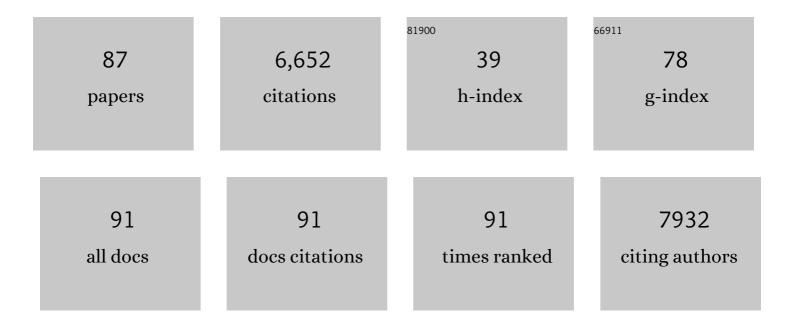
Joseph F Cubells

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Association of <emph type="ital">FKBP5</emph> Polymorphisms and Childhood Abuse With Risk of Posttraumatic Stress Disorder Symptoms in Adults. JAMA - Journal of the American Medical Association, 2008, 299, 1291.	7.4	1,190
2	Influence of Child Abuse on Adult Depression. Archives of General Psychiatry, 2008, 65, 190.	12.3	583
3	Trauma exposure and stress-related disorders in inner city primary care patients. General Hospital Psychiatry, 2009, 31, 505-514.	2.4	401
4	Serotonin transporter protein (SLC6A4) allele and haplotype frequencies and linkage disequilibria in African- and European-American and Japanese populations and in alcohol-dependent subjects. Human Genetics, 1997, 101, 243-246.	3.8	393
5	Differential immune system DNA methylation and cytokine regulation in postâ€ŧraumatic stress disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 700-708.	1.7	294
6	SPARK: A US Cohort of 50,000 Families to Accelerate Autism Research. Neuron, 2018, 97, 488-493.	8.1	265
7	A Quantitative-Trait Analysis of Human Plasma–Dopamine β-Hydroxylase Activity: Evidence for a Major Functional Polymorphism at the DBH Locus. American Journal of Human Genetics, 2001, 68, 515-522.	6.2	253
8	An epigenetic clock for gestational age at birth based on blood methylation data. Genome Biology, 2016, 17, 206.	8.8	193
9	Diagnostic reliability of the Semi-structured Assessment for Drug Dependence and Alcoholism (SSADDA). Drug and Alcohol Dependence, 2005, 80, 303-312.	3.2	180
10	Dopamine β-hydroxylase: two polymorphisms in linkage disequilibrium at the structural gene DBH associate with biochemical phenotypic variation. Human Genetics, 1998, 102, 533-540.	3.8	127
11	Reliability of DSM-IV diagnostic criteria using the semi-structured assessment for drug dependence and alcoholism (SSADDA). Drug and Alcohol Dependence, 2007, 91, 85-90.	3.2	124
12	Population genetics of a functional variant of the dopamine β-hydroxylase gene (DBH). American Journal of Medical Genetics Part A, 1997, 74, 374-379.	2.4	104
13	Neonatal DNA methylation patterns associate with gestational age. Epigenetics, 2011, 6, 1498-1504.	2.7	95
14	A Previously Undescribed Intron and Extensive 5′ Upstream Sequence, but Not Phox2a-mediated Transactivation, Are Necessary for High Level Cell Type-specific Expression of the Human Norepinephrine Transporter Gene. Journal of Biological Chemistry, 1999, 274, 6507-6518.	3.4	93
15	Using common genetic variation to examine phenotypic expression and risk prediction in 22q11.2 deletion syndrome. Nature Medicine, 2020, 26, 1912-1918.	30.7	90
16	Mutations in the dopamine ?-hydroxylase gene are associated with human norepinephrine deficiency. American Journal of Medical Genetics Part A, 2002, 108, 140-147.	2.4	88
17	Genetic contributors to risk of schizophrenia in the presence of a 22q11.2 deletion. Molecular Psychiatry, 2021, 26, 4496-4510.	7.9	87
18	Polymorphisms in <i>CRHR1</i> and the serotonin transporter loci: Gene × Gene × Enviror interactions on depressive symptoms. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 812-824.	nment 1.7	83

#	Article	IF	CITATIONS
19	The Structure of Linkage Disequilibrium at the DBH Locus Strongly Influences the Magnitude of Association between Diallelic Markers and Plasma Dopamine β-Hydroxylase Activity. American Journal of Human Genetics, 2003, 72, 1389-1400.	6.2	81
20	A genotype-controlled analysis of plasma dopamine Î ² -hydroxylase in healthy and alcoholic subjects: evidence for alcohol-related differences in noradrenergic function. Biological Psychiatry, 2002, 52, 1151-1158.	1.3	75
21	Inducible cAMP Early Repressor Can Modulate Tyrosine Hydroxylase Gene Expression after Stimulation of cAMP Synthesis. Journal of Biological Chemistry, 1996, 271, 25375-25381.	3.4	73
22	Association study of theCNR1 gene exon 3 alternative promoter region polymorphisms and substance dependence. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2006, 141B, 499-503.	1.7	71
23	DNA methylation in neonates born to women receiving psychiatric care. Epigenetics, 2012, 7, 409-414.	2.7	68
24	Epigenetic modification of the oxytocin receptor gene: implications for autism symptom severity and brain functional connectivity. Neuropsychopharmacology, 2020, 45, 1150-1158.	5.4	62
25	Randomized, double blind, placebo-controlled trial of disulfiram for the treatment of cocaine dependence in methadone-stabilized patients. Drug and Alcohol Dependence, 2011, 113, 184-191.	3.2	61
26	Cyclic AMP-Dependent Protein Kinase Regulates Basal and Cyclic AMP-Stimulated but Not Phorbol Ester-Stimulated Transcription of the Tyrosine Hydroxylase Gene. Journal of Neurochemistry, 2002, 63, 834-842.	3.9	59
27	Genotype-controlled analysis of plasma dopamine β-hydroxylase activity in psychotic unipolar major depression. Biological Psychiatry, 2002, 51, 358-364.	1.3	58
28	Dopamine β-Hydroxylase Gene (DβH) -1021C→T Influences Self-Reported Paranoia during Cocaine Self-Administration. Biological Psychiatry, 2007, 61, 1310-1313.	1.3	58
29	Differential In Vivo Regulation of mRNA Encoding the Norepinephrine Transporter and Tyrosine Hydroxylase in Rat Adrenal Medulla and Locus Ceruleus. Journal of Neurochemistry, 1995, 65, 502-509.	3.9	57
30	Dopamine betaâ€hydroxylase (DBH) activity and â€1021C/T polymorphism of <i>DBH</i> gene in combatâ€related postâ€traumatic stress disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 1087-1089.	1.7	57
31	Rating the severity and character of transient cocaine-induced delusions and hallucinations with a new instrument, the Scale for Assessment of Positive Symptoms for Cocaine-Induced Psychosis (SAPS-CIP). Drug and Alcohol Dependence, 2005, 80, 23-33.	3.2	55
32	A review of neurocognitive and behavioral profiles associated with 22q11 deletion syndrome: Implications for clinical evaluation and treatment. Current Psychiatry Reports, 2007, 9, 148-158.	4.5	49
33	A serotonin transporter gene polymorphism predicts peripartum depressive symptoms in an at-risk psychiatric cohort. Journal of Psychiatric Research, 2010, 44, 640-646.	3.1	49
34	Subthreshold Psychosis in 22q11.2 Deletion Syndrome: Multisite Naturalistic Study. Schizophrenia Bulletin, 2017, 43, 1079-1089.	4.3	47
35	Comorbid Psychiatric Diagnoses and Their Association with Cocaine-Induced Psychosis in Cocaine-Dependent Subjects. American Journal on Addictions, 2007, 16, 343-351.	1.4	46
36	Pharmaco-genetically guided treatment of recurrent rage outbursts in an adult male with 15q13.3		46

deletion syndrome. , 2011, 155, 805-810.

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#	Article	IF	CITATIONS
37	Linkage analysis of plasma dopamine β-hydroxylase activity in families of patients with schizophrenia. Human Genetics, 2011, 130, 635-643.	3.8	45
38	Dopamine Beta-Hydroxylase (DBH) gene and schizophrenia phenotypic variability: A genetic association study. American Journal of Medical Genetics Part A, 2003, 117B, 33-38.	2.4	43
39	Genotypic and haplotypic associations of the DBH gene with plasma dopamine β-hydroxylase activity in African Americans. European Journal of Human Genetics, 2007, 15, 878-883.	2.8	43
40	A Single Nucleotide Polymorphism at DBH, Possibly Associated with Attention-Deficit/Hyperactivity Disorder, Associates with Lower Plasma Dopamine Î ² -Hydroxylase Activity and is in Linkage Disequilibrium with Two Putative Functional Single Nucleotide Polymorphisms. Biological Psychiatry, 2006, 60, 1034-1038.	1.3	42
41	Study protocol for The Emory 3q29 Project: evaluation of neurodevelopmental, psychiatric, and medical symptoms in 3q29 deletion syndrome. BMC Psychiatry, 2018, 18, 183.	2.6	40
42	Self-reported paranoia during laboratory "binge―cocaine self-administration in humans. Pharmacology Biochemistry and Behavior, 2006, 83, 249-256.	2.9	39
43	Socialâ€adaptive and psychological functioning of patients affected by Fabry disease. Journal of Inherited Metabolic Disease, 2010, 33, 73-81.	3.6	33
44	Mutations in the dopamine beta-hydroxylase gene are associated with human norepinephrine deficiency. American Journal of Medical Genetics Part A, 2002, 108, 140-7.	2.4	33
45	Sex dependent influence of a functional polymorphism in steroid 5â€Î±â€reductase type 2 (<i>SRD5A2</i>) on postâ€traumatic stress symptoms. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 283-292.	1.7	32
46	Deep phenotyping in 3q29 deletion syndrome: recommendations for clinical care. Genetics in Medicine, 2021, 23, 872-880.	2.4	32
47	Haplotypeâ€controlled analysis of the association of a nonâ€synonymous single nucleotide polymorphism at DBH (+ 1603C → T) with plasma dopamine βâ€hydroxylase activity. American Journal of Medical Part B: Neuropsychiatric Genetics, 2005, 139B, 88-90.	¢മnetics	31
48	Predictors of neonatal hypothalamic–pituitary–adrenal axis activity at delivery. Clinical Endocrinology, 2011, 75, 90-95.	2.4	30
49	Association between polymorphisms in catecholâ€ <i>O</i> â€methyltransferase (<i>COMT</i>) and cocaineâ€induced paranoia in Europeanâ€American and Africanâ€American populations. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 651-660.	1.7	30
50	Randomized clinical trial of disulfiram for cocaine dependence or abuse during buprenorphine treatment. Drug and Alcohol Dependence, 2014, 136, 36-42.	3.2	27
51	Pharmacogenetic Association of the Galanin Receptor (GALR1) SNP rs2717162 with Smoking Cessation. Neuropsychopharmacology, 2012, 37, 1683-1688.	5.4	26
52	Ciliary neurotrophic factor null allele frequencies in schizophrenia, affective disorders, and Alzheimer's disease. American Journal of Medical Genetics Part A, 1997, 74, 497-500.	2.4	24
53	The Galanin Receptor 1 Gene Associates with Tobacco Craving in Smokers Seeking Cessation Treatment. Neuropsychopharmacology, 2011, 36, 1412-1420.	5.4	23
54	Transient Cocaine-Associated Behavioral Symptoms Rated with a New Instrument, the Scale for Assessment of Positive Symptoms for Cocaine-Induced Psychosis (SAPS-CIP). American Journal on Addictions, 2009, 18, 339-345.	1.4	23

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#	Article	IF	CITATIONS
55	Association of Variants in MANEA With Cocaine-Related Behaviors. Archives of General Psychiatry, 2009, 66, 267.	12.3	22
56	Cross-Disorder Comparison of Four Neuropsychiatric CNV Loci. Current Genetic Medicine Reports, 2014, 2, 151-161.	1.9	22
57	Substance Use Disorders Assessed Using the Kreek–McHugh–Schluger–Kellogg (KMSK) Scale in an Urban Lowâ€Income and Predominantly African American Sample of Primary Care Patients. American Journal on Addictions, 2011, 20, 292-299.	1.4	21
58	Regulatory Polymorphisms in Human <i>DBH</i> Affect Peripheral Gene Expression and Sympathetic Activity. Circulation Research, 2014, 115, 1017-1025.	4.5	21
59	Subtypes of major depression in substance dependence. Addiction, 2009, 104, 1700-1709.	3.3	20
60	Analysis of variations in the tryptophan hydroxylase-2 (TPH2) gene in cocaine dependence. Addiction Biology, 2006, 11, 76-83.	2.6	18
61	Functional variants atCYP2A6: New genotyping methods, population genetics, and relevance to studies of tobacco dependence. American Journal of Medical Genetics Part A, 2000, 96, 638-645.	2.4	16
62	Dopamine βâ€hydroxylase gene associates with stroop colorâ€word task performance in Han Chinese children with attention deficit/hyperactivity disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 730-736.	1.7	16
63	Genotype-controlled analysis of serum dopamine β-hydroxylase activity in civilian post-traumatic stress disorder. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2010, 34, 1396-1401.	4.8	15
64	Innervation-independent changes in the mRNAs encoding tyrosine hydroxylase and the norepinephrine transporter in rat adrenal medulla after high-dose reserpine. Neuroscience Letters, 1995, 193, 189-192.	2.1	13
65	Variations in the dopamine ?-hydroxylase gene are not associated with the autonomic disorders, pure autonomic failure, or multiple system atrophy. American Journal of Medical Genetics Part A, 2003, 120A, 234-236.	2.4	13
66	Prodromal and autistic symptoms in schizotypal personality disorder and 22q11.2 deletion syndrome Journal of Abnormal Psychology, 2013, 122, 238-249.	1.9	13
67	Brief Report: Relationship Between ADOS-2, Module 4 Calibrated Severity Scores (CSS) and Social and Non-Social Standardized Assessment Measures in Adult Males with Autism Spectrum Disorder (ASD). Journal of Autism and Developmental Disorders, 2017, 47, 4018-4024.	2.7	13
68	Human Bacterial Artificial Chromosome (BAC) Transgenesis Fully Rescues Noradrenergic Function in Dopamine β-Hydroxylase Knockout Mice. PLoS ONE, 2016, 11, e0154864.	2.5	12
69	Associations between the DBH gene, plasma dopamine β-hydroxylase activity and cognitive measures in Han Chinese patients with schizophrenia. Schizophrenia Research, 2018, 193, 58-63.	2.0	12
70	Beliefs in vaccine as causes of autism among SPARK cohort caregivers. Vaccine, 2020, 38, 1794-1803.	3.8	12
71	Comprehensive phenotyping of neuropsychiatric traits in a multiplex 3q29 deletion family: a case report. BMC Psychiatry, 2020, 20, 184.	2.6	12
72	Convergent and distributed effects of the 3q29 deletion on the human neural transcriptome. Translational Psychiatry, 2021, 11, 357.	4.8	12

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73	A revised allele frequency estimate and haplotype analysis of the DBH deficiency mutation IVS1+2T → C African- and European-Americans. , 2003, 123A, 190-192.	Cin	11
74	Copy Number Variants: A New Molecular Frontier in Clinical Psychiatry. Current Psychiatry Reports, 2011, 13, 129-137.	4.5	11
75	Sources of Unreliability in the Diagnosis of Substance Dependence. Journal of Studies on Alcohol and Drugs, 2009, 70, 475-481.	1.0	9
76	Meaning in Measurement: Evaluating Young Autistic Adults' Active Engagement and Expressed Interest in Quality-of-Life Goals. Autism in Adulthood, 2020, 2, 227-242.	6.9	9
77	Prevalence of autism spectrum disorders in China. Shanghai Archives of Psychiatry, 2013, 25, 176-7.	0.7	5
78	Clinical trials report. Current Psychiatry Reports, 2007, 9, 131-134.	4.5	3
79	Clinical genomic psychiatry comes of age in the evaluation and treatment of developmental disabilities: Is our nation prepared to make the benefits available to all who need them?. Current Psychiatry Reports, 2007, 9, 81-82.	4.5	2
80	Concerns Over Participant Suicides Prematurely Abort a Clinical Trial of Potentially Significant Impact on Public Health: How Will We Make Progress in Timid Times?. Current Psychiatry Reports, 2011, 13, 80-81.	4.5	1
81	Ciliary neurotrophic factor null allele frequencies in schizophrenia, affective disorders, and Alzheimer's disease. , 1997, 74, 497.		1
82	Beyond irritability and aggressive behavior: does risperidone improve adaptive behavior in autistic spectrum disorders?. Current Psychiatry Reports, 2007, 9, 132-3.	4.5	1
83	A Head-to-Head Comparison of Risperidone and Divalproex for Treatment of Pediatric Bipolar Disorder. Current Psychiatry Reports, 2011, 13, 82-83.	4.5	0
84	Genetics of stimulant dependence. , 0, , 306-315.		0
85	Molecular Genetic Analysis of Plasma Dopamine β-Hydroxylase in Depression. Advances in Behavioral Biology, 2002, , 423-426.	0.2	0
86	22q11.2 Deletion Syndrome: A Paradigmatic Copy-Number-Variant (CNV) Disorder. , 2016, , 723-730.		0
87	Acetylcholine and cognition in schizophrenia: effects of galantamine. Current Psychiatry Reports, 2007, 9, 133-4.	4.5	0