Michael B Miller

List of Publications by Year in descending order

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50 5,889 papers citations

9 30
ns h-index

52 g-index

53 all docs

53 docs citations 53 times ranked 11072 citing authors

#	Article	IF	CITATIONS
1	Polygenic prediction of educational attainment within and between families from genome-wide association analyses in 3 million individuals. Nature Genetics, 2022, 54, 437-449.	9.4	215
2	Brain Somatic Mutation in Aging and Alzheimer's Disease. Annual Review of Genomics and Human Genetics, 2021, 22, 239-256.	2.5	32
3	Genetic evidence of assortative mating in humans. Nature Human Behaviour, 2017, 1, .	6.2	242
4	Genetic variants associated with subjective well-being, depressive symptoms, and neuroticism identified through genome-wide analyses. Nature Genetics, 2016, 48, 624-633.	9.4	870
5	Identification of Common Genetic Variants Influencing Spontaneous Dizygotic Twinning and Female Fertility. American Journal of Human Genetics, 2016, 98, 898-908.	2.6	89
6	Genome-wide association study identifies 74 loci associated with educational attainment. Nature, 2016, 533, 539-542.	13.7	1,204
7	Genetic variants linked to education predict longevity. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 13366-13371.	3.3	110
8	Concept and design of a genome-wide association genotyping array tailored for transplantation-specific studies. Genome Medicine, 2015, 7, 90.	3.6	49
9	Rare variant genotype imputation with thousands of study-specific whole-genome sequences: implications for cost-effective study designs. European Journal of Human Genetics, 2015, 23, 975-983.	1.4	92
10	A Structural and Functional Comparison Between Infectious and Non-Infectious Autocatalytic Recombinant PrP Conformers. PLoS Pathogens, 2015, 11, e1005017.	2.1	38
11	Heritability and molecular genetic basis of electrodermal activity: A genomeâ€wide association study. Psychophysiology, 2014, 51, 1259-1271.	1.2	18
12	Heritability and molecular genetic basis of antisaccade eye tracking error rate: A genomeâ€wide association study. Psychophysiology, 2014, 51, 1272-1284.	1.2	20
13	Heritability and molecular genetic basis of acoustic startle eye blink and affectively modulated startle response: A genomeâ€wide association study. Psychophysiology, 2014, 51, 1285-1299.	1.2	35
14	Premorbid risk factors for major depressive disorder: Are they associated with early onset and recurrent course?. Development and Psychopathology, 2014, 26, 1477-1493.	1.4	54
15	Common genetic variants associated with cognitive performance identified using the proxy-phenotype method. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 13790-13794.	3.3	244
16	Gammaâ€Aminobutyric Acid System Genes—No Evidence for a Role in Alcohol Use and Abuse in a Communityâ€Based Sample. Alcoholism: Clinical and Experimental Research, 2014, 38, 938-947.	1.4	14
17	Heritability and molecularâ€genetic basis of resting <scp>EEG</scp> activity: A genomeâ€wide association study. Psychophysiology, 2014, 51, 1225-1245.	1.2	46
18	Heritability and molecularâ€genetic basis of the <scp>P</scp> 3 eventâ€related brain potential: A genomeâ€wide association study. Psychophysiology, 2014, 51, 1246-1258.	1.2	32

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19	Low-frequency copy-number variants and general cognitive ability: No evidence of association. Intelligence, 2014, 42, 98-106.	1.6	10
20	Rare Nonsynonymous Exonic Variants in Addiction and Behavioral Disinhibition. Biological Psychiatry, 2014, 75, 783-789.	0.7	41
21	Results of a "GWAS Plus:―General Cognitive Ability Is Substantially Heritable and Massively Polygenic. PLoS ONE, 2014, 9, e112390.	1.1	41
22	Cofactor Molecules Induce Structural Transformation during Infectious Prion Formation. Structure, 2013, 21, 2061-2068.	1.6	64
23	GWAS of 126,559 Individuals Identifies Genetic Variants Associated with Educational Attainment. Science, 2013, 340, 1467-1471.	6.0	750
24	A Rapid Gene-Based Genome-Wide Association Test with Multivariate Traits. Human Heredity, 2013, 76, 53-63.	0.4	17
25	Three Mutually Informative Ways to Understand the Genetic Relationships Among Behavioral Disinhibition, Alcohol Use, Drug Use, Nicotine Use/Dependence, and Their Co-occurrence: Twin Biometry, GCTA, and Genome-Wide Scoring. Behavior Genetics, 2013, 43, 97-107.	1.4	91
26	A Genome-Wide Association Study of Behavioral Disinhibition. Behavior Genetics, 2013, 43, 363-373.	1.4	119
27	Assumptions in studies of heritability and genotype–phenotype association. Behavioral and Brain Sciences, 2012, 35, 372-373.	0.4	7
28	The Minnesota Center for Twin and Family Research Genome-Wide Association Study. Twin Research and Human Genetics, 2012, 15, 767-774.	0.3	70
29	Superparamagnetic Nanoparticle Capture of Prions for Amplification. Journal of Virology, 2011, 85, 2813-2817.	1.5	19
30	A Rapid Generalized Least Squares Model for a Genome-Wide Quantitative Trait Association Analysis in Families. Human Heredity, 2011, 71, 67-82.	0.4	27
31	An Assessment of the Individual and Collective Effects of Variants on Height Using Twins and a Developmentally Informative Study Design. PLoS Genetics, 2011, 7, e1002413.	1.5	11
32	Dissociation of Infectivity from Seeding Ability in Prions with Alternate Docking Mechanism. PLoS Pathogens, 2011, 7, e1002128.	2.1	43
33	Genotyping Errors and Their Impact on Genetic Analysis. Advances in Genetics, 2008, 60, 141-152.	0.8	7
34	Genome scan of glomerular filtration rate and albuminuria: the HyperGEN study. Nephrology Dialysis Transplantation, 2007, 22, 763-771.	0.4	34
35	Genotype-by-Sex Interaction on Fasting Insulin Concentration: The HyperGEN Study. Diabetes, 2007, 56, 137-142.	0.3	19
36	Genetic Analysis Workshop 15: simulation of a complex genetic model for rheumatoid arthritis in nuclear families including a dense SNP map with linkage disequilibrium between marker loci and trait loci. BMC Proceedings, 2007, 1, S4.	1.8	31

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37	A Whole Genome Scan for Pulse Pressure/Stroke Volume Ratio in African Americans: The HyperGEN Study. American Journal of Hypertension, 2007, 20, 398-402.	1.0	25
38	Sex-specific effects of ACE I/D and AGT-M235T on pulse pressure: the HyperGEN Study. Human Genetics, 2007, 122, 33-40.	1.8	6
39	Genome-Wide Linkage Analysis for Loci Affecting Pulse Pressure. Hypertension, 2005, 46, 1286-1293.	1.3	42
40	Further Evidence of a Quantitative Trait Locus on Chromosome 18 Influencing Postural Change in Systolic Blood Pressure: The Hypertension Genetic Epidemiology Network (HyperGEN) Study. American Journal of Hypertension, 2005, 18, 672-678.	1.0	34
41	Linkage Analysis of Diabetes Status Among Hypertensive Families: The Hypertension Genetic Epidemiology Network Study. Diabetes, 2004, 53, 3307-3312.	0.3	19
42	Evidence for a Gene on Chromosome 13 Influencing Postural Systolic Blood Pressure Change and Body Mass Index. Hypertension, 2004, 43, 780-784.	1.3	26
43	Refined Mapping of Suggestive Linkage to Renal Function in African Americans: The HyperGEN Study. American Journal of Human Genetics, 2002, 71, 204-205.	2.6	14
44	A family history study of male sexual orientation using three independent samples. Behavior Genetics, 1999, 29, 79-86.	1.4	80
45	The use of likelihood-based confidence intervals in genetic models. Behavior Genetics, 1997, 27, 113-120.	1.4	284
46	Genomic scanning and the transmission/disequilibrium test: Analysis of error rates. Genetic Epidemiology, 1997, 14, 854-856.	0.6	2
47	Task difficulty and cognitive deficits in schizophrenia Journal of Abnormal Psychology, 1995, 104, 251-258.	2.0	61
48	Computing conditional recombination probabilities given marker information. Genetic Epidemiology, 1995, 12, 883-888.	0.6	0
49	Coefficient alpha: A basic introduction from the perspectives of classical test theory and structural equation modeling. Structural Equation Modeling, 1995, 2, 255-273.	2.4	289
50	Do Children and the Elderly Show Heightened Semantic Priming? How to Answer the Question. Developmental Review, 1994, 14, 159-185.	2.6	165