Ann E Pulver

List of Publications by Year
in descending order

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33 papers	3,715 citations	331670 21 h-index	454955 30 g-index
34 all docs	34 docs citations	34 times ranked	5802 citing authors

#	Article	IF	Citations
1	The benefit of diagnostic whole genome sequencing in schizophrenia and other psychotic disorders. Molecular Psychiatry, 2022, 27, 1435-1447.	7.9	12
2	Mapping genomic loci implicates genes and synaptic biology in schizophrenia. Nature, 2022, 604, 502-508.	27.8	929
3	De novo variation in bipolar disorder. Molecular Psychiatry, 2021, 26, 4127-4136.	7.9	18
4	S184. IN SILICO PREDICTION OF T-CELL-MEDIATED MOLECULAR MIMICRY IN TOXOPLASMOSIS AND SCHIZOPHRENIA. Schizophrenia Bulletin, 2020, 46, S108-S108.	4.3	0
5	New insights into tardive dyskinesia genetics: Implementation of whole-exome sequencing approach. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2019, 94, 109659.	4.8	9
6	Genome-wide association study in two populations to determine genetic variants associated with Toxoplasma gondii infection and relationship to schizophrenia risk. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2019, 92, 133-147.	4.8	26
7	HLA typing using genome wide data reveals susceptibility types for infections in a psychiatric disease enriched sample. Brain, Behavior, and Immunity, 2018, 70, 203-213.	4.1	10
8	Cognitive and functional deficits in bipolar disorder and schizophrenia as a function of the presence and history of psychosis. Bipolar Disorders, 2018, 20, 604-613.	1.9	31
9	Insights into the genetic epidemiology of Crohn's and rare diseases in the Ashkenazi Jewish population. PLoS Genetics, 2018, 14, e1007329.	3.5	66
10	Revisiting the prevalence of nonclassic congenital adrenal hyperplasia in US Ashkenazi Jews and Caucasians. Genetics in Medicine, 2017, 19, 1276-1279.	2.4	90
11	Thorase variants are associated with defects in glutamatergic neurotransmission that can be rescued by Perampanel. Science Translational Medicine, 2017, 9, .	12.4	20
12	Contribution of copy number variants to schizophrenia from a genome-wide study of 41,321 subjects. Nature Genetics, 2017, 49, 27-35.	21.4	838
13	ADHD and executive functioning deficits in OCD youths who hoard. Journal of Psychiatric Research, 2016, 82, 141-148.	3.1	24
14	Neuregulin 3 Knockout Mice Exhibit Behaviors Consistent with Psychotic Disorders. Molecular Neuropsychiatry, 2016, 2, 79-87.	2.9	27
15	Common genetic variation and schizophrenia polygenic risk influence neurocognitive performance in young adulthood. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2015, 168, 392-401.	1.7	52
16	Identification and Functional Studies of Regulatory Variants Responsible for the Association of <i>NRG3</i> with a Delusion Phenotype in Schizophrenia. Molecular Neuropsychiatry, 2015, 1, 36-46.	2.9	14
17	Genomeâ€wide association study of schizophrenia in Ashkenazi Jews. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2015, 168, 649-659.	1.7	203
18	Improving the understanding of the link between cognition and functional capacity in schizophrenia and bipolar disorder. Schizophrenia Research, 2015, 169, 121-127.	2.0	13

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19	Infection and Inflammation in Schizophrenia and Bipolar Disorder: A Genome Wide Study for Interactions with Genetic Variation. PLoS ONE, 2015, 10, e0116696.	2.5	92
20	Exome Sequencing in 53 Sporadic Cases of Schizophrenia Identifies 18 Putative Candidate Genes. PLoS ONE, 2014, 9, e112745.	2.5	79
21	Reciprocal Duplication of the Williams-Beuren Syndrome Deletion on Chromosome 7q11.23 Is Associated with Schizophrenia. Biological Psychiatry, 2014, 75, 371-377.	1.3	66
22	Hoarding in children and adolescents with obsessiveâ€“compulsive disorder. Journal of Obsessive-Compulsive and Related Disorders, 2014, 3, 325-331.	1.5	31
23	An indirect test of the new mutation hypothesis associating advanced paternal age with the etiology of schizophrenia. American Journal of Medical Genetics Part A, 2004, 124B, 6-9.	2.4	22
24	Genomewide Linkage Scan for Schizophrenia Susceptibility Loci among Ashkenazi Jewish Families Shows Evidence of Linkage on Chromosome 10q22. American Journal of Human Genetics, 2003, 73, 601-611.	6.2	99
25	No evidence for linkage between schizophrenia and markers at chromosome 15q13-14. , 1999, 88, 109-112.		59
26	Chromosome workshop: Chromosomes 11, 14, and 15. American Journal of Medical Genetics Part A, 1999, 88, 244-254.	2.4	53
27	Lack of linkage or association between schizophrenia and the polymorphic trinucleotide repeat within the KCNN3 gene on chromosome 1q21. , 1999, 88, 348-351.		33
28	Schizophrenia susceptibility loci on chromosomes 13q32 and 8p21. Nature Genetics, 1998, 20, 70-73.	21.4	506
29	Schizophrenia susceptibility and chromosome 6p24â€“22. Nature Genetics, 1995, 11, 235-236.	21.4	123
30	Report from the Maryland epidemiology schizophrenia linkage study: No evidence for linkage between schizophrenia and a number of candidate and other genomic regions using a complex dominant model. American Journal of Medical Genetics Part A, 1994, 54, 345-353.	2.4	27
31	Estimating effects of proband characteristics on familial risk: II. The association between age at onset and familial risk in the Maryland schizophrenia sample. Genetic Epidemiology, 1991, 8, 339-350.	1.3	45
32	Availability of schizophrenic patients and their families for genetic linkage studies: Findings from the Maryland epidemiology sample. Genetic Epidemiology, 1989, 6, 671-680.	1.3	35
33	Risk Factors in Schizophrenia: Season of Birth in Maryland, USA. British Journal of Psychiatry, 1983, 143, 389-396.	2.8	52