Joseph F Cubells

List of Publications by Year
in descending order

87 papers	6,652 citations	87723 38 h-index	66788 78 g-index
91 all docs	91 docs citations	91 times ranked	7932 citing authors

#	Article	IF	Citations
1	Deep phenotyping in 3q29 deletion syndrome: recommendations for clinical care. Genetics in Medicine, 2021, 23, 872-880.	1.1	32
2	Convergent and distributed effects of the 3q29 deletion on the human neural transcriptome. Translational Psychiatry, 2021, 11, 357.	2.4	12
3	Genetic contributors to risk of schizophrenia in the presence of a 22q11.2 deletion. Molecular Psychiatry, 2021, 26, 4496-4510.	4.1	87
4	Using common genetic variation to examine phenotypic expression and risk prediction in 22q11.2 deletion syndrome. Nature Medicine, 2020, 26, 1912-1918.	15.2	90
5	Meaning in Measurement: Evaluating Young Autistic Adults' Active Engagement and Expressed Interest in Quality-of-Life Goals. Autism in Adulthood, 2020, 2, 227-242.	4.0	9
6	Epigenetic modification of the oxytocin receptor gene: implications for autism symptom severity and brain functional connectivity. Neuropsychopharmacology, 2020, 45, 1150-1158.	2.8	62
7	Beliefs in vaccine as causes of autism among SPARK cohort caregivers. Vaccine, 2020, 38, 1794-1803.	1.7	12
8	Comprehensive phenotyping of neuropsychiatric traits in a multiplex 3q29 deletion family: a case report. BMC Psychiatry, 2020, 20, 184.	1.1	12
9	SPARK: A US Cohort of 50,000 Families to Accelerate Autism Research. Neuron, 2018, 97, 488-493.	3.8	265
10	Associations between the DBH gene, plasma dopamine Î²-hydroxylase activity and cognitive measures in Han Chinese patients with schizophrenia. Schizophrenia Research, 2018, 193, 58-63.	1.1	12
11	Study protocol for The Emory 3q29 Project: evaluation of neurodevelopmental, psychiatric, and medical symptoms in 3q29 deletion syndrome. BMC Psychiatry, 2018, 18, 183.	1.1	40
12	Subthreshold Psychosis in 22q11.2 Deletion Syndrome: Multisite Naturalistic Study. Schizophrenia Bulletin, 2017, 43, 1079-1089.	2.3	47
13	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.	1.7	13
14	Human Bacterial Artificial Chromosome (BAC) Transgenesis Fully Rescues Noradrenergic Function in Dopamine Î²-Hydroxylase Knockout Mice. PLoS ONE, 2016, 11, e0154864.	1.1	12
15	An epigenetic clock for gestational age at birth based on blood methylation data. Genome Biology, 2016, 17, 206.	3.8	193
16	22q11.2 Deletion Syndrome: A Paradigmatic Copy-Number-Variant (CNV) Disorder. , 2016, , 723-730.		0
17	Regulatory Polymorphisms in Human DBH Affect Peripheral Gene Expression and Sympathetic Activity. Circulation Research, 2014, 115, 1017-1025.	2.0	21
18	Cross-Disorder Comparison of Four Neuropsychiatric CNV Loci. Current Genetic Medicine Reports, 2014, 2, 151-161.	1.9	22

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19	Randomized clinical trial of disulfiram for cocaine dependence or abuse during buprenorphine treatment. Drug and Alcohol Dependence, 2014, 136, 36-42.	1.6	27
20	Prodromal and autistic symptoms in schizotypal personality disorder and 22q11.2 deletion syndrome.. Journal of Abnormal Psychology, 2013, 122, 238-249.	2.0	13
21	Sex dependent influence of a functional polymorphism in steroid 5â€Î±â€reductase type 2 (SRD5A2) on postâ€traumatic stress symptoms. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 283-292.	1.1	32
22	Prevalence of autism spectrum disorders in China. Shanghai Archives of Psychiatry, 2013, 25, 176-7.	0.7	5
23	Pharmacogenetic Association of the Galanin Receptor (GALR1) SNP rs2717162 with Smoking Cessation. Neuropsychopharmacology, 2012, 37, 1683-1688.	2.8	26
24	DNA methylation in neonates born to women receiving psychiatric care. Epigenetics, 2012, 7, 409-414.	1.3	68
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.	1.6	61
26	Predictors of neonatal hypothalamicâ€“pituitaryâ€“adrenal axis activity at delivery. Clinical Endocrinology, 2011, 75, 90-95.	1.2	30
27	Substance Use Disorders Assessed Using the Kreekâ€“McHughâ€“Schlugerâ€“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.	1.3	21
28	Linkage analysis of plasma dopamine Î²-hydroxylase activity in families of patients with schizophrenia. Human Genetics, 2011, 130, 635-643.	1.8	45
29	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.	2.1	1
30	A Head-to-Head Comparison of Risperidone and Divalproex for Treatment of Pediatric Bipolar Disorder. Current Psychiatry Reports, 2011, 13, 82-83.	2.1	0
31	Copy Number Variants: A New Molecular Frontier in Clinical Psychiatry. Current Psychiatry Reports, 2011, 13, 129-137.	2.1	11
32	Pharmaco-genetically guided treatment of recurrent rage outbursts in an adult male with 15q13.3 deletion syndrome. , 2011, 155, 805-810.		46
33	Association between polymorphisms in catecholâ€Oâ€methyltransferase (COMT) and cocaineâ€induced paranoia in Europeanâ€American and Africanâ€American populations. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 651-660.	1.1	30
34	Differential immune system DNA methylation and cytokine regulation in postâ€traumatic stress disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 700-708.	1.1	294
35	Dopamine Î²â€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.	1.1	16
36	Neonatal DNA methylation patterns associate with gestational age. Epigenetics, 2011, 6, 1498-1504.	1.3	95

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37	The Galanin Receptor 1 Gene Associates with Tobacco Craving in Smokers Seeking Cessation Treatment. Neuropsychopharmacology, 2011, 36, 1412-1420.	2.8	23
38	Polymorphisms in CRHR1 and the serotonin transporter loci: Geneâ€‰Ã—â€‰Geneâ€‰Ã—â€‰Environment interactions on depressive symptoms. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 812-824.	1.1	83
39	A serotonin transporter gene polymorphism predicts peripartum depressive symptoms in an at-risk psychiatric cohort. Journal of Psychiatric Research, 2010, 44, 640-646.	1.5	49
40	Socialâ€adaptive and psychological functioning of patients affected by Fabry disease. Journal of Inherited Metabolic Disease, 2010, 33, 73-81.	1.7	33
41	Genotype-controlled analysis of serum dopamine Î²-hydroxylase activity in civilian post-traumatic stress disorder. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2010, 34, 1396-1401.	2.5	15
42	Sources of Unreliability in the Diagnosis of Substance Dependence. Journal of Studies on Alcohol and Drugs, 2009, 70, 475-481.	0.6	9
43	Trauma exposure and stress-related disorders in inner city primary care patients. General Hospital Psychiatry, 2009, 31, 505-514.	1.2	401
44	Subtypes of major depression in substance dependence. Addiction, 2009, 104, 1700-1709.	1.7	20
45	Association of Variants in MANEA With Cocaine-Related Behaviors. Archives of General Psychiatry, 2009, 66, 267.	13.8	22
46	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.	1.3	23
47	Influence of Child Abuse on Adult Depression. Archives of General Psychiatry, 2008, 65, 190.	13.8	583
48	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.	3.8	1,190
49	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.	1.6	124
50	Dopamine Î²-Hydroxylase Gene (DÎ²H) -1021Câ†'T Influences Self-Reported Paranoia during Cocaine Self-Administration. Biological Psychiatry, 2007, 61, 1310-1313.	0.7	58
51	Dopamine betaâ€hydroxylase (DBH) activity and â€1021C/T polymorphism of DBH gene in combatâ€related postâ€traumatic stress disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 1087-1089.	1.1	57
52	Genotypic and haplotypic associations of the DBH gene with plasma dopamine Î²-hydroxylase activity in African Americans. European Journal of Human Genetics, 2007, 15, 878-883.	1.4	43
53	Comorbid Psychiatric Diagnoses and Their Association with Cocaine-Induced Psychosis in Cocaine-Dependent Subjects. American Journal on Addictions, 2007, 16, 343-351.	1.3	46
54	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.	2.1	2

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55	Clinical trials report. Current Psychiatry Reports, 2007, 9, 131-134.	2.1	3
56	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.	2.1	49
57	Beyond irritability and aggressive behavior: does risperidone improve adaptive behavior in autistic spectrum disorders?. Current Psychiatry Reports, 2007, 9, 132-3.	2.1	1
58	Acetylcholine and cognition in schizophrenia: effects of galantamine. Current Psychiatry Reports, 2007, 9, 133-4.	2.1	0
59	A Single Nucleotide Polymorphism at DBH, Possibly Associated with Attention-Deficit/Hyperactivity Disorder, Associates with Lower Plasma Dopamine Î²-Hydroxylase Activity and is in Linkage Disequilibrium with Two Putative Functional Single Nucleotide Polymorphisms. Biological Psychiatry, 2006, 60, 1034-1038.	0.7	42
60	Analysis of variations in the tryptophan hydroxylase-2 (TPH2) gene in cocaine dependence. Addiction Biology, 2006, 11, 76-83.	1.4	18
61	Self-reported paranoia during laboratory â€œbingeâ€•cocaine self-administration in humans. Pharmacology Biochemistry and Behavior, 2006, 83, 249-256.	1.3	39
62	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.	1.1	71
63	Haplotypeâ€•controlled analysis of the association of a nonâ€•synonymous single nucleotide polymorphism at DBH (+â€‰1603Câ€‰â†’â€‰T) with plasma dopamine Î²â€•hydroxylase activity. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2005, 139B, 88-90.	1.1	31
64	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.	1.6	55
65	Diagnostic reliability of the Semi-structured Assessment for Drug Dependence and Alcoholism (SSADDA). Drug and Alcohol Dependence, 2005, 80, 303-312.	1.6	180
66	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
67	A revised allele frequency estimate and haplotype analysis of the DBH deficiency mutation IVS1+2Tâ€‰â†’â€‰C in African- and European-Americans. , 2003, 123A, 190-192.		11
68	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
69	The Structure of Linkage Disequilibrium at the DBH Locus Strongly Influences the Magnitude of Association between Diallelic Markers and Plasma Dopamine Î²-Hydroxylase Activity. American Journal of Human Genetics, 2003, 72, 1389-1400.	2.6	81
70	Genotype-controlled analysis of plasma dopamine Î²-hydroxylase activity in psychotic unipolar major depression. Biological Psychiatry, 2002, 51, 358-364.	0.7	58
71	A genotype-controlled analysis of plasma dopamine Î²-hydroxylase in healthy and alcoholic subjects: evidence for alcohol-related differences in noradrenergic function. Biological Psychiatry, 2002, 52, 1151-1158.	0.7	75
72	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

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73	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.	2.1	59
74	Molecular Genetic Analysis of Plasma Dopamine Î²-Hydroxylase in Depression. Advances in Behavioral Biology, 2002, , 423-426.	0.2	0
75	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
76	A Quantitative-Trait Analysis of Human Plasmaâ€“Dopamine Î²-Hydroxylase Activity: Evidence for a Major Functional Polymorphism at the DBH Locus. American Journal of Human Genetics, 2001, 68, 515-522.	2.6	253
77	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
78	A Previously Undescribed Intron and Extensive 5â€² 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.	1.6	93
79	Dopamine Î²-hydroxylase: two polymorphisms in linkage disequilibrium at the structural gene DBH associate with biochemical phenotypic variation. Human Genetics, 1998, 102, 533-540.	1.8	127
80	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.	1.8	393
81	Population genetics of a functional variant of the dopamine Î²-hydroxylase gene (DBH). , 1997, 74, 374-379.		104
82	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
83	Ciliary neurotrophic factor null allele frequencies in schizophrenia, affective disorders, and Alzheimer's disease. , 1997, 74, 497.		1
84	Inducible cAMP Early Repressor Can Modulate Tyrosine Hydroxylase Gene Expression after Stimulation of cAMP Synthesis. Journal of Biological Chemistry, 1996, 271, 25375-25381.	1.6	73
85	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.	1.0	13
86	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.	2.1	57
87	Genetics of stimulant dependence. , 0, , 306-315.		0