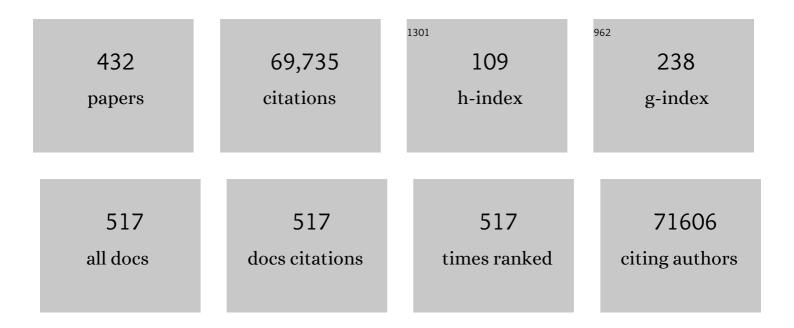
Philip L De Jager

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Integrative analysis of 111 reference human epigenomes. Nature, 2015, 518, 317-330.	27.8	5,653
2	Meta-analysis of 74,046 individuals identifies 11 new susceptibility loci for Alzheimer's disease. Nature Genetics, 2013, 45, 1452-1458.	21.4	3,741
3	Genetic risk and a primary role for cell-mediated immune mechanisms in multiple sclerosis. Nature, 2011, 476, 214-219.	27.8	2,400
4	Genetics of rheumatoid arthritis contributes to biology and drug discovery. Nature, 2014, 506, 376-381.	27.8	1,974
5	Genetic meta-analysis of diagnosed Alzheimer's disease identifies new risk loci and implicates Aβ, tau, immunity and lipid processing. Nature Genetics, 2019, 51, 414-430.	21.4	1,962
6	Common variants at MS4A4/MS4A6E, CD2AP, CD33 and EPHA1 are associated with late-onset Alzheimer's disease. Nature Genetics, 2011, 43, 436-441.	21.4	1,676
7	Genetic and epigenetic fine mapping of causal autoimmune disease variants. Nature, 2015, 518, 337-343.	27.8	1,669
8	Risk Alleles for Multiple Sclerosis Identified by a Genomewide Study. New England Journal of Medicine, 2007, 357, 851-862.	27.0	1,529
9	Analysis of immune-related loci identifies 48 new susceptibility variants for multiple sclerosis. Nature Genetics, 2013, 45, 1353-1360.	21.4	1,213
10	Genome-wide association study identifies 74 loci associated with educational attainment. Nature, 2016, 533, 539-542.	27.8	1,204
11	Charting a dynamic DNA methylation landscape of the human genome. Nature, 2013, 500, 477-481.	27.8	1,168
12	Genome-wide association study meta-analysis identifies seven new rheumatoid arthritis risk loci. Nature Genetics, 2010, 42, 508-514.	21.4	1,132
13	Modeling Linkage Disequilibrium Increases Accuracy of Polygenic Risk Scores. American Journal of Human Genetics, 2015, 97, 576-592.	6.2	1,098
14	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	12.6	1,085
15	Alterations of the human gut microbiome in multiple sclerosis. Nature Communications, 2016, 7, 12015.	12.8	957
16	Gene expression elucidates functional impact of polygenic risk for schizophrenia. Nature Neuroscience, 2016, 19, 1442-1453.	14.8	952
17	Genetic variants associated with subjective well-being, depressive symptoms, and neuroticism identified through genome-wide analyses. Nature Genetics, 2016, 48, 624-633.	21.4	870
18	Alzheimer's disease: early alterations in brain DNA methylation at ANK1, BIN1, RHBDF2 and other loci. Nature Neuroscience, 2014, 17, 1156-1163.	14.8	800

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19	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. Nature Genetics, 2017, 49, 1373-1384.	21.4	783
20	GWAS of 126,559 Individuals Identifies Genetic Variants Associated with Educational Attainment. Science, 2013, 340, 1467-1471.	12.6	750
21	Meta-analysis of genome scans and replication identify CD6, IRF8 and TNFRSF1A as new multiple sclerosis susceptibility loci. Nature Genetics, 2009, 41, 776-782.	21.4	729
22	Multiple sclerosis genomic map implicates peripheral immune cells and microglia in susceptibility. Science, 2019, 365, .	12.6	710
23	Parkinson's Disease: Genetics and Pathogenesis. Annual Review of Pathology: Mechanisms of Disease, 2011, 6, 193-222.	22.4	654
24	Large-scale proteomic analysis of Alzheimer's disease brain and cerebrospinal fluid reveals early changes in energy metabolism associated with microglia and astrocyte activation. Nature Medicine, 2020, 26, 769-780.	30.7	547
25	Pervasive Sharing of Genetic Effects in Autoimmune Disease. PLoS Genetics, 2011, 7, e1002254.	3.5	540
26	Temporal Tracking of Microglia Activation in Neurodegeneration at Single-Cell Resolution. Cell Reports, 2017, 21, 366-380.	6.4	538
27	The transcriptional landscape of age in human peripheral blood. Nature Communications, 2015, 6, 8570.	12.8	533
28	Neurodegeneration in Lurcher mice caused by mutation in δ2 glutamate receptor gene. Nature, 1997, 388, 769-773.	27.8	522
29	Defining the Role of the MHC in Autoimmunity: A Review and Pooled Analysis. PLoS Genetics, 2008, 4, e1000024.	3.5	488
30	Methylomic profiling implicates cortical deregulation of ANK1 in Alzheimer's disease. Nature Neuroscience, 2014, 17, 1164-1170.	14.8	488
31	CD33 Alzheimer's disease locus: altered monocyte function and amyloid biology. Nature Neuroscience, 2013, 16, 848-850.	14.8	485
32	Study of 300,486 individuals identifies 148 independent genetic loci influencing general cognitive function. Nature Communications, 2018, 9, 2098.	12.8	484
33	Polarization of the Effects of Autoimmune and Neurodegenerative Risk Alleles in Leukocytes. Science, 2014, 344, 519-523.	12.6	480
34	Genome-wide Chromatin State Transitions Associated with Developmental and Environmental Cues. Cell, 2013, 152, 642-654.	28.9	473
35	A molecular network of the aging human brain provides insights into the pathology and cognitive decline of Alzheimer's disease. Nature Neuroscience, 2018, 21, 811-819.	14.8	422
36	A High-Density Admixture Map for Disease Gene Discovery in African Americans. American Journal of Human Genetics, 2004, 74, 1001-1013.	6.2	416

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37	Altered bile acid profile associates with cognitive impairment in Alzheimer's disease—An emerging role for gut microbiome. Alzheimer's and Dementia, 2019, 15, 76-92.	0.8	396
38	Common Genetic Variants Modulate Pathogen-Sensing Responses in Human Dendritic Cells. Science, 2014, 343, 1246980.	12.6	391
39	An xQTL map integrates the genetic architecture of the human brain's transcriptome and epigenome. Nature Neuroscience, 2017, 20, 1418-1426.	14.8	377
40	A transcriptomic atlas of aged human microglia. Nature Communications, 2018, 9, 539.	12.8	375
41	De novo copy number variants identify new genes and loci in isolated sporadic tetralogy of Fallot. Nature Genetics, 2009, 41, 931-935.	21.4	373
42	Single cell RNA sequencing of human microglia uncovers a subset associated with Alzheimer's disease. Nature Communications, 2020, 11, 6129.	12.8	371
43	Variants in the ATP-Binding Cassette Transporter (ABCA7), Apolipoprotein E ϵ4, and the Risk of Late-Onset Alzheimer Disease in African Americans. JAMA - Journal of the American Medical Association, 2013, 309, 1483.	7.4	360
44	A multi-omic atlas of the human frontal cortex for aging and Alzheimer's disease research. Scientific Data, 2018, 5, 180142.	5.3	357
45	Automated high-dimensional flow cytometric data analysis. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 8519-8524.	7.1	355
46	The power of genetic diversity in genome-wide association studies of lipids. Nature, 2021, 600, 675-679.	27.8	353
47	GWAS of Cerebrospinal Fluid Tau Levels Identifies Risk Variants for Alzheimer's Disease. Neuron, 2013, 78, 256-268.	8.1	344
48	Self-antigen tetramers discriminate between myelin autoantibodies to native or denatured protein. Nature Medicine, 2007, 13, 211-217.	30.7	342
49	Blood Kidney Injury Molecule-1 Is a Biomarker of Acute and Chronic Kidney Injury and Predicts Progression to ESRD in Type I Diabetes. Journal of the American Society of Nephrology: JASN, 2014, 25, 2177-2186.	6.1	341
50	The Influence of Age and Sex on Genetic Associations with Adult Body Size and Shape: A Large-Scale Genome-Wide Interaction Study. PLoS Genetics, 2015, 11, e1005378.	3.5	331
51	Genomeâ€wide metaâ€analysis identifies novel multiple sclerosis susceptibility loci. Annals of Neurology, 2011, 70, 897-912.	5.3	314
52	Class II HLA interactions modulate genetic risk for multiple sclerosis. Nature Genetics, 2015, 47, 1107-1113.	21.4	312
53	Integrative transcriptome analyses of the aging brain implicate altered splicing in Alzheimer's disease susceptibility. Nature Genetics, 2018, 50, 1584-1592.	21.4	307
54	Genome-Wide Association Meta-analysis of Neuropathologic Features of Alzheimer's Disease and Related Dementias. PLoS Genetics, 2014, 10, e1004606.	3.5	305

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55	Infection-Triggered Familial or Recurrent Cases of Acute Necrotizing Encephalopathy Caused by Mutations in a Component of the Nuclear Pore, RANBP2. American Journal of Human Genetics, 2009, 84, 44-51.	6.2	291
56	A High-Density Screen for Linkage in Multiple Sclerosis. American Journal of Human Genetics, 2005, 77, 454-467.	6.2	268
57	A novel Alzheimer disease locus located near the gene encoding tau protein. Molecular Psychiatry, 2016, 21, 108-117.	7.9	260
58	Fine-Mapping the Genetic Association of the Major Histocompatibility Complex in Multiple Sclerosis: HLA and Non-HLA Effects. PLoS Genetics, 2013, 9, e1003926.	3.5	250
59	GWAS of Longevity in CHARGE Consortium Confirms APOE and FOXO3 Candidacy. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2015, 70, 110-118.	3.6	250
60	Parsing the Interferon Transcriptional Network and Its Disease Associations. Cell, 2016, 164, 564-578.	28.9	250
61	Novel genetic loci associated with hippocampal volume. Nature Communications, 2017, 8, 13624.	12.8	250
62	Exceptionally low likelihood of Alzheimer's dementia in APOE2 homozygotes from a 5,000-person neuropathological study. Nature Communications, 2020, 11, 667.	12.8	246
63	A whole-genome admixture scan finds a candidate locus for multiple sclerosis susceptibility. Nature Genetics, 2005, 37, 1113-1118.	21.4	243
64	Life Extension Factor Klotho Enhances Cognition. Cell Reports, 2014, 7, 1065-1076.	6.4	243
65	Association of Brain DNA Methylation in <i>SORL1</i> , <i>ABCA7</i> , <i>HLA-DRB5</i> , <i>SLC24A4</i> , and <i>BIN1</i> With Pathological Diagnosis of Alzheimer Disease. JAMA Neurology, 2015, 72, 15.	9.0	239
66	Integration of genetic risk factors into a clinical algorithm for multiple sclerosis susceptibility: a weighted genetic risk score. Lancet Neurology, The, 2009, 8, 1111-1119.	10.2	233
67	Mapping of multiple susceptibility variants within the MHC region for 7 immune-mediated diseases. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 18680-18685.	7.1	231
68	Association of CR1, CLU and PICALM with Alzheimer's disease in a cohort of clinically characterized and neuropathologically verified individuals. Human Molecular Genetics, 2010, 19, 3295-3301.	2.9	223
69	Sex-Specific Association of Apolipoprotein E With Cerebrospinal Fluid Levels of Tau. JAMA Neurology, 2018, 75, 989.	9.0	223
70	Intersection of population variation and autoimmunity genetics in human T cell activation. Science, 2014, 345, 1254665.	12.6	218
71	Tau Activates Transposable Elements in Alzheimer's Disease. Cell Reports, 2018, 23, 2874-2880.	6.4	216
72	Novel genetic loci underlying human intracranial volume identified through genome-wide association. Nature Neuroscience, 2016, 19, 1569-1582.	14.8	213

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73	Common variants at 12q14 and 12q24 are associated with hippocampal volume. Nature Genetics, 2012, 44, 545-551.	21.4	212
74	IL2RA Genetic Heterogeneity in Multiple Sclerosis and Type 1 Diabetes Susceptibility and Soluble Interleukin-2 Receptor Production. PLoS Genetics, 2009, 5, e1000322.	3.5	210
75	Genome-wide Association Study in a High-Risk Isolate for Multiple Sclerosis Reveals Associated Variants in STAT3 Gene. American Journal of Human Genetics, 2010, 86, 285-291.	6.2	210
76	Limited statistical evidence for shared genetic effects of eQTLs and autoimmune-disease-associated loci in three major immune-cell types. Nature Genetics, 2017, 49, 600-605.	21.4	205
77	Large-scale deep multi-layer analysis of Alzheimer's disease brain reveals strong proteomic disease-related changes not observed at the RNA level. Nature Neuroscience, 2022, 25, 213-225.	14.8	202
78	Genetic Analysis of Human Traits In Vitro: Drug Response and Gene Expression in Lymphoblastoid Cell Lines. PLoS Genetics, 2008, 4, e1000287.	3.5	200
79	Genome-wide association study identifies four novel loci associated with Alzheimer's endophenotypes and disease modifiers. Acta Neuropathologica, 2017, 133, 839-856.	7.7	199
80	Meta-Analysis of the Alzheimer's Disease Human Brain Transcriptome and Functional Dissection in Mouse Models. Cell Reports, 2020, 32, 107908.	6.4	199
81	Genetic architecture of subcortical brain structures in 38,851 individuals. Nature Genetics, 2019, 51, 1624-1636.	21.4	192
82	Epigenome-wide study uncovers large-scale changes in histone acetylation driven by tau pathology in aging and Alzheimer's human brains. Nature Neuroscience, 2019, 22, 37-46.	14.8	188
83	The role of the <i>CD58</i> locus in multiple sclerosis. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 5264-5269.	7.1	185
84	Convergent genetic and expression data implicate immunity in Alzheimer's disease. Alzheimer's and Dementia, 2015, 11, 658-671.	0.8	173
85	Higher brain <i>BDNF</i> gene expression is associated with slower cognitive decline in older adults. Neurology, 2016, 86, 735-741.	1.1	170
86	Normalization of Plasma 25-Hydroxy Vitamin D Is Associated with Reduced Risk of Surgery in Crohn's Disease. Inflammatory Bowel Diseases, 2013, 19, 1.	1.9	168
87	Effects of Multiple Genetic Loci on Age at Onset in Late-Onset Alzheimer Disease. JAMA Neurology, 2014, 71, 1394.	9.0	166
88	Transethnic genomeâ€wide scan identifies novel Alzheimer's disease loci. Alzheimer's and Dementia, 2017, 13, 727-738.	0.8	166
89	Evidence for Polygenic Susceptibility to Multiple Sclerosis—The Shape of Things to Come. American Journal of Human Genetics, 2010, 86, 621-625.	6.2	162
90	Multiethnic Genome-Wide Association Study of Cerebral White Matter Hyperintensities on MRI. Circulation: Cardiovascular Genetics, 2015, 8, 398-409.	5.1	162

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91	Admixture Mapping of an Allele Affecting Interleukin 6 Soluble Receptor and Interleukin 6 Levels. American Journal of Human Genetics, 2007, 80, 716-726.	6.2	160
92	A genome-wide scan for common variants affecting the rate of age-related cognitive decline. Neurobiology of Aging, 2012, 33, 1017.e1-1017.e15.	3.1	160
93	Polygenic risk of Alzheimer disease is associated with early- and late-life processes. Neurology, 2016, 87, 481-488.	1.1	159
94	Integrating human brain proteomes with genome-wide association data implicates new proteins in Alzheimer's disease pathogenesis. Nature Genetics, 2021, 53, 143-146.	21.4	158
95	Gene-Wide Analysis Detects Two New Susceptibility Genes for Alzheimer's Disease. PLoS ONE, 2014, 9, e94661.	2.5	155
96	Genome-wide meta-analysis identifies multiple novel associations and ethnic heterogeneity of psoriasis susceptibility. Nature Communications, 2015, 6, 6916.	12.8	154
97	Functionally defective germline variants of sialic acid acetylesterase in autoimmunity. Nature, 2010, 466, 243-247.	27.8	150
98	A Genome-Wide Association Study of Depressive Symptoms. Biological Psychiatry, 2013, 73, 667-678.	1.3	149
99	<i>CR1</i> is associated with amyloid plaque burden and ageâ€related cognitive decline. Annals of Neurology, 2011, 69, 560-569.	5.3	148
100	Genomeâ€wide association study of the rate of cognitive decline in Alzheimer's disease. Alzheimer's and Dementia, 2014, 10, 45-52.	0.8	147
101	Functional screening in Drosophila identifies Alzheimer's disease susceptibility genes and implicates Tau-mediated mechanisms. Human Molecular Genetics, 2014, 23, 870-877.	2.9	147
102	Large meta-analysis of genome-wide association studies identifies five loci for lean body mass. Nature Communications, 2017, 8, 80.	12.8	147
103	Genome-Wide Association Study and Gene Expression Analysis Identifies CD84 as a Predictor of Response to Etanercept Therapy in Rheumatoid Arthritis. PLoS Genetics, 2013, 9, e1003394.	3.5	146
104	Novel late-onset Alzheimer disease loci variants associate with brain gene expression. Neurology, 2012, 79, 221-228.	1.1	144
105	Novel Alzheimer Disease Risk Loci and Pathways in African American Individuals Using the African Genome Resources Panel. JAMA Neurology, 2021, 78, 102.	9.0	144
106	Genetic Susceptibility for Alzheimer Disease Neuritic Plaque Pathology. JAMA Neurology, 2013, 70, 1150.	9.0	143
107	Genetic determinants of co-accessible chromatin regions in activated T cells across humans. Nature Genetics, 2018, 50, 1140-1150.	21.4	139
108	Elevated DNA methylation across a 48â€kb region spanning the <i>HOXA</i> gene cluster is associated with Alzheimer's disease neuropathology. Alzheimer's and Dementia, 2018, 14, 1580-1588.	0.8	138

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109	Soluble IL-2RA Levels in Multiple Sclerosis Subjects and the Effect of Soluble IL-2RA on Immune Responses. Journal of Immunology, 2009, 182, 1541-1547.	0.8	136
110	CD33 modulates TREM2: convergence of Alzheimer loci. Nature Neuroscience, 2015, 18, 1556-1558.	14.8	134
111	Multicolored stain-free histopathology with coherent Raman imaging. Laboratory Investigation, 2012, 92, 1492-1502.	3.7	130
112	Identification of additional risk loci for stroke and small vessel disease: a meta-analysis of genome-wide association studies. Lancet Neurology, The, 2016, 15, 695-707.	10.2	130
113	CD33: increased inclusion of exon 2 implicates the Ig V-set domain in Alzheimer's disease susceptibility. Human Molecular Genetics, 2014, 23, 2729-2736.	2.9	128
114	Neuropathological correlates and genetic architecture of microglial activation in elderly human brain. Nature Communications, 2019, 10, 409.	12.8	121
115	Tau-Mediated Disruption of the Spliceosome Triggers Cryptic RNA Splicing and Neurodegeneration in Alzheimer's Disease. Cell Reports, 2019, 29, 301-316.e10.	6.4	118
116	GWAS of longitudinal amyloid accumulation on ¹⁸ F-florbetapir PET in Alzheimer's disease implicates microglial activation gene <i>IL1RAP</i> . Brain, 2015, 138, 3076-3088.	7.6	117
117	Nuclei multiplexing with barcoded antibodies for single-nucleus genomics. Nature Communications, 2019, 10, 2907.	12.8	117
118	Replication analysis identifies TYK2 as a multiple sclerosis susceptibility factor. European Journal of Human Genetics, 2009, 17, 1309-1313.	2.8	115
119	Low-Frequency and Rare-Coding Variation Contributes to Multiple Sclerosis Risk. Cell, 2018, 175, 1679-1687.e7.	28.9	115
120	Gut Microbiome in Progressive Multiple Sclerosis. Annals of Neurology, 2021, 89, 1195-1211.	5.3	115
121	Interindividual variation in human T regulatory cells. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, E1111-20.	7.1	112
122	Genetic variants linked to education predict longevity. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 13366-13371.	7.1	110
123	Comprehensive follow-up of the first genome-wide association study of multiple sclerosis identifies KIF21B and TMEM39A as susceptibility loci. Human Molecular Genetics, 2010, 19, 953-962.	2.9	108
124	A second X chromosome contributes to resilience in a mouse model of Alzheimer's disease. Science Translational Medicine, 2020, 12, .	12.4	107
125	A human microglia-like cellular model for assessing the effects of neurodegenerative disease gene variants. Science Translational Medicine, 2017, 9, .	12.4	106
126	NMNAT2:HSP90 Complex Mediates Proteostasis in Proteinopathies. PLoS Biology, 2016, 14, e1002472.	5.6	105

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127	Targeted brain proteomics uncover multiple pathways to Alzheimer's dementia. Annals of Neurology, 2018, 84, 78-88.	5.3	102
128	Evaluation of TDP-43 proteinopathy and hippocampal sclerosis in relation to APOE ε4 haplotype status: a community-based cohort study. Lancet Neurology, The, 2018, 17, 773-781.	10.2	101
129	Common Risk Alleles for Inflammatory Diseases Are Targets of Recent Positive Selection. American Journal of Human Genetics, 2013, 92, 517-529.	6.2	100
130	Epigenomics of Alzheimer's disease. Translational Research, 2015, 165, 200-220.	5.0	97
131	Genetic variants in Alzheimer disease — molecular and brain network approaches. Nature Reviews Neurology, 2016, 12, 413-427.	10.1	97
132	Association of APOE with tau-tangle pathology with and without Î ² -amyloid. Neurobiology of Aging, 2016, 37, 19-25.	3.1	97
133	A method for high-throughput, sensitive analysis of IgG Fc and Fab glycosylation by capillary electrophoresis. Journal of Immunological Methods, 2015, 417, 34-44.	1.4	95
134	Modification of Multiple Sclerosis Phenotypes by African Ancestry at HLA. Archives of Neurology, 2009, 66, 226-33.	4.5	92
135	A coding variant in CR1 interacts with APOE-É>4 to influence cognitive decline. Human Molecular Genetics, 2012, 21, 2377-2388.	2.9	90
136	Cerebral small vessel disease genomics and its implications across the lifespan. Nature Communications, 2020, 11, 6285.	12.8	89
137	Identification of genes associated with dissociation of cognitive performance and neuropathological burden: Multistep analysis of genetic, epigenetic, and transcriptional data. PLoS Medicine, 2017, 14, e1002287.	8.4	88
138	Sex-specific genetic predictors of Alzheimer's disease biomarkers. Acta Neuropathologica, 2018, 136, 857-872.	7.7	87
139	Human Herpesvirus 6 Detection in Alzheimer's Disease Cases and Controls across Multiple Cohorts. Neuron, 2020, 105, 1027-1035.e2.	8.1	87
140	Neurodegeneration in <i>Lurcher</i> Mice Occurs via Multiple Cell Death Pathways. Journal of Neuroscience, 2000, 20, 3687-3694.	3.6	86
141	The <i>CD6</i> Multiple Sclerosis Susceptibility Allele Is Associated with Alterations in CD4+ T Cell Proliferation. Journal of Immunology, 2011, 187, 3286-3291.	0.8	85
142	Brain proteome-wide association study implicates novel proteins in depression pathogenesis. Nature Neuroscience, 2021, 24, 810-817.	14.8	85
143	An Analysis of Two Genome-wide Association Meta-analyses Identifies a New Locus for Broad Depression Phenotype. Biological Psychiatry, 2017, 82, 322-329.	1.3	84
144	TIGAR: An Improved Bayesian Tool for Transcriptomic Data Imputation Enhances Gene Mapping of Complex Traits. American Journal of Human Genetics, 2019, 105, 258-266.	6.2	84

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145	Associations of autozygosity with a broad range of human phenotypes. Nature Communications, 2019, 10, 4957.	12.8	84
146	PhIP-Seq characterization of autoantibodies from patients with multiple sclerosis, type 1 diabetes and rheumatoid arthritis. Journal of Autoimmunity, 2013, 43, 1-9.	6.5	83
147	A genomeâ€wide profiling of brain DNA hydroxymethylation in Alzheimer's disease. Alzheimer's and Dementia, 2017, 13, 674-688.	0.8	83
148	Circadian alterations during early stages of Alzheimer's disease are associated with aberrant cycles of DNA methylation in BMAL1. Alzheimer's and Dementia, 2017, 13, 689-700.	0.8	83
149	Functional Screening of Alzheimer Pathology Genome-wide Association Signals in Drosophila. American Journal of Human Genetics, 2011, 88, 232-238.	6.2	81
150	24-Hour Rhythms of DNA Methylation and Their Relation with Rhythms of RNA Expression in the Human Dorsolateral Prefrontal Cortex. PLoS Genetics, 2014, 10, e1004792.	3.5	80
151	Single-Cell Detection of Secreted Aβ and sAPPα from Human IPSC-Derived Neurons and Astrocytes. Journal of Neuroscience, 2016, 36, 1730-1746.	3.6	80
152	Genome-wide Comparison of African-Ancestry Populations from CARe and Other Cohorts Reveals Signals of Natural Selection. American Journal of Human Genetics, 2011, 89, 368-381.	6.2	79
153	Evaluation of a Genetic Risk Score to Improve Risk Prediction for Alzheimer's Disease. Journal of Alzheimer's Disease, 2016, 53, 921-932.	2.6	77
154	APOE and cerebral amyloid angiopathy in community-dwelling older persons. Neurobiology of Aging, 2015, 36, 2946-2953.	3.1	76
155	Multiple sclerosis risk loci and disease severity in 7,125 individuals from 10 studies. Neurology: Genetics, 2016, 2, e87.	1.9	76
156	Large eQTL meta-analysis reveals differing patterns between cerebral cortical and cerebellar brain regions. Scientific Data, 2020, 7, 340.	5.3	75
157	Genome-wide meta-analysis of muscle weakness identifies 15 susceptibility loci in older men and women. Nature Communications, 2021, 12, 654.	12.8	75
158	Stem cell-derived neurons reflect features of protein networks, neuropathology, and cognitive outcome of their aged human donors. Neuron, 2021, 109, 3402-3420.e9.	8.1	75
159	Cytometric profiling in multiple sclerosis uncovers patient population structure and a reduction of CD8low cells. Brain, 2008, 131, 1701-1711.	7.6	73
160	Early Elevation of Serum Tumor Necrosis Factor-α is Associated with Poor Outcome in Subarachnoid Hemorrhage. Journal of Investigative Medicine, 2012, 60, 1054-1058.	1.6	72
161	Similar Risk of Depression and Anxiety Following Surgery or Hospitalization for Crohn's Disease and Ulcerative Colitis. American Journal of Gastroenterology, 2013, 108, 594-601.	0.4	72
162	A meta-analysis of epigenome-wide association studies in Alzheimer's disease highlights novel differentially methylated loci across cortex. Nature Communications, 2021, 12, 3517.	12.8	72

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163	Alzheimer Disease Susceptibility Loci: Evidence for a Protein Network under Natural Selection. American Journal of Human Genetics, 2012, 90, 720-726.	6.2	71
164	The <i>TMEM106B</i> locus and TDP-43 pathology in older persons without FTLD. Neurology, 2015, 84, 927-934.	1.1	71
165	Brain expression of the vascular endothelial growth factor gene family in cognitive aging and alzheimer's disease. Molecular Psychiatry, 2021, 26, 888-896.	7.9	71
166	Dissecting the role of non-coding RNAs in the accumulation of amyloid and tau neuropathologies in Alzheimer's disease. Molecular Neurodegeneration, 2017, 12, 51.	10.8	70
167	Cortical Proteins Associated With Cognitive Resilience in Community-Dwelling Older Persons. JAMA Psychiatry, 2020, 77, 1172.	11.0	70
168	A <scp><i>TREM</i></scp> <i>1</i> variant alters the accumulation of Alzheimerâ€related amyloid pathology. Annals of Neurology, 2015, 77, 469-477.	5.3	69
169	Relation of genomic variants for Alzheimer disease dementia to common neuropathologies. Neurology, 2016, 87, 489-496.	1.1	68
170	Integrating Gene and Protein Expression Reveals Perturbed Functional Networks in Alzheimer's Disease. Cell Reports, 2019, 28, 1103-1116.e4.	6.4	67
171	Modeling Disease Severity in Multiple Sclerosis Using Electronic Health Records. PLoS ONE, 2013, 8, e78927.	2.5	67
172	Genetic analysis of isoform usage in the human anti-viral response reveals influenza-specific regulation of <i>ERAP2</i> transcripts under balancing selection. Genome Research, 2018, 28, 1812-1825.	5.5	66
173	Deconvolving the contributions of cell-type heterogeneity on cortical gene expression. PLoS Computational Biology, 2020, 16, e1008120.	3.2	66
174	Enhancing linkage analysis of complex disorders: an evaluation of high-density genotyping. Human Molecular Genetics, 2004, 13, 1943-1949.	2.9	65
175	Evaluation of an Online Platform for Multiple Sclerosis Research: Patient Description, Validation of Severity Scale, and Exploration of BMI Effects on Disease Course. PLoS ONE, 2013, 8, e59707.	2.5	65
176	Sex differences in the genetic predictors of Alzheimer's pathology. Brain, 2019, 142, 2581-2589.	7.6	65
177	Exploration of changes in disability after menopause in a longitudinal multiple sclerosis cohort. Multiple Sclerosis Journal, 2016, 22, 935-943.	3.0	64
178	Genome-wide interaction analysis of pathological hallmarks in Alzheimer's disease. Neurobiology of Aging, 2020, 93, 61-68.	3.1	63
179	A Major Histocompatibility Class I Locus Contributes to Multiple Sclerosis Susceptibility Independently from HLA-DRB1*15:01. PLoS ONE, 2010, 5, e11296.	2.5	60
180	High-dimensional immunomonitoring models of HIV-1–specific CD8 T-cell responses accurately identify subjects achieving spontaneous viral control. Blood, 2013, 121, 801-811.	1.4	60

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181	5′RNA-Seq identifies Fhl1 as a genetic modifier in cardiomyopathy. Journal of Clinical Investigation, 2014, 124, 1364-1370.	8.2	58
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