

Aiden P Corvin

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.

187 papers	20,752 citations	56 h-index	143 g-index
202 ext. papers	25,973 ext. citations	10.2 avg, IF	5.24 L-index

#	Paper	IF	Citations
187	Sex-Dependent Shared and Nonshared Genetic Architecture Across Mood and Psychotic Disorders. <i>Biological Psychiatry</i> , 2022 , 91, 102-117	7.9	11
186	Schizophrenia genomics 2022 , 17-41		
185	Mapping genomic loci implicates genes and synaptic biology in schizophrenia.. <i>Nature</i> , 2022 ,	50.4	35
184	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
183	Minding metabolism: targeted interventions to improve cardio-metabolic monitoring across early and chronic psychosis. <i>Irish Journal of Medical Science</i> , 2021 , 1	1.9	1
182	Changes in Default-Mode Network Associated With Childhood Trauma in Schizophrenia. <i>Schizophrenia Bulletin</i> , 2021 , 47, 1482-1494	1.3	3
181	Converting single nucleotide variants between genome builds: from cautionary tale to solution. <i>Briefings in Bioinformatics</i> , 2021 , 22,	13.4	1
180	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
179	Identifying nootropic drug targets via large-scale cognitive GWAS and transcriptomics. <i>Neuropsychopharmacology</i> , 2021 , 46, 1788-1801	8.7	1
178	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
177	DNA methylation meta-analysis reveals cellular alterations in psychosis and markers of treatment-resistant schizophrenia. <i>ELife</i> , 2021 , 10,	8.9	15
176	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
175	Characterisation of age and polarity at onset in bipolar disorder.. <i>British Journal of Psychiatry</i> , 2021 , 219, 659-669	5.4	2
174	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
173	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
172	Dissecting the Shared Genetic Architecture of Suicide Attempt, Psychiatric Disorders, and Known Risk Factors. <i>Biological Psychiatry</i> , 2021 ,	7.9	11
171	Investigating rare pathogenic/likely pathogenic exonic variation in bipolar disorder. <i>Molecular Psychiatry</i> , 2021 , 26, 5239-5250	15.1	3

170	The Relationship Between Polygenic Risk Scores and Cognition in Schizophrenia. <i>Schizophrenia Bulletin</i> , 2020 , 46, 336-344	1.3	38
169	Prevalence of N-Methyl-d-Aspartate Receptor antibody (NMDAR-Ab) encephalitis in patients with first episode psychosis and treatment resistant schizophrenia on clozapine, a population based study. <i>Schizophrenia Research</i> , 2020 , 222, 455-461	3.6	8
168	Deficit not bias: A quantifiable neuropsychological model of delusions. <i>Schizophrenia Research</i> , 2020 , 222, 496-498	3.6	1
167	The genetic architecture of the human cerebral cortex. <i>Science</i> , 2020 , 367,	33.3	156
166	Neuroharmony: A new tool for harmonizing volumetric MRI data from unseen scanners. <i>NeuroImage</i> , 2020 , 220, 117127	7.9	14
165	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	7
164	Autoantibodies and Psychosis. <i>Current Topics in Behavioral Neurosciences</i> , 2020 , 44, 85-123	3.4	4
163	Methyl-CpG-binding protein 2 mediates overlapping mechanisms across brain disorders. <i>Scientific Reports</i> , 2020 , 10, 22255	4.9	5
162	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
161	Integrating machine learning and multimodal neuroimaging to detect schizophrenia at the level of the individual. <i>Human Brain Mapping</i> , 2020 , 41, 1119-1135	5.9	23
160	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
159	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
158	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
157	Detecting schizophrenia at the level of the individual: relative diagnostic value of whole-brain images, connectome-wide functional connectivity and graph-based metrics. <i>Psychological Medicine</i> , 2020 , 50, 1852-1861	6.9	32
156	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
155	Cortical patterning of abnormal morphometric similarity in psychosis is associated with brain expression of schizophrenia-related genes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 9604-9609	11.5	70
154	Genome-wide association study identifies 30 loci associated with bipolar disorder. <i>Nature Genetics</i> , 2019 , 51, 793-803	36.3	662
153	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

152	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
151	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
150	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
149	A randomized controlled trial of cognitive remediation for a national cohort of forensic patients with schizophrenia or schizoaffective disorder. <i>BMC Psychiatry</i> , 2019 , 19, 27	4.2	20
148	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
147	Moral cognition and homicide amongst forensic patients with schizophrenia and schizoaffective disorder: A cross-sectional cohort study. <i>Schizophrenia Research</i> , 2018 , 193, 468-469	3.6	3
146	Analysis of shared heritability in common disorders of the brain. <i>Science</i> , 2018 , 360,	33.3	666
145	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
144	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
143	Study of 300,486 individuals identifies 148 independent genetic loci influencing general cognitive function. <i>Nature Communications</i> , 2018 , 9, 2098	17.4	254
142	Computerised working memory-based cognitive remediation therapy does not affect Reading the Mind in The Eyes test performance or neural activity during a Facial Emotion Recognition test in psychosis. <i>European Journal of Neuroscience</i> , 2018 , 48, 1691	3.5	0
141	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
140	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
139	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
138	Novel genetic loci associated with hippocampal volume. <i>Nature Communications</i> , 2017 , 8, 13624	17.4	173
137	Occurrence and co-occurrence of hallucinations by modality in schizophrenia-spectrum disorders. <i>Psychiatry Research</i> , 2017 , 252, 154-160	9.9	62
136	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
135	Cognitive Characterization of Schizophrenia Risk Variants Involved in Synaptic Transmission: Evidence of CACNA1C Role in Working Memory. <i>Neuropsychopharmacology</i> , 2017 , 42, 2612-2622	8.7	16

134	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
133	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
132	Fluorescent nanodiamond tracking reveals intraneuronal transport abnormalities induced by brain-disease-related genetic risk factors. <i>Nature Nanotechnology</i> , 2017 , 12, 322-328	28.7	79
131	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
130	Genetics of Schizophrenia: Ready to Translate?. <i>Current Psychiatry Reports</i> , 2017 , 19, 61	9.1	36
129	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
128	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
127	Polymorphism in a lincRNA Associates with a Doubled Risk of Pneumococcal Bacteremia in Kenyan Children. <i>American Journal of Human Genetics</i> , 2016 , 98, 1092-1100	11	30
126	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
125	Psychiatric genetics: what's new in 2015?. <i>Lancet Psychiatry</i> , 2016 , 3, 10-12	23.3	2
124	What Next in Schizophrenia Genetics for the Psychiatric Genomics Consortium?. <i>Schizophrenia Bulletin</i> , 2016 , 42, 538-41	1.3	19
123	Expression of nuclear Methyl-CpG binding protein 2 (Mecp2) is dependent on neuronal stimulation and application of Insulin-like growth factor 1. <i>Neuroscience Letters</i> , 2016 , 621, 111-116	3.3	10
122	Data science for mental health: a UK perspective on a global challenge. <i>Lancet Psychiatry</i> , 2016 , 3, 993-998	23.3	34
121	Novel genetic loci underlying human intracranial volume identified through genome-wide association. <i>Nature Neuroscience</i> , 2016 , 19, 1569-1582	25.5	147
120	Common genetic variants influence human subcortical brain structures. <i>Nature</i> , 2015 , 520, 224-9	50.4	601
119	MIR137HG risk variant rs1625579 genotype is related to corpus callosum volume in schizophrenia. <i>Neuroscience Letters</i> , 2015 , 602, 44-9	3.3	16
118	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	
117	Modeling Linkage Disequilibrium Increases Accuracy of Polygenic Risk Scores. <i>American Journal of Human Genetics</i> , 2015 , 97, 576-92	11	649

116	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
115	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
114	Biomarkers for Psychosis: the Molecular Genetics of Psychosis. <i>Current Behavioral Neuroscience Reports</i> , 2015 , 2, 112-118	1.7	1
113	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
112	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
111	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
110	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
109	Unlocking the treasure trove: from genes to schizophrenia biology. <i>Schizophrenia Bulletin</i> , 2014 , 40, 492-63	13	
108	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
107	Polygenic dissection of diagnosis and clinical dimensions of bipolar disorder and schizophrenia. <i>Molecular Psychiatry</i> , 2014 , 19, 1017-1024	15.1	258
106	Effects of a novel schizophrenia risk variant rs7914558 at CNNM2 on brain structure and attributional style. <i>British Journal of Psychiatry</i> , 2014 , 204, 115-21	5.4	25
105	The correlation between reading and mathematics ability at age twelve has a substantial genetic component. <i>Nature Communications</i> , 2014 , 5, 4204	17.4	54
104	The phenotypic manifestations of rare CNVs in schizophrenia. <i>Schizophrenia Research</i> , 2014 , 158, 255-60	3.6	16
103	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
102	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
101	A genome-wide association analysis of a broad psychosis phenotype identifies three loci for further investigation. <i>Biological Psychiatry</i> , 2014 , 75, 386-97	7.9	36
100	Altered medial prefrontal activity during dynamic face processing in schizophrenia spectrum patients. <i>Schizophrenia Research</i> , 2014 , 157, 225-30	3.6	24
99	Genome-wide association studies: findings at the major histocompatibility complex locus in psychosis. <i>Biological Psychiatry</i> , 2014 , 75, 276-83	7.9	95

98	Repeated insulin-like growth factor 1 treatment in a patient with rett syndrome: a single case study. <i>Frontiers in Pediatrics</i> , 2014 , 2, 52	3.4	24
97	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
96	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
95	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
94	Effects of MIR137 on fronto-amygdala functional connectivity. <i>NeuroImage</i> , 2014 , 90, 189-95	7.9	36
93	Genetic modifiers and subtypes in schizophrenia: investigations of age at onset, severity, sex and family history. <i>Schizophrenia Research</i> , 2014 , 154, 48-53	3.6	49
92	Genome-wide association analysis identifies 13 new risk loci for schizophrenia. <i>Nature Genetics</i> , 2013 , 45, 1150-9	36.3	1153
91	Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. <i>Nature Genetics</i> , 2013 , 45, 984-94	36.3	1628
90	Implication of a rare deletion at distal 16p11.2 in schizophrenia. <i>JAMA Psychiatry</i> , 2013 , 70, 253-60	14.5	56
89	Mosaic copy number variation in schizophrenia. <i>European Journal of Human Genetics</i> , 2013 , 21, 1007-11	5.3	10
88	Common variants in the HLA-DRB1-HLA-DQA1 HLA class II region are associated with susceptibility to visceral leishmaniasis. <i>Nature Genetics</i> , 2013 , 45, 208-13	36.3	76
87	Social dysfunction in schizophrenia: an investigation of the GAF scale's sensitivity to deficits in social cognition. <i>Schizophrenia Research</i> , 2013 , 146, 363-5	3.6	21
86	A comprehensive family-based replication study of schizophrenia genes. <i>JAMA Psychiatry</i> , 2013 , 70, 573-81	14.5	115
85	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
84	Genome-wide association study of intraocular pressure identifies the GLCCI1/ICA1 region as a glaucoma susceptibility locus. <i>Human Molecular Genetics</i> , 2013 , 22, 4653-60	5.6	24
83	Schizophrenia at a genetics crossroads: where to now?. <i>Schizophrenia Bulletin</i> , 2013 , 39, 490-5	1.3	12
82	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
81	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

80	Genome-wide association study identifies a variant in HDAC9 associated with large vessel ischemic stroke. <i>Nature Genetics</i> , 2012 , 44, 328-33	36.3	314
79	Identification of common variants associated with human hippocampal and intracranial volumes. <i>Nature Genetics</i> , 2012 , 44, 552-61	36.3	498
78	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
77	Social cognition in bipolar disorder versus schizophrenia: comparability in mental state decoding deficits. <i>Bipolar Disorders</i> , 2012 , 14, 743-8	3.8	37
76	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
75	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
74	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
73	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
72	Common variants at the MHC locus and at chromosome 16q24.1 predispose to Barrett's esophagus. <i>Nature Genetics</i> , 2012 , 44, 1131-6	36.3	139
71	Psychiatric genetics 2012 , 35-53		0
70	A neuropsychological investigation of the genome wide associated schizophrenia risk variant NRGN rs12807809. <i>Schizophrenia Research</i> , 2011 , 125, 304-6	3.6	20
69	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
68	High frequencies of de novo CNVs in bipolar disorder and schizophrenia. <i>Neuron</i> , 2011 , 72, 951-63	13.9	240
67	Duplications of the neuropeptide receptor gene VIPR2 confer significant risk for schizophrenia. <i>Nature</i> , 2011 , 471, 499-503	50.4	257
66	Genetic classification of populations using supervised learning. <i>PLoS ONE</i> , 2011 , 6, e14802	3.7	14
65	Common variants near ATM are associated with glycemic response to metformin in type 2 diabetes. <i>Nature Genetics</i> , 2011 , 43, 117-20	36.3	319
64	Interaction between ERAP1 and HLA-B27 in ankylosing spondylitis implicates peptide handling in the mechanism for HLA-B27 in disease susceptibility. <i>Nature Genetics</i> , 2011 , 43, 761-7	36.3	646
63	Two patients walk into a clinic...a genomics perspective on the future of schizophrenia. <i>BMC Biology</i> , 2011 , 9, 77	7.3	10

62	Genetic risk and a primary role for cell-mediated immune mechanisms in multiple sclerosis. <i>Nature</i> , 2011 , 476, 214-9	50.4	1948
61	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
60	Multiplex target enrichment using DNA indexing for ultra-high throughput SNP detection. <i>DNA Research</i> , 2011 , 18, 31-8	4.5	35
59	Dissection of the genetics of Parkinson's disease identifies an additional association of SNCA and multiple associated haplotypes at 17q21. <i>Human Molecular Genetics</i> , 2011 , 20, 345-53	5.6	178
58	Mutation of Semaphorin-6A disrupts limbic and cortical connectivity and models neurodevelopmental psychopathology. <i>PLoS ONE</i> , 2011 , 6, e26488	3.7	32
57	The shock of the new: progress in schizophrenia genomics. <i>Current Genomics</i> , 2011 , 12, 516-24	2.6	15
56	Population structure and genome-wide patterns of variation in Ireland and Britain. <i>European Journal of Human Genetics</i> , 2010 , 18, 1248-54	5.3	36
55	Genes predict village of origin in rural Europe. <i>European Journal of Human Genetics</i> , 2010 , 18, 1269-70	5.3	19
54	A genome-wide association study identifies new psoriasis susceptibility loci and an interaction between HLA-C and ERAP1. <i>Nature Genetics</i> , 2010 , 42, 985-90	36.3	773
53	Evidence for cis-acting regulation of ANK3 and CACNA1C gene expression. <i>Bipolar Disorders</i> , 2010 , 12, 440-5	3.8	26
52	Psychosis susceptibility gene ZNF804A and cognitive performance in schizophrenia. <i>Archives of General Psychiatry</i> , 2010 , 67, 692-700		120
51	Reduced occipital and prefrontal brain volumes in dysbindin-associated schizophrenia. <i>Neuropsychopharmacology</i> , 2010 , 35, 368-73	8.7	25
50	Genetic differences between five European populations. <i>Human Heredity</i> , 2010 , 70, 141-9	1.1	24
49	Neuronal cell adhesion genes: Key players in risk for schizophrenia, bipolar disorder and other neurodevelopmental brain disorders?. <i>Cell Adhesion and Migration</i> , 2010 , 4, 511-4	3.2	23
48	The psychosis susceptibility gene ZNF804A: associations, functions, and phenotypes. <i>Schizophrenia Bulletin</i> , 2010 , 36, 904-9	1.3	44
47	Replicated genetic evidence supports a role for HOMER2 in schizophrenia. <i>Neuroscience Letters</i> , 2010 , 468, 229-33	3.3	18
46	Is "clinical" insight the same as "cognitive" insight in schizophrenia?. <i>Journal of the International Neuropsychological Society</i> , 2009 , 15, 471-5	3.1	27
45	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

44	The SNP ratio test: pathway analysis of genome-wide association datasets. <i>Bioinformatics</i> , 2009 , 25, 2762-3	7.3	116
43	Copy-number variants in neurodevelopmental disorders: promises and challenges. <i>Trends in Genetics</i> , 2009 , 25, 536-44	8.5	102
42	Genome-wide association study of ulcerative colitis identifies three new susceptibility loci, including the HNF4A region. <i>Nature Genetics</i> , 2009 , 41, 1330-4	36.3	411
41	Mood-incongruent psychosis in bipolar disorder: conditional linkage analysis shows genome-wide suggestive linkage at 1q32.3, 7p13 and 20q13.31. <i>Bipolar Disorders</i> , 2009 , 11, 610-20	3.8	20
40	Does the ability to sustain attention underlie symptom severity in schizophrenia?. <i>Schizophrenia Research</i> , 2009 , 107, 319-23	3.6	24
39	Identification of loci associated with schizophrenia by genome-wide association and follow-up. <i>Nature Genetics</i> , 2008 , 40, 1053-5	36.3	877
38	Collaborative genome-wide association analysis supports a role for ANK3 and CACNA1C in bipolar disorder. <i>Nature Genetics</i> , 2008 , 40, 1056-8	36.3	949
37	An assessment of the Irish population for large-scale genetic mapping studies involving epilepsy and other complex diseases. <i>European Journal of Human Genetics</i> , 2008 , 16, 176-83	5.3	4
36	Familial patterns and the origins of individual differences in synaesthesia. <i>Cognition</i> , 2008 , 106, 871-93	3.5	123
35	Mental state decoding v. mental state reasoning as a mediator between cognitive and social function in psychosis. <i>British Journal of Psychiatry</i> , 2008 , 193, 77-8	5.4	58
34	Dysbindin (DTNBP1) and the biogenesis of lysosome-related organelles complex 1 (BLOC-1): main and epistatic gene effects are potential contributors to schizophrenia susceptibility. <i>Biological Psychiatry</i> , 2008 , 63, 24-31	7.9	51
33	Early visual processing deficits in dysbindin-associated schizophrenia. <i>Biological Psychiatry</i> , 2008 , 63, 484-9	7.9	59
32	Chitinase-3-like 1 (CHI3L1) gene and schizophrenia: genetic association and a potential functional mechanism. <i>Biological Psychiatry</i> , 2008 , 64, 98-103	7.9	25
31	A dysbindin risk haplotype associated with less severe manic-type symptoms in psychosis. <i>Neuroscience Letters</i> , 2008 , 431, 146-9	3.3	29
30	Are relational style and neuropsychological performance predictors of social attributions in chronic schizophrenia?. <i>Psychiatry Research</i> , 2008 , 161, 19-27	9.9	31
29	Physical health and attendance at primary care in people with schizophrenia. <i>Irish Journal of Psychological Medicine</i> , 2008 , 25, 57-60	3	2
28	Variance in facial recognition performance associated with BDNF in schizophrenia. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2007 , 144B, 578-9	3.5	6
27	Variance in neurocognitive performance is associated with dysbindin-1 in schizophrenia: a preliminary study. <i>Neuropsychologia</i> , 2007 , 45, 454-8	3.2	101

26	D-amino acid oxidase (DAO) genotype and mood symptomatology in schizophrenia. <i>Neuroscience Letters</i> , 2007 , 426, 97-100	3.3	25
25	Functional genomics and schizophrenia: endophenotypes and mutant models. <i>Psychiatric Clinics of North America</i> , 2007 , 30, 365-99	3.1	38
24	Evidence that interaction between neuregulin 1 and its receptor erbB4 increases susceptibility to schizophrenia. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2006 , 141B, 96-101	3.5	143
23	Evidence that specific executive functions predict symptom variance among schizophrenia patients with a predominantly negative symptom profile. <i>Cognitive Neuropsychiatry</i> , 2006 , 11, 13-32	2	41
22	Evaluation of a susceptibility gene for schizophrenia: genotype based meta-analysis of RGS4 polymorphisms from thirteen independent samples. <i>Biological Psychiatry</i> , 2006 , 60, 152-62	7.9	80
21	Neurocognition and suicidal behaviour in an Irish population with major psychotic disorders. <i>Schizophrenia Research</i> , 2006 , 85, 196-200	3.6	53
20	Are deficits in executive sub-processes simply reflecting more general cognitive decline in schizophrenia?. <i>Schizophrenia Research</i> , 2006 , 85, 168-73	3.6	22
19	Do antisaccade deficits in schizophrenia provide evidence of a specific inhibitory function?. <i>Journal of the International Neuropsychological Society</i> , 2006 , 12, 901-6	3.1	14
18	Investigation of the apolipoprotein-L (APOL) gene family and schizophrenia using a novel DNA pooling strategy for public database SNPs. <i>Schizophrenia Research</i> , 2005 , 76, 231-8	3.6	12
17	Are the cognitive deficits associated with impaired insight in schizophrenia specific to executive task performance?. <i>Journal of Nervous and Mental Disease</i> , 2005 , 193, 803-8	1.8	37
16	Genomewide linkage scan in schizoaffective disorder: significant evidence for linkage at 1q42 close to DISC1, and suggestive evidence at 22q11 and 19p13. <i>Archives of General Psychiatry</i> , 2005 , 62, 1081-8		164
15	Confirming RGS4 as a susceptibility gene for schizophrenia. <i>American Journal of Medical Genetics Part A</i> , 2004 , 125B, 50-3		114
14	No evidence for association of the dysbindin gene [DTNBP1] with schizophrenia in an Irish population-based study. <i>Schizophrenia Research</i> , 2003 , 60, 167-72	3.6	74
13	The attitudes of Irish trainees to their training and its supervision: a five-year follow up study. <i>Irish Journal of Psychological Medicine</i> , 2001 , 18, 120-125	3	6
12	Cigarette smoking and psychotic symptoms in bipolar affective disorder. <i>British Journal of Psychiatry</i> , 2001 , 179, 35-8	5.4	77
11	Proof of concept: Molecular prediction of schizophrenia risk		1
10	Genome-wide Association Study of Clinical Features in the Schizophrenia Psychiatric Genomics Consortium: Confirmation of Polygenic Effect on Negative Symptoms		3
9	Genome-wide association study identifies 30 Loci Associated with Bipolar Disorder		28

8	GWAS meta-analysis (N=279,930) identifies new genes and functional links to intelligence	9
7	Large-scale analysis of DNA methylation identifies cellular alterations in blood from psychosis patients and molecular biomarkers of treatment-resistant schizophrenia	1
6	Genome-wide association study of over 40,000 bipolar disorder cases provides new insights into the underlying biology	11
5	Dissecting the shared genetic architecture of suicide attempt, psychiatric disorders and known risk factors	2
4	Association between schizophrenia and both loss of function and missense mutations in paralog conserved sites of voltage-gated sodium channels	2
3	Pleiotropic Meta-Analysis of Cognition, Education, and Schizophrenia Differentiates Roles of Early Neurodevelopmental and Adult Synaptic Pathways	2
2	Characterization of Age and Polarity at Onset in Bipolar Disorder	1
1	Overlapping Etiology of Neurodevelopmental Disorders ²⁹⁻⁴⁸	4