

# Dorothy E Grice

## List of Publications by Year in descending order

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Version: 2024-02-01

59  
papers

10,697  
citations

117571

34  
h-index

161767

54  
g-index

68  
all docs

68  
docs citations

68  
times ranked

15428  
citing authors

#	ARTICLE	IF	CITATIONS
1	Systematic review and meta-analysis: relationships between attention-deficit/hyperactivity disorder and urinary symptoms in children. <i>European Child and Adolescent Psychiatry</i> , 2022, 31, 663-670.	2.8	10
2	The Genetic Architecture of Obsessive-Compulsive Disorder: Contribution of Liability to OCD From Alleles Across the Frequency Spectrum. <i>American Journal of Psychiatry</i> , 2022, 179, 216-225.	4.0	16
3	Psychometric properties of the Swedish translation of the Obsessive-Compulsive Inventory-Revised and the population characteristics of the symptom dimensions of OCD. <i>Social Psychiatry and Psychiatric Epidemiology</i> , 2022, , 1.	1.6	0
4	Synaptic processes and immune-related pathways implicated in Tourette syndrome. <i>Translational Psychiatry</i> , 2021, 11, 56.	2.4	31
5	Prospective and detailed behavioral phenotyping in DDX3X syndrome. <i>Molecular Autism</i> , 2021, 12, 36.	2.6	25
6	Developmental and Behavioral Phenotypes in a Mouse Model of DDX3X Syndrome. <i>Biological Psychiatry</i> , 2021, 90, 742-755.	0.7	21
7	Investigation of gene-environment interactions in relation to tic severity. <i>Journal of Neural Transmission</i> , 2021, 128, 1757-1765.	1.4	2
8	Systematic review and meta-analysis identify significant relationships between clinical anxiety and lower urinary tract symptoms. <i>Brain and Behavior</i> , 2021, 11, e2268.	1.0	12
9	Prevalence and phenotypic impact of rare potentially damaging variants in autism spectrum disorder. <i>Molecular Autism</i> , 2021, 12, 65.	2.6	22
10	Cohort profile: Epidemiology and Genetics of Obsessive-compulsive disorder and chronic tic disorders in Sweden (EGOS). <i>Social Psychiatry and Psychiatric Epidemiology</i> , 2020, 55, 1383-1393.	1.6	13
11	Episignatures Stratifying Helsmoortel-Van Der Aa Syndrome Show Modest Correlation with Phenotype. <i>American Journal of Human Genetics</i> , 2020, 107, 555-563.	2.6	32
12	Don't Worry, the Genetics of Obsessive-Compulsive Disorder Is Finally Catching Up. <i>Biological Psychiatry</i> , 2020, 87, 1017-1018.	0.7	2
13	Maternal Effects as Causes of Risk for Obsessive-Compulsive Disorder. <i>Biological Psychiatry</i> , 2020, 87, 1045-1051.	0.7	18
14	Parental Age and Differential Estimates of Risk for Neuropsychiatric Disorders: Findings From the Danish Birth Cohort. <i>Journal of the American Academy of Child and Adolescent Psychiatry</i> , 2019, 58, 618-627.	0.3	24
15	Genomic Relationships, Novel Loci, and Pleiotropic Mechanisms across Eight Psychiatric Disorders. <i>Cell</i> , 2019, 179, 1469-1482.e11.	13.5	935
16	Dr Nurnberger and Colleagues Reply. <i>Journal of Clinical Psychiatry</i> , 2019, 80, .	1.1	0
17	Investigation of previously implicated genetic variants in chronic tic disorders: a transmission disequilibrium test approach. <i>European Archives of Psychiatry and Clinical Neuroscience</i> , 2018, 268, 301-316.	1.8	23
18	Heritable Variation, With Little or No Maternal Effect, Accounts for Recurrence Risk to Autism Spectrum Disorder in Sweden. <i>Biological Psychiatry</i> , 2018, 83, 589-597.	0.7	38

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19	De Novo Sequence and Copy Number Variants Are Strongly Associated with Tourette Disorder and Implicate Cell Polarity in Pathogenesis. <i>Cell Reports</i> , 2018, 24, 3441-3454.e12.	2.9	91
20	Identification of rare de novo epigenetic variations in congenital disorders. <i>Nature Communications</i> , 2018, 9, 2064.	5.8	82
21	What Should a Psychiatrist Know About Genetics?. <i>Journal of Clinical Psychiatry</i> , 2018, 80, .	1.1	40
22	“Just Right” Transitioning to College with Obsessive-Compulsive Disorder. , 2018, , 161-168.		0
23	Polygenic transmission disequilibrium confirms that common and rare variation act additively to create risk for autism spectrum disorders. <i>Nature Genetics</i> , 2017, 49, 978-985.	9.4	401
24	De Novo Coding Variants Are Strongly Associated with Tourette Disorder. <i>Neuron</i> , 2017, 94, 486-499.e9.	3.8	155
25	Parental Age And Differential Risk For Asd, Adhd, Ocd And Tic Disorders: Data From A Large National Cohort. <i>European Neuropsychopharmacology</i> , 2017, 27, S492.	0.3	0
26	Diagnostic validity of early-onset obsessive-compulsive disorder in the Danish Psychiatric Central Register: findings from a cohort sample. <i>BMJ Open</i> , 2017, 7, e017172.	0.8	18
27	Prenatal Maternal Smoking and Increased Risk for Tourette Syndrome and Chronic Tic Disorders. <i>Journal of the American Academy of Child and Adolescent Psychiatry</i> , 2016, 55, 784-791.	0.3	43
28	Pre- and perinatal complications in relation to Tourette syndrome and co-occurring obsessive-compulsive disorder and attention-deficit/hyperactivity disorder. <i>Journal of Psychiatric Research</i> , 2016, 82, 126-135.	1.5	36
29	Familial Clustering of Tic Disorders and Obsessive-Compulsive Disorder. <i>JAMA Psychiatry</i> , 2015, 72, 359.	6.0	67
30	Joint Analysis of Psychiatric Disorders Increases Accuracy of Risk Prediction for Schizophrenia, Bipolar Disorder, and Major Depressive Disorder. <i>American Journal of Human Genetics</i> , 2015, 96, 283-294.	2.6	225
31	Psychiatric genome-wide association study analyses implicate neuronal, immune and histone pathways. <i>Nature Neuroscience</i> , 2015, 18, 199-209.	7.1	701
32	Insights into Autism Spectrum Disorder Genomic Architecture and Biology from 71 Risk Loci. <i>Neuron</i> , 2015, 87, 1215-1233.	3.8	1,219
33	A Genome-wide Association Study of Autism Using the Simons Simplex Collection: Does Reducing Phenotypic Heterogeneity in Autism Increase Genetic Homogeneity?. <i>Biological Psychiatry</i> , 2015, 77, 775-784.	0.7	133
34	The increasing prevalence of reported diagnoses of childhood psychiatric disorders: a descriptive multinational comparison. <i>European Child and Adolescent Psychiatry</i> , 2015, 24, 173-183.	2.8	201
35	Modest Impact on Risk for Autism Spectrum Disorder of Rare Copy Number Variants at 15q11.2, Specifically Breakpoints 1 to 2. <i>Autism Research</i> , 2014, 7, 355-362.	2.1	59
36	Genetics of Obsessive-Compulsive Disorder and Related Disorders. <i>Psychiatric Clinics of North America</i> , 2014, 37, 319-335.	0.7	62

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37	Obsessive-Compulsive Disorder. <i>Psychiatric Clinics of North America</i> , 2014, 37, 257-267.	0.7	126
38	Adjusting Head Circumference for Covariates in Autism: Clinical Correlates of a Highly Heritable Continuous Trait. <i>Biological Psychiatry</i> , 2013, 74, 576-584.	0.7	70
39	Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. <i>Nature Genetics</i> , 2013, 45, 984-994.	9.4	2,067
40	Characterization of SLITRK1 Variation in Obsessive-Compulsive Disorder. <i>PLoS ONE</i> , 2013, 8, e70376.	1.1	47
41	Common genetic variants, acting additively, are a major source of risk for autism. <i>Molecular Autism</i> , 2012, 3, 9.	2.6	357
42	Rare structural variation of synapse and neurotransmission genes in autism. <i>Molecular Psychiatry</i> , 2012, 17, 402-411.	4.1	151
43	Multiple Recurrent De Novo CNVs, Including Duplications of the 7q11.23 Williams Syndrome Region, Are Strongly Associated with Autism. <i>Neuron</i> , 2011, 70, 863-885.	3.8	1,146
44	SLITRK1 Binds 14-3-3 and Regulates Neurite Outgrowth in a Phosphorylation-Dependent Manner. <i>Biological Psychiatry</i> , 2009, 66, 918-925.	0.7	58
45	Identification of three mouse $\mu$ -opioid receptor (MOR) gene (Oprm1) splice variants containing a newly identified alternatively spliced exon. <i>Gene</i> , 2007, 388, 135-147.	1.0	30
46	Identification of five mouse $\mu$ -opioid receptor (MOR) gene (Oprm1) splice variants containing a newly identified alternatively spliced exon. <i>Gene</i> , 2007, 395, 98-107.	1.0	38
47	Transcriptional profiling of C57 and DBA strains of mice in the absence and presence of morphine. <i>BMC Genomics</i> , 2007, 8, 76.	1.2	39
48	The Genetics of Autism Spectrum Disorders. <i>NeuroMolecular Medicine</i> , 2006, 8, 451-460.	1.8	38
49	The genetic architecture of autism and related disorders. <i>Clinical Neuroscience Research</i> , 2006, 6, 161-168.	0.8	4
50	Candidate genes for anorexia nervosa in the 1p33-36 linkage region: serotonin 1D and delta opioid receptor loci exhibit significant association to anorexia nervosa. <i>Molecular Psychiatry</i> , 2003, 8, 397-406.	4.1	132
51	Genomewide Linkage Analyses of Bipolar Disorder: A New Sample of 250 Pedigrees from the National Institute of Mental Health Genetics Initiative. <i>American Journal of Human Genetics</i> , 2003, 73, 107-114.	2.6	202
52	A genetic association study of the mu opioid receptor and severe opioid dependence. <i>Psychiatric Genetics</i> , 2003, 13, 169-173.	0.6	152
53	Evidence for a Susceptibility Gene for Anorexia Nervosa on Chromosome 1. <i>American Journal of Human Genetics</i> , 2002, 70, 787-792.	2.6	199
54	Genetics of Mental Disorders: A Guide for Students, Clinicians and Researchers.. <i>Journal of the American Academy of Child and Adolescent Psychiatry</i> , 2001, 40, 1238-1239.	0.3	0

#	ARTICLE	IF	CITATIONS
55	Case control and family-based studies of tryptophan hydroxylase gene A218C polymorphism and suicidality in adolescents. American Journal of Medical Genetics Part A, 2001, 105, 451-457.	2.4	45
56	Symptoms of obsessive-compulsive disorder. American Journal of Psychiatry, 1997, 154, 911-917.	4.0	674
57	Tic-related vs. non-tic-related obsessive compulsive disorder. Anxiety, 1994, 1, 208-15.	0.5	113
58	Gender differences in substance use disorders. American Journal of Psychiatry, 1993, 150, 1707-1711.	4.0	206
59	De Novo Sequence and Copy Number Variants are Strongly Associated with Tourette Disorder and Implicate Cell Polarity in Pathogenesis. SSRN Electronic Journal, 0, , .	0.4	0