

# David A Collier

## List of Publications by Citations

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281  
papers

26,599  
citations

76  
h-index

156  
g-index

307  
ext. papers

30,428  
ext. citations

8.7  
avg, IF

5.95  
L-index

#	Paper	IF	Citations
281	Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. <i>Nature Genetics</i> , <b>2013</b> , 45, 984-94	36.3	1628
280	Large recurrent microdeletions associated with schizophrenia. <i>Nature</i> , <b>2008</b> , 455, 232-6	50.4	1427
279	Common variants conferring risk of schizophrenia. <i>Nature</i> , <b>2009</b> , 460, 744-7	50.4	1350
278	Association scan of 14,500 nonsynonymous SNPs in four diseases identifies autoimmunity variants. <i>Nature Genetics</i> , <b>2007</b> , 39, 1329-37	36.3	1130
277	Collaborative genome-wide association analysis supports a role for ANK3 and CACNA1C in bipolar disorder. <i>Nature Genetics</i> , <b>2008</b> , 40, 1056-8	36.3	949
276	Common schizophrenia alleles are enriched in mutation-intolerant genes and in regions under strong background selection. <i>Nature Genetics</i> , <b>2018</b> , 50, 381-389	36.3	787
275	The UK10K project identifies rare variants in health and disease. <i>Nature</i> , <b>2015</b> , 526, 82-90	50.4	776
274	The same sequence variant on 9p21 associates with myocardial infarction, abdominal aortic aneurysm and intracranial aneurysm. <i>Nature Genetics</i> , <b>2008</b> , 40, 217-24	36.3	596
273	Contribution of copy number variants to schizophrenia from a genome-wide study of 41,321 subjects. <i>Nature Genetics</i> , <b>2017</b> , 49, 27-35	36.3	530
272	The chemistry and biology of unusual DNA structures adopted by oligopurine ̳oligopyrimidine sequences. <i>FASEB Journal</i> , <b>1988</b> , 2, 2939-2949	0.9	484
271	A novel functional polymorphism within the promoter of the serotonin transporter gene: possible role in susceptibility to affective disorders. <i>Molecular Psychiatry</i> , <b>1996</b> , 1, 453-60	15.1	452
270	Disruption of the neurexin 1 gene is associated with schizophrenia. <i>Human Molecular Genetics</i> , <b>2009</b> , 18, 988-96	5.6	376
269	Genomic Dissection of Bipolar Disorder and Schizophrenia, Including 28 Subphenotypes. <i>Cell</i> , <b>2018</b> , 173, 1705-1715.e16	56.2	360
268	The frequency and distribution of thiopurine methyltransferase alleles in Caucasian and Asian populations. <i>Pharmacogenetics and Genomics</i> , <b>1999</b> , 9, 37-42		314
267	Common variants near CAV1 and CAV2 are associated with primary open-angle glaucoma. <i>Nature Genetics</i> , <b>2010</b> , 42, 906-9	36.3	303
266	Genetic and functional analyses of SHANK2 mutations suggest a multiple hit model of autism spectrum disorders. <i>PLoS Genetics</i> , <b>2012</b> , 8, e1002521	6	297
265	Short-term effects of antipsychotic treatment on cerebral function in drug-naive first-episode schizophrenia revealed by "resting state" functional magnetic resonance imaging. <i>Archives of General Psychiatry</i> , <b>2010</b> , 67, 783-92		294

264	Pharmacogenetic prediction of clozapine response. <i>Lancet, The</i> , <b>2000</b> , 355, 1615-6	40	293
263	Genome-wide association identifies a common variant in the reelin gene that increases the risk of schizophrenia only in women. <i>PLoS Genetics</i> , <b>2008</b> , 4, e28	6	270
262	Meta-analysis of the association of urbanicity with schizophrenia. <i>Schizophrenia Bulletin</i> , <b>2012</b> , 38, 1118-23		265
261	Meta-analysis shows strong positive association of the neuregulin 1 (NRG1) gene with schizophrenia. <i>Human Molecular Genetics</i> , <b>2006</b> , 15, 1995-2002	5.6	239
260	Meta-analysis of studies on genetic variation in 5-HT2A receptors and clozapine response. <i>Schizophrenia Research</i> , <b>1998</b> , 32, 93-9	3.6	229
259	A genome-wide association study of anorexia nervosa. <i>Molecular Psychiatry</i> , <b>2014</b> , 19, 1085-94	15.1	224
258	Association of cerebral deficits with clinical symptoms in antipsychotic-naive first-episode schizophrenia: an optimized voxel-based morphometry and resting state functional connectivity study. <i>American Journal of Psychiatry</i> , <b>2009</b> , 166, 196-205	11.9	217
257	Common variants on 8p12 and 1q24.2 confer risk of schizophrenia. <i>Nature Genetics</i> , <b>2011</b> , 43, 1224-7	36.3	201
256	Set shifting in anorexia nervosa: an examination before and after weight gain, in full recovery and relationship to childhood and adult OCPD traits. <i>Journal of Psychiatric Research</i> , <b>2004</b> , 38, 545-52	5.2	201
255	An integrated genetic-epigenetic analysis of schizophrenia: evidence for co-localization of genetic associations and differential DNA methylation. <i>Genome Biology</i> , <b>2016</b> , 17, 176	18.3	189
254	Neurexin 1 (NRXN1) deletions in schizophrenia. <i>Schizophrenia Bulletin</i> , <b>2009</b> , 35, 851-4	1.3	183
253	Evidence for association between polymorphisms in the promoter and coding regions of the 5-HT2A receptor gene and response to clozapine. <i>Molecular Psychiatry</i> , <b>1998</b> , 3, 61-6	15.1	183
252	Prevalence and heritability of compulsive hoarding: a twin study. <i>American Journal of Psychiatry</i> , <b>2009</b> , 166, 1156-61	11.9	175
251	Cognitive flexibility in anorexia nervosa and bulimia nervosa. <i>Journal of the International Neuropsychological Society</i> , <b>2004</b> , 10, 513-20	3.1	172
250	The serotonin transporter is a potential susceptibility factor for bipolar affective disorder. <i>NeuroReport</i> , <b>1996</b> , 7, 1675-9	1.7	170
249	Association of BDNF with anorexia, bulimia and age of onset of weight loss in six European populations. <i>Human Molecular Genetics</i> , <b>2004</b> , 13, 1205-12	5.6	168
248	Common variants at VRK2 and TCF4 conferring risk of schizophrenia. <i>Human Molecular Genetics</i> , <b>2011</b> , 20, 4076-81	5.6	162
247	Systematic meta-analyses and field synopsis of genetic association studies of violence and aggression. <i>Molecular Psychiatry</i> , <b>2014</b> , 19, 471-7	15.1	158

246	Population-based and family-based studies on the serotonin transporter gene polymorphisms and bipolar disorder: a systematic review and meta-analysis. <i>Molecular Psychiatry</i> , <b>2005</b> , 10, 771-81	15.1	157
245	Rare deletions at the neurexin 3 locus in autism spectrum disorder. <i>American Journal of Human Genetics</i> , <b>2012</b> , 90, 133-41	11	155
244	Genome-wide association study identifies a sequence variant within the DAB2IP gene conferring susceptibility to abdominal aortic aneurysm. <i>Nature Genetics</i> , <b>2010</b> , 42, 692-7	36.3	155
243	Low activity allele of catechol-O-methyltransferase gene associated with rapid cycling bipolar disorder. <i>Molecular Psychiatry</i> , <b>1998</b> , 3, 342-5	15.1	150
242	Association between 5-HT2A gene promoter polymorphism and anorexia nervosa. <i>Lancet, The</i> , <b>1997</b> , 350, 412	40	146
241	Association between BDNF val66 met genotype and episodic memory. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2005</b> , 134B, 73-5	3.5	143
240	Identification of a novel neuregulin 1 at-risk haplotype in Han schizophrenia Chinese patients, but no association with the Icelandic/Scottish risk haplotype. <i>Molecular Psychiatry</i> , <b>2004</b> , 9, 698-704	15.1	142
239	5-HT2A and 5-HT2C receptor polymorphisms and psychopathology in late onset Alzheimer's disease. <i>Human Molecular Genetics</i> , <b>1998</b> , 7, 1507-9	5.6	136
238	Copy number variations in neurodevelopmental disorders. <i>Progress in Neurobiology</i> , <b>2012</b> , 99, 81-91	10.9	129
237	Expanding the range of ZNF804A variants conferring risk of psychosis. <i>Molecular Psychiatry</i> , <b>2011</b> , 16, 59-66	15.1	129
236	Allelic association between a Ser-9-Gly polymorphism in the dopamine D3 receptor gene and schizophrenia. <i>Human Genetics</i> , <b>1996</b> , 97, 714-9	6.3	126
235	Meta-analysis of genome-wide linkage studies in BMI and obesity. <i>Obesity</i> , <b>2007</b> , 15, 2263-75	8	122
234	Therapygenetics: the 5HTTLPR and response to psychological therapy. <i>Molecular Psychiatry</i> , <b>2012</b> , 17, 236-7	15.1	119
233	Penetrance for copy number variants associated with schizophrenia. <i>Human Molecular Genetics</i> , <b>2010</b> , 19, 3477-81	5.6	117
232	Preferential transmission of the high activity allele of COMT in schizophrenia. <i>Psychiatric Genetics</i> , <b>1996</b> , 6, 131-3	2.9	117
231	Influence of X chromosome and hormones on human brain development: a magnetic resonance imaging and proton magnetic resonance spectroscopy study of Turner syndrome. <i>Biological Psychiatry</i> , <b>2006</b> , 59, 273-83	7.9	116
230	Association of BDNF with restricting anorexia nervosa and minimum body mass index: a family-based association study of eight European populations. <i>European Journal of Human Genetics</i> , <b>2005</b> , 13, 428-34	5.3	115
229	Family-based linkage disequilibrium mapping using SNP marker haplotypes: application to a potential locus for schizophrenia at chromosome 22q11. <i>Molecular Psychiatry</i> , <b>2000</b> , 5, 77-84	15.1	111

228	High and low activity alleles of catechol-O-methyltransferase gene: ethnic difference and possible association with Parkinson's disease. <i>Neuroscience Letters</i> , <b>1997</b> , 221, 202-4	3.3	105
227	Genetics of behavioural domains across the neuropsychiatric spectrum; of mice and men. <i>Molecular Psychiatry</i> , <b>2007</b> , 12, 324-30	15.1	103
226	Effect of BDNF val(66)met polymorphism on declarative memory and its neural substrate: a meta-analysis. <i>Neuroscience and Biobehavioral Reviews</i> , <b>2012</b> , 36, 2165-77	9	101
225	At-risk variant in TCF7L2 for type II diabetes increases risk of schizophrenia. <i>Biological Psychiatry</i> , <b>2011</b> , 70, 59-63	7.9	101
224	No evidence for an association of affective disorders with high- or low-activity allele of catechol-o-methyltransferase gene. <i>Biological Psychiatry</i> , <b>1997</b> , 42, 282-5	7.9	97
223	Individual-specific risk factors for anorexia nervosa: a pilot study using a discordant sister-pair design. <i>Psychological Medicine</i> , <b>2001</b> , 31, 317-29	6.9	97
222	The quantification of COMT mRNA in post mortem cerebellum tissue: diagnosis, genotype, methylation and expression. <i>BMC Medical Genetics</i> , <b>2006</b> , 7, 10	2.1	95
221	Strong genetic evidence for a selective influence of GABAA receptors on a component of the bipolar disorder phenotype. <i>Molecular Psychiatry</i> , <b>2010</b> , 15, 146-53	15.1	94
220	Catechol-O-methyltransferase polymorphisms and schizophrenia: a transmission disequilibrium study in multiply affected families. <i>Psychiatric Genetics</i> , <b>1997</b> , 7, 97-101	2.9	93
219	Male-biased autosomal effect of 16p13.11 copy number variation in neurodevelopmental disorders. <i>PLoS ONE</i> , <b>2013</b> , 8, e61365	3.7	88
218	Translating genome-wide association findings into new therapeutics for psychiatry. <i>Nature Neuroscience</i> , <b>2016</b> , 19, 1392-1396	25.5	86
217	Performance deficit of alpha7 nicotinic receptor knockout mice in a delayed matching-to-place task suggests a mild impairment of working/episodic-like memory. <i>Genes, Brain and Behavior</i> , <b>2006</b> , 5, 433-40 <sup>3.6</sup>	3.6	86
216	Sensorimotor gating depends on polymorphisms of the serotonin-2A receptor and catechol-O-methyltransferase, but not on neuregulin-1 Arg38Gln genotype: a replication study. <i>Biological Psychiatry</i> , <b>2009</b> , 66, 614-20	7.9	85
215	Catechol-O-methyltransferase Val158Met polymorphism: frequency analysis in Han Chinese subjects and allelic association of the low activity allele with bipolar affective disorder. <i>Pharmacogenetics and Genomics</i> , <b>1997</b> , 7, 349-53		85
214	Association analysis of the DRD4 and COMT genes in methamphetamine abuse. <i>American Journal of Medical Genetics Part A</i> , <b>2004</b> , 129B, 120-4		85
213	Association analysis of the 5-HT5A gene in depression, psychosis and antipsychotic response. <i>NeuroReport</i> , <b>2000</b> , 11, 2017-20	1.7	85
212	Maternally derived microduplications at 15q11-q13: implication of imprinted genes in psychotic illness. <i>American Journal of Psychiatry</i> , <b>2011</b> , 168, 408-17	11.9	84
211	Association analysis of the dopamine D4 gene exon III VNTR and heroin abuse in Chinese subjects. <i>Molecular Psychiatry</i> , <b>1997</b> , 2, 413-6	15.1	84

210	Evaluation of a susceptibility gene for schizophrenia: genotype based meta-analysis of RGS4 polymorphisms from thirteen independent samples. <i>Biological Psychiatry</i> , <b>2006</b> , 60, 152-62	7.9	80
209	Aberrant intrinsic brain activity and cognitive deficit in first-episode treatment-naive patients with schizophrenia. <i>Psychological Medicine</i> , <b>2013</b> , 43, 769-80	6.9	79
208	Serotonin transporter gene and risk for bipolar affective disorder: an association study in Spanish population. <i>Biological Psychiatry</i> , <b>1998</b> , 43, 843-7	7.9	79
207	No association between Parkinson's disease and low-activity alleles of catechol O-methyltransferase. <i>Biochemical and Biophysical Research Communications</i> , <b>1996</b> , 228, 780-4	3.4	77
206	Genetic utility of broadly defined bipolar schizoaffective disorder as a diagnostic concept. <i>British Journal of Psychiatry</i> , <b>2009</b> , 195, 23-9	5.4	76
205	Analysis of Intellectual Disability Copy Number Variants for Association With Schizophrenia. <i>JAMA Psychiatry</i> , <b>2016</b> , 73, 963-969	14.5	75
204	Hypermethylation in the ZBTB20 gene is associated with major depressive disorder. <i>Genome Biology</i> , <b>2014</b> , 15, R56	18.3	73
203	Epistasis between the DAT 3' UTR VNTR and the COMT Val158Met SNP on cortical function in healthy subjects and patients with schizophrenia. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2009</b> , 106, 13600-5	11.5	72
202	Supercoiling and integration host factor change the DNA conformation and alter the flow of convergent transcription in phage Mu. <i>Journal of Biological Chemistry</i> , <b>1989</b> , 264, 3035-42	5.4	72
201	The schizophrenia risk allele C of the TCF4 rs9960767 polymorphism disrupts sensorimotor gating in schizophrenia spectrum and healthy volunteers. <i>Journal of Neuroscience</i> , <b>2011</b> , 31, 6684-91	6.6	70
200	The genetics of schizophrenia: glutamate not dopamine?. <i>European Journal of Pharmacology</i> , <b>2003</b> , 480, 177-84	5.3	70
199	Stereological estimation of the total number of neurons in the murine hippocampus using the optical disector. <i>Journal of Comparative Neurology</i> , <b>1999</b> , 408, 560-6	3.4	69
198	White-matter microstructure in previously drug-naive patients with schizophrenia after 6 weeks of treatment. <i>Psychological Medicine</i> , <b>2013</b> , 43, 2301-9	6.9	68
197	Evidence for association between novel polymorphisms in the PRODH gene and schizophrenia in a Chinese population. <i>American Journal of Medical Genetics Part A</i> , <b>2004</b> , 129B, 13-5		68
196	Allelic association analysis of the dopamine D2, D3, 5-HT2A, and GABA(A)gamma2 receptors and serotonin transporter genes with heroin abuse in Chinese subjects. <i>American Journal of Medical Genetics Part A</i> , <b>2002</b> , 114, 329-35		68
195	Common variant at 16p11.2 conferring risk of psychosis. <i>Molecular Psychiatry</i> , <b>2014</b> , 19, 108-14	15.1	67
194	A study of chromosome 4p markers and dopamine D5 receptor gene in schizophrenia and bipolar disorder. <i>Molecular Psychiatry</i> , <b>1998</b> , 3, 310-20	15.1	67
193	Lack of association between a polymorphism in the promoter region of the dopamine-2 receptor gene and clozapine response. <i>Pharmacogenetics and Genomics</i> , <b>1998</b> , 8, 481-4		67

192	Sensorimotor gating is associated with CHRNA3 polymorphisms in schizophrenia and healthy volunteers. <i>Neuropsychopharmacology</i> , <b>2010</b> , 35, 1429-39	8.7	65
191	Dopaminergic and brain-derived neurotrophic factor signalling in inbred mice exposed to a restricted feeding schedule. <i>Genes, Brain and Behavior</i> , <b>2008</b> , 7, 552-9	3.6	65
190	Difference in susceptibility to activity-based anorexia in two inbred strains of mice. <i>European Neuropsychopharmacology</i> , <b>2007</b> , 17, 199-205	1.2	65
189	Identification of increased genetic risk scores for schizophrenia in treatment-resistant patients. <i>Molecular Psychiatry</i> , <b>2015</b> , 20, 150-1	15.1	63
188	Association study of nonsynonymous single nucleotide polymorphisms in schizophrenia. <i>Biological Psychiatry</i> , <b>2012</b> , 71, 169-77	7.9	63
187	Non-B right-handed DNA conformations of homopurine.homopyrimidine sequences in the murine immunoglobulin C alpha switch region. <i>Journal of Biological Chemistry</i> , <b>1988</b> , 263, 7397-405	5.4	63
186	Site-specific intercalation at the triplex-duplex junction induces a conformational change which is detectable by hypersensitivity to diethylpyrocarbonate. <i>Nucleic Acids Research</i> , <b>1991</b> , 19, 4219-24	20.1	62
185	Analysis of clozapine response and polymorphisms of the dopamine D4 receptor gene (DRD4) in schizophrenic patients. <i>American Journal of Medical Genetics Part A</i> , <b>1995</b> , 60, 541-5		61
184	Genetic Markers of Human Evolution Are Enriched in Schizophrenia. <i>Biological Psychiatry</i> , <b>2016</b> , 80, 284-292	7.9	60
183	Impairment of inhibitory control processing related to acute psychotomimetic effects of cannabis. <i>European Neuropsychopharmacology</i> , <b>2015</b> , 25, 26-37	1.2	60
182	Association study of CHRFAM7A copy number and 2 bp deletion polymorphisms with schizophrenia and bipolar affective disorder. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2006</b> , 141B, 571-5	3.5	60
181	Cytochrome P4502D6 genotype does not determine response to clozapine. <i>British Journal of Clinical Pharmacology</i> , <b>1995</b> , 39, 417-20	3.8	60
180	The effects of neuregulin1 on brain function in controls and patients with schizophrenia and bipolar disorder. <i>NeuroImage</i> , <b>2008</b> , 42, 817-26	7.9	59
179	No association between (AAT)n repeats in the cannabinoid receptor gene (CNR1) and heroin abuse in a Chinese population. <i>Molecular Psychiatry</i> , <b>2000</b> , 5, 128-30	15.1	57
178	Synthesis, molecular modeling, DNA binding, and antitumor properties of some substituted amidoanthraquinones. <i>Journal of Medicinal Chemistry</i> , <b>1988</b> , 31, 847-57	8.3	57
177	Opposite effects of catechol-O-methyltransferase Val158Met on cortical function in healthy subjects and patients with schizophrenia. <i>Biological Psychiatry</i> , <b>2009</b> , 65, 473-80	7.9	56
176	Brain-derived neurotrophic factor (BDNF) and set-shifting in currently ill and recovered anorexia nervosa (AN) patients. <i>Psychological Medicine</i> , <b>2009</b> , 39, 1029-35	6.9	55
175	An examination of decision making in bulimia nervosa. <i>Journal of Clinical and Experimental Neuropsychology</i> , <b>2009</b> , 31, 455-61	2.1	55

174	Consensus paper of the WFSBP Task Force on Genetics: Genetics, epigenetics and gene expression markers of major depressive disorder and antidepressant response. <i>World Journal of Biological Psychiatry</i> , <b>2017</b> , 18, 5-28	3.8	54
173	Gene-environment interaction in anorexia nervosa: relevance of non-shared environment and the serotonin transporter gene. <i>Molecular Psychiatry</i> , <b>2011</b> , 16, 590-2	15.1	52
172	5-HT2A receptor and bipolar affective disorder: association studies in affected patients. <i>Neuroscience Letters</i> , <b>1997</b> , 224, 95-8	3.3	52
171	Identifying potential risk haplotypes for schizophrenia at the DTNBP1 locus in Han Chinese and Scottish populations. <i>Molecular Psychiatry</i> , <b>2005</b> , 10, 1037-44	15.1	52
170	An association study of the neuregulin 1 gene, bipolar affective disorder and psychosis. <i>Psychiatric Genetics</i> , <b>2009</b> , 19, 113-6	2.9	51
169	Reaction time of the Continuous Performance Test is an endophenotypic marker for schizophrenia: a study of first-episode neuroleptic-naive schizophrenia, their non-psychotic first-degree relatives and healthy population controls. <i>Schizophrenia Research</i> , <b>2007</b> , 89, 293-8	3.6	51
168	Failure to find linkage between a functional polymorphism in the dopamine D4 receptor gene and schizophrenia. <i>American Journal of Medical Genetics Part A</i> , <b>1994</b> , 54, 8-11		51
167	Catechol-O-methyltransferase (COMT) val158met genotype is associated with BOLD response as a function of task characteristic. <i>Neuropsychopharmacology</i> , <b>2008</b> , 33, 3046-57	8.7	50
166	Neurocognitive deficits in first-episode schizophrenic patients and their first-degree relatives. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2007</b> , 144B, 407-16	3.5	50
165	The impact of CACNA1C allelic variation on effective connectivity during emotional processing in bipolar disorder. <i>Molecular Psychiatry</i> , <b>2013</b> , 18, 526-7	15.1	49
164	The impact of the Val158Met catechol-O-methyltransferase genotype on neural correlates of sad facial affect processing in patients with bipolar disorder and their relatives. <i>Psychological Medicine</i> , <b>2011</b> , 41, 779-88	6.9	49
163	Interspecies trait genetics reveals association of Adcy8 with mouse avoidance behavior and a human mood disorder. <i>Biological Psychiatry</i> , <b>2009</b> , 66, 1123-30	7.9	49
162	Effect of disrupted-in-schizophrenia-1 on pre-frontal cortical function. <i>Molecular Psychiatry</i> , <b>2008</b> , 13, 915-7, 909	15.1	49
161	Behavioral, physiological, and molecular differences in response to dietary restriction in three inbred mouse strains. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , <b>2006</b> , 291, E574-81 <sup>6</sup>		49
160	Left-handed Z-DNA and intramolecular triplex formation at the site of an unequal sister chromatid exchange. <i>Journal of Biological Chemistry</i> , <b>1990</b> , 265, 1352-9	5.4	49
159	Genetic targeting of NRXN2 in mice unveils role in excitatory cortical synapse function and social behaviors. <i>Frontiers in Synaptic Neuroscience</i> , <b>2015</b> , 7, 3	3.5	47
158	Volume increases in putamen associated with positive symptom reduction in previously drug-naive schizophrenia after 6 weeks antipsychotic treatment. <i>Psychological Medicine</i> , <b>2012</b> , 42, 1475-83	6.9	47
157	Epilepsy-associated GRIN2A mutations reduce NMDA receptor trafficking and agonist potency - molecular profiling and functional rescue. <i>Scientific Reports</i> , <b>2017</b> , 7, 66	4.9	46



156	Assessment of white matter abnormalities in paranoid schizophrenia and bipolar mania patients. <i>Psychiatry Research - Neuroimaging</i> , <b>2011</b> , 194, 347-353	2.9	46
155	The association of white matter volume in psychotic disorders with genotypic variation in NRG1, MOG and CNP: a voxel-based analysis in affected individuals and their unaffected relatives. <i>Translational Psychiatry</i> , <b>2012</b> , 2, e167	8.6	46
154	Genome-wide common and rare variant analysis provides novel insights into clozapine-associated neutropenia. <i>Molecular Psychiatry</i> , <b>2017</b> , 22, 1502-1508	15.1	45
153	Ohnologs are overrepresented in pathogenic copy number mutations. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2014</b> , 111, 361-6	11.5	45
152	Association analysis between dopamine receptor genes and bipolar affective disorder. <i>Psychiatry Research</i> , <b>1999</b> , 86, 193-201	9.9	45
151	PACAP and PAC1 receptor in brain development and behavior. <i>Neuropeptides</i> , <b>2013</b> , 47, 421-30	3.3	44
150	Eating disorders and obesity: two sides of the same coin?. <i>Epidemiologia E Psichiatria Sociale</i> , <b>2009</b> , 18, 96-100		43
149	Association study of dysbindin gene with clinical and outcome measures in a representative cohort of Italian schizophrenic patients. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2007</b> , 144B, 647-59	3.5	43
148	Association of the 5-HT2c gene with susceptibility and minimum body mass index in anorexia nervosa. <i>NeuroReport</i> , <b>2003</b> , 14, 781-3	1.7	43
147	Two novel variants in the DOPA decarboxylase gene: association with bipolar affective disorder. <i>Molecular Psychiatry</i> , <b>1999</b> , 4, 545-51	15.1	43
146	Linkage of Wolfram syndrome to chromosome 4p16.1 and evidence for heterogeneity. <i>American Journal of Human Genetics</i> , <b>1996</b> , 59, 855-63	11	43
145	A large replication study and meta-analysis in European samples provides further support for association of AHI1 markers with schizophrenia. <i>Human Molecular Genetics</i> , <b>2010</b> , 19, 1379-86	5.6	42
144	Neurotrophic gene polymorphisms and response to psychological therapy. <i>Translational Psychiatry</i> , <b>2012</b> , 2, e108	8.6	42
143	Effect of length, supercoiling, and pH on intramolecular triplex formation. Multiple conformers at pur.pyr mirror repeats. <i>Journal of Biological Chemistry</i> , <b>1990</b> , 265, 10652-8	5.4	42
142	Apolipoprotein E: depressive illness, depressive symptoms, and Alzheimer's disease. <i>Biological Psychiatry</i> , <b>1998</b> , 43, 159-64	7.9	39
141	A quantitative association study between schizotypal traits and COMT, PRODH and BDNF genes in a healthy Chinese population. <i>Psychiatry Research</i> , <b>2007</b> , 153, 7-15	9.9	39
140	Personality disorders and personality dimensions in anorexia nervosa. <i>Journal of Personality Disorders</i> , <b>2003</b> , 17, 73-85	2.6	39
139	Expression-based drug screening of neural progenitor cells from individuals with schizophrenia. <i>Nature Communications</i> , <b>2018</b> , 9, 4412	17.4	39

138	The serotonin transporter and clozapine response. <i>Molecular Psychiatry</i> , <b>2000</b> , 5, 124-5	15.1	38
137	TCF4 (e2-2; ITF2): a schizophrenia-associated gene with pleiotropic effects on human disease. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2013</b> , 162B, 1-16	3.5	37
136	Neuregulin-1 and the P300 waveform--a preliminary association study using a psychosis endophenotype. <i>Schizophrenia Research</i> , <b>2008</b> , 103, 178-85	3.6	37
135	Case-control and combined family trios analysis of three polymorphisms in the ghrelin gene in European patients with anorexia and bulimia nervosa. <i>Psychiatric Genetics</i> , <b>2006</b> , 16, 51-2	2.9	37
134	Combined family trio and case-control analysis of the COMT Val158Met polymorphism in European patients with anorexia nervosa. <i>American Journal of Medical Genetics Part A</i> , <b>2004</b> , 124B, 68-72		37
133	Molecular models for the interaction of the anti-tumour drug nogalamycin with DNA. <i>Biochemical Pharmacology</i> , <b>1984</b> , 33, 2877-80	6	37
132	StemBANCC: Governing Access to Material and Data in a Large Stem Cell Research Consortium. <i>Stem Cell Reviews and Reports</i> , <b>2015</b> , 11, 681-7	6.4	36
131	A genome-wide association analysis of a broad psychosis phenotype identifies three loci for further investigation. <i>Biological Psychiatry</i> , <b>2014</b> , 75, 386-97	7.9	36
130	Convergent lines of evidence support CAMKK2 as a schizophrenia susceptibility gene. <i>Molecular Psychiatry</i> , <b>2014</b> , 19, 774-83	15.1	36
129	Association analysis of the RGS4 gene in Han Chinese and Scottish populations with schizophrenia. <i>Genes, Brain and Behavior</i> , <b>2005</b> , 4, 444-8	3.6	36
128	Paraoxonase in Persian Gulf War veterans. <i>Journal of Occupational and Environmental Medicine</i> , <b>2003</b> , 45, 668-75	2	35
127	Clozapine pharmacokinetics and pharmacodynamics studied with Cyp1A2-null mice. <i>Journal of Psychopharmacology</i> , <b>2000</b> , 14, 353-9	4.6	35
126	Chromosome 22 markers demonstrate transmission disequilibrium with schizophrenia. <i>Psychiatric Genetics</i> , <b>1995</b> , 5, 127-30	2.9	35
125	Mapping genomic loci implicates genes and synaptic biology in schizophrenia.. <i>Nature</i> , <b>2022</b> ,	50.4	35
124	Protein kinase B (AKT1) genotype mediates sensitivity to cannabis-induced impairments in psychomotor control. <i>Psychological Medicine</i> , <b>2014</b> , 44, 3315-28	6.9	34
123	Altered effect of dopamine transporter 3'UTR VNTR genotype on prefrontal and striatal function in schizophrenia. <i>Archives of General Psychiatry</i> , <b>2009</b> , 66, 1162-72		34
122	Association analysis of polymorphisms in the DRD4 gene and heroin abuse in Chinese subjects. <i>American Journal of Medical Genetics Part A</i> , <b>2000</b> , 96, 616-21		34
121	Case-control, haplotype relative risk and transmission disequilibrium analysis of a dopamine D2 receptor functional promoter polymorphism in schizophrenia. <i>Schizophrenia Research</i> , <b>1998</b> , 32, 87-92	3.6	34

120	Genetics of schizophrenia: A consensus paper of the WFSBP Task Force on Genetics. <i>World Journal of Biological Psychiatry</i> , <b>2017</b> , 18, 492-505	3.8	33
119	Genetic overlap between episodic memory deficits and schizophrenia: results from the Maudsley Twin Study. <i>Psychological Medicine</i> , <b>2011</b> , 41, 521-32	6.9	33
118	Abnormalities in connectivity of white-matter tracts in patients with familial and non-familial schizophrenia. <i>Psychological Medicine</i> , <b>2011</b> , 41, 1691-700	6.9	33
117	Systematic screening of the 14-3-3 eta ( $\eta$ ) chain gene for polymorphic variants and case-control analysis in schizophrenia. <i>American Journal of Medical Genetics Part A</i> , <b>2000</b> , 96, 736-743		33
116	Transmission disequilibrium analysis of HLA class II DRB1, DQA1, DQB1 and DPB1 polymorphisms in schizophrenia using family trios from a Han Chinese population. <i>Schizophrenia Research</i> , <b>2001</b> , 49, 73-8	3.6	33
115	Overlapping clusters of gray matter deficits in paranoid schizophrenia and psychotic bipolar mania with family history. <i>Neuroscience Letters</i> , <b>2011</b> , 489, 94-8	3.3	32
114	Analysis of CAG/CTG repeat size in Chinese subjects with schizophrenia and bipolar affective disorder using the repeat expansion detection method. <i>Biological Psychiatry</i> , <b>1998</b> , 44, 1160-5	7.9	32
113	A computational and experimental study of the bending induced at a double-triple helix junction. <i>Biophysical Chemistry</i> , <b>1992</b> , 45, 143-52	3.5	32
112	Microdeletions of ELP4 Are Associated with Language Impairment, Autism Spectrum Disorder, and Mental Retardation. <i>Human Mutation</i> , <b>2015</b> , 36, 842-50	4.7	31
111	Advanced paternal age increases the risk of schizophrenia and obsessive-compulsive disorder in a Chinese Han population. <i>Psychiatry Research</i> , <b>2012</b> , 198, 353-9	9.9	31
110	The genetics of eating disorders. <i>Current Topics in Behavioral Neurosciences</i> , <b>2011</b> , 6, 157-75	3.4	31
109	Identification of novel polymorphisms in the 5' flanking region of CYP1A2, characterization of interethnic variability, and investigation of their functional significance. <i>Pharmacogenetics and Genomics</i> , <b>2000</b> , 10, 695-704		31
108	Association between clozapine response and allelic variation in the 5-HT <sub>2C</sub> receptor gene. <i>NeuroReport</i> , <b>1995</b> , 7, 169-72	1.7	31
107	Using ancestry-informative markers to identify fine structure across 15 populations of European origin. <i>European Journal of Human Genetics</i> , <b>2014</b> , 22, 1190-200	5.3	30
106	Possible parent-of-origin effect of Dopa decarboxylase in susceptibility to bipolar affective disorder <b>2003</b> , 117B, 18-22		30
105	Association analysis of polymorphisms in the $\beta$ opioid gene and heroin abuse in Chinese subjects. <i>Addiction Biology</i> , <b>2000</b> , 5, 181-6	4.6	30
104	Transmission disequilibrium analysis of a triplet repeat within the hKCa3 gene using family trios with schizophrenia. <i>Biochemical and Biophysical Research Communications</i> , <b>1998</b> , 251, 662-5	3.4	30
103	Advances in multidisciplinary and cross-species approaches to examine the neurobiology of psychiatric disorders. <i>European Neuropsychopharmacology</i> , <b>2011</b> , 21, 532-44	1.2	29

102	Voxel-based morphometric analysis on the volume of gray matter in bipolar I disorder. <i>Psychiatry Research - Neuroimaging</i> , <b>2011</b> , 191, 92-7	2.9	29
101	Prenatal exposure to maternal obesity leads to hyperactivity in offspring. <i>Molecular Psychiatry</i> , <b>2012</b> , 17, 1159-60	15.1	29
100	Systematic screening of the LDL-PLA2 gene for polymorphic variants and case-control analysis in schizophrenia. <i>Biochemical and Biophysical Research Communications</i> , <b>1997</b> , 241, 630-5	3.4	28
99	Novel polymorphisms in the somatostatin receptor 5 (SSTR5) gene associated with bipolar affective disorder. <i>Molecular Psychiatry</i> , <b>2002</b> , 7, 745-54	15.1	28
98	NRXN1 deletions identified by array comparative genome hybridisation in a clinical case series - further understanding of the relevance of NRXN1 to neurodevelopmental disorders. <i>Journal of Molecular Psychiatry</i> , <b>2013</b> , 1, 4		27
97	Serum glutamine, set-shifting ability and anorexia nervosa. <i>Annals of General Psychiatry</i> , <b>2010</b> , 9, 29	3.4	27
96	Failure to respond to treatment with typical antipsychotics is not associated with CYP2D6 ultrarapid hydroxylation. <i>British Journal of Clinical Pharmacology</i> , <b>1999</b> , 48, 388-94	3.8	27
95	A genome-wide association study for quantitative traits in schizophrenia in China. <i>Genes, Brain and Behavior</i> , <b>2011</b> , 10, 734-9	3.6	26
94	Evidence of association of KIBRA genotype with episodic memory in families of psychotic patients and controls. <i>Journal of Psychiatric Research</i> , <b>2010</b> , 44, 795-8	5.2	26
93	No association between polymorphisms of methylenetetrahydrofolate reductase gene and schizophrenia in both Chinese and Scottish populations. <i>Molecular Psychiatry</i> , <b>2004</b> , 9, 1063-5	15.1	26
92	Genetic variation in the 5-HT5A receptor gene in patients with bipolar disorder and major depression. <i>Neuroscience Letters</i> , <b>2001</b> , 303, 111-4	3.3	26
91	Expression analysis in a rat psychosis model identifies novel candidate genes validated in a large case-control sample of schizophrenia. <i>Translational Psychiatry</i> , <b>2015</b> , 5, e656	8.6	25
90	An epidemiological survey of alcohol use disorders in a Tibetan population. <i>Psychiatry Research</i> , <b>2008</b> , 159, 56-66	9.9	25
89	Pre-morbid psychiatric morbidity, comorbidity and personality in patients with anorexia nervosa compared to their healthy sisters. <i>European Eating Disorders Review</i> , <b>2002</b> , 10, 255-270	5.3	25
88	Analysis of a structural polymorphism in the 5-HT2A receptor and clinical response to clozapine. <i>Neuroscience Letters</i> , <b>1996</b> , 217, 177-8	3.3	24
87	Extensive brain structural network abnormality in first-episode treatment-naive patients with schizophrenia: morphometrical and covariation study. <i>Psychological Medicine</i> , <b>2014</b> , 44, 2489-501	6.9	23
86	Premorbid tobacco smoking is associated with later age at onset in schizophrenia. <i>Psychiatry Research</i> , <b>2010</b> , 178, 461-6	9.9	23
85	Differential effects of DAAO on regional activation and functional connectivity in schizophrenia, bipolar disorder and controls. <i>NeuroImage</i> , <b>2011</b> , 56, 2283-91	7.9	22

84	Chromosomal mapping of excessive physical activity in mice in response to a restricted feeding schedule. <i>European Neuropsychopharmacology</i> , <b>2010</b> , 20, 317-26	1.2	22
83	Is neuregulin 1 involved in determining cerebral volumes in schizophrenia? Preliminary results showing a decrease in superior temporal gyrus volume. <i>Neuropsychobiology</i> , <b>2012</b> , 65, 119-25	4	22
82	Correlation and familial aggregation of dimensions of psychosis in affected sibling pairs from China. <i>British Journal of Psychiatry</i> , <b>2008</b> , 193, 305-10	5.4	22
81	Mutation screening and association analysis of the parkin gene in Parkinson's disease patients from South-West China. <i>European Neurology</i> , <b>2003</b> , 49, 85-9	2.1	22
80	Genome-Wide Association of Heroin Dependence in Han Chinese. <i>PLoS ONE</i> , <b>2016</b> , 11, e0167388	3.7	22
79	Genetic modulation of neural response during working memory in healthy individuals: interaction of glucocorticoid receptor and dopaminergic genes. <i>Molecular Psychiatry</i> , <b>2013</b> , 18, 174-82	15.1	21
78	Identification of increased genetic risk scores for schizophrenia in treatment-resistant patients. <i>Molecular Psychiatry</i> , <b>2015</b> , 20, 913	15.1	20
77	High-throughput DNA methylation analysis in anorexia nervosa confirms TNXB hypermethylation. <i>World Journal of Biological Psychiatry</i> , <b>2018</b> , 19, 187-199	3.8	20
76	Interspecies comparisons of functional genetic variations and their implications in neuropsychiatry. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2009</b> , 150B, 309-17	3.5	20
75	A COMT gene haplotype associated with methamphetamine abuse. <i>Pharmacogenetics and Genomics</i> , <b>2011</b> , 21, 731-40	1.9	20
74	Common schizophrenia alleles are enriched in mutation-intolerant genes and maintained by background selection		20
73	Microduplications at the pseudoautosomal SHOX locus in autism spectrum disorders and related neurodevelopmental conditions. <i>Journal of Medical Genetics</i> , <b>2016</b> , 53, 536-47	5.8	19
72	Gene-set analysis based on the pharmacological profiles of drugs to identify repurposing opportunities in schizophrenia. <i>Journal of Psychopharmacology</i> , <b>2016</b> , 30, 826-30	4.6	19
71	Quantitative promoter DNA methylation analysis of four candidate genes in anorexia nervosa: a pilot study. <i>Journal of Psychiatric Research</i> , <b>2013</b> , 47, 280-2	5.2	19
70	Factor structures of the neurocognitive assessments and familial analysis in first-episode schizophrenia patients, their relatives and controls. <i>Australian and New Zealand Journal of Psychiatry</i> , <b>2010</b> , 44, 109-19	2.6	19
69	Autosome search for schizophrenia susceptibility genes in multiply affected families. <i>Molecular Psychiatry</i> , <b>1999</b> , 4, 353-9	15.1	19
68	Aberrant spontaneous neural activity and correlation with evoked-brain potentials in first-episode, treatment-naïve patients with deficit and non-deficit schizophrenia. <i>Psychiatry Research - Neuroimaging</i> , <b>2017</b> , 261, 9-19	2.9	18
67	Random or stochastic monoallelic expressed genes are enriched for neurodevelopmental disorder candidate genes. <i>PLoS ONE</i> , <b>2013</b> , 8, e85093	3.7	18

66	No association of Disrupted-in-Schizophrenia-1 variation with prefrontal function in patients with schizophrenia and bipolar disorder. <i>Genes, Brain and Behavior</i> , <b>2011</b> , 10, 276-85	3.6	18
65	Normative data on a battery of neuropsychological tests in the Han Chinese population. <i>Journal of Neuropsychology</i> , <b>2011</b> , 5, 126-42	2.6	18
64	Association of Neuregulin 1 rs3924999 genotype with antisaccades and smooth pursuit eye movements. <i>Genes, Brain and Behavior</i> , <b>2010</b> , 9, 621-7	3.6	18
63	Dissecting the many genetic faces of schizophrenia. <i>Epidemiologia E Psichiatria Sociale</i> , <b>2009</b> , 18, 91-95		18
62	Accuracy and sensitivity of DNA pooling with microsatellite repeats using capillary electrophoresis. <i>Molecular and Cellular Probes</i> , <b>1999</b> , 13, 359-65	3.3	18
61	A genome-wide association study in individuals of African ancestry reveals the importance of the Duffy-null genotype in the assessment of clozapine-related neutropenia. <i>Molecular Psychiatry</i> , <b>2019</b> , 24, 328-337	15.1	18
60	A dopamine D4 receptor exon 3 VNTR allele protecting against migraine without aura. <i>Annals of Neurology</i> , <b>2007</b> , 61, 574-8	9.4	17
59	A Comparison of Ten Polygenic Score Methods for Psychiatric Disorders Applied Across Multiple Cohorts. <i>Biological Psychiatry</i> , <b>2021</b> , 90, 611-620	7.9	17
58	Establishing the characteristics of an effective pharmacogenetic test for clozapine-induced agranulocytosis. <i>Pharmacogenomics Journal</i> , <b>2015</b> , 15, 461-6	3.5	16
57	Translational neuroscience of schizophrenia: seeking a meeting of minds between mouse and man. <i>Science Translational Medicine</i> , <b>2011</b> , 3, 102mr3	17.5	16
56	A linkage study of schizophrenia with DNA markers from chromosome 8p21-p22 in 25 multiplex families. <i>Schizophrenia Research</i> , <b>1996</b> , 22, 61-8	3.6	16
55	Linkage studies in bipolar affective disorder with markers on chromosome 21. <i>Journal of Affective Disorders</i> , <b>1996</b> , 41, 217-21	6.6	16
54	<sup>123</sup> I-iododexetimide preferentially binds to the muscarinic receptor subtype M1 in vivo. <i>Journal of Nuclear Medicine</i> , <b>2015</b> , 56, 317-22	8.9	15
53	Using genetic findings in autism for the development of new pharmaceutical compounds. <i>Psychopharmacology</i> , <b>2014</b> , 231, 1063-78	4.7	15
52	Effect of D-amino acid oxidase activator (DAOA; G72) on brain function during verbal fluency. <i>Human Brain Mapping</i> , <b>2012</b> , 33, 143-53	5.9	15
51	Replication study and meta-analysis in European samples supports association of the 3p21.1 locus with bipolar disorder. <i>Biological Psychiatry</i> , <b>2012</b> , 72, 645-50	7.9	15
50	Sensorimotor gating and D2 receptor signalling: evidence from a molecular genetic approach. <i>International Journal of Neuropsychopharmacology</i> , <b>2012</b> , 15, 1427-40	5.8	14
49	Bipolar 1 disorder is not associated with the RGS4, PRODH, COMT and GRK3 genes. <i>Psychiatric Genetics</i> , <b>2006</b> , 16, 229-30	2.9	14

48	Fronto-parietal white matter microstructural deficits are linked to performance IQ in a first-episode schizophrenia Han Chinese sample. <i>Psychological Medicine</i> , <b>2013</b> , 43, 2047-56	6.9	13
47	Association analysis of STX1A gene variants in common forms of migraine. <i>Cephalalgia</i> , <b>2012</b> , 32, 203-126.1		12
46	Analysis of microsatellite markers at the UCP2/UCP3 locus on chromosome 11q13 in anorexia nervosa. <i>Molecular Psychiatry</i> , <b>2002</b> , 7, 276-7	15.1	12
45	Linkage analysis of the fragile X gene FMR-1 and schizophrenia: no evidence for linkage but report of a family with schizophrenia and an unstable triplet repeat. <i>Psychiatric Genetics</i> , <b>1996</b> , 6, 81-6	2.9	12
44	PRODH gene is associated with executive function in schizophrenic families. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2008</b> , 147B, 654-7	3.5	11
43	Linkage analysis between bipolar affective disorder and markers on chromosome X. <i>Psychiatric Genetics</i> , <b>1998</b> , 8, 183-6	2.9	11
42	Pharmacogenetics in psychosis. <i>Drug News and Perspectives</i> , <b>2003</b> , 16, 159-65		11
41	Sex-Dependent Shared and Nonshared Genetic Architecture Across Mood and Psychotic Disorders. <i>Biological Psychiatry</i> , <b>2022</b> , 91, 102-117	7.9	11
40	Detection of metabolites in the white matter of frontal lobes and hippocampus with proton in first-episode treatment-naïve schizophrenia patients. <i>Microbial Biotechnology</i> , <b>2012</b> , 6, 166-75	3.3	10
39	CNTF and psychiatric disorders. <i>Nature Genetics</i> , <b>1996</b> , 13, 143-4	36.3	10
38	Advances in the genetics of schizophrenia: toward a network and pathway view for drug discovery. <i>Annals of the New York Academy of Sciences</i> , <b>2016</b> , 1366, 61-75	6.5	10
37	Cross-species genetics converge to TLL2 for mouse avoidance behavior and human bipolar disorder. <i>Genes, Brain and Behavior</i> , <b>2013</b> , 12, 653-7	3.6	9
36	The androgen receptor gene polyglycine repeat polymorphism is associated with memory performance in healthy Chinese individuals. <i>Psychoneuroendocrinology</i> , <b>2009</b> , 34, 947-52	5	9
35	Schizophrenia: the polygene princess and the pea. <i>Psychological Medicine</i> , <b>2008</b> , 38, 1687-91; discussion 1818-20	6.9	9
34	Family-based analysis of serotonin transporter gene polymorphisms in migraine with and without aura. <i>Cephalalgia</i> , <b>2007</b> , 27, 773-80	6.1	9
33	Affected sibling pair linkage analysis of qualitative and quantitative traits for schizophrenia on chromosome 22 in a Chinese population. <i>American Journal of Medical Genetics Part A</i> , <b>2001</b> , 105, 321-7		9
32	Perception of autonomy and connectedness prior to the onset of anorexia nervosa and bulimia nervosa. <i>Zeitschrift für Kinder- Und Jugendpsychiatrie Und Psychotherapie</i> , <b>2012</b> , 40, 61-8	1.8	9
31	Clinical utility gene card for: 16p13.11 microdeletion syndrome. <i>European Journal of Human Genetics</i> , <b>2014</b> , 22,	5.3	8

30	Clinical utility gene card for: 15q13.3 microdeletion syndrome. <i>European Journal of Human Genetics</i> , <b>2014</b> , 22,	5.3	8
29	Opening Pandora's box in the UK: a hypothetical pharmacogenetic test for clozapine. <i>Pharmacogenomics</i> , <b>2013</b> , 14, 1907-14	2.6	8
28	No association between a promoter polymorphism in the noradrenaline transporter gene and anorexia nervosa. <i>Psychiatric Genetics</i> , <b>2007</b> , 17, 247-8	2.9	8
27	Understanding the genetic predisposition to anorexia nervosa. <i>European Eating Disorders Review</i> , <b>1999</b> , 7, 96-102	5.3	8
26	The DISC1 Ser704Cys polymorphism is associated with prefrontal function in healthy individuals. <i>Molecular Psychiatry</i> , <b>2008</b> , 13, 909-909	15.1	6
25	The functional MMP-9 microsatellite marker is not associated with episodic memory in humans. <i>Psychiatric Genetics</i> , <b>2008</b> , 18, 252	2.9	6
24	Family-based association analysis of functional VNTR polymorphisms in the dopamine transporter gene in migraine with and without aura. <i>Journal of Neural Transmission</i> , <b>2008</b> , 115, 91-5	4.3	6
23	The genetics of cannabis involvement in humans: a genetic epidemiological perspective. <i>Addiction</i> , <b>2006</b> , 101, 780-1; author reply 781-2	4.6	6
22	The 5-HT2A rs438G/A polymorphism in anorexia nervosa: a combined analysis of 316 trios from six European centres		6
21	Interaction among genes influencing ethanol metabolism and sex is association with alcohol use disorders in a Tibet population. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2010</b> , 153B, 561-569	3.5	5
20	Full-length transcript sequencing of human and mouse cerebral cortex identifies widespread isoform diversity and alternative splicing. <i>Cell Reports</i> , <b>2021</b> , 37, 110022	10.6	5
19	BBGRE: brain and body genetic resource exchange. <i>Database: the Journal of Biological Databases and Curation</i> , <b>2013</b> , 2013, bat067	5	4
18	Gender-specific interactions between alcohol metabolism genes and severity of quantitative alcohol-related-traits in a Tibetan population. <i>Neuroscience Letters</i> , <b>2011</b> , 495, 22-5	3.3	4
17	No association between bipolar disorder risk polymorphisms in ANK3 and CACNA1C and common migraine. <i>Headache</i> , <b>2011</b> , 51, 796-803	4.2	4
16	Genetic mapping of 14 short tandem repeat polymorphisms on human chromosome 22. <i>Human Genetics</i> , <b>1994</b> , 93, 688-90	6.3	4
15	Advances in the genetics of schizophrenia: will high-risk copy number variants be useful in clinical genetics or diagnostics?. <i>F1000 Medicine Reports</i> , <b>2009</b> , 1,		3
14	The Spectrum of Eating Disorders in Humans <b>2001</b> , 19-49		3
13	HLA-DQB1 6672G>C (rs113332494) is associated with clozapine-induced neutropenia and agranulocytosis in individuals of European ancestry. <i>Translational Psychiatry</i> , <b>2021</b> , 11, 214	8.6	3



12	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> , <b>2019</b> , 180, 223-231	3.5	2
11	Association between the COMT gene and neurological abnormalities and poorer executive function in psychosis. <i>Psychiatry Research</i> , <b>2015</b> , 230, 742-3	9.9	2
10	No evidence of linkage disequilibrium between a CAG repeat in the SCA1 gene and schizophrenia in Caucasian and Chinese schizophrenic subjects. <i>Psychiatric Genetics</i> , <b>1999</b> , 9, 123-7	2.9	2
9	Genetic underpinnings of sociability in the general population. <i>Neuropsychopharmacology</i> , <b>2021</b> , 46, 1628-1634	3.1	2
8	Autosome search for schizophrenia susceptibility genes in multiply affected families		1
7	Avances en enfoques multidisciplinarios y en diversas especies para el examen de la neurobiología de los trastornos psiquiátricos. <i>Psiquiatría Biológica</i> , <b>2012</b> , 19, 9-20	0.2	0
6	Highly pleiotropic variants of human traits are enriched in genomic regions with strong background selection. <i>Human Genetics</i> , <b>2021</b> , 140, 1343-1351	6.3	0
5	[O2306]: ASSESSING TREM2 FUNCTION IN ALZHEIMER'S DISEASE WITH RNA-SEQ <b>2017</b> , 13, P590		
4	Clozapine response and genetic variation in neurotransmitter receptor targets <b>2002</b> , 217-244		
3	Pharmacogenetic methods in schizophrenia. <i>International Review of Psychiatry</i> , <b>2001</b> , 13, 47-49	3.6	
2	Assignment of the neuronal cochaperone, HSJ1, to human chromosome bands 2q32-->q34 between D2S295 and D2S339 by in situ hybridization and somatic cell and radiation hybrids. <i>Cytogenetic and Genome Research</i> , <b>1999</b> , 86, 62-3	1.9	
1	Functional polymorphism of CYP2E1 gene and alcohol use disorders in a Tibetan population. <i>Journal of Central South University (Medical Sciences)</i> , <b>2008</b> , 33, 284-92	0.4	