Edwin H Cook

List of Publications by Year
in descending order

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196 papers	36,726 citations	13332 70 h-index	4305 179 g-index
203 all docs	203 docs citations	203 times ranked	34854 citing authors

#	Article	IF	Citations
1	Case Report: Association of Comorbid Psychiatric Disorders and Sigmoid Prolapse with de novo POGZ Mutation. Journal of Autism and Developmental Disorders, 2022, 52, 1408-1411.	1.7	2
2	Recurrent de novo missense variants in GNB2 can cause syndromic intellectual disability. Journal of Medical Genetics, 2022, 59, 511-516.	1.5	4
3	Maternal Duplication 15q11-13 Syndrome with Autism Spectrum Disorder: Mood Stabilization by Carbamazepine. Journal of Child and Adolescent Psychopharmacology, 2022, 32, 122-126.	0.7	3
4	Elevated Polygenic Burden for Autism Spectrum Disorder Is Associated With the Broad Autism Phenotype in Mothers of Individuals With Autism Spectrum Disorder. Biological Psychiatry, 2021, 89, 476-485.	0.7	32
5	Contextualizing genetic risk score for disease screening and rare variant discovery. Nature Communications, 2021, 12, 4418.	5.8	11
6	Behavioral characterization of dup15q syndrome: Toward meaningful endpoints for clinical trials. American Journal of Medical Genetics, Part A, 2020, 182, 71-84.	0.7	21
7	Clinical and neurocognitive issues associated with Boschâ€Boonstraâ€Schaaf optic atrophy syndrome: A case study. American Journal of Medical Genetics, Part A, 2020, 182, 213-218.	0.7	11
8	Properties of beta oscillations in Dup15q syndrome. Journal of Neurodevelopmental Disorders, 2020, 12, 22.	1.5	7
9	Large-Scale Exome Sequencing Study Implicates Both Developmental and Functional Changes in the Neurobiology of Autism. Cell, 2020, 180, 568-584.e23.	13.5	1,422
10	A framework for an evidence-based gene list relevant to autism spectrum disorder. Nature Reviews Genetics, 2020, 21, 367-376.	7.7	83
11	Lovastatin Treatment of a Patient with a De Novo SYNGAP1 Protein Truncating Variant. Journal of Child and Adolescent Psychopharmacology, 2019, 29, 321-322.	0.7	5
12	A Bayesian framework that integrates multi-omics data and gene networks predicts risk genes from schizophrenia GWAS data. Nature Neuroscience, 2019, 22, 691-699.	7.1	118
13	Whole Blood Serotonin Levels and Platelet 5-HT2A Binding in Autism Spectrum Disorder. Journal of Autism and Developmental Disorders, 2019, 49, 2417-2425.	1.7	10
14	Vocabulary comprehension in adults with fragile X syndrome (FXS). Journal of Neurodevelopmental Disorders, 2019, 11, 25.	1.5	5
15	Familiality of behavioral flexibility and response inhibition deficits in autism spectrum disorder (ASD). Molecular Autism, 2019, 10, 47.	2.6	20
16	Phenotypic association of 15q11.2 CNVs of the region of breakpoints 1â€“2 (BP1â€“BP2) in a large cohort of samples referred for genetic diagnosis. Journal of Human Genetics, 2019, 64, 253-255.	1.1	9
17	Maternal Serotonin Levels Are Associated With Cognitive Ability and Core Symptoms in Autism Spectrum Disorder. Journal of the American Academy of Child and Adolescent Psychiatry, 2018, 57, 867-875.	0.3	24
18	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	6.0	1,085

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19	Pharmacogenomics of autism spectrum disorder. Pharmacogenomics, 2017, 18, 403-414.	0.6	17
20	Whole genome sequencing resource identifies 18 new candidate genes for autism spectrum disorder. Nature Neuroscience, 2017, 20, 602-611.	7.1	691
21	De novo unbalanced translocation (4p duplication/8p deletion) in a patient with autism, OCD, and overgrowth syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 1656-1662.	0.7	9
22	Is there sexual dimorphism of hyperserotonemia in autism spectrum disorder?. Autism Research, 2017, 10, 1417-1423.	2.1	24
23	Does MAOA increase susceptibility to prenatal stress in young children?. Neurotoxicology and Teratology, 2017, 61, 82-91.	1.2	7
24	Leveraging blood serotonin as an endophenotype to identify de novo and rare variants involved in autism. Molecular Autism, 2017, 8, 14.	2.6	50
25	Arbaclofen in Children and Adolescents with Autism Spectrum Disorder: A Randomized, Controlled, Phase 2 Trial. Neuropsychopharmacology, 2017, 42, 1390-1398.	2.8	112
26	Variants in Adjacent Oxytocin/Vasopressin Gene Region and Associations with ASD Diagnosis and Other Autism Related Endophenotypes. Frontiers in Neuroscience, 2016, 10, 195.	1.4	21
27	ASD and Genetic Associations with Receptors for Oxytocin and Vasopressinâ€”AVPR1A, AVPR1B, and OXTR. Frontiers in Neuroscience, 2016, 10, 516.	1.4	38
28	Mortality in isodicentric chromosome 15 syndrome: The role of SUDEP. Epilepsy and Behavior, 2016, 61, 1-5.	0.9	24
29	Confirmation of the Factor Structure and Measurement Invariance of the Children's Scale of Hostility and Aggression: Reactive/Proactive in Clinic-Referred Children With and Without Autism Spectrum Disorder. Journal of Child and Adolescent Psychopharmacology, 2016, 26, 10-18.	0.7	12
30	Commentary on â€œPlatelet Studies in Autism Spectrum Disorder Patients and First-Degree Relativesâ€. Molecular Autism, 2016, 7, 20.	2.6	4
31	Constance E. Lieber, Theodore R. Stanley, and the Enduring Impact of Philanthropy on Psychiatry Research. Biological Psychiatry, 2016, 80, 84-86.	0.7	2
32	The impact of genotype calling errors on family-based studies. Scientific Reports, 2016, 6, 28323.	1.6	12
33	STAMS: STRING-assisted module search for genome wide association studies and application to autism. Bioinformatics, 2016, 32, 3815-3822.	1.8	17
34	Separating Family-Level and Direct Exposure Effects of Smoking During Pregnancy on Offspring Externalizing Symptoms: Bridging the Behavior Genetic and Behavior Teratologic Divide. Behavior Genetics, 2016, 46, 389-402.	1.4	40
35	Identification of a distinct developmental and behavioral profile in children with Dup15q syndrome. Journal of Neurodevelopmental Disorders, 2016, 8, 19.	1.5	47
36	CYP2A6Longitudinal Effects in Young Smokers. Nicotine and Tobacco Research, 2016, 18, 196-203.	1.4	7

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37	Significant neuronal soma volume deficit in the limbic system in subjects with 15q11.2-q13 duplications. Acta Neuropathologica Communications, 2015, 3, 63.	2.4	11
38	Escitalopram pharmacogenetics. Pharmacogenetics and Genomics, 2015, 25, 548-554.	0.7	22
39	Integrin Î²3 Haploinsufficiency Modulates Serotonin Transport and Antidepressant-Sensitive Behavior in Mice. Neuropsychopharmacology, 2015, 40, 2015-2024.	2.8	26
40	UGT1A and UGT2B Genetic Variation Alters Nicotine and Nitrosamine Glucuronidation in European and African American Smokers. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 94-104.	1.1	27
41	An ontology for Autism Spectrum Disorder (ASD) to infer ASD phenotypes from Autism Diagnostic Interview-Revised data. Journal of Biomedical Informatics, 2015, 56, 333-347.	2.5	14
42	Consensus Genotyper for Exome Sequencing (CGES): improving the quality of exome variant genotypes. Bioinformatics, 2015, 31, 187-193.	1.8	18
43	A haplotype-based framework for group-wise transmission/disequilibrium tests for rare variant association analysis. Bioinformatics, 2015, 31, 1452-1459.	1.8	14
44	Insights into Autism Spectrum Disorder Genomic Architecture and Biology from 71 Risk Loci. Neuron, 2015, 87, 1215-1233.	3.8	1,219
45	Pharmacogenetic Study of Serotonin Transporter and 5HT2A Genotypes in Autism. Journal of Child and Adolescent Psychopharmacology, 2015, 25, 467-474.	0.7	11
46	Preliminary evidence for the interaction of the oxytocin receptor gene (oxtr) and face processing in differentiating prenatal smoking patterns. Neuroscience Letters, 2015, 584, 259-264.	1.0	14
47	A Genome-wide Association Study of Autism Using the Simons Simplex Collection: Does Reducing Phenotypic Heterogeneity in Autism Increase Genetic Homogeneity?. Biological Psychiatry, 2015, 77, 775-784.	0.7	133
48	Cognitive Set Shifting Deficits and Their Relationship to Repetitive Behaviors in Autism Spectrum Disorder. Journal of Autism and Developmental Disorders, 2015, 45, 805-815.	1.7	95
49	Reducing Auditory Hypersensitivities in Autistic Spectrum Disorder: Preliminary Findings Evaluating the Listening Project Protocol. Frontiers in Pediatrics, 2014, 2, 80.	0.9	24
50	Environmental and State-Level Regulatory Factors Affect the Incidence of Autism and Intellectual Disability. PLoS Computational Biology, 2014, 10, e1003518.	1.5	48
51	Dopamine Transporter Genotype and Stimulant Dose-Response in Youth with Attention-Deficit/Hyperactivity Disorder. Journal of Child and Adolescent Psychopharmacology, 2014, 24, 238-244.	0.7	14
52	Self-injury in autism spectrum disorder: An effect of serotonin transporter gene promoter variants. Psychiatry Research, 2014, 220, 987-990.	1.7	11
53	A survey of seizures and current treatments in 15q duplication syndrome. Epilepsia, 2014, 55, 396-402.	2.6	80
54	Saccadic eye movement abnormalities in autism spectrum disorder indicate dysfunctions in cerebellum and brainstem. Molecular Autism, 2014, 5, 47.	2.6	131

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55	The Autism Simplex Collection: an international, expertly phenotyped autism sample for genetic and phenotypic analyses. Molecular Autism, 2014, 5, 34.	2.6	31
56	Convergence of Genes and Cellular Pathways Dysregulated in Autism Spectrum Disorders. American Journal of Human Genetics, 2014, 94, 677-694.	2.6	819
57	Synaptic, transcriptional and chromatin genes disrupted in autism. Nature, 2014, 515, 209-215.	13.7	2,254
58	A framework for the interpretation of de novo mutation in human disease. Nature Genetics, 2014, 46, 944-950.	9.4	943
59	Epigenetic Mechanisms in Autism Spectrum Disorder. International Review of Neurobiology, 2014, 115, 203-244.	0.9	41
60	Genetic factors affecting gene transcription and catalytic activity of UDP-glucuronosyltransferases in human liver. Human Molecular Genetics, 2014, 23, 5558-5569.	1.4	50
61	Expression of microRNAs and Other Small RNAs in Prefrontal Cortex in Schizophrenia, Bipolar Disorder and Depressed Subjects. PLoS ONE, 2014, 9, e86469.	1.1	166
62	Coâ€œoccurrence of autism, childhood psychosis, and intellectual disability associated with a de novo 3q29 microdeletion. American Journal of Medical Genetics, Part A, 2013, 161, 845-849.	0.7	48
63	Gene Ã— Environment effects of serotonin transporter, dopamine receptor D4, and monoamine oxidase A genes with contextual and parenting risk factors on symptoms of oppositional defiant disorder, anxiety, and depression in a community sample of 4-year-old children. Development and Psychopathology, 2013, 25, 555-575.	1.4	50
64	Parental Broader Autism Subphenotypes in <scp>ASD</scp> Affected Families: Relationship to Gender, Child's Symptoms, <scp>SSRI</scp> Treatment, and Platelet Serotonin. Autism Research, 2013, 6, 621-630.	2.1	16
65	Reduced behavioral flexibility in autism spectrum disorders.. Neuropsychology, 2013, 27, 152-160.	1.0	207
66	Clinicopathological Stratification of Idiopathic Autism and Autism with 15q11.2â€“q13 Duplications. , 2013, , 347-359.		0
67	Individual common variants exert weak effects on the risk for autism spectrum disorders. Human Molecular Genetics, 2012, 21, 4781-4792.	1.4	334
68	Differences Between the Pattern of Developmental Abnormalities in Autism Associated With Duplications 15q11.2-q13 and Idiopathic Autism. Journal of Neuropathology and Experimental Neurology, 2012, 71, 382-397.	0.9	78
69	Loci nominally associated with autism from genome-wide analysis show enrichment of brain expression quantitative trait loci but not lymphoblastoid cell line expression quantitative trait loci. Molecular Autism, 2012, 3, 3.	2.6	38
70	Examining Autism Spectrum Disorders by Biomarkers: Example From the Oxytocin and Serotonin Systems. Journal of the American Academy of Child and Adolescent Psychiatry, 2012, 51, 712-721.e1.	0.3	65
71	Common genetic variants, acting additively, are a major source of risk for autism. Molecular Autism, 2012, 3, 9.	2.6	357
72	Abnormal Intracellular Accumulation and Extracellular AÎ² Deposition in Idiopathic and Dup15q11.2-q13 Autism Spectrum Disorders. PLoS ONE, 2012, 7, e35414.	1.1	48

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73	Maternal depressive history, teen 5HTTLPR genotype, and the processing of emotional faces: Exploring mechanisms of risk. Behaviour Research and Therapy, 2011, 49, 80-84.	1.6	20
74	Mitochondrial small RNAs that are up-regulated in hippocampus during olfactory discrimination training in mice. Mitochondrion, 2011, 11, 994-995.	1.6	21
75	Multiple Recurrent De Novo CNVs, Including Duplications of the 7q11.23 Williams Syndrome Region, Are Strongly Associated with Autism. Neuron, 2011, 70, 863-885.	3.8	1,146
76	Family-based association testing of glutamate transporter genes in autism. Psychiatric Genetics, 2011, 21, 212-213.	0.6	17
77	A genotype resource for postmortem brain samples from the Autism Tissue Program. Autism Research, 2011, 4, 89-97.	2.1	23
78	De novo autosomal dominant mutation in SYNGAP1. Autism Research, 2011, 4, 155-156.	2.1	9
79	Repetitive behavior profiles: Consistency across autism spectrum disorder cohorts and divergence from Praderâ€“Willi syndrome. Journal of Neurodevelopmental Disorders, 2011, 3, 316-324.	1.5	22
80	A quantitative association study of SLC25A12 and restricted repetitive behavior traits in autism spectrum disorders. Molecular Autism, 2011, 2, 8.	2.6	25
81	Effect of dopamine transporter genotype on caudate volume in childhood ADHD and controls. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 28-35.	1.1	28
82	Parent-of-origin effects of the serotonin transporter gene associated with autism. , 2011, 156, 139-144.		41
83	Endogenous siRNAs and noncoding RNA-derived small RNAs are expressed in adult mouse hippocampus and are up-regulated in olfactory discrimination training. Rna, 2011, 17, 166-181.	1.6	59
84	Interactions between early parenting and a polymorphism of the child's dopamine transporter gene in predicting future child conduct disorder symptoms.. Journal of Abnormal Psychology, 2011, 120, 33-45.	2.0	59
85	Molecular Genetics of Childhood and Adolescent Onset Psychiatric Disorders. , 2011, , 1173-1186.		0
86	A pharmacogenetic study of escitalopram in autism spectrum disorders. Autism Research, 2010, 3, 1-7.	2.1	45
87	Reduction of increased repetitive selfâ€grooming in ASD mouse model by metabotropic 5 glutamate receptor antagonism; randomized controlled trial of early start denver model. Autism Research, 2010, 3, 40-42.	2.1	5
88	Decreased brain serotonin transporter binding in ASD; intranasal oxytocin administration: effects on computerized social discrimination task and eyeâ€tracking. Autism Research, 2010, 3, 94-95.	2.1	1
89	Clinical genetic microarray testing; ASD neuropathology. Autism Research, 2010, 3, 142-143.	2.1	0
90	Autism spectrum disorder: unbroken mirror neurons; rare copy number variants. Autism Research, 2010, 3, 196-197.	2.1	1

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91	Multiparameter Classification Approach to Structural Neuroimaging Data; Heterogeneity of 16p11.2 Microdeletion Clinical Presentation. Autism Research, 2010, 3, 286-287.	2.1	0
92	Functional impact of global rare copy number variation in autism spectrum disorders. Nature, 2010, 466, 368-372.	13.7	1,803
93	A genome-wide scan for common alleles affecting risk for autism. Human Molecular Genetics, 2010, 19, 4072-4082.	1.4	538
94	Disruption at the PTCHD1 Locus on Xp22.11 in Autism Spectrum Disorder and Intellectual Disability. Science Translational Medicine, 2010, 2, 49ra68.	5.8	178
95	Dopamine transporter gene variation modulates activation of striatum in youth with ADHD. NeuroImage, 2010, 53, 935-942.	2.1	62
96	Neural response to working memory load varies by dopamine transporter genotype in children. NeuroImage, 2010, 53, 970-977.	2.1	45
97	A de novo 1p34.2 microdeletion identifies the synaptic vesicle gene RIMS3 as a novel candidate for autism. Journal of Medical Genetics, 2010, 47, 81-90.	1.5	52
98	Association and Mutation Analyses of 16p11.2 Autism Candidate Genes. PLoS ONE, 2009, 4, e4582.	1.1	80
99	Dopamine Transporter Genotype and Stimulant Side Effect Factors in Youth Diagnosed with Attention-Deficit/Hyperactivity Disorder. Journal of Child and Adolescent Psychopharmacology, 2009, 19, 233-239.	0.7	30
100	Genome-Wide Analyses of Exonic Copy Number Variants in a Family-Based Study Point to Novel Autism Susceptibility Genes. PLoS Genetics, 2009, 5, e1000536.	1.5	374
101	Autism genome-wide copy number variation reveals ubiquitin and neuronal genes. Nature, 2009, 459, 569-573.	13.7	1,270
102	Common genetic variants on 5p14.1 associate with autism spectrum disorders. Nature, 2009, 459, 528-533.	13.7	912
103	Literature review: Similar prevalence of ASD across the life span; amygdala enlargement in young children. Autism Research, 2009, 2, 365-366.	2.1	0
104	Literature Review: Mitchondrial disorders; Circumscribed interests. Autism Research, 2009, 2, 60-61.	2.1	0
105	Habituation to neutral faces; Language regression in ASD. Autism Research, 2009, 2, 119-120.	2.1	0
106	Literature Review: Preference for physical compared to biological motion; Genomeâ€wide association study. Autism Research, 2009, 2, 178-179.	2.1	1
107	Citalopram efficacy study; interstitial 15q11â€q13 duplication mouse. Autism Research, 2009, 2, 238-240.	2.1	1
108	Gastrointestinal symptoms in ASD, brain structure of identical twins with ASD. Autism Research, 2009, 2, 285-286.	2.1	1

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109	Transmission disequilibrium testing of the chromosome 15q11â€q13 region in autism. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 1116-1125.	1.1	26
110	Familyâ€based association testing of OCDâ€associated SNPs of SLC1A1 in an autism sample. Autism Research, 2008, 1, 108-113.	2.1	21
111	Literature review: adult structural MRI, postmortem GABAâ€A protein, 1q21.1 microdeletion/microduplication. Autism Research, 2008, 1, 308-310.	2.1	1
112	Literature Review: Overlapping genetic association in developmental language disorder and autism; Grey matter in high functioning autism and Asperger's syndrome; Brain activation in selfâ€other face discrimination. Autism Research, 2008, 1, 370-371.	2.1	0
113	Copy-number variations associated with neuropsychiatric conditions. Nature, 2008, 455, 919-923.	13.7	587
114	Novel Submicroscopic Chromosomal Abnormalities Detected in Autism Spectrum Disorder. Biological Psychiatry, 2008, 63, 1111-1117.	0.7	268
115	A Common Genetic Variant in the Neurexin Superfamily Member CNTNAP2 Increases Familial Risk of Autism. American Journal of Human Genetics, 2008, 82, 160-164.	2.6	566
116	Molecular Genetics of the Platelet Serotonin System in First-Degree Relatives of Patients with Autism. Neuropsychopharmacology, 2008, 33, 353-360.	2.8	57
117	GAD1 Single Nucleotide Polymorphism Is in Linkage Disequilibrium with a Child Bipolar I Disorder Phenotype. Journal of Child and Adolescent Psychopharmacology, 2008, 18, 25-29.	0.7	20
118	Single nucleotide polymorphism discovery and functional assessment of variation in the UDP-glucuronosyltransferase 2B7 gene. Pharmacogenetics and Genomics, 2008, 18, 683-697.	0.7	73
119	Interactions between integrin Î±IIbÎ²3 and the serotonin transporter regulate serotonin transport and platelet aggregation in mice and humans. Journal of Clinical Investigation, 2008, 118, 1544-1552.	3.9	164
120	Recurrent 16p11.2 microdeletions in autism. Human Molecular Genetics, 2007, 17, 628-638.	1.4	614
121	Association of the oxytocin receptor gene (OXTR) in Caucasian children and adolescents with autism. Neuroscience Letters, 2007, 417, 6-9.	1.0	409
122	Using the Autism Diagnostic Interviewâ€"Revised to Increase Phenotypic Homogeneity in Genetic Studies of Autism. Biological Psychiatry, 2007, 61, 438-448.	0.7	153
123	Association Studies of Serotonin System Candidate Genes in Early-onset Obsessive-Compulsive Disorder. Biological Psychiatry, 2007, 61, 322-329.	0.7	81
124	Norepinephrine Transporter Gene Variation Modulates Acute Response to d-Amphetamine. Biological Psychiatry, 2007, 61, 1296-1305.	0.7	39
125	Association of dopamine transporter genotype with disruptive behavior disorders in an eight-year longitudinal study of children and adolescents. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 310-317.	1.1	41
126	Cytogenetic and molecular characterization of A2BP1 / FOX1 as a candidate gene for autism. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 869-876.	1.1	192

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127	Between a ROC and a hard place: decision making and making decisions about using the SCQ. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2007, 48, 932-940.	3.1	258
128	Mapping autism risk loci using genetic linkage and chromosomal rearrangements. Nature Genetics, 2007, 39, 319-328.	9.4	1,272
129	Combining Information From Multiple Sources in the Diagnosis of Autism Spectrum Disorders. Journal of the American Academy of Child and Adolescent Psychiatry, 2006, 45, 1094-1103.	0.3	481
130	High frequency of neurexin 1Î² signal peptide structural variants in patients with autism. Neuroscience Letters, 2006, 409, 10-13.	1.0	285
131	The Autistic Spectrum Disorders. , 2006, , 371-389.		1
132	5-HTTLPR Genotype-Specific Phenotype in Children and Adolescents With Autism. American Journal of Psychiatry, 2006, 163, 2148-2156.	4.0	158
133	Variation in ITGB3 is associated with whole-blood serotonin level and autism susceptibility. European Journal of Human Genetics, 2006, 14, 923-931.	1.4	82
134	Serotonin Transporter Genotype and Acute Subjective Response to Amphetamine. American Journal on Addictions, 2006, 15, 327-335.	1.3	28
135	ITGB3 shows genetic and expression interaction with SLC6A4. Human Genetics, 2006, 120, 93-100.	1.8	44
136	An association study of the brain-derived neurotrophic factor Val66Met polymorphism and amphetamine response. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2006, 141B, 576-583.	1.1	35
137	Association Testing of the Positional and Functional Candidate Gene SLC1A1/EAAC1 in Early-Onset Obsessive-compulsive Disorder. Archives of General Psychiatry, 2006, 63, 778.	13.8	252
138	Are the arginine vasopressin V1a receptor microsatellites related to hypersexuality in children with a prepubertal and early adolescent bipolar disorder phenotype?. Bipolar Disorders, 2005, 7, 610-616.	1.1	7
139	Family-based association study of the serotonin transporter gene polymorphisms in Korean ADHD trios. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2005, 139B, 14-18.	1.1	27
140	Interindividual variation in anxiety response to amphetamine: Possible role for adenosine A2Areceptor gene variants. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2005, 139B, 42-44.	1.1	50
141	Variation in ITGB3 has sex-specific associations with plasma lipoprotein(a) and whole blood serotonin levels in a population-based sample. Human Genetics, 2005, 117, 81-87.	1.8	27
142	Dopamine Transporter Gene Associated with Diminished Subjective Response to Amphetamine. Neuropsychopharmacology, 2005, 30, 602-609.	2.8	139
143	EPS or Stereotypies?. Journal of Child and Adolescent Psychopharmacology, 2005, 15, 150-151.	0.7	0
144	Dopamine Transporter Genotype and Methylphenidate Dose Response in Children with ADHD. Neuropsychopharmacology, 2005, 30, 1374-1382.	2.8	137

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145	Sex-Specific Genetic Architecture of Whole Blood Serotonin Levels. American Journal of Human Genetics, 2005, 76, 33-41.	2.6	122
146	An Open-Label Trial of Escitalopram in Pervasive Developmental Disorders. Journal of the American Academy of Child and Adolescent Psychiatry, 2005, 44, 343-348.	0.3	98
147	Linkage Disequilibrium of the Brain-Derived Neurotrophic FactorVal66MetPolymorphism in Children With a Prepubertal and Early Adolescent Bipolar Disorder Phenotype. American Journal of Psychiatry, 2004, 161, 1698-1700.	4.0	176
148	AUTISM AS A PARADIGMATIC COMPLEX GENETIC DISORDER. Annual Review of Genomics and Human Genetics, 2004, 5, 379-405.	2.5	237
149	Genome-wide association study identifies ITGB3 as a QTL for whole blood serotonin. European Journal of Human Genetics, 2004, 12, 949-954.	1.4	65
150	Pharmacogenetics of Outcome in Children with Acute Lymphoblastic Leukemia.. Blood, 2004, 104, 163-163.	0.6	2
151	Pharmacogenetics of Minimal Residual Disease Response in Children with Acute Lymphoblastic Leukemia (ALL).. Blood, 2004, 104, 451-451.	0.6	0
152	A pharmacogenetic study of uridine diphosphateâ€“glucuronosyltransferase 2B7 in patients receiving morphine. Clinical Pharmacology and Therapeutics, 2003, 73, 566-574.	2.3	149
153	Genetics of Childhood Disorders: XLVI. Autism, Part 5: Genetics of Autism. Journal of the American Academy of Child and Adolescent Psychiatry, 2003, 42, 116-118.	0.3	59
154	Remission Status After Long-Term Sertraline Treatment of Pediatric Obsessive-Compulsive Disorder. Journal of Child and Adolescent Psychopharmacology, 2003, 13, 53-60.	0.7	51
155	Case Series: AdderallÂ® Augmentation of Serotonin Reuptake Inhibitors in Childhood-Onset Obsessive Compulsive Disorder. Journal of Child and Adolescent Psychopharmacology, 2002, 12, 165-171.	0.7	10
156	Serotonin transporter promoter polymorphism, peripheral indexes of serotonin function, and personality measures in families with alcoholism. American Journal of Medical Genetics Part A, 2002, 114, 230-234.	2.4	126
157	Transmission disequilibrium studies of the serotonin 5-HT2A receptor gene (HTR2A) in autism. American Journal of Medical Genetics Part A, 2002, 114, 277-283.	2.4	76
158	Genome-wide linkage analysis of families with obsessive-compulsive disorder ascertained through pediatric probands. American Journal of Medical Genetics Part A, 2002, 114, 541-552.	2.4	223
159	No evidence for linkage of liability to autism toHOXA1 in a sample from the CPEA network. American Journal of Medical Genetics Part A, 2002, 114, 667-672.	2.4	33
160	Long-Term Sertraline Treatment of Children and Adolescents With Obsessive-Compulsive Disorder. Journal of the American Academy of Child and Adolescent Psychiatry, 2001, 40, 1175-1181.	0.3	166
161	Multisite, Double-Blind, Placebo-Controlled Trial of Porcine Secretin in Autism. Journal of the American Academy of Child and Adolescent Psychiatry, 2001, 40, 1293-1299.	0.3	118
162	Risperidone-Induced Prolonged Erections Following the Addition of Lithium. Journal of Child and Adolescent Psychopharmacology, 2001, 11, 441-442.	0.7	28

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163	Double-Blind, Placebo-Controlled Study of Amantadine Hydrochloride in the Treatment of Children With Autistic Disorder. Journal of the American Academy of Child and Adolescent Psychiatry, 2001, 40, 658-665.	0.3	181
164	Genetics of Autism. Child and Adolescent Psychiatric Clinics of North America, 2001, 10, 333-350.	1.0	76
165	Self-injurious behavior: Gene-brain-behavior relationships. Mental Retardation and Developmental Disabilities Research Reviews, 2001, 7, 3-12.	3.5	119
166	Serotonin Transporter Promoter Polymorphism Genotype Is Associated With Behavioral Disinhibition and Negative Affect in Children of Alcoholics. Alcoholism: Clinical and Experimental Research, 2001, 25, 953-959.	1.4	56
167	Genetics of Psychiatric Disorders: Where Have We Been and Where Are We Going?. American Journal of Psychiatry, 2000, 157, 1039-1040.	4.0	8
168	Novelde novo nonsense mutation ofMECP2 in a patient with Rett syndrome. Human Mutation, 2000, 15, 382-383.	1.1	37
169	Serotonergic Function, Behavioral Disinhibition, and Negative Affect in Children of Alcoholics: The Moderating Effects of Puberty. Alcoholism: Clinical and Experimental Research, 2000, 24, 972-979.	1.4	19
170	Pharmacogenetics and the serotonin system: initial studies and future directions. European Journal of Pharmacology, 2000, 410, 165-181.	1.7	236
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