions. Nature Neuroscience, 2019, 22, 343-352.	7.1	1,589
7	Genome-wide association analysis identifies 13 new risk loci for schizophrenia. Nature Genetics, 2013, 45, 1150-1159.	9.4	1,395
8	Association scan of 14,500 nonsynonymous SNPs in four diseases identifies autoimmunity variants. Nature Genetics, 2007, 39, 1329-1337.	9.4	1,298
9	Genome-wide association study identifies 30 loci associated with bipolar disorder. Nature Genetics, 2019, 51, 793-803.	9.4	1,191
10	Multiancestry genome-wide association study of 520,000 subjects identifies 32 loci associated with stroke and stroke subtypes. Nature Genetics, 2018, 50, 524-537.	9.4	1,124
11	PRSice: Polygenic Risk Score software. Bioinformatics, 2015, 31, 1466-1468.	1.8	1,109
12	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	6.0	1,085
13	Genome Scan Meta-Analysis of Schizophrenia and Bipolar Disorder, Part II: Schizophrenia. American Journal of Human Genetics, 2003, 73, 34-48.	2.6	1,072
14	Association between insertion mutation in NOD2 gene and Crohn's disease in German and British populations. Lancet, The, 2001, 357, 1925-1928.	6.3	1,071
15	Sequence variants in the autophagy gene IRGM and multiple other replicating loci contribute to Crohn's disease susceptibility. Nature Genetics, 2007, 39, 830-832.	9.4	1,063
16	A mega-analysis of genome-wide association studies for major depressive disorder. Molecular Psychiatry, 2013, 18, 497-511.	4.1	1,002
17	Genomic Relationships, Novel Loci, and Pleiotropic Mechanisms across Eight Psychiatric Disorders. Cell, 2019, 179, 1469-1482.e11.	13.5	935
18	Suberoylanilide hydroxamic acid, a histone deacetylase inhibitor, ameliorates motor deficits in a mouse model of Huntington's disease. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 2041-2046.	3.3	805

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19	Psychiatric genome-wide association study analyses implicate neuronal, immune and histone pathways. Nature Neuroscience, 2015, 18, 199-209.	7.1	701
20	The contribution of NOD2 gene mutations to the risk and site of disease in inflammatory bowel disease. Gastroenterology, 2002, 122, 867-874.	0.6	670
21	Polygenic risk scores: from research tools to clinical instruments. Genome Medicine, 2020, 12, 44.	3.6	646
22	Genome-wide association study of more than 40,000 bipolar disorder cases provides new insights into the underlying biology. Nature Genetics, 2021, 53, 817-829.	9.4	629
23	Meta-analysis of the Association Between the Level of Cannabis Use and Risk of Psychosis. Schizophrenia Bulletin, 2016, 42, 1262-1269.	2.3	615
24	The contribution of cannabis use to variation in the incidence of psychotic disorder across Europe (EU-GEI): a multicentre case-control study. Lancet Psychiatry,the, 2019, 6, 427-436.	3.7	528
25	Assignment of a locus for familial melanoma, MLM, to chromosome 9p13-p22. Science, 1992, 258, 1148-1152.	6.0	506
26	Genome-wide association analyses identify new risk variants and the genetic architecture of amyotrophic lateral sclerosis. Nature Genetics, 2016, 48, 1043-1048.	9.4	494
27	Genome-wide association study of depression phenotypes in UK Biobank identifies variants in excitatory synaptic pathways. Nature Communications, 2018, 9, 1470.	5.8	415
28	Genome-wide association study of major depressive disorder: new results, meta-analysis, and lessons learned. Molecular Psychiatry, 2012, 17, 36-48.	4.1	405
29	Genetic determinants of ulcerative colitis include the ECM1 locus and five loci implicated in Crohn's disease. Nature Genetics, 2008, 40, 710-712.	9.4	403
30	Genome Scan Meta-Analysis of Schizophrenia and Bipolar Disorder, Part III: Bipolar Disorder. American Journal of Human Genetics, 2003, 73, 49-62.	2.6	400
31	Genetic association studies: Design, analysis and interpretation. Briefings in Bioinformatics, 2002, 3, 146-153.	3.2	392
32	EULAR recommendations for terminology and research in individuals at risk of rheumatoid arthritis: report from the Study Group for Risk Factors for Rheumatoid Arthritis. Annals of the Rheumatic Diseases, 2012, 71, 638-641.	0.5	354
33	Meta-Analysis of the Association of Urbanicity With Schizophrenia. Schizophrenia Bulletin, 2012, 38, 1118-1123.	2.3	349
34	Association analysis in over 329,000 individuals identifies 116 independent variants influencing neuroticism. Nature Genetics, 2018, 50, 6-11.	9.4	327
35	Genome-Wide Pharmacogenetics of Antidepressant Response in the GENDEP Project. American Journal of Psychiatry, 2010, 167, 555-564.	4.0	314
36	Adverse drug reactions to azathioprine therapy are associated with polymorphism in the gene encoding inosine triphosphate pyrophosphatase (ITPase). Pharmacogenetics and Genomics, 2004, 14, 181-187.	5.7	305

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37	Environmental enrichment slows disease progression in R6/2 Huntington's disease mice. Annals of Neurology, 2002, 51, 235-242.	2.8	303
38	Fecundity of Patients With Schizophrenia, Autism, Bipolar Disorder, Depression, Anorexia Nervosa, or Substance Abuse vs Their Unaffected Siblings. JAMA Psychiatry, 2013, 70, 22.	6.0	284
39	A genome-wide association study of anorexia nervosa. Molecular Psychiatry, 2014, 19, 1085-1094.	4.1	282
40	A Nonsynonymous SNP in ATG16L1 Predisposes to Ileal Crohn's Disease and Is Independent of CARD15 and IBD5. Gastroenterology, 2007, 132, 1665-1671.	0.6	268
41	An additional k-means clustering step improves the biological features of WGCNA gene co-expression networks. BMC Systems Biology, 2017, 11, 47.	3.0	253
42	Meta-analysis of 32 genome-wide linkage studies of schizophrenia. Molecular Psychiatry, 2009, 14, 774-785.	4.1	235
43	Meta-analysis of genome scans of age-related macular degeneration. Human Molecular Genetics, 2005, 14, 2257-2264.	1.4	224
44	Genome-Wide Association Study of Major Recurrent Depression in the U.K. Population. American Journal of Psychiatry, 2010, 167, 949-957.	4.0	221
45	Inflammatory bowel disease susceptibility loci defined by genome scan meta-analysis of 1952 affected relative pairs. Human Molecular Genetics, 2004, 13, 763-770.	1.4	219
46	Common Genetic Variation and Antidepressant Efficacy in Major Depressive Disorder: A Meta-Analysis of Three Genome-Wide Pharmacogenetic Studies. American Journal of Psychiatry, 2013, 170, 207-217.	4.0	216
47	Minimal phenotyping yields genome-wide association signals of low specificity for major depression. Nature Genetics, 2020, 52, 437-447.	9.4	207
48	Meta-analysis of genome searches. Annals of Human Genetics, 1999, 63, 263-272.	0.3	205
49	Chromosome 9p21 in sporadic amyotrophic lateral sclerosis in the UK and seven other countries: a genome-wide association study. Lancet Neurology, The, 2010, 9, 986-994.	4.9	205
50	Contribution of Common Genetic Variants to Antidepressant Response. Biological Psychiatry, 2013, 73, 679-682.	0.7	199
51	Investigation of Crohn's Disease Risk Loci in Ulcerative Colitis Further Defines Their Molecular Relationship. Gastroenterology, 2009, 136, 523-529.e3.	0.6	198
52	The Neuronal Transporter Gene SLC6A15 Confers Risk to Major Depression. Neuron, 2011, 70, 252-265.	3.8	189
53	Genetic predictors of response to antidepressants in the GENDEP project. Pharmacogenomics Journal, 2009, 9, 225-233.	0.9	188
54	GWAS of Suicide Attempt in Psychiatric Disorders and Association With Major Depression Polygenic Risk Scores. American Journal of Psychiatry, 2019, 176, 651-660.	4.0	186

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55	Polygenic interactions with environmental adversity in the aetiology of major depressive disorder. Psychological Medicine, 2016, 46, 759-770.	2.7	176
56	An Examination of Polygenic Score Risk Prediction in Individuals With First-Episode Psychosis. Biological Psychiatry, 2017, 81, 470-477.	0.7	176
57	Genome-wide Association for Major Depression Through Age at Onset Stratification: Major Depressive Disorder Working Group of the Psychiatric Genomics Consortium. Biological Psychiatry, 2017, 81, 325-335.	0.7	175
58	Genetic Association of Major Depression With Atypical Features and Obesity-Related Immunometabolic Dysregulations. JAMA Psychiatry, 2017, 74, 1214.	6.0	174
59	Cost-effectiveness of pharmacogenetic-guided treatment: are we there yet?. Pharmacogenomics Journal, 2017, 17, 395-402.	0.9	173
60	IL23R Variation Determines Susceptibility But Not Disease Phenotype in Inflammatory Bowel Disease. Gastroenterology, 2007, 132, 1657-1664.	0.6	170
61	Metaâ€analysis of genomeâ€wide linkage scans of attention deficit hyperactivity disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 1392-1398.	1.1	160
62	Using genetics to understand the causal influence of higher BMI on depression. International Journal of Epidemiology, 2019, 48, 834-848.	0.9	156
63	Prospects for using risk scores in polygenic medicine. Genome Medicine, 2017, 9, 96.	3.6	153
64	Prospective evaluation of the pharmacogenetics of azathioprine in the treatment of inflammatory bowel disease. Alimentary Pharmacology and Therapeutics, 2008, 28, 973-983.	1.9	152
65	Combining clinical variables to optimize prediction of antidepressant treatment outcomes. Journal of Psychiatric Research, 2016, 78, 94-102.	1.5	149
66	Genome Scan Meta-Analysis of Schizophrenia and Bipolar Disorder, Part I: Methods and Power Analysis. American Journal of Human Genetics, 2003, 73, 17-33.	2.6	147
67	Meta-Analysis of Genome-Wide Scans Provides Evidence for Sex- and Site-Specific Regulation of Bone Mass. Journal of Bone and Mineral Research, 2006, 22, 173-183.	3.1	144
68	MicroRNAs 146a/b-5 and 425-3p and 24-3p are markers of antidepressant response and regulate MAPK/Wnt-system genes. Nature Communications, 2017, 8, 15497.	5.8	144
69	Introduction to Genetic Association Studies. Cold Spring Harbor Protocols, 2012, 2012, pdb.top068163.	0.2	140
70	Pharmacogenetic variants in the DPYD, TYMS, CDA and MTHFR genes are clinically significant predictors of fluoropyrimidine toxicity. British Journal of Cancer, 2013, 108, 2505-2515.	2.9	139
71	Metaâ€Analysis of Genomeâ€wide Linkage Studies in BMI and Obesity. Obesity, 2007, 15, 2263-2275	1.5	138
72	Therapygenetics: the 5HTTLPR and response to psychological therapy. Molecular Psychiatry, 2012, 17, 236-237.	4.1	135

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73	Penetrance for copy number variants associated with schizophrenia. Human Molecular Genetics, 2010, 19, 3477-3481.	1.4	132
74	Standardization and statistical approaches to therapeutic trials in the R6/2 mouse. Brain Research Bulletin, 2003, 61, 469-479.	1.4	129
75	Expression quantitative trait loci in the developing human brain and their enrichment in neuropsychiatric disorders. Genome Biology, 2018, 19, 194.	3.8	126
76	The ?174G allele of the interleukin-6 gene confers susceptibility to systemic arthritis in children: A multicenter study using simplex and multiplex juvenile idiopathic arthritis families. Arthritis and Rheumatism, 2003, 48, 3202-3206.	6.7	123
77	A genome-wide association meta-analysis identifies a novel locus at 17q11.2 associated with sporadic amyotrophic lateral sclerosis. Human Molecular Genetics, 2014, 23, 2220-2231.	1.4	123
78	Genome-wide gene-environment analyses of major depressive disorder and reported lifetime traumatic experiences in UK Biobank. Molecular Psychiatry, 2020, 25, 1430-1446.	4.1	116
79	Response to radiation therapy and prognosis in breast cancer patients with BRCA1 and BRCA2 mutations. Radiotherapy and Oncology, 1998, 47, 129-136.	0.3	114
80	Comorbid medical illness in bipolar disorder. British Journal of Psychiatry, 2014, 205, 465-472.	1.7	113
81	Uncovering the Genetic Architecture of Major Depression. Neuron, 2019, 102, 91-103.	3.8	113
82	Genetic Evidence for Interaction of the 5q31 Cytokine Locus and the CARD15 Gene in Crohn Disease. American Journal of Human Genetics, 2003, 72, 1018-1022.	2.6	111
83	Genetic Predictors of Response to Serotonergic and Noradrenergic Antidepressants in Major Depressive Disorder: A Genome-Wide Analysis of Individual-Level Data and a Meta-Analysis. PLoS Medicine, 2012, 9, e1001326.	3.9	110
84	Elevated C-Reactive Protein in Patients With Depression, Independent of Genetic, Health, and Psychosocial Factors: Results From the UK Biobank. American Journal of Psychiatry, 2021, 178, 522-529.	4.0	110
85	A multiobserver, population-based analysis of histologic dysplasia in melanocytic nevi. Journal of the American Academy of Dermatology, 1994, 30, 707-714.	0.6	108
86	Correlation of SMNt and SMNc gene copy number with age of onset and survival in spinal muscular atrophy. European Journal of Human Genetics, 1998, 6, 467-474.	1.4	108
87	Genetics of inflammatory bowel disease: progress and prospects. Human Molecular Genetics, 2004, 13, 161R-168.	1.4	106
88	Genetic Predictors of Increase in Suicidal Ideation During Antidepressant Treatment in the GENDEP Project. Neuropsychopharmacology, 2009, 34, 2517-2528.	2.8	105
89	Polymorphisms in Folate, Pyrimidine, and Purine Metabolism Are Associated with Efficacy and Toxicity of Methotrexate in Psoriasis. Journal of Investigative Dermatology, 2007, 127, 1860-1867.	0.3	104
90	Genetics of Depression: Progress at Last. Current Psychiatry Reports, 2017, 19, 43.	2.1	101

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91	Genetic architecture of 11 major psychiatric disorders at biobehavioral, functional genomic and molecular genetic levels of analysis. Nature Genetics, 2022, 54, 548-559.	9.4	101
92	Incidence of multiple primary cancers in a cohort of women diagnosed with breast cancer in southeast England. British Journal of Cancer, 2001, 84, 435-440.	2.9	99
93	Genetic relationships between suicide attempts, suicidal ideation and major psychiatric disorders: A genomeâ€wide association and polygenic scoring study. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2014, 165, 428-437.	1.1	99
94	Evaluation of polygenic prediction methodology within a reference-standardized framework. PLoS Genetics, 2021, 17, e1009021.	1.5	99
95	Genomewide Association Scan of Suicidal Thoughts and Behaviour in Major Depression. PLoS ONE, 2011, 6, e20690.	1.1	98
96	The risk of subsequent primary cancers after colorectal cancer in southeast England. Gut, 2002, 50, 647-652.	6.1	97
97	Genetic basis of lacunar stroke: a pooled analysis of individual patient data and genome-wide association studies. Lancet Neurology, The, 2021, 20, 351-361.	4.9	95
98	Independent and population-specific association of risk variants at the IRGM locus with Crohn's disease. Human Molecular Genetics, 2010, 19, 1828-1839.	1.4	93
99	Modelling the Effects of Penetrance and Family Size on Rates of Sporadic and Familial Disease. Human Heredity, 2011, 71, 281-288.	0.4	93
100	Genome-wide association study of MRI markers of cerebral small vessel disease in 42,310 participants. Nature Communications, 2020, 11, 2175.	5.8	93
101	Genome-wide association study of increasing suicidal ideation during antidepressant treatment in the GENDEP project. Pharmacogenomics Journal, 2012, 12, 68-77.	0.9	92
102	Genome-wide meta-analysis of cerebral white matter hyperintensities in patients with stroke. Neurology, 2016, 86, 146-153.	1.5	91
103	The protective effect of alcohol on developing rheumatoid arthritis: a systematic review and meta-analysis. Rheumatology, 2013, 52, 856-867.	0.9	89
104	A Genome-Wide Test of the Differential Susceptibility Hypothesis Reveals a Genetic Predictor of Differential Response to Psychological Treatments for Child Anxiety Disorders. Psychotherapy and Psychosomatics, 2016, 85, 146-158.	4.0	89
105	Molecular Characterization of CTNS Deletions in Nephropathic Cystinosis: Development of a PCR-Based Detection Assay. American Journal of Human Genetics, 1999, 65, 353-359.	2.6	88
106	The Genetic Architecture of Depression in Individuals of East Asian Ancestry. JAMA Psychiatry, 2021, 78, 1258.	6.0	88
107	Associations of allelic variants of the multidrug resistance gene (ABCB1 or MDR1) and Inflammatory Bowel Disease and their effects on disease behavior: A case-control and meta-analysis study. Inflammatory Bowel Diseases, 2006, 12, 263-271.	0.9	87
108	Does Childhood Trauma Moderate Polygenic Risk for Depression? A Meta-analysis of 5765 Subjects From the Psychiatric Genomics Consortium. Biological Psychiatry, 2018, 84, 138-147.	0.7	87

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109	An Analysis of Two Genome-wide Association Meta-analyses Identifies a New Locus for Broad Depression Phenotype. Biological Psychiatry, 2017, 82, 322-329.	0.7	84
110	Relationship Between p53 Codon 72 Polymorphism and Susceptibility to Sunburn and Skin Cancer. Journal of Investigative Dermatology, 2002, 119, 84-90.	0.3	83
111	Controlling misdiagnosis errors in preimplantation genetic diagnosis: a comprehensive model encompassing extrinsic and intrinsic sources of error. Human Reproduction, 2001, 16, 43-50.	0.4	81
112	Novel pharmacogenetic markers for treatment outcome in azathioprineâ€ŧreated inflammatory bowel disease. Alimentary Pharmacology and Therapeutics, 2009, 30, 375-384.	1.9	78
113	Joint genome-wide association study of progressive supranuclear palsy identifies novel susceptibility loci and genetic correlation to neurodegenerative diseases. Molecular Neurodegeneration, 2018, 13, 41.	4.4	77
114	Two Families with Familial Amyotrophic Lateral Sclerosis Are Linked to a Novel Locus on Chromosome 16q. American Journal of Human Genetics, 2003, 73, 390-396.	2.6	76
115	Meta-Analysis of 4 Coronary Heart Disease Genome-Wide Linkage Studies Confirms a Susceptibility Locus on Chromosome 3q. Arteriosclerosis, Thrombosis, and Vascular Biology, 2003, 23, 1863-1868.	1.1	76
116	A bidirectional relationship between depression and the autoimmune disorders – New perspectives from the National Child Development Study. PLoS ONE, 2017, 12, e0173015.	1.1	76
117	A Novel MMP12 Locus Is Associated with Large Artery Atherosclerotic Stroke Using a Genome-Wide Age-at-Onset Informed Approach. PLoS Genetics, 2014, 10, e1004469.	1.5	75
118	EPIBLASTER-fast exhaustive two-locus epistasis detection strategy using graphical processing units. European Journal of Human Genetics, 2011, 19, 465-471.	1.4	74
119	Evidence of causal effect of major depression on alcohol dependence: findings from the psychiatric genomics consortium. Psychological Medicine, 2019, 49, 1218-1226.	2.7	74
120	Novel IL10 gene family associations with systemic juvenile idiopathic arthritis. Arthritis Research and Therapy, 2006, 8, R148.	1.6	73
121	Genetic variation at 16q24.2 is associated with small vessel stroke. Annals of Neurology, 2017, 81, 383-394.	2.8	73
122	Meta-analysis of four rheumatoid arthritis genome-wide linkage studies: Confirmation of a susceptibility locus on chromosome 16. Arthritis and Rheumatism, 2003, 48, 1200-1206.	6.7	72
123	Depressive disorder moderates the effect of the FTO gene on body mass index. Molecular Psychiatry, 2012, 17, 604-611.	4.1	72
124	ACPA-positive and ACPA-negative rheumatoid arthritis differ in their requirements for combination DMARDs and corticosteroids: secondary analysis of a randomized controlled trial. Arthritis Research and Therapy, 2014, 16, R13.	1.6	72
125	The correlation between reading and mathematics ability at age twelve has a substantial genetic component. Nature Communications, 2014, 5, 4204.	5.8	72
126	Pharmacogenetics of antidepressant response: A polygenic approach. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2017, 75, 128-134.	2.5	71

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127	Association of DLG5 R30Q variant with inflammatory bowel disease. European Journal of Human Genetics, 2005, 13, 835-839.	1.4	70
128	Transdiagnostic dimensions of psychopathology at first episode psychosis: findings from the multinational EU-GEI study. Psychological Medicine, 2019, 49, 1378-1391.	2.7	69
129	Polygenic scores in biomedical research. Nature Reviews Genetics, 2022, 23, 524-532.	7.7	69
130	Predicting outcomes following cognitive behaviour therapy in child anxiety disorders: the influence of genetic, demographic and clinical information. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2013, 54, 1086-1094.	3.1	68
131	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.	0.3	68
132	Genetic and clinical characteristics of treatment-resistant depression using primary care records in two UK cohorts. Molecular Psychiatry, 2021, 26, 3363-3373.	4.1	66
133	A candidate gene analysis of three related photosensitivity disorders: cutaneous lupus erythematosus, polymorphic light eruption and actinic prurigo. British Journal of Dermatology, 2001, 145, 229-236.	1.4	65
134	Predicting treatment response in psoriasis using serum levels of adalimumab and etanercept: a single-centre, cohort study. British Journal of Dermatology, 2013, 169, 306-313.	1.4	65
135	Genetic effects influencing risk for major depressive disorder in China and Europe. Translational Psychiatry, 2017, 7, e1074-e1074.	2.4	64
136	Effect of cytochrome CYP2C19 metabolizing activity on antidepressant response and side effects: Meta-analysis of data from genome-wide association studies. European Neuropsychopharmacology, 2018, 28, 945-954.	0.3	64
137	Delineating the Genetic Component of Gene Expression in Major Depression. Biological Psychiatry, 2021, 89, 627-636.	0.7	63
138	Relationship between obesity and the risk of clinically significant depression: Mendelian randomisation study. British Journal of Psychiatry, 2014, 205, 24-28.	1.7	62
139	From SNPs to Genes: Disease Association at the Gene Level. PLoS ONE, 2011, 6, e20133.	1.1	61
140	Allelic differences between Europeans and Chinese for CREB1 SNPs and their implications in gene expression regulation, hippocampal structure and function, and bipolar disorder susceptibility. Molecular Psychiatry, 2014, 19, 452-461.	4.1	61
141	Sex-Dependent Shared and Nonshared Genetic Architecture Across Mood and Psychotic Disorders. Biological Psychiatry, 2022, 91, 102-117.	0.7	61
142	Genetic modelling of dizygotic twinning in pedigrees of spontaneous dizygotic twins. , 1996, 61, 258-263.		60
143	Genetic risk score analysis indicates migraine with and without comorbid depression are genetically different disorders. Human Genetics, 2014, 133, 173-186.	1.8	60
144	The association between lower educational attainment and depression owing to shared genetic effects? Results in ~25 000 subjects. Molecular Psychiatry, 2015, 20, 735-743.	4.1	59

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145	Pooled Sequencing of 531 Genes in Inflammatory Bowel Disease Identifies an Associated Rare Variant in BTNL2 and Implicates Other Immune Related Genes. PLoS Genetics, 2015, 11, e1004955.	1.5	59
146	Meta-analysis of genome-wide scans for hypertension and blood pressure in Caucasians shows evidence of susceptibility regions on chromosomes 2 and 3. Human Molecular Genetics, 2004, 13, 2325-2332.	1.4	58
147	Comprehensive association study of genetic variants in the IL-1 gene family in systemic juvenile idiopathic arthritis. Genes and Immunity, 2008, 9, 349-357.	2.2	58
148	Meta-analysis of linkage studies for Alzheimer's disease—A web resource. Neurobiology of Aging, 2009, 30, 1037-1047.	1.5	58
149	Mutation in theITPAGene Predicts Intolerance to Azathioprine. Nucleosides, Nucleotides and Nucleic Acids, 2004, 23, 1393-1397.	0.4	57
150	Association of a Locus in the <i>CAMTA1</i> Gene With Survival in Patients With Sporadic Amyotrophic Lateral Sclerosis. JAMA Neurology, 2016, 73, 812.	4.5	57
151	A polygenic risk score analysis of psychosis endophenotypes across brain functional, structural, and cognitive domains. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2018, 177, 21-34.	1.1	57
152	Meta-analysis of genome-wide linkage studies of systemic lupus erythematosus. Genes and Immunity, 2006, 7, 609-614.	2.2	56
153	A genetic risk score combining 32 SNPs is associated with body mass index and improves obesity prediction in people with major depressive disorder. BMC Medicine, 2015, 13, 86.	2.3	56
154	Shared genetic contribution to ischemic stroke and Alzheimer's disease. Annals of Neurology, 2016, 79, 739-747.	2.8	56
155	Genetic Associations Between Childhood Psychopathology and Adult Depression and Associated Traits in 42â€ <sup></sup> 998 Individuals. JAMA Psychiatry, 2020, 77, 715.	6.0	56
156	Localization of the 9p Melanoma Susceptibility Locus (MLM) to a 2-cM Region between D9S736 and D9S171. Genomics, 1994, 23, 265-268.	1.3	55
157	Genetic determinants of the thiopurine methyltransferase intermediate activity phenotype in British Asians and Caucasians. Pharmacogenetics and Genomics, 2003, 13, 97-105.	5.7	55
158	Methylâ€CpGâ€binding protein 2 polymorphisms and vulnerability to autism. Genes, Brain and Behavior, 2008, 7, 754-760.	1.1	55
159	COMT but not serotoninâ€related genes modulates the influence of childhood abuse on anger traits. Genes, Brain and Behavior, 2010, 9, 193-202.	1.1	55
160	The genetics of affective disorder and suicide. European Psychiatry, 2010, 25, 275-277.	0.1	55
161	Predicting the Risk of Rheumatoid Arthritis and Its Age of Onset through Modelling Genetic Risk Variants with Smoking. PLoS Genetics, 2013, 9, e1003808.	1.5	55
162	Genomeâ€wide geneâ€environment interaction in depression: A systematic evaluation of candidate genes. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2018, 177, 40-49.	1.1	55

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163	Genetic Modeling of Abnormal Photosensitivity in Families with Polymorphic Light Eruption and Actinic Prurigo. Journal of Investigative Dermatology, 2000, 115, 471-476.	0.3	54
164	Meta-analysis of 20 genome-wide linkage studies evidenced new regions linked to asthma and atopy. European Journal of Human Genetics, 2010, 18, 700-706.	1.4	54
165	Mutation Rates in Humans. I. Overall and Sex-Specific Rates Obtained from a Population Study of Hemophilia B. American Journal of Human Genetics, 1999, 65, 1572-1579.	2.6	53
166	A common Fanconi anemia mutation in black populations of sub-Saharan Africa. Blood, 2005, 105, 3542-3544.	0.6	53
167	Visual analysis of geocoded twin data puts nature and nurture on the map. Molecular Psychiatry, 2012, 17, 867-874.	4.1	52
168	A genome-wide association study of a sustained pattern of antidepressant response. Journal of Psychiatric Research, 2013, 47, 1157-1165.	1.5	52
169	Interobserver concordance in discriminating clinical atypia of melanocytic nevi, and correlations with histologic atypia. Journal of the American Academy of Dermatology, 1996, 34, 618-625.	0.6	51
170	A Genome-Wide Significant Linkage for Severe Depression on Chromosome 3: The Depression Network Study. American Journal of Psychiatry, 2011, 168, 840-847.	4.0	51
171	Antidepressant drug-specific prediction of depression treatment outcomes from genetic and clinical variables. Scientific Reports, 2018, 8, 5530.	1.6	51
172	Treatment response classes in major depressive disorder identified by model-based clustering and validated by clinical prediction models. Translational Psychiatry, 2019, 9, 187.	2.4	51
173	Sex stratification of an inflammatory bowel disease genome search shows male-specific linkage to the HLA region of chromosome 6. European Journal of Human Genetics, 2002, 10, 259-265.	1.4	49
174	Interaction between the <i>FTO</i> gene, body mass index and depression: meta-analysis of 13701 individuals. British Journal of Psychiatry, 2017, 211, 70-76.	1.7	49
175	Genetic contribution to DZ twinning. , 1996, 61, 237-246.		48
176	Familial clustering of polymorphic light eruption in relatives of patients with lupus erythematosus: evidence of a shared pathogenesis. British Journal of Dermatology, 2001, 144, 334-338.	1.4	47
177	Direct or indirect association in a complex disease: the role ofSLC22A4 andSLC22A5 functional variants in Crohn disease. Human Mutation, 2006, 27, 778-785.	1.1	47
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