Thomas V Fernandez

List of Publications by Year
in descending order

62	6,478	126907 33	168389 53
papers	citations	h-index	g-index

67	67	67	10344
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Wholeâ€exome DNA sequencing in childhood anxiety disorders identifies rare de novo damaging coding variants. Depression and Anxiety, 2022, 39, 474-484.	4.1	5
2	Synaptic processes and immune-related pathways implicated in Tourette syndrome. Translational Psychiatry, 2021, 11, 56.	4.8	31
3	Early developmental asymmetries in cell lineage trees in living individuals. Science, 2021, 371, 1245-1248.	12.6	39
4	Whole-exome sequencing identifies genes associated with Touretteâ€™s disorder in multiplex families. Molecular Psychiatry, 2021, , .	7.9	16
5	Investigation of geneâ€“environment interactions in relation to tic severity. Journal of Neural Transmission, 2021, 128, 1757-1765.	2.8	2
6	De Novo Damaging DNA Coding Mutations Are Associated With Obsessive-Compulsive Disorder and Overlap With Touretteâ€™s Disorder and Autism. Biological Psychiatry, 2020, 87, 1035-1044.	1.3	59
7	Leveraging aggression risk gene expression in the developing and adult human brain to guide future precision interventions. Molecular Psychiatry, 2020, 26, 2680-2682.	7.9	0
8	Empiric Recurrence Risk Estimates for Chronic Tic Disorders: Implications for Genetic Counseling. Frontiers in Neurology, 2020, 11, 770.	2.4	7
9	Interrogating the Genetic Determinants of Touretteâ€™s Syndrome and Other Tic Disorders Through Genome-Wide Association Studies. American Journal of Psychiatry, 2019, 176, 217-227.	7.2	242
10	Investigation of previously implicated genetic variants in chronic tic disorders: a transmission disequilibrium test approach. European Archives of Psychiatry and Clinical Neuroscience, 2018, 268, 301-316.	3.2	23
11	De Novo Sequence and Copy Number Variants Are Strongly Associated with Tourette Disorder and Implicate Cell Polarity in Pathogenesis. Cell Reports, 2018, 24, 3441-3454.e12.	6.4	91
12	Genetic Insights Into ADHD Biology. Frontiers in Psychiatry, 2018, 9, 251.	2.6	28
13	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	12.6	1,085
14	Genetic susceptibility in obsessive-compulsive disorder. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 148, 767-781.	1.8	16
15	Tourette disorder and other tic disorders. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 147, 343-354.	1.8	24
16	Neurogenetic analysis of childhood disintegrative disorder. Molecular Autism, 2017, 8, 19.	4.9	19
17	De Novo Coding Variants Are Strongly Associated with Tourette Disorder. Neuron, 2017, 94, 486-499.e9.	8.1	155
18	Extended-Release Guanfacine Does Not Show a Large Effect on Tic Severity in Children with Chronic Tic Disorders. Journal of Child and Adolescent Psychopharmacology, 2017, 27, 762-770.	1.3	51

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19	Motor Stereotypies: A Pathophysiological Review. Frontiers in Neuroscience, 2017, 11, 171.	2.8	57
20	Transcriptome Analysis of the Human Striatum in Tourette Syndrome. Biological Psychiatry, 2016, 79, 372-382.	1.3	160
21	Whole-exome sequencing in obsessive-compulsive disorder identifies rare mutations in immunological and neurodevelopmental pathways. Translational Psychiatry, 2016, 6, e764-e764.	4.8	59
22	Prenatal and Perinatal Risk Factors and the Promise of Birth Cohort Studies. JAMA Psychiatry, 2016, 73, 1117.	11.0	3
23	The Origins of Tourette Syndrome: Prenatal Risk Factors and the Promise of Birth Cohort Studies. Journal of the American Academy of Child and Adolescent Psychiatry, 2016, 55, 751-753.	0.5	5
24	Pre- and perinatal complications in relation to Tourette syndrome and co-occurring obsessive-compulsive disorder and attention-deficit/hyperactivity disorder. Journal of Psychiatric Research, 2016, 82, 126-135.	3.1	36
25	What Makes You Tic? A New Lead in Tourette Syndrome Genetics. Biological Psychiatry, 2016, 79, 341-342.	1.3	2
26	Tourette Syndrome: Bridging the Gap between Genetics and Biology. Molecular Neuropsychiatry, 2015, 1, 156-164.	2.9	9
27	Cross-Disorder Genome-Wide Analyses Suggest a Complex Genetic Relationship Between Touretteâ€™s Syndrome and OCD. American Journal of Psychiatry, 2015, 172, 82-93.	7.2	117
28	No Evidence for Association of Autism with Rare Heterozygous Point Mutations in Contactin-Associated Protein-Like 2 (CNTNAP2), or in Other Contactin-Associated Proteins or Contactins. PLoS Genetics, 2015, 11, e1004852.	3.5	47
29	Touretteâ€™s Syndrome and Translational Clinical Science. Journal of the American Academy of Child and Adolescent Psychiatry, 2015, 54, 6-8.	0.5	1
30	Modeling non-syndromic autism and the impact of TRPC6 disruption in human neurons. Molecular Psychiatry, 2015, 20, 1350-1365.	7.9	175
31	The Tourette International Collaborative Genetics (TIC Genetics) study, finding the genes causing Tourette syndrome: objectives and methods. European Child and Adolescent Psychiatry, 2015, 24, 141-151.	4.7	41
32	Genetics of Tourette Syndrome. , 2015, , 169-189.		1
33	The inheritance of Tourette Disorder: A review. Journal of Obsessive-Compulsive and Related Disorders, 2014, 3, 380-385.	1.5	70
34	Copy Number Variation in Obsessive-Compulsive Disorder and Tourette Syndrome: A Cross-Disorder Study. Journal of the American Academy of Child and Adolescent Psychiatry, 2014, 53, 910-919.	0.5	111
35	Rare deleterious mutations of the gene EFR3A in autism spectrum disorders. Molecular Autism, 2014, 5, 31.	4.9	27
36	Genome-wide association study of Tourette's syndrome. Molecular Psychiatry, 2013, 18, 721-728.	7.9	161

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37	Gene variants associated with antisocial behaviour: a latent variable approach. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2013, 54, 1074-1085.	5.2	16
38	Transcranial Magnetic Stimulation of Wernickeâ€™s and Right Homologous Sites to Curtail â€œVoicesâ€: A Randomized Trial. Biological Psychiatry, 2013, 73, 1008-1014.	1.3	70
39	Genetic Susceptibility and Neurotransmitters in Tourette Syndrome. International Review of Neurobiology, 2013, 112, 155-177.	2.0	48
40	Partitioning the Heritability of Tourette Syndrome and Obsessive Compulsive Disorder Reveals Differences in Genetic Architecture. PLoS Genetics, 2013, 9, e1003864.	3.5	241
41	Data Blitz Debuts at the AACAP Annual Meeting. Journal of Child and Adolescent Psychopharmacology, 2013, 23, 306-307.	1.3	0
42	CNTN4: Contactin 4. , 2013, , 681-682.		1
43	Genetic Susceptibility in Tourette Syndrome. , 2013, , 137-155.		2
44	Rare Copy Number Variants in Tourette Syndrome Disrupt Genes in Histaminergic Pathways and Overlap with Autism. Biological Psychiatry, 2012, 71, 392-402.	1.3	167
45	Elevated Functional Connectivity Along a Corticostriatal Loop and the Mechanism of Auditory/Verbal Hallucinations in Patients with Schizophrenia. Biological Psychiatry, 2011, 69, 407-414.	1.3	145
46	Multiple Recurrent De Novo CNVs, Including Duplications of the 7q11.23 Williams Syndrome Region, Are Strongly Associated with Autism. Neuron, 2011, 70, 863-885.	8.1	1,146
47	Reply to 3p deletion and (skewed) literature review. American Journal of Medical Genetics, Part A, 2010, 152A, 1060-1060.	1.2	0
48	L-Histidine Decarboxylase and Tourette's Syndrome. New England Journal of Medicine, 2010, 362, 1901-1908.	27.0	304
49	Molecular characterization of a patient with 3p deletion syndrome and a review of the literature. American Journal of Medical Genetics, Part A, 2008, 146A, 2746-2752.	1.2	38
50	Disruption of Contactin 4 (CNTN4) Results in Developmental Delay and Other Features of 3p Deletion Syndrome. American Journal of Human Genetics, 2008, 82, 1385.	6.2	51
51	Disruption of Contactin 4 (CNTN4) Results in Developmental Delay and Other Features of 3p Deletion Syndrome. American Journal of Human Genetics, 2004, 74, 1286-1293.	6.2	162
52	Premorbid Speech and Language Impairments in Childhood-Onset Schizophrenia: Association With Risk Factors. American Journal of Psychiatry, 2000, 157, 794-800.	7.2	128
53	Lessons from childhood-onset schizophrenia. Brain Research Reviews, 2000, 31, 147-156.	9.0	90
54	Obstetrical Complications and Childhood-Onset Schizophrenia. American Journal of Psychiatry, 1999, 156, 1650-1652.	7.2	30

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55	Apolipoprotein E alleles in childhood-onset schizophrenia. , 1999, 88, 211-213.		24
56	Childhood-onset schizophrenia: progressive brain changes during adolescence. Biological Psychiatry, 1999, 46, 892-898.	1.3	202
57	Progressive Cortical Change During Adolescence in Childhood-Onset Schizophrenia. Archives of General Psychiatry, 1999, 56, 649.	12.3	361
58	Clinical and Neurobiological Correlates of Cytogenetic Abnormalities in Childhood-Onset Schizophrenia. American Journal of Psychiatry, 1999, 156, 1575-1579.	7.2	59
59	Brief Report: Association of Sex Chromosome Anomalies With Childhood-Onset Psychotic Disorders. Journal of the American Academy of Child and Adolescent Psychiatry, 1998, 37, 292-296.	0.5	51
60	Functional Differentiation Along the Anterior-Posterior Axis of the Hippocampus in Monkeys. Journal of Neurophysiology, 1998, 80, 1002-1005.	1.8	149
61	Genetics of Tourette syndrome and related disorders. , 0, , 336-346.		0
62	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