## Sarah E Bergen

List of Publications by Year in descending order

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SADAH F REDCEN

#	Article	IF	CITATIONS
1	Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. Nature Genetics, 2013, 45, 984-994.	9.4	2,067
2	Genome-wide association study identifies five new schizophrenia loci. Nature Genetics, 2011, 43, 969-976.	9.4	1,758
3	Genome-wide association analysis identifies 13 new risk loci for schizophrenia. Nature Genetics, 2013, 45, 1150-1159.	9.4	1,395
4	A polygenic burden of rare disruptive mutations in schizophrenia. Nature, 2014, 506, 185-190.	13.7	1,305
5	Large-scale genome-wide association analysis of bipolar disorder identifies a new susceptibility locus near ODZ4. Nature Genetics, 2011, 43, 977-983.	9.4	1,283
6	Genome-wide association study identifies 30 loci associated with bipolar disorder. Nature Genetics, 2019, 51, 793-803.	9.4	1,191
7	Modeling Linkage Disequilibrium Increases Accuracy of Polygenic Risk Scores. American Journal of Human Genetics, 2015, 97, 576-592.	2.6	1,098
8	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	6.0	1,085
9	Genomic Relationships, Novel Loci, and Pleiotropic Mechanisms across Eight Psychiatric Disorders. Cell, 2019, 179, 1469-1482.e11.	13.5	935
10	Mapping genomic loci implicates genes and synaptic biology in schizophrenia. Nature, 2022, 604, 502-508.	13.7	929
11	Contribution of copy number variants to schizophrenia from a genome-wide study of 41,321 subjects. Nature Genetics, 2017, 49, 27-35.	9.4	838
12	Psychiatric genome-wide association study analyses implicate neuronal, immune and histone pathways. Nature Neuroscience, 2015, 18, 199-209.	7.1	701
13	Genome-wide association study of more than 40,000 bipolar disorder cases provides new insights into the underlying biology. Nature Genetics, 2021, 53, 817-829.	9.4	629
14	Genomic Dissection of Bipolar Disorder and Schizophrenia, Including 28 Subphenotypes. Cell, 2018, 173, 1705-1715.e16.	13.5	623
15	Partitioning Heritability of Regulatory and Cell-Type-Specific Variants across 11 Common Diseases. American Journal of Human Genetics, 2014, 95, 535-552.	2.6	569
16	Discovery and Statistical Genotyping of Copy-Number Variation from Whole-Exome Sequencing Depth. American Journal of Human Genetics, 2012, 91, 597-607.	2.6	513
17	Age-Related Changes in Heritability of Behavioral Phenotypes Over Adolescence and Young Adulthood: A Meta-Analysis. Twin Research and Human Genetics, 2007, 10, 423-433.	0.3	398
18	Copy Number Variants in Schizophrenia: Confirmation of Five Previous Findings and New Evidence for 3q29 Microdeletions and VIPR2 Duplications. American Journal of Psychiatry, 2011, 168, 302-316.	4.0	398

Sarah E Bergen

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19	Relationship of Brain-Derived Neurotrophic Factor and Its Receptor TrkB to Altered Inhibitory Prefrontal Circuitry in Schizophrenia. Journal of Neuroscience, 2005, 25, 372-383.	1.7	390
20	Genomic and drug target evaluation of 90 cardiovascular proteins in 30,931 individuals. Nature Metabolism, 2020, 2, 1135-1148.	5.1	327
21	Copy number variation in schizophrenia in Sweden. Molecular Psychiatry, 2014, 19, 762-773.	4.1	257
22	Extremely low-coverage sequencing and imputation increases power for genome-wide association studies. Nature Genetics, 2012, 44, 631-635.	9.4	239
23	Genome-wide association study in a Swedish population yields support for greater CNV and MHC involvement in schizophrenia compared with bipolar disorder. Molecular Psychiatry, 2012, 17, 880-886.	4.1	230
24	Joint Analysis of Psychiatric Disorders Increases Accuracy of Risk Prediction for Schizophrenia, Bipolar Disorder, and Major Depressive Disorder. American Journal of Human Genetics, 2015, 96, 283-294.	2.6	225
25	Genetic influences on schizophrenia and subcortical brain volumes: large-scale proof of concept. Nature Neuroscience, 2016, 19, 420-431.	7.1	204
26	Genome-wide association study of 40,000 individuals identifies two novel loci associated with bipolar disorder. Human Molecular Genetics, 2016, 25, 3383-3394.	1.4	182
27	Evidence for genetic heterogeneity between clinical subtypes of bipolar disorder. Translational Psychiatry, 2017, 7, e993-e993.	2.4	162
28	Genome-wide association study of borderline personality disorder reveals genetic overlap with bipolar disorder, major depression and schizophrenia. Translational Psychiatry, 2017, 7, e1155-e1155.	2.4	150
29	The Genetics of the Mood Disorder Spectrum: Genome-wide Association Analyses of More Than 185,000 Cases and 439,000 Controls. Biological Psychiatry, 2020, 88, 169-184.	0.7	137
30	Bipolar disorder and its relation to major psychiatric disorders: a familyâ€based study in the <scp>S</scp> wedish population. Bipolar Disorders, 2015, 17, 184-193.	1.1	119
31	Estimation of Genetic Correlation via Linkage Disequilibrium Score Regression and Genomic Restricted Maximum Likelihood. American Journal of Human Genetics, 2018, 102, 1185-1194.	2.6	119
32	Validation of Electronic Health Record Phenotyping of Bipolar Disorder Cases and Controls. American Journal of Psychiatry, 2015, 172, 363-372.	4.0	116
33	Genome-Wide Association Study of Clinical Dimensions of Schizophrenia: Polygenic Effect on Disorganized Symptoms. American Journal of Psychiatry, 2012, 169, 1309-1317.	4.0	112
34	High density methylation QTL analysis in human blood via next-generation sequencing of the methylated genomic DNA fraction. Genome Biology, 2015, 16, 291.	3.8	112
35	Common DISC1 Polymorphisms Disrupt Wnt/GSK3β Signaling and Brain Development. Neuron, 2011, 72, 545-558.	3.8	110
36	Schizophrenia and subsequent neighborhood deprivation: revisiting the social drift hypothesis using population, twin and molecular genetic data. Translational Psychiatry, 2016, 6, e796-e796.	2.4	110

SARAH E BERGEN

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37	Joint Contributions of Rare Copy Number Variants and Common SNPs to Risk for Schizophrenia. American Journal of Psychiatry, 2019, 176, 29-35.	4.0	104
38	A Comparison of Ten Polygenic Score Methods for Psychiatric Disorders Applied Across Multiple Cohorts. Biological Psychiatry, 2021, 90, 611-620.	0.7	103
39	Suicidal Behavior During Lithium and Valproate Treatment: A Within-Individual 8-Year Prospective Study of 50,000 Patients With Bipolar Disorder. American Journal of Psychiatry, 2017, 174, 795-802.	4.0	98
40	Cerebrospinal fluid metabolomics identifies a key role of isocitrate dehydrogenase in bipolar disorder: evidence in support of mitochondrial dysfunction hypothesis. Molecular Psychiatry, 2016, 21, 1504-1510.	4.1	95
41	Pyramidal cell size reduction in schizophrenia: evidence for involvement of auditory feedforward circuits. Biological Psychiatry, 2004, 55, 1128-1137.	0.7	87
42	Anatomical Evidence of Impaired Feedforward Auditory Processing in Schizophrenia. Biological Psychiatry, 2007, 61, 854-864.	0.7	73
43	Cis-acting regulation of brain-specific ANK3 gene expression by a genetic variant associated with bipolar disorder. Molecular Psychiatry, 2013, 18, 922-929.	4.1	73
44	Genome-wide association studies of schizophrenia. Current Opinion in Psychiatry, 2012, 25, 76-82.	3.1	72
45	A genome-wide association study of kynurenic acid in cerebrospinal fluid: implications for psychosis and cognitive impairment in bipolar disorder. Molecular Psychiatry, 2016, 21, 1342-1350.	4.1	71
46	Implication of a Rare Deletion at Distal 16p11.2 in Schizophrenia. JAMA Psychiatry, 2013, 70, 253.	6.0	69
47	Genome-wide association study identifies SESTD1 as a novel risk gene for lithium-responsive bipolar disorder. Molecular Psychiatry, 2016, 21, 1290-1297.	4.1	69
48	Genetic modifiers and subtypes in schizophrenia: Investigations of age at onset, severity, sex and family history. Schizophrenia Research, 2014, 154, 48-53.	1.1	68
49	Allelic differences between Europeans and Chinese for CREB1 SNPs and their implications in gene expression regulation, hippocampal structure and function, and bipolar disorder susceptibility. Molecular Psychiatry, 2014, 19, 452-461.	4.1	61
50	Sex-Dependent Shared and Nonshared Genetic Architecture Across Mood and Psychotic Disorders. Biological Psychiatry, 2022, 91, 102-117.	0.7	61
51	Environmental Risk Factors for Schizophrenia and Bipolar Disorder and Their Relationship to Genetic Risk: Current Knowledge and Future Directions. Frontiers in Genetics, 2021, 12, 686666.	1.1	61
52	The protocadherin 17 gene affects cognition, personality, amygdala structure and function, synapse development and risk of major mood disorders. Molecular Psychiatry, 2018, 23, 400-412.	4.1	60
53	A principal component approach to improve association testing with polygenic risk scores. Genetic Epidemiology, 2020, 44, 676-686.	0.6	56
54	Longitudinal Cortical Thickness Changes in Bipolar Disorder and the Relationship to Genetic Risk, Mania, and Lithium Use. Biological Psychiatry, 2020, 87, 271-281.	0.7	46

SARAH E BERGEN

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55	Contribution of Rare Copy Number Variants toÂBipolar Disorder Risk Is Limited to Schizoaffective Cases. Biological Psychiatry, 2019, 86, 110-119.	0.7	45
56	Polymorphisms in SLC6A4, PAH, GABRB3, and MAOB and modification of psychotic disorder features. Schizophrenia Research, 2009, 109, 94-97.	1.1	38
57	The association between Darier disease, bipolar disorder, and schizophrenia revisited: a population-based family study. Bipolar Disorders, 2015, 17, 340-344.	1.1	37
58	Specificity in Etiology of Subtypes of Bipolar Disorder: Evidence From a Swedish Population-Based Family Study. Biological Psychiatry, 2018, 84, 810-816.	0.7	37
59	Association of Youth Depression With Subsequent Somatic Diseases and Premature Death. JAMA Psychiatry, 2021, 78, 302.	6.0	35
60	A Loss-of-Function Variant in a Minor Isoform of ANK3 Protects Against Bipolar Disorder and Schizophrenia. Biological Psychiatry, 2016, 80, 323-330.	0.7	31
61	Socioeconomic Status and Social Support Following Illicit Drug Use: Causal Pathways or Common Liability?. Twin Research and Human Genetics, 2008, 11, 266-274.	0.3	29
62	Analysis of schizophrenia-related genes and electrophysiological measures reveals ZNF804A association with amplitude of P300b elicited by novel sounds. Translational Psychiatry, 2014, 4, e346-e346.	2.4	29
63	Elevated expression of a minor isoform of ANK3 is a risk factor for bipolar disorder. Translational Psychiatry, 2018, 8, 210.	2.4	24
64	Modifiers and Subtype-Specific Analyses in Whole-Genome Association Studies: A Likelihood Framework. Human Heredity, 2011, 72, 10-20.	0.4	20
65	Convergent Lines of Evidence Support LRP8 as a Susceptibility Gene for Psychosis. Molecular Neurobiology, 2016, 53, 6608-6619.	1.9	20
66	The role of ADHD genetic risk in mid-to-late life somatic health conditions. Translational Psychiatry, 2022, 12, 152.	2.4	20
67	Lack of Support for the Genes by Early Environment Interaction Hypothesis in the Pathogenesis of Schizophrenia. Schizophrenia Bulletin, 2022, 48, 20-26.	2.3	19
68	Technological readiness and implementation of genomicâ€driven precision medicine for complex diseases. Journal of Internal Medicine, 2021, 290, 602-620.	2.7	18
69	Familial co-aggregation of schizophrenia and eating disorders in Sweden and Denmark. Molecular Psychiatry, 2021, 26, 5389-5397.	4.1	17
70	Polygenic risk for anxiety influences anxiety comorbidity and suicidal behavior in bipolar disorder. Translational Psychiatry, 2020, 10, 298.	2.4	16
71	Combined Whole Methylome and Genomewide Association Study Implicates <i>CNTN4</i> in Alcohol Use. Alcoholism: Clinical and Experimental Research, 2015, 39, 1396-1405.	1.4	15
72	Investigating rare pathogenic/likely pathogenic exonic variation in bipolar disorder. Molecular Psychiatry, 2021, 26, 5239-5250.	4.1	15

SARAH E BERGEN

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73	Association Study of 167 Candidate Genes for Schizophrenia Selected by a Multi-Domain Evidence-Based Prioritization Algorithm and Neurodevelopmental Hypothesis. PLoS ONE, 2013, 8, e67776.	1.1	15
74	Acute intermittent porphyria: Comorbidity and shared familial risks with schizophrenia and bipolar disorder in Sweden. British Journal of Psychiatry, 2015, 207, 556-557.	1.7	14
75	Association of family history of schizophrenia and clinical outcomes in individuals with eating disorders. Psychological Medicine, 2021, , 1-8.	2.7	14
76	Novel disease associations with schizophrenia genetic risk revealed in ~400,000 UK Biobank participants. Molecular Psychiatry, 2022, 27, 1448-1454.	4.1	13
77	Impact of a <i>cis</i> -associated gene expression SNP on chromosome 20q11.22 on bipolar disorder susceptibility, hippocampal structure and cognitive performance. British Journal of Psychiatry, 2016, 208, 128-137.	1.7	11
78	Genes, biomarkers, and clinical features associated with the course of bipolar disorder. European Neuropsychopharmacology, 2019, 29, 1152-1160.	0.3	11
79	Genetic variation in 117 myelination-related genes in schizophrenia: Replication of association to lipid biosynthesis genes. Scientific Reports, 2018, 8, 6915.	1.6	10
80	Characterization of Single Gene Copy Number Variants in Schizophrenia. Biological Psychiatry, 2020, 87, 736-744.	0.7	10
81	Genetic risk for bipolar disorder and schizophrenia predicts structure and function of the ventromedial prefrontal cortex. Journal of Psychiatry and Neuroscience, 2021, 46, E441-E450.	1.4	10
82	Novel gene-brain structure relationships in psychotic disorder revealed using parallel independent component analyses. Schizophrenia Research, 2017, 182, 74-83.	1.1	9
83	Identification of a Bipolar Disorder Vulnerable Gene CHDH at 3p21.1. Molecular Neurobiology, 2017, 54, 5166-5176.	1.9	9
84	Crossâ€sex shifts in two brain imaging phenotypes and their relation to polygenic scores for sameâ€sex sexual behavior: A study of 18,645 individuals from the UK Biobank. Human Brain Mapping, 2021, 42, 2292-2304.	1.9	8
85	Genome-wide study of immune biomarkers in cerebrospinal fluid and serum from patients with bipolar disorder and controls. Translational Psychiatry, 2020, 10, 58.	2.4	8
86	Nationalâ€scale precision medicine for psychiatric disorders in Sweden. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2018, 177, 630-634.	1.1	7
87	Detection of susceptibility genes as modifiers due to subgroup differences in complex disease. European Journal of Human Genetics, 2010, 18, 960-964.	1.4	6
88	Polygenic association with severity and long-term outcome in eating disorder cases. Translational Psychiatry, 2022, 12, 61.	2.4	6
89	No association of dysbindin with symptom factors of schizophrenia in an Irish case–control sample. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 700-705.	1.1	5
90	Chronicity and Sex Affect Genetic Risk Prediction in Schizophrenia. Frontiers in Psychiatry, 2020, 11, 313.	1.3	5

Sarah E Bergen

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91	Improving lithium dose prediction using population pharmacokinetics and pharmacogenomics: a cohort genome-wide association study in Sweden. Lancet Psychiatry,the, 2022, 9, 447-457.	3.7	4
92	A loop-counting method for covariate-corrected low-rank biclustering of gene-expression and genome-wide association study data. PLoS Computational Biology, 2018, 14, e1006105.	1.5	3
93	Association of severe childhood infections with depression and intentional self-harm in adolescents and young adults. Brain, Behavior, and Immunity, 2022, 99, 247-255.	2.0	3
94	Populationâ€based identityâ€byâ€descent mapping combined with exome sequencing to detect rare risk variants for schizophrenia. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2019, 180, 223-231.	1.1	2
95	Overview of CAPICE—Childhood and Adolescence Psychopathology: unravelling the complex etiology by a large Interdisciplinary Collaboration in Europe—an EU Marie SkÅ,odowska-Curie International Training Network. European Child and Adolescent Psychiatry, 2021, , 1.	2.8	2
96	Genetic Modifiers and Subtypes in Schizophrenia. Current Behavioral Neuroscience Reports, 2014, 1, 197-205.	0.6	1
97	Polygenic Risk for Anxiety Influences Anxiety Comorbidity and Suicidal Behavior in Bipolar Disorder. SSRN Electronic Journal, 0, , .	0.4	1
98	Parsing psychosis subtypes through investigations of rare genetic variants. EBioMedicine, 2016, 6, 16-17.	2.7	0