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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	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
2	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
3	The psychiatric phenotypes of 1q21 distal deletion and duplication. Translational Psychiatry, 2021, 11, 105.	4.8	6
4	Neurodevelopmental Trajectories and Psychiatric Morbidity: Lessons Learned From the 22q11.2 Deletion Syndrome. Current Psychiatry Reports, 2021, 23, 13.	4.5	20
5	Clinical evaluation of patients with a neuropsychiatric risk copy number variant. Current Opinion in Genetics and Development, 2021, 68, 26-34.	3.3	12
6	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
7	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
8	Copy number variants (CNVs): a powerful tool for iPSC-based modelling of ASD. Molecular Autism, 2020, 11, 42.	4.9	14
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	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
11	Psychiatric disorders in children with 16p11.2 deletion and duplication. Translational Psychiatry, 2019, 9, 8.	4.8	93
12	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
13	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
14	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
15	O4.8. VULNERABLE PERIODS FOR COGNITIVE DEVELOPMENT IN INDIVIDUALS AT HIGH GENOMIC RISK OF SCHIZOPHRENIA. Schizophrenia Bulletin, 2018, 44, S86-S86.	4.3	0
16	Childhood cognitive development in 22q11.2 deletion syndrome: Caseâ€“control study. British Journal of Psychiatry, 2017, 211, 223-230.	2.8	33
17	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
18	16p11.2 Locus modulates response to satiety before the onset of obesity. International Journal of Obesity, 2016, 40, 870-876.	3.4	31

#	Article	IF	Citations
19	Defining the Effect of the 16p11.2 Duplication on Cognition, Behavior, and Medical Comorbidities. JAMA Psychiatry, 2016, 73, 20.	11.0	195