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List of Publications by Year in descending order

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24
papers

5,601
citations

331670

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610901

24
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25
all docs

25
docs citations

25
times ranked

11796
citing authors

#	ARTICLE	IF	CITATIONS
1	Dissecting the Shared Genetic Architecture of Suicide Attempt, Psychiatric Disorders, and Known Risk Factors. <i>Biological Psychiatry</i> , 2022, 91, 313-327.	1.3	114
2	Rare coding variants in ten genes confer substantial risk for schizophrenia. <i>Nature</i> , 2022, 604, 509-516.	27.8	326
3	Genome-wide association study of more than 40,000 bipolar disorder cases provides new insights into the underlying biology. <i>Nature Genetics</i> , 2021, 53, 817-829.	21.4	629
4	GWAS of Suicide Attempt in Psychiatric Disorders and Association With Major Depression Polygenic Risk Scores. <i>American Journal of Psychiatry</i> , 2019, 176, 651-660.	7.2	186
5	Population-based identity-by-descent mapping combined with exome sequencing to detect rare risk variants for schizophrenia. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2019, 180, 223-231.	1.7	2
6	Estimation of Genetic Correlation via Linkage Disequilibrium Score Regression and Genomic Restricted Maximum Likelihood. <i>American Journal of Human Genetics</i> , 2018, 102, 1185-1194.	6.2	119
7	Analysis of shared heritability in common disorders of the brain. <i>Science</i> , 2018, 360, .	12.6	1,085
8	Contribution of copy number variants to schizophrenia from a genome-wide study of 41,321 subjects. <i>Nature Genetics</i> , 2017, 49, 27-35.	21.4	838
9	Evidence for Genetic Overlap Between Schizophrenia and Age at First Birth in Women. <i>JAMA Psychiatry</i> , 2016, 73, 497.	11.0	51
10	The use of actigraphy in the monitoring of sleep and activity in ADHD: A meta-analysis. <i>Sleep Medicine Reviews</i> , 2016, 26, 9-20.	8.5	91
11	Modeling Linkage Disequilibrium Increases Accuracy of Polygenic Risk Scores. <i>American Journal of Human Genetics</i> , 2015, 97, 576-592.	6.2	1,098
12	The use of actigraphy in the monitoring of methylphenidate versus placebo in ADHD: a meta-analysis. <i>ADHD Attention Deficit and Hyperactivity Disorders</i> , 2014, 6, 49-58.	1.7	41
13	Partitioning Heritability of Regulatory and Cell-Type-Specific Variants across 11 Common Diseases. <i>American Journal of Human Genetics</i> , 2014, 95, 535-552.	6.2	569
14	Case-case genome-wide association analysis shows markers differentially associated with schizophrenia and bipolar disorder and implicates calcium channel genes. <i>Psychiatric Genetics</i> , 2011, 21, 1-4.	1.1	70
15	Evidence for the association of the DAOA (G72) gene with schizophrenia and bipolar disorder but not for the association of the DAO gene with schizophrenia. <i>Behavioral and Brain Functions</i> , 2009, 5, 28.	3.3	40
16	Confirmation of the genetic association between the U2AF homology motif (UHM) kinase 1 (UHMK1) gene and schizophrenia on chromosome 1q23.3. <i>European Journal of Human Genetics</i> , 2008, 16, 1275-1282.	2.8	18
17	A Genetic Association Study of Chromosome 11q22-24 in Two Different Samples Implicates the FXYP6 Gene, Encoding Phosphohippolin, in Susceptibility to Schizophrenia. <i>American Journal of Human Genetics</i> , 2007, 80, 664-672.	6.2	32
18	Failure to confirm allelic and haplotypic association between markers at the chromosome 6p22.3 dystrobrevin-binding protein 1 (DTNBP1) locus and schizophrenia. <i>Behavioral and Brain Functions</i> , 2007, 3, 50.	3.3	20

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19	Fine Mapping by Genetic Association Implicates the Chromosome 1q23.3 Gene UHMK1, Encoding a Serine/Threonine Protein Kinase, as a Novel Schizophrenia Susceptibility Gene. <i>Biological Psychiatry</i> , 2007, 61, 873-879.	1.3	35
20	Failure to Confirm Allelic Association Between Markers at the CAPON Gene Locus and Schizophrenia in a British Sample. <i>Biological Psychiatry</i> , 2006, 59, 195-197.	1.3	28
21	Genetic Association and Brain Morphology Studies and the Chromosome 8p22 Pericentriolar Material 1 (PCM1) Gene in Susceptibility to Schizophrenia. <i>Archives of General Psychiatry</i> , 2006, 63, 844.	12.3	82
22	Failure to confirm genetic association between schizophrenia and markers on chromosome 1q23.3 in the region of the gene encoding the regulator of G-protein signaling 4 protein (RGS4). <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2006, 141B, 296-300.	1.7	32
23	The Epsin 4 Gene on Chromosome 5q, Which Encodes the Clathrin-Associated Protein Enthoprotin, Is Involved in the Genetic Susceptibility to Schizophrenia. <i>American Journal of Human Genetics</i> , 2005, 76, 902-907.	6.2	62
24	Genetic association studies of schizophrenia using the 8p21-22 genes: prepronociceptin (PNOC), neuronal nicotinic cholinergic receptor alpha polypeptide 2 (CHRNA2) and arylamine N-acetyltransferase 1 (NAT1). <i>European Journal of Human Genetics</i> , 2001, 9, 469-472.	2.8	29