Samuel J R A Chawner

List of Publications by Year
in descending order

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19 papers	763 citations	759233 12 h-index	794594 19 g-index
26 all docs	26 docs citations	26 times ranked	1346 citing authors

#	Article	IF	Citations
1	Defining the Effect of the 16p11.2 Duplication on Cognition, Behavior, and Medical Comorbidities. JAMA Psychiatry, 2016, 73, 20.	11.0	195
2	Psychiatric disorders in children with 16p11.2 deletion and duplication. Translational Psychiatry, 2019, 9, 8.	4.8	93
3	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
4	Genotypeâ€“phenotype associations in children with copy number variants associated with high neuropsychiatric risk in the UK (IMAGINE-ID): a case-control cohort study. Lancet Psychiatry,the, 2019, 6, 493-505.	7.4	87
5	A Genetics-First Approach to Dissecting the Heterogeneity of Autism: Phenotypic Comparison of Autism Risk Copy Number Variants. American Journal of Psychiatry, 2021, 178, 77-86.	7.2	62
6	Childhood cognitive development in 22q11.2 deletion syndrome: Caseâ€“control study. British Journal of Psychiatry, 2017, 211, 223-230.	2.8	33
7	16p11.2 Locus modulates response to satiety before the onset of obesity. International Journal of Obesity, 2016, 40, 870-876.	3.4	31
8	Genes To Mental Health (G2MH): A Framework to Map the Combined Effects of Rare and Common Variants on Dimensions of Cognition and Psychopathology. American Journal of Psychiatry, 2022, 179, 189-203.	7.2	29
9	Cognitive deficits in childhood, adolescence and adulthood in 22q11.2 deletion syndrome and association with psychopathology. Translational Psychiatry, 2020, 10, 53.	4.8	28
10	The emergence of psychotic experiences in the early adolescence of 22q11.2 Deletion Syndrome. Journal of Psychiatric Research, 2019, 109, 10-17.	3.1	21
11	Neurodevelopmental Trajectories and Psychiatric Morbidity: Lessons Learned From the 22q11.2 Deletion Syndrome. Current Psychiatry Reports, 2021, 23, 13.	4.5	20
12	Attention deficit hyperactivity disorder symptoms as antecedents of later psychotic outcomes in 22q11.2 deletion syndrome. Schizophrenia Research, 2019, 204, 320-325.	2.0	19
13	Copy number variants (CNVs): a powerful tool for iPSC-based modelling of ASD. Molecular Autism, 2020, 11, 42.	4.9	14
14	Clinical evaluation of patients with a neuropsychiatric risk copy number variant. Current Opinion in Genetics and Development, 2021, 68, 26-34.	3.3	12
15	A GENETIC FIRST APPROACH TO DISSECTING THE HETEROGENEITY OF AUTISM: PHENOTYPIC COMPARISON OF AUTISM RISK COPY NUMBER VARIANTS. European Neuropsychopharmacology, 2019, 29, S783-S784.	0.7	6
16	The psychiatric phenotypes of 1q21 distal deletion and duplication. Translational Psychiatry, 2021, 11, 105.	4.8	6
17	Pan-european landscape of research into neurodevelopmental copy number variants: A survey by the MINDDS consortium. European Journal of Medical Genetics, 2020, 63, 104093.	1.3	2
18	Summaries of plenary, symposia, and oral sessions at the XXII World Congress of Psychiatric Genetics, Copenhagen, Denmark, 12â€“16 October 2014. Psychiatric Genetics, 2016, 26, 1-47.	1.1	0

#	Article	IF	Citations
19	O4.8. VULNERABLE PERIODS FOR COGNITIVE DEVELOPMENT IN INDIVIDUALS AT HIGH GENOMIC RISK OF SCHIZOPHRENIA. Schizophrenia Bulletin, 2018, 44, S86-S86.	4.3	0