

Anjene M Addington

List of Publications by Year in descending order

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Version: 2024-02-01

17
papers

3,753
citations

706676

14
h-index

993246

17
g-index

19
all docs

19
docs citations

19
times ranked

6948
citing authors

#	ARTICLE	IF	CITATIONS
1	Introduction to special section on Leveraging Electronic Health Records for psychiatric genetic research. , 2018, 177, 599-600.		0
2	The Open Translational Science in Schizophrenia (OPTICS) project: an open-science project bringing together Janssen clinical trial and NIMH data. NPJ Schizophrenia, 2018, 4, 14.	2.0	1
3	Whole genome sequencing in psychiatric disorders: the WGSPD consortium. Nature Neuroscience, 2017, 20, 1661-1668.	7.1	122
4	Convergence of Advances in Genomics, Team Science, and Repositories as Drivers of Progress in Psychiatric Genomics. Biological Psychiatry, 2015, 77, 6-14.	0.7	18
5	Microduplications disrupting the MYT1L gene (2p25.3) are associated with schizophrenia. Psychiatric Genetics, 2012, 22, 206-209.	0.6	42
6	Annual Research Review: Impact of advances in genetics in understanding developmental psychopathology. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2012, 53, 510-518.	3.1	32
7	A Novel Microduplication in the Neurodevelopmental Gene <i>SRGAP3</i> That Segregates with Psychotic Illness in the Family of a COS Proband. Case Reports in Genetics, 2011, 2011, 1-5.	0.1	20
8	Direct Measure of the De Novo Mutation Rate in Autism and Schizophrenia Cohorts. American Journal of Human Genetics, 2010, 87, 316-324.	2.6	222
9	De novo mutations in the gene encoding the synaptic scaffolding protein <i>SHANK3</i> in patients ascertained for schizophrenia. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 7863-7868.	3.3	361
10	The genetics of childhood-onset schizophrenia: When madness strikes the prepubescent. Current Psychiatry Reports, 2009, 11, 156-161.	2.1	79
11	Microduplications of 16p11.2 are associated with schizophrenia. Nature Genetics, 2009, 41, 1223-1227.	9.4	646
12	Sequencing and Analyzing the t(1;7) Reciprocal Translocation Breakpoints Associated with a Case of Childhood-onset Schizophrenia/Autistic Disorder. Journal of Autism and Developmental Disorders, 2008, 38, 668-677.	1.7	6
13	Rare Structural Variants Disrupt Multiple Genes in Neurodevelopmental Pathways in Schizophrenia. Science, 2008, 320, 539-543.	6.0	1,654
14	Molecular genetic studies of ADHD: 1991 to 2004. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2006, 141B, 551-565.	1.1	60
15	Support for association between ADHD and two candidate genes: <i>NET1</i> and <i>DRD1</i> . American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2005, 134B, 67-72.	1.1	180
16	Pervasive developmental disorder and childhood-onset schizophrenia: comorbid disorder or a phenotypic variant of a very early onset illness?. Biological Psychiatry, 2004, 55, 989-994.	0.7	164
17	Polymorphisms in the 13q33.2 gene <i>G72/G30</i> are associated with childhood-onset schizophrenia and psychosis not otherwise specified. Biological Psychiatry, 2004, 55, 976-980.	0.7	143