

Jason L Stein

List of Publications by Year in descending order

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Version: 2024-02-01

89
papers

13,338
citations

53794

45
h-index

48315

88
g-index

118
all docs

118
docs citations

118
times ranked

19396
citing authors

#	ARTICLE	IF	CITATIONS
1	Ten years of enhancing <scp>neuroimaging</scp> genetics through <scp>meta-analysis</scp>: An overview from the <scp>ENIGMA Genetics Working Group</scp>. Human Brain Mapping, 2022, 43, 292-299.	3.6	19
2	Evaluating brain structure traits as endophenotypes using polygenicity and discoverability. Human Brain Mapping, 2022, 43, 329-340.	3.6	19
3	<scp>ToxCast</scp> chemical library Wnt screen identifies diethanolamine as an activator of neural progenitor proliferation. FASEB BioAdvances, 2022, 4, 441-453.	2.4	1
4	Genetic variants associated with longitudinal changes in brain structure across the lifespan. Nature Neuroscience, 2022, 25, 421-432.	14.8	75
5	Common variants contribute to intrinsic human brain functional networks. Nature Genetics, 2022, 54, 508-517.	21.4	37
6	The Evolutionary History of Common Genetic Variants Influencing Human Cortical Surface Area. Cerebral Cortex, 2021, 31, 1873-1887.	2.9	21
7	MRLocus: Identifying causal genes mediating a trait through Bayesian estimation of allelic heterogeneity. PLoS Genetics, 2021, 17, e1009455.	3.5	24
8	From base pair to brain. Nature Neuroscience, 2021, 24, 619-621.	14.8	4
9	Cell-type-specific effects of genetic variation on chromatin accessibility during human neuronal differentiation. Nature Neuroscience, 2021, 24, 941-953.	14.8	47
10	Segmentor: a tool for manual refinement of 3D microscopy annotations. BMC Bioinformatics, 2021, 22, 260.	2.6	11
11	Common genetic variation influencing human white matter microstructure. Science, 2021, 372, .	12.6	106
12	Brain-trait-associated variants impact cell-type-specific gene regulation during neurogenesis. American Journal of Human Genetics, 2021, 108, 1647-1668.	6.2	36
13	NuMorph: Tools for cortical cellular phenotyping in tissue-cleared whole-brain images. Cell Reports, 2021, 37, 109802.	6.4	8
14	Genetic correlations and genome-wide associations of cortical structure in general population samples of 22,824 adults. Nature Communications, 2020, 11, 4796.	12.8	61
15	Cas9 gene therapy for Angelman syndrome traps Ube3a-ATS long non-coding RNA. Nature, 2020, 587, 281-284.	27.8	92
16	Common genetic risk variants identified in the SPARK cohort support DDHD2 as a candidate risk gene for autism. Translational Psychiatry, 2020, 10, 265.	4.8	56
17	ENIGMA and global neuroscience: A decade of large-scale studies of the brain in health and disease across more than 40 countries. Translational Psychiatry, 2020, 10, 100.	4.8	365
18	The genetic architecture of the human cerebral cortex. Science, 2020, 367, .	12.6	450

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19	A Single-Cell Transcriptomic Atlas of Human Neocortical Development during Mid-gestation. <i>Neuron</i> , 2019, 103, 785-801.e8.	8.1	361
20	Memo1-Mediated Tiling of Radial Glial Cells Facilitates Cerebral Cortical Development. <i>Neuron</i> , 2019, 103, 836-852.e5.	8.1	46
21	Genetic Control of Expression and Splicing in Developing Human Brain Informs Disease Mechanisms. <i>Cell</i> , 2019, 179, 750-771.e22.	28.9	174
22	Genetic Markers of ADHD-Related Variations in Intracranial Volume. <i>American Journal of Psychiatry</i> , 2019, 176, 228-238.	7.2	68
23	Mapping causal pathways from genetics to neuropsychiatric disorders using genome-wide imaging genetics: Current status and future directions. <i>Psychiatry and Clinical Neurosciences</i> , 2019, 73, 357-369.	1.8	22
24	Genetic architecture of subcortical brain structures in 38,851 individuals. <i>Nature Genetics</i> , 2019, 51, 1624-1636.	21.4	192
25	The Dynamic Landscape of Open Chromatin during Human Cortical Neurogenesis. <i>Cell</i> , 2018, 172, 289-304.e18.	28.9	281
26	Human neural stem cell-derived cultures in three-dimensional substrates form spontaneously functional neuronal networks. <i>Journal of Tissue Engineering and Regenerative Medicine</i> , 2017, 11, 1022-1033.	2.7	20
27	ENIGMA and the individual: Predicting factors that affect the brain in 35 countries worldwide. <i>NeuroImage</i> , 2017, 145, 389-408.	4.2	173
28	Novel genetic loci associated with hippocampal volume. <i>Nature Communications</i> , 2017, 8, 13624.	12.8	250
29	Default Patterning Produces Pan-cortical Glutamatergic and CGE/LGE-like GABAergic Neurons from Human Pluripotent Stem Cells. <i>Stem Cell Reports</i> , 2017, 9, 1463-1476.	4.8	14
30	Advancing the understanding of autism disease mechanisms through genetics. <i>Nature Medicine</i> , 2016, 22, 345-361.	30.7	684
31	Circuit-wide Transcriptional Profiling Reveals Brain Region-Specific Gene Networks Regulating Depression Susceptibility. <i>Neuron</i> , 2016, 90, 969-983.	8.1	272
32	Transcriptomic signatures of neuronal differentiation and their association with risk genes for autism spectrum and related neuropsychiatric disorders. <i>Translational Psychiatry</i> , 2016, 6, e864-e864.	4.8	22
33	Novel genetic loci underlying human intracranial volume identified through genome-wide association. <i>Nature Neuroscience</i> , 2016, 19, 1569-1582.	14.8	213
34	Imaging Genomics and ENIGMA. , 2016, , 101-115.		3
35	Chromosome conformation elucidates regulatory relationships in developing human brain. <i>Nature</i> , 2016, 538, 523-527.	27.8	507
36	Early developmental gene enhancers affect subcortical volumes in the adult human brain. <i>Human Brain Mapping</i> , 2016, 37, 1788-1800.	3.6	6

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37	Genetic influences on schizophrenia and subcortical brain volumes: large-scale proof of concept. <i>Nature Neuroscience</i> , 2016, 19, 420-431.	14.8	204
38	Genetics of the Connectome and the ENIGMA Project. <i>Research and Perspectives in Neurosciences</i> , 2016, , 147-164.	0.4	10
39	Genome engineering of isogenic human ES cells to model autism disorders. <i>Nucleic Acids Research</i> , 2015, 43, e65-e65.	14.5	15
40	Common genetic variants influence human subcortical brain structures. <i>Nature</i> , 2015, 520, 224-229.	27.8	772
41	Genome-wide interaction analysis reveals replicated epistatic effects on brain structure. <i>Neurobiology of Aging</i> , 2015, 36, S151-S158.	3.1	22
42	Copy number variation and brain structure: lessons learned from chromosome 16p11.2. <i>Genome Medicine</i> , 2015, 7, 13.	8.2	6
43	JAKMIP1, a Novel Regulator of Neuronal Translation, Modulates Synaptic Function and Autistic-like Behaviors in Mouse. <i>Neuron</i> , 2015, 88, 1173-1191.	8.1	34
44	Joint genetic analysis of hippocampal size in mouse and human identifies a novel gene linked to neurodegenerative disease. <i>BMC Genomics</i> , 2014, 15, 850.	2.8	59
45	The ENIGMA Consortium: large-scale collaborative analyses of neuroimaging and genetic data. <i>Brain Imaging and Behavior</i> , 2014, 8, 153-182.	2.1	696
46	A Quantitative Framework to Evaluate Modeling of Cortical Development by Neural Stem Cells. <i>Neuron</i> , 2014, 83, 69-86.	8.1	184
47	Allelic differences between Europeans and Chinese for CREB1 SNPs and their implications in gene expression regulation, hippocampal structure and function, and bipolar disorder susceptibility. <i>Molecular Psychiatry</i> , 2014, 19, 452-461.	7.9	61
48	Genome-wide association identifies genetic variants associated with lentiform nucleus volume in N=1345 young and elderly subjects. <i>Brain Imaging and Behavior</i> , 2013, 7, 102-115.	2.1	26
49	Rare Inherited Variation in Autism: Beginning to See the Forest and a Few Trees. <i>Neuron</i> , 2013, 77, 209-211.	8.1	56
50	Hippocampal structure and human cognition: Key role of spatial processing and evidence supporting the efficiency hypothesis in females. <i>Intelligence</i> , 2013, 41, 129-140.	3.0	40
51	Exhaustive Search of the SNP-SNP Interactome Identifies Epistatic Effects on Brain Volume in Two Cohorts. <i>Lecture Notes in Computer Science</i> , 2013, 16, 600-607.	1.3	9
52	Genetic Clustering on the Hippocampal Surface for Genome-Wide Association Studies. <i>Lecture Notes in Computer Science</i> , 2013, 16, 690-697.	1.3	7
53	Investigation of Anatomical Thalamo-Cortical Connectivity and fMRI Activation in Schizophrenia. <i>Neuropsychopharmacology</i> , 2012, 37, 499-507.	5.4	133
54	Predicting White Matter Integrity from Multiple Common Genetic Variants. <i>Neuropsychopharmacology</i> , 2012, 37, 2012-2019.	5.4	49

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55	Predicting temporal lobe volume on MRI from genotypes using L<sup>1</sup>-L<sup>2</sup>-regularized regression. , 2012, , 1160-1163.		23
56	Common variants at 12q14 and 12q24 are associated with hippocampal volume. Nature Genetics, 2012, 44, 545-551.	21.4	212
57	Neuroimaging and Genetics: Exploring, Searching, and Finding. Twin Research and Human Genetics, 2012, 15, 267-272.	0.6	7
58	EnigmaVis: Online Interactive Visualization of Genome-Wide Association Studies of the Enhancing Neuroimaging Genetics through Meta-Analysis (ENIGMA) Consortium. Twin Research and Human Genetics, 2012, 15, 414-418.	0.6	34
59	Alzheimer's Disease Risk Gene, <i>GAB2</i>, is Associated with Regional Brain Volume Differences in 755 Young Healthy Twins. Twin Research and Human Genetics, 2012, 15, 286-295.	0.6	16
60	Relationship of a Variant in the <i>NTRK1</i> Gene to White Matter Microstructure in Young Adults. Journal of Neuroscience, 2012, 32, 5964-5972.	3.6	40
61	Identification of common variants associated with human hippocampal and intracranial volumes. Nature Genetics, 2012, 44, 552-561.	21.4	594
62	Sparse reduced-rank regression detects genetic associations with voxel-wise longitudinal phenotypes in Alzheimer's disease. NeuroImage, 2012, 60, 700-716.	4.2	121
63	Structural and functional neuroimaging phenotypes in dysbindin mutant mice. NeuroImage, 2012, 62, 120-129.	4.2	19
64	Common folate gene variant, MTHFR C677T, is associated with brain structure in two independent cohorts of people with mild cognitive impairment. NeuroImage: Clinical, 2012, 1, 179-187.	2.7	29
65	Brain structure in healthy adults is related to serum transferrin and the H63D polymorphism in the <i>HFE</i> gene. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, E851-9.	7.1	83
66	Discovery and replication of gene influences on brain structure using LASSO regression. Frontiers in Neuroscience, 2012, 6, 115.	2.8	91
67	De novo mutations revealed by whole-exome sequencing are strongly associated with autism. Nature, 2012, 485, 237-241.	27.8	1,863
68	Principal components regression: Multivariate, gene-based tests in imaging genomics. , 2011, , .		2
69	Genome-wide association reveals dopamine-related genetic effects on caudate volume. Molecular Psychiatry, 2011, 16, 881-881.	7.9	8
70	Voxelwise gene-wide association study (vGeneWAS): Multivariate gene-based association testing in 731 elderly subjects. NeuroImage, 2011, 56, 1875-1891.	4.2	116
71	Multilocus Genetic Analysis of Brain Images. Frontiers in Genetics, 2011, 2, 73.	2.3	53
72	Genetic Analysis of Cortical Thickness and Fractional Anisotropy of Water Diffusion in the Brain. Frontiers in Neuroscience, 2011, 5, 120.	2.8	52

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73	The contribution of genes to cortical thickness and volume. <i>NeuroReport</i> , 2011, 22, 101-105.	1.2	84
74	Hippocampal volume is related to body mass index in Alzheimer's disease. <i>NeuroReport</i> , 2011, 22, 10-14.	1.2	35
75	Discovery and replication of dopamine-related gene effects on caudate volume in young and elderly populations (N=1198) using genome-wide search. <i>Molecular Psychiatry</i> , 2011, 16, 927-937.	7.9	52
76	The effects of physical activity, education, and body mass index on the aging brain. <i>Human Brain Mapping</i> , 2011, 32, 1371-1382.	3.6	117
77	Boosting power to detect genetic associations in imaging using multi-locus, genome-wide scans and ridge regression. , 2011, , .		9
78	Common Alzheimer's Disease Risk Variant Within the <i>CLU</i> Gene Affects White Matter Microstructure in Young Adults. <i>Journal of Neuroscience</i> , 2011, 31, 6764-6770.	3.6	157
79	Vasopressin modulates social recognition-related activity in the left temporoparietal junction in humans. <i>Translational Psychiatry</i> , 2011, 1, e3-e3.	4.8	52
80	Comparing 3 T and 1.5 T MRI for tracking Alzheimer's disease progression with tensor-based morphometry. <i>Human Brain Mapping</i> , 2010, 31, 499-514.	3.6	66
81	A commonly carried allele of the obesity-related <i>FTO</i> gene is associated with reduced brain volume in the healthy elderly. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 8404-8409.	7.1	227
82	Vasopressin Modulates Medial Prefrontal Cortex–Amygdala Circuitry during Emotion Processing in Humans. <i>Journal of Neuroscience</i> , 2010, 30, 7017-7022.	3.6	118
83	Voxelwise genome-wide association study (vGWAS). <i>NeuroImage</i> , 2010, 53, 1160-1174.	4.2	239
84	Genome-wide analysis reveals novel genes influencing temporal lobe structure with relevance to neurodegeneration in Alzheimer's disease. <i>NeuroImage</i> , 2010, 51, 542-554.	4.2	141
85	Obesity is linked with lower brain volume in 700 AD and MCI patients. <i>Neurobiology of Aging</i> , 2010, 31, 1326-1339.	3.1	170
86	Alzheimer's Disease Neuroimaging Initiative biomarkers as quantitative phenotypes: Genetics core aims, progress, and plans. <i>Alzheimer's and Dementia</i> , 2010, 6, 265-273.	0.8	378
87	Active fibers: Matching deformable tract templates to diffusion tensor images. <i>NeuroImage</i> , 2009, 47, T82-T89.	4.2	18
88	Know Your Place: Neural Processing of Social Hierarchy in Humans. <i>Neuron</i> , 2008, 58, 273-283.	8.1	516
89	A validated network of effective amygdala connectivity. <i>NeuroImage</i> , 2007, 36, 736-745.	4.2	360