

Derek Morris

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.

212
papers

30,458
citations

61
h-index

174
g-index

248
ext. papers

37,510
ext. citations

9.9
avg, IF

7.99
L-index

#	Paper	IF	Citations
212	Normalization of impaired emotion inhibition in bipolar disorder mediated by cholinergic neurotransmission in the cingulate cortex.. <i>Neuropsychopharmacology</i> , 2022 ,	8.7	1
211	Genetic variants associated with longitudinal changes in brain structure across the lifespan.. <i>Nature Neuroscience</i> , 2022 , 25, 421-432	25.5	1
210	Major Depressive Disorder: Existing Hypotheses about Pathophysiological Mechanisms and New Genetic Findings.. <i>Genes</i> , 2022 , 13,	4.2	5
209	Mapping genomic loci implicates genes and synaptic biology in schizophrenia.. <i>Nature</i> , 2022 ,	50.4	35
208	Rare coding variants in ten genes confer substantial risk for schizophrenia.. <i>Nature</i> , 2022 ,	50.4	16
207	Interleukin 6 predicts increased neural response during face processing in a sample of individuals with schizophrenia and healthy participants: A functional magnetic resonance imaging study. <i>NeuroImage: Clinical</i> , 2021 , 32, 102851	5.3	0
206	SATB2-LEMD2 interaction links nuclear shape plasticity to regulation of cognition-related genes. <i>EMBO Journal</i> , 2021 , 40, e103701	13	2
205	1q21.1 distal copy number variants are associated with cerebral and cognitive alterations in humans. <i>Translational Psychiatry</i> , 2021 , 11, 182	8.6	6
204	Changes in Default-Mode Network Associated With Childhood Trauma in Schizophrenia. <i>Schizophrenia Bulletin</i> , 2021 , 47, 1482-1494	1.3	3
203	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
202	Identifying nootropic drug targets via large-scale cognitive GWAS and transcriptomics. <i>Neuropsychopharmacology</i> , 2021 , 46, 1788-1801	8.7	1
201	Childhood trauma, parental bonding, and social cognition in patients with schizophrenia and healthy adults. <i>Journal of Clinical Psychology</i> , 2021 , 77, 241-253	2.8	10
200	Genes influenced by MEF2C contribute to neurodevelopmental disease via gene expression changes that affect multiple types of cortical excitatory neurons. <i>Human Molecular Genetics</i> , 2021 , 30, 961-970	5.6	7
199	Contamination of domestic groundwater systems by verotoxigenic escherichia coli (VTEC), 2003-2019: A global scoping review. <i>Water Research</i> , 2021 , 188, 116496	12.5	2
198	DNA methylation meta-analysis reveals cellular alterations in psychosis and markers of treatment-resistant schizophrenia. <i>ELife</i> , 2021 , 10,	8.9	15
197	Early life Adversity, functional connectivity and cognitive performance in Schizophrenia: The mediating role of IL-6. <i>Brain, Behavior, and Immunity</i> , 2021 , 98, 388-396	16.6	2
196	Rare Copy Number Variants Are Associated With Poorer Cognition in Schizophrenia. <i>Biological Psychiatry</i> , 2021 , 90, 28-34	7.9	5

195	Microglial-expressed genetic risk variants, cognitive function and brain volume in patients with schizophrenia and healthy controls. <i>Translational Psychiatry</i> , 2021 , 11, 490	8.6	1
194	A meta-analysis of deep brain structural shape and asymmetry abnormalities in 2,833 individuals with schizophrenia compared with 3,929 healthy volunteers via the ENIGMA Consortium. <i>Human Brain Mapping</i> , 2021 ,	5.9	7
193	Controlling for background genetic effects using polygenic scores improves the power of genome-wide association studies. <i>Scientific Reports</i> , 2021 , 11, 19571	4.9	0
192	Investigating rare pathogenic/likely pathogenic exonic variation in bipolar disorder. <i>Molecular Psychiatry</i> , 2021 , 26, 5239-5250	15.1	3
191	The Relationship Between Polygenic Risk Scores and Cognition in Schizophrenia. <i>Schizophrenia Bulletin</i> , 2020 , 46, 336-344	1.3	38
190	The genetic architecture of the human cerebral cortex. <i>Science</i> , 2020 , 367,	33.3	156
189	The Relationship Between White Matter Microstructure and General Cognitive Ability in Patients With Schizophrenia and Healthy Participants in the ENIGMA Consortium. <i>American Journal of Psychiatry</i> , 2020 , 177, 537-547	11.9	21
188	Neuroharmony: A new tool for harmonizing volumetric MRI data from unseen scanners. <i>NeuroImage</i> , 2020 , 220, 117127	7.9	14
187	Identifying schizophrenia patients who carry pathogenic genetic copy number variants using standard clinical assessment: retrospective cohort study. <i>British Journal of Psychiatry</i> , 2020 , 216, 275-279 ^{5.4}	5.4	7
186	Cognitive Genomics: Recent Advances and Current Challenges. <i>Current Psychiatry Reports</i> , 2020 , 22, 2	9.1	2
185	Diagnostic yield of a custom-designed multi-gene cancer panel in Irish patients with breast cancer. <i>Irish Journal of Medical Science</i> , 2020 , 189, 849-864	1.9	1
184	Dynamics of Brain Structure and its Genetic Architecture over the Lifespan 2020 ,		7
183	The Genetics of the Mood Disorder Spectrum: Genome-wide Association Analyses of More Than 185,000 Cases and 439,000 Controls. <i>Biological Psychiatry</i> , 2020 , 88, 169-184	7.9	57
182	Association of Copy Number Variation of the 15q11.2 BP1-BP2 Region With Cortical and Subcortical Morphology and Cognition. <i>JAMA Psychiatry</i> , 2020 , 77, 420-430	14.5	24
181	Altered gene regulation as a candidate mechanism by which ciliopathy gene SDCCAG8 contributes to schizophrenia and cognitive function. <i>Human Molecular Genetics</i> , 2020 , 29, 407-417	5.6	1
180	Effects of early life adversity on immune function and cognitive performance: results from the ALSPAC cohort. <i>Social Psychiatry and Psychiatric Epidemiology</i> , 2020 , 55, 723-733	4.5	6
179	Effects of complement gene-set polygenic risk score on brain volume and cortical measures in patients with psychotic disorders and healthy controls. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2020 , 183, 445-453	3.5	1
178	Childhood trauma, brain structure and emotion recognition in patients with schizophrenia and healthy participants. <i>Social Cognitive and Affective Neuroscience</i> , 2020 , 15, 1336-1350	4	5

177	Genes regulated by BCL11B during T-cell development are enriched for de novo mutations found in schizophrenia patients. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2020 , 183, 370-379	3.5	
176	Dose response of the 16p11.2 distal copy number variant on intracranial volume and basal ganglia. <i>Molecular Psychiatry</i> , 2020 , 25, 584-602	15.1	24
175	Genome-wide association study identifies 30 loci associated with bipolar disorder. <i>Nature Genetics</i> , 2019 , 51, 793-803	36.3	662
174	T91. NOVEL INFLUENCE OF EARLY-LIFE ADVERSITY ACROSS FUNCTIONAL NETWORKS DURING WORKING MEMORY IN SCHIZOPHRENIA. <i>Schizophrenia Bulletin</i> , 2019 , 45, S239-S239	1.3	0
173	Population-based identity-by-descent mapping combined with exome sequencing to detect rare risk variants for schizophrenia. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2019 , 180, 223-231	3.5	2
172	Genes encoding SATB2-interacting proteins in adult cerebral cortex contribute to human cognitive ability. <i>PLoS Genetics</i> , 2019 , 15, e1007890	6	5
171	Beyond C4: Analysis of the complement gene pathway shows enrichment for IQ in patients with psychotic disorders and healthy controls. <i>Genes, Brain and Behavior</i> , 2019 , 18, e12602	3.6	8
170	Pleiotropic Meta-Analysis of Cognition, Education, and Schizophrenia Differentiates Roles of Early Neurodevelopmental and Adult Synaptic Pathways. <i>American Journal of Human Genetics</i> , 2019 , 105, 334-350	11.1	37
169	The effect of breed and diet type on the global transcriptome of hepatic tissue in beef cattle divergent for feed efficiency. <i>BMC Genomics</i> , 2019 , 20, 525	4.5	13
168	Genomic Relationships, Novel Loci, and Pleiotropic Mechanisms across Eight Psychiatric Disorders. <i>Cell</i> , 2019 , 179, 1469-1482.e11	56.2	402
167	Targeted Sequencing of 10,198 Samples Confirms Abnormalities in Neuronal Activity and Implicates Voltage-Gated Sodium Channels in Schizophrenia Pathogenesis. <i>Biological Psychiatry</i> , 2019 , 85, 554-562	7.9	21
166	Effects of MiR-137 genetic risk score on brain volume and cortical measures in patients with schizophrenia and controls. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2018 , 177, 369-376	3.5	6
165	Genes regulated by SATB2 during neurodevelopment contribute to schizophrenia and educational attainment. <i>PLoS Genetics</i> , 2018 , 14, e1007515	6	17
164	Analysis of shared heritability in common disorders of the brain. <i>Science</i> , 2018 , 360,	33.3	666
163	Multi-Trait Analysis of GWAS and Biological Insights Into Cognition: A Response to Hill (2018). <i>Twin Research and Human Genetics</i> , 2018 , 21, 394-397	2.2	2
162	Activated charcoal as a capture material for silver nanoparticles in environmental water samples. <i>Science of the Total Environment</i> , 2018 , 645, 356-362	10.2	8
161	The Genetics of Endophenotypes of Neurofunction to Understand Schizophrenia (GENUS) consortium: A collaborative cognitive and neuroimaging genetics project. <i>Schizophrenia Research</i> , 2018 , 195, 306-317	3.6	14
160	GWAS and eQTL analysis identifies a SNP associated with both residual feed intake and GFRA2 expression in beef cattle. <i>Scientific Reports</i> , 2018 , 8, 14301	4.9	27

159	Study of 300,486 individuals identifies 148 independent genetic loci influencing general cognitive function. <i>Nature Communications</i> , 2018 , 9, 2098	17.4	254
158	Cortical Brain Abnormalities in 4474 Individuals With Schizophrenia and 5098 Control Subjects via the Enhancing Neuro Imaging Genetics Through Meta Analysis (ENIGMA) Consortium. <i>Biological Psychiatry</i> , 2018 , 84, 644-654	7.9	325
157	Shiga toxigenic Escherichia coli incidence is related to small area variation in cattle density in a region in Ireland. <i>Science of the Total Environment</i> , 2018 , 637-638, 865-870	10.2	6
156	DNA methylation differences at the glucocorticoid receptor gene in depression are related to functional alterations in hypothalamic-pituitary-adrenal axis activity and to early life emotional abuse. <i>Psychiatry Research</i> , 2018 , 265, 341-348	9.9	85
155	Genome-wide association meta-analysis in 269,867 individuals identifies new genetic and functional links to intelligence. <i>Nature Genetics</i> , 2018 , 50, 912-919	36.3	475
154	GWAS meta-analysis reveals novel loci and genetic correlates for general cognitive function: a report from the COGENT consortium. <i>Molecular Psychiatry</i> , 2017 , 22, 336-345	15.1	123
153	Novel genetic loci associated with hippocampal volume. <i>Nature Communications</i> , 2017 , 8, 13624	17.4	173
152	Occurrence and co-occurrence of hallucinations by modality in schizophrenia-spectrum disorders. <i>Psychiatry Research</i> , 2017 , 252, 154-160	9.9	62
151	Further evidence of alerted default network connectivity and association with theory of mind ability in schizophrenia. <i>Schizophrenia Research</i> , 2017 , 184, 52-58	3.6	14
150	Cognitive Characterization of Schizophrenia Risk Variants Involved in Synaptic Transmission: Evidence of CACNA1CS Role in Working Memory. <i>Neuropsychopharmacology</i> , 2017 , 42, 2612-2622	8.7	16
149	ltn2a silencing rescues lamin A mediated inhibition of 3T3-L1 adipocyte differentiation. <i>Adipocyte</i> , 2017 , 6, 259-276	3.2	2
148	Large-Scale Cognitive GWAS Meta-Analysis Reveals Tissue-Specific Neural Expression and Potential Nootropic Drug Targets. <i>Cell Reports</i> , 2017 , 21, 2597-2613	10.6	71
147	Human subcortical brain asymmetries in 15,847 people worldwide reveal effects of age and sex. <i>Brain Imaging and Behavior</i> , 2017 , 11, 1497-1514	4.1	87
146	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
145	Silver nanoparticles in the environment: Sources, detection and ecotoxicology. <i>Science of the Total Environment</i> , 2017 , 575, 231-246	10.2	308
144	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
143	Common polygenic variation in coeliac disease and confirmation of ZNF335 and NIFA as disease susceptibility loci. <i>European Journal of Human Genetics</i> , 2016 , 24, 291-7	5.3	18
142	Dissemination of clonally related multidrug-resistant Klebsiella pneumoniae in Ireland. <i>Epidemiology and Infection</i> , 2016 , 144, 443-8	4.3	11

141	Cognitive analysis of schizophrenia risk genes that function as epigenetic regulators of gene expression. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2016 , 171, 1170-1179 ^{3.5}	25
140	Evidence for Genetic Overlap Between Schizophrenia and Age at First Birth in Women. <i>JAMA Psychiatry</i> , 2016 , 73, 497-505	14.5 40
139	No Reliable Association between Runs of Homozygosity and Schizophrenia in a Well-Powered Replication Study. <i>PLoS Genetics</i> , 2016 , 12, e1006343	6 15
138	Fertility and genomics: comparison of gene expression in contrasting reproductive tissues of female cattle. <i>Reproduction, Fertility and Development</i> , 2016 , 28, 11-24	1.8 6
137	P.3.018 Schizophrenia susceptibility gene CACNA1C associates with microstructure of fornix. <i>European Neuropsychopharmacology</i> , 2016 , 26, S61-S62	1.2
136	Novel genetic loci underlying human intracranial volume identified through genome-wide association. <i>Nature Neuroscience</i> , 2016 , 19, 1569-1582	25.5 147
135	Common genetic variants influence human subcortical brain structures. <i>Nature</i> , 2015 , 520, 224-9	50.4 601
134	MIR137HG risk variant rs1625579 genotype is related to corpus callosum volume in schizophrenia. <i>Neuroscience Letters</i> , 2015 , 602, 44-9	3.3 16
133	Greater number of older siblings is associated with decreased theory of mind ability in psychosis. <i>Schizophrenia Research</i> , 2015 , 165, 247-8	3.6
132	BDNF Val66Met polymorphism in patterns of neural activation in individuals with MDD and healthy controls. <i>Journal of Affective Disorders</i> , 2015 , 184, 239-44	6.6 13
131	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
130	Dense genotyping of immune-related susceptibility loci reveals new insights into the genetics of psoriatic arthritis. <i>Nature Communications</i> , 2015 , 6, 6046	17.4 103
129	Characterization of methicillin-resistant <i>Staphylococcus aureus</i> from residents and the environment in a long-term care facility. <i>Epidemiology and Infection</i> , 2015 , 143, 2985-8	4.3 4
128	Mutational analysis of the insulin-like growth factor 1 receptor tyrosine kinase domain in non-small cell lung cancer patients. <i>Molecular and Clinical Oncology</i> , 2015 , 3, 1073-1079	1.6 2
127	Independent evidence for an association between general cognitive ability and a genetic locus for educational attainment. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2015 , 168B, 363-73	3.5 21
126	DNA methylation of the serotonin transporter gene in peripheral cells and stress-related changes in hippocampal volume: a study in depressed patients and healthy controls. <i>PLoS ONE</i> , 2015 , 10, e0119061 ^{3.7}	110
125	AuthorsSreply. <i>British Journal of Psychiatry</i> , 2015 , 206, 344	5.4
124	Genome-wide comparative analysis of atopic dermatitis and psoriasis gives insight into opposing genetic mechanisms. <i>American Journal of Human Genetics</i> , 2015 , 96, 104-20	11 113

123	DNA methylation of the serotonin transporter gene (SLC6A4) is associated with brain function involved in processing emotional stimuli. <i>Journal of Psychiatry and Neuroscience</i> , 2015 , 40, 296-305	4.5	56
122	Translation of 5S leaders is pervasive in genes resistant to eIF2 repression. <i>ELife</i> , 2015 , 4, e03971	8.9	203
121	No evidence that runs of homozygosity are associated with schizophrenia in an Irish genome-wide association dataset. <i>Schizophrenia Research</i> , 2014 , 154, 79-82	3.6	13
120	Identification of mechanosensitive genes during skeletal development: alteration of genes associated with cytoskeletal rearrangement and cell signalling pathways. <i>BMC Genomics</i> , 2014 , 15, 48	4.5	58
119	Variability in working memory performance explained by epistasis vs polygenic scores in the ZNF804A pathway. <i>JAMA Psychiatry</i> , 2014 , 71, 778-785	14.5	24
118	Evidence that duplications of 22q11.2 protect against schizophrenia. <i>Molecular Psychiatry</i> , 2014 , 19, 37-40	45.1	130
117	Molecular genetic evidence for overlap between general cognitive ability and risk for schizophrenia: a report from the Cognitive Genomics consortium (COGENT). <i>Molecular Psychiatry</i> , 2014 , 19, 168-74	15.1	142
116	The ENIGMA Consortium: large-scale collaborative analyses of neuroimaging and genetic data. <i>Brain Imaging and Behavior</i> , 2014 , 8, 153-82	4.1	539
115	Biological insights from 108 schizophrenia-associated genetic loci. <i>Nature</i> , 2014 , 511, 421-7	50.4	5249
114	Effects of a novel schizophrenia risk variant rs7914558 at CNM2 on brain structure and attributional style. <i>British Journal of Psychiatry</i> , 2014 , 204, 115-21	5.4	25
113	De novo mutations in schizophrenia implicate chromatin remodeling and support a genetic overlap with autism and intellectual disability. <i>Molecular Psychiatry</i> , 2014 , 19, 652-8	15.1	263
112	Excess of rare novel loss-of-function variants in synaptic genes in schizophrenia and autism spectrum disorders. <i>Molecular Psychiatry</i> , 2014 , 19, 872-9	15.1	131
111	BDNF Val66Met genotype interacts with childhood adversity and influences the formation of hippocampal subfields. <i>Human Brain Mapping</i> , 2014 , 35, 5776-83	5.9	53
110	Genome-wide schizophrenia variant at MIR137 does not impact white matter microstructure in healthy participants. <i>Neuroscience Letters</i> , 2014 , 574, 6-10	3.3	15
109	Analysis of the hexanucleotide repeat expansion and founder haplotype at C9ORF72 in an Irish psychosis case-control sample. <i>Neurobiology of Aging</i> , 2014 , 35, 1510.e1-5	5.6	15
108	Altered medial prefrontal activity during dynamic face processing in schizophrenia spectrum patients. <i>Schizophrenia Research</i> , 2014 , 157, 225-30	3.6	24
107	Genome-wide association studies: findings at the major histocompatibility complex locus in psychosis. <i>Biological Psychiatry</i> , 2014 , 75, 276-83	7.9	95
106	Allelic differences between Europeans and Chinese for CREB1 SNPs and their implications in gene expression regulation, hippocampal structure and function, and bipolar disorder susceptibility. <i>Molecular Psychiatry</i> , 2014 , 19, 452-61	15.1	52

105	Next-generation sequencing of the mitochondrial genome and association with IgA nephropathy in a renal transplant population. <i>Scientific Reports</i> , 2014 , 4, 7379	4.9	11
104	Effects of ZNF804A on auditory P300 response in schizophrenia. <i>Translational Psychiatry</i> , 2014 , 4, e345	8.6	16
103	The miR-137 schizophrenia susceptibility variant rs1625579 does not predict variability in brain volume in a sample of schizophrenic patients and healthy individuals. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2014 , 165B, 467-71	3.5	15
102	An inherited duplication at the gene p21 Protein-Activated Kinase 7 (PAK7) is a risk factor for psychosis. <i>Human Molecular Genetics</i> , 2014 , 23, 3316-26	5.6	32
101	The one and the many: effects of the cell adhesion molecule pathway on neuropsychological function in psychosis. <i>Psychological Medicine</i> , 2014 , 44, 2177-87	6.9	12
100	CNV analysis in a large schizophrenia sample implicates deletions at 16p12.1 and SLC1A1 and duplications at 1p36.33 and CGNL1. <i>Human Molecular Genetics</i> , 2014 , 23, 1669-76	5.6	61
99	Common variant at 16p11.2 conferring risk of psychosis. <i>Molecular Psychiatry</i> , 2014 , 19, 108-14	15.1	67
98	Effects of MIR137 on fronto-amygdala functional connectivity. <i>NeuroImage</i> , 2014 , 90, 189-95	7.9	36
97	Genome-wide association analysis identifies 13 new risk loci for schizophrenia. <i>Nature Genetics</i> , 2013 , 45, 1150-9	36.3	1153
96	Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. <i>Nature Genetics</i> , 2013 , 45, 984-94	36.3	1628
95	Mood congruent psychotic symptoms and specific cognitive deficits in carriers of the novel schizophrenia risk variant at MIR-137. <i>Neuroscience Letters</i> , 2013 , 532, 33-8	3.3	55
94	Implication of a rare deletion at distal 16p11.2 in schizophrenia. <i>JAMA Psychiatry</i> , 2013 , 70, 253-60	14.5	56
93	Neuropsychological effects of the CSMD1 genome-wide associated schizophrenia risk variant rs10503253. <i>Genes, Brain and Behavior</i> , 2013 , 12, 203-9	3.6	41
92	Social dysfunction in schizophrenia: an investigation of the GAF scales' sensitivity to deficits in social cognition. <i>Schizophrenia Research</i> , 2013 , 146, 363-5	3.6	21
91	Genome-wide significant associations in schizophrenia to ITIH3/4, CACNA1C and SDCCAG8, and extensive replication of associations reported by the Schizophrenia PGC. <i>Molecular Psychiatry</i> , 2013 , 18, 708-12	15.1	184
90	The role of the major histocompatibility complex region in cognition and brain structure: a schizophrenia GWAS follow-up. <i>American Journal of Psychiatry</i> , 2013 , 170, 877-85	11.9	51
89	Brain-derived neurotrophic factor Val66Met polymorphism and early life adversity affect hippocampal volume. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2013 , 162B, 183-90	3.5	71
88	Neural effects of the CSMD1 genome-wide associated schizophrenia risk variant rs10503253. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2013 , 162B, 530-7	3.5	26

87	Delineating the genetic heterogeneity of ALS using targeted high-throughput sequencing. <i>Journal of Medical Genetics</i> , 2013 , 50, 776-83	5.8	129
86	Relationship between the COMT-Val158Met and BDNF-Val66Met polymorphisms, childhood trauma and psychotic experiences in an adolescent general population sample. <i>PLoS ONE</i> , 2013 , 8, e79747	2.7	23
85	Development of strategies for SNP detection in RNA-seq data: application to lymphoblastoid cell lines and evaluation using 1000 Genomes data. <i>PLoS ONE</i> , 2013 , 8, e58815	3.7	94
84	Promoter polymorphisms in two overlapping 6p25 genes implicate mitochondrial proteins in cognitive deficit in schizophrenia. <i>Molecular Psychiatry</i> , 2012 , 17, 1328-39	15.1	14
83	ZNF804A and social cognition in patients with schizophrenia and healthy controls. <i>Molecular Psychiatry</i> , 2012 , 17, 118-9	15.1	18
82	Identification of common variants associated with human hippocampal and intracranial volumes. <i>Nature Genetics</i> , 2012 , 44, 552-61	36.3	498
81	Insulin-like growth factor 1 (IGF1) and its active peptide (1-3)IGF1 enhance the expression of synaptic markers in neuronal circuits through different cellular mechanisms. <i>Neuroscience Letters</i> , 2012 , 520, 51-6	3.3	42
80	Global endometrial transcriptomic profiling: transient immune activation precedes tissue proliferation and repair in healthy beef cows. <i>BMC Genomics</i> , 2012 , 13, 489	4.5	22
79	Social cognition in bipolar disorder versus schizophrenia: comparability in mental state decoding deficits. <i>Bipolar Disorders</i> , 2012 , 14, 743-8	3.8	37
78	Neurotrophic tyrosine kinase polymorphism impacts white matter connections in patients with major depressive disorder. <i>Biological Psychiatry</i> , 2012 , 72, 663-70	7.9	45
77	The NOS1 variant rs6490121 is associated with variation in prefrontal function and grey matter density in healthy individuals. <i>NeuroImage</i> , 2012 , 60, 614-22	7.9	22
76	Preserved cognitive function is associated with suicidal ideation and single suicide attempts in schizophrenia. <i>Schizophrenia Research</i> , 2012 , 140, 232-6	3.6	59
75	Avian resistance to <i>Campylobacter jejuni</i> colonization is associated with an intestinal immunogene expression signature identified by mRNA sequencing. <i>PLoS ONE</i> , 2012 , 7, e40409	3.7	39
74	A NOS1 variant implicated in cognitive performance influences evoked neural responses during a high density EEG study of early visual perception. <i>Human Brain Mapping</i> , 2012 , 33, 1202-11	5.9	17
73	Reduced fractional anisotropy in the uncinate fasciculus in patients with major depression carrying the met-allele of the Val66Met brain-derived neurotrophic factor genotype. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2012 , 159B, 537-48	3.5	64
72	Effect of genetic variant in BICC1 on functional and structural brain changes in depression. <i>Neuropsychopharmacology</i> , 2012 , 37, 2855-62	8.7	37
71	The effect of the neurogranin schizophrenia risk variant rs12807809 on brain structure and function. <i>Twin Research and Human Genetics</i> , 2012 , 15, 296-303	2.2	26
70	Functional investigation of a schizophrenia GWAS signal at the CDC42 gene. <i>World Journal of Biological Psychiatry</i> , 2012 , 13, 550-4	3.8	12

69	Genome-wide linkage analysis of 972 bipolar pedigrees using single-nucleotide polymorphisms. <i>Molecular Psychiatry</i> , 2012 , 17, 818-26	15.1	24
68	Genome-wide association study identifies five new schizophrenia loci. <i>Nature Genetics</i> , 2011 , 43, 969-76	36.3	1508
67	A neuropsychological investigation of the genome wide associated schizophrenia risk variant NRGN rs12807809. <i>Schizophrenia Research</i> , 2011 , 125, 304-6	3.6	20
66	ZNF804A risk allele is associated with relatively intact gray matter volume in patients with schizophrenia. <i>NeuroImage</i> , 2011 , 54, 2132-7	7.9	74
65	Duplications of the neuropeptide receptor gene VIPR2 confer significant risk for schizophrenia. <i>Nature</i> , 2011 , 471, 499-503	50.4	257
64	Genetic classification of populations using supervised learning. <i>PLoS ONE</i> , 2011 , 6, e14802	3.7	14
63	Assessment of inactivating stop codon mutations in forty <i>Saccharomyces cerevisiae</i> strains: implications for [PSI] prion-mediated phenotypes. <i>PLoS ONE</i> , 2011 , 6, e28684	3.7	13
62	Fine mapping of ZNF804A and genome-wide significant evidence for its involvement in schizophrenia and bipolar disorder. <i>Molecular Psychiatry</i> , 2011 , 16, 429-41	15.1	221
61	Molecular pathways involved in neuronal cell adhesion and membrane scaffolding contribute to schizophrenia and bipolar disorder susceptibility. <i>Molecular Psychiatry</i> , 2011 , 16, 286-92	15.1	175
60	GWA study data mining and independent replication identify cardiomyopathy-associated 5 (CMYA5) as a risk gene for schizophrenia. <i>Molecular Psychiatry</i> , 2011 , 16, 1117-29	15.1	58
59	Functional assessment of a promoter polymorphism in S100B, a putative risk variant for bipolar disorder. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2011 , 156B, 691-9	3.5	15
58	First implication of STRA6 mutations in isolated anophthalmia, microphthalmia, and coloboma: a new dimension to the STRA6 phenotype. <i>Human Mutation</i> , 2011 , 32, 1417-26	4.7	62
57	Large-scale genome-wide association analysis of bipolar disorder identifies a new susceptibility locus near ODZ4. <i>Nature Genetics</i> , 2011 , 43, 977-83	36.3	1094
56	Allelic expression imbalance of the schizophrenia susceptibility gene CHI3L1: evidence of cis-acting variation and tissue specific regulation. <i>Psychiatric Genetics</i> , 2011 , 21, 281-6	2.9	1
55	Multiplex target enrichment using DNA indexing for ultra-high throughput SNP detection. <i>DNA Research</i> , 2011 , 18, 31-8	4.5	35
54	Mutation of Semaphorin-6A disrupts limbic and cortical connectivity and models neurodevelopmental psychopathology. <i>PLoS ONE</i> , 2011 , 6, e26488	3.7	32
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43	Influence of NOS1 on verbal intelligence and working memory in both patients with schizophrenia and healthy control subjects. <i>Archives of General Psychiatry</i> , 2009 , 66, 1045-54		42
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24	D-amino acid oxidase (DAO) genotype and mood symptomatology in schizophrenia. <i>Neuroscience Letters</i> , 2007 , 426, 97-100	3.3	25
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22	Analysis of high-resolution HapMap of DTNBP1 (Dysbindin) suggests no consistency between reported common variant associations and schizophrenia. <i>American Journal of Human Genetics</i> , 2006 , 79, 903-9	11	103
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12	Universal, robust, highly quantitative SNP allele frequency measurement in DNA pools. <i>Human Genetics</i> , 2002 , 110, 471-8	6.3	152
11	Molecular genetic evidence for overlap between general cognitive ability and risk for schizophrenia: a report from the Cognitive Genomics consortium (COGENT)		1
10	Genome-wide association study identifies 30 Loci Associated with Bipolar Disorder		28
9	Ninety-nine independent genetic loci influencing general cognitive function include genes associated with brain health and structure (N = 280,360)		6
8	GWAS meta-analysis (N=279,930) identifies new genes and functional links to intelligence		9
7	Genes influenced by MEF2C contribute to neurodevelopmental disease via gene expression changes that affect multiple types of cortical excitatory neurons		1
6	Large-scale analysis of DNA methylation identifies cellular alterations in blood from psychosis patients and molecular biomarkers of treatment-resistant schizophrenia		1
5	Controlling for background genetic effects using polygenic scores improves the power of genome-wide association studies		1
4	Genome-wide association study of over 40,000 bipolar disorder cases provides new insights into the underlying biology		11
3	Association between schizophrenia and both loss of function and missense mutations in paralog conserved sites of voltage-gated sodium channels		2
2	The genetic architecture of the human cerebral cortex		12
1	Pleiotropic Meta-Analysis of Cognition, Education, and Schizophrenia Differentiates Roles of Early Neurodevelopmental and Adult Synaptic Pathways		2