List of Publications by Year in descending order

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IOHN F POWELL

#	Article	IF	CITATIONS
1	Mapping genomic loci implicates genes and synaptic biology in schizophrenia. Nature, 2022, 604, 502-508.	27.8	929
2	Genetic copy number variants, cognition and psychosis: a meta-analysis and a family study. Molecular Psychiatry, 2021, 26, 5307-5319.	7.9	18
3	A Comparison of Ten Polygenic Score Methods for Psychiatric Disorders Applied Across Multiple Cohorts. Biological Psychiatry, 2021, 90, 611-620.	1.3	103
4	Gene-based analysis in HRC imputed genome wide association data identifies three novel genes for Alzheimer's disease. PLoS ONE, 2019, 14, e0218111.	2.5	23
5	Pattern of Altered Plasma Elemental Phosphorus, Calcium, Zinc, and Iron in Alzheimer's Disease. Scientific Reports, 2019, 9, 3147.	3.3	25
6	Telomere length is greater in ALS than in controls: a whole genome sequencing study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 229-234.	1.7	18
7	A plasma protein classifier for predicting amyloid burden for preclinical Alzheimer's disease. Science Advances, 2019, 5, eaau7220.	10.3	59
8	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
9	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
10	Mendelian adult-onset leukodystrophy genes in Alzheimer's disease: critical influence of CSF1R and NOTCH3. Neurobiology of Aging, 2018, 66, 179.e17-179.e29.	3.1	32
11	A polygenic risk score analysis of psychosis endophenotypes across brain functional, structural, and cognitive domains. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2018, 177, 21-34.	1.7	57
12	Red blood cell indices and anaemia as causative factors for cognitive function deficits and for Alzheimer's disease. Genome Medicine, 2018, 10, 51.	8.2	46
13	Use of schizophrenia and bipolar disorder polygenic risk scores to identify psychotic disorders. British Journal of Psychiatry, 2018, 213, 535-541.	2.8	37
14	ATXN2 trinucleotide repeat length correlates with risk of ALS. Neurobiology of Aging, 2017, 51, 178.e1-178.e9.	3.1	86
15	Mitochondrial genes are altered in blood early in Alzheimer's disease. Neurobiology of Aging, 2017, 53, 36-47.	3.1	132
16	Association Between Telomere Length and Risk of Cancer and Non-Neoplastic Diseases. JAMA Oncology, 2017, 3, 636.	7.1	376
17	Association between Plasma Ceramides and Phosphatidylcholines and Hippocampal Brain Volume in Late Onset Alzheimer's Disease. Journal of Alzheimer's Disease, 2017, 60, 809-817.	2.6	72
18	No Genetic Overlap Between Circulating Iron Levels and Alzheimer's Disease. Journal of Alzheimer's Disease, 2017, 59, 85-99.	2.6	10

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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	An Examination of Polygenic Score Risk Prediction in Individuals With First-Episode Psychosis. Biological Psychiatry, 2017, 81, 470-477.	1.3	176
21	Association of blood lipids with Alzheimer's disease: AÂcomprehensiveÂlipidomics analysis. Alzheimer's and Dementia, 2017, 13, 140-151.	0.8	144
22	Contribution of copy number variants to schizophrenia from a genome-wide study of 41,321 subjects. Nature Genetics, 2017, 49, 27-35.	21.4	838
23	Shared genetic contribution to ischemic stroke and Alzheimer's disease. Annals of Neurology, 2016, 79, 739-747.	5.3	56
24	Rare genetic variation in UNC13A may modify survival in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2016, 17, 593-599.	1.7	22
25	Genome-wide association analyses identify new risk variants and the genetic architecture of amyotrophic lateral sclerosis. Nature Genetics, 2016, 48, 1043-1048.	21.4	494
26	ABCA7 p.G215S as potential protective factor for Alzheimer's disease. Neurobiology of Aging, 2016, 46, 235.e1-235.e9.	3.1	37
27	Association of a Locus in the <i>CAMTA1</i> Gene With Survival in Patients With Sporadic Amyotrophic Lateral Sclerosis. JAMA Neurology, 2016, 73, 812.	9.0	57
28	Genome-wide analysis of genetic correlation in dementia with Lewy bodies, Parkinson's and Alzheimer's diseases. Neurobiology of Aging, 2016, 38, 214.e7-214.e10.	3.1	78
29	The effect of increased genetic risk for Alzheimer's disease on hippocampal and amygdala volume. Neurobiology of Aging, 2016, 40, 68-77.	3.1	115
30	A novel Alzheimer disease locus located near the gene encoding tau protein. Molecular Psychiatry, 2016, 21, 108-117.	7.9	260
31	Influence of Coding Variability in APP-Aβ Metabolism Genes in Sporadic Alzheimer's Disease. PLoS ONE, 2016, 11, e0150079.	2.5	34
32	Interaction between DRD2 and AKT1 genetic variations on risk of psychosis in cannabis users: a case–control study. NPJ Schizophrenia, 2015, 1, 15025.	3.6	29
33	Associations between Potentially Modifiable Risk Factors and Alzheimer Disease: A Mendelian Randomization Study. PLoS Medicine, 2015, 12, e1001841.	8.4	153
34	Plasma lipidomics analysis finds long chain cholesteryl esters to be associated with Alzheimer's disease. Translational Psychiatry, 2015, 5, e494-e494.	4.8	105
35	Proportion of patients in south London with first-episode psychosis attributable to use of high potency cannabis: a case-control study. Lancet Psychiatry,the, 2015, 2, 233-238.	7.4	429
36	Stratified gene expression analysis identifies major amyotrophic lateral sclerosis genes. Neurobiology of Aging, 2015, 36, 2006.e1-2006.e9.	3.1	22

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37	Interaction Between Functional Genetic Variation of DRD2 and Cannabis Use on Risk of Psychosis. Schizophrenia Bulletin, 2015, 41, 1171-1182.	4.3	73
38	Modeling Linkage Disequilibrium Increases Accuracy of Polygenic Risk Scores. American Journal of Human Genetics, 2015, 97, 576-592.	6.2	1,098
39	Common polygenic variation enhances risk prediction for Alzheimer's disease. Brain, 2015, 138, 3673-3684.	7.6	359
40	Convergent genetic and expression data implicate immunity in Alzheimer's disease. Alzheimer's and Dementia, 2015, 11, 658-671.	0.8	173
41	Genetic Predisposition to Increased Blood Cholesterol and Triglyceride Lipid Levels and Risk of Alzheimer Disease: A Mendelian Randomization Analysis. PLoS Medicine, 2014, 11, e1001713.	8.4	75
42	Investigating the role of rare coding variability in Mendelian dementia genes (APP , PSEN1 , PSEN2 , GRN) Tj ETQ	q0_0.0 rgE	BT /Qverlock
43	Heritability of Amyotrophic Lateral Sclerosis. JAMA Neurology, 2014, 71, 1579.	9.0	0
44	Role of Environmental Confounding in the Association between FKBP5 and First-Episode Psychosis. Frontiers in Psychiatry, 2014, 5, 84.	2.6	17
45	Exome sequencing identifies 2 novel presenilin 1 mutations (p.L166V and p.S230R) in British early-onset Alzheimer's disease. Neurobiology of Aging, 2014, 35, 2422.e13-2422.e16.	3.1	28
46	Rare coding variants in the phospholipase D3 gene confer risk for Alzheimer's disease. Nature, 2014, 505, 550-554.	27.8	425
47	A genome-wide association meta-analysis identifies a novel locus at 17q11.2 associated with sporadic amyotrophic lateral sclerosis. Human Molecular Genetics, 2014, 23, 2220-2231.	2.9	123
48	Alleles that increase risk for type 2 diabetes mellitus are not associated with increased risk for Alzheimer's disease. Neurobiology of Aging, 2014, 35, 2883.e3-2883.e10.	3.1	9
49	Partitioning Heritability of Regulatory and Cell-Type-Specific Variants across 11 Common Diseases. American Journal of Human Genetics, 2014, 95, 535-552.	6.2	569
50	Methylomic profiling implicates cortical deregulation of ANK1 in Alzheimer's disease. Nature Neuroscience, 2014, 17, 1164-1170.	14.8	488
51	Missense variant in TREML2 protects against Alzheimer's disease. Neurobiology of Aging, 2014, 35, 1510.e19-1510.e26.	3.1	110
52	Alzheimer's disease susceptibility variants in the MS4A6A gene are associated with altered levels of MS4A6A expression in blood. Neurobiology of Aging, 2014, 35, 279-290.	3.1	56
53	A Genome-wide Association Analysis of a Broad Psychosis Phenotype Identifies Three Loci for Further Investigation. Biological Psychiatry, 2014, 75, 386-397.	1.3	44
54	Gene-Wide Analysis Detects Two New Susceptibility Genes for Alzheimer's Disease. PLoS ONE, 2014, 9, e94661.	2.5	155

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55	Genome-wide association analysis identifies 13 new risk loci for schizophrenia. Nature Genetics, 2013, 45, 1150-1159.	21.4	1,395
56	Residual association at C9orf72 suggests an alternative amyotrophic lateral sclerosis-causing hexanucleotide repeat. Neurobiology of Aging, 2013, 34, 2234.e1-2234.e7.	3.1	22
57	Smell identification function as a severity and progression marker in Alzheimer's disease. International Psychogeriatrics, 2013, 25, 1157-1166.	1.0	68
58	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
59	<i>TREM2</i> Variants in Alzheimer's Disease. New England Journal of Medicine, 2013, 368, 117-127.	27.0	2,385
60	The C9ORF72 expansion mutation is a common cause of ALS+/â^'FTD in Europe and has a single founder. European Journal of Human Genetics, 2013, 21, 102-108.	2.8	201
61	The Role of ABCA1 Gene Sequence Variants on Risk of Alzheimer's Disease. Journal of Alzheimer's Disease, 2013, 38, 897-906.	2.6	45
62	Effect of DISC1 on the P300 Waveform in Psychosis. Schizophrenia Bulletin, 2013, 39, 161-167.	4.3	19
63	Entorhinal Cortex Thickness Predicts Cognitive Decline in Alzheimer's Disease. Journal of Alzheimer's Disease, 2013, 33, 755-766.	2.6	105
64	Credibility Analysis of Putative Disease-Causing Genes Using Bioinformatics. PLoS ONE, 2013, 8, e64899.	2.5	13
65	Development of a Smartphone App for a Genetics Website: The Amyotrophic Lateral Sclerosis Online Genetics Database (ALSoD). JMIR MHealth and UHealth, 2013, 1, e18.	3.7	51
66	Evidence for a role of the rare p.A152T variant in MAPT in increasing the risk for FTD-spectrum and Alzheimer's diseases. Human Molecular Genetics, 2012, 21, 3500-3512.	2.9	198
67	Plasma Transthyretin as a Candidate Marker for Alzheimer's Disease. Journal of Alzheimer's Disease, 2012, 28, 369-375.	2.6	86
68	The Role of Variation at AβPP, PSEN1, PSEN2, and MAPT in Late Onset Alzheimer's Disease. Journal of Alzheimer's Disease, 2012, 28, 377-387.	2.6	53
69	Alzheimer's disease and age-related macular degeneration have different genetic models for complement gene variation. Neurobiology of Aging, 2012, 33, 1843.e9-1843.e17.	3.1	24
70	Association of serotonin and dopamine gene pathways with behavioral subphenotypes in dementia. Neurobiology of Aging, 2012, 33, 791-803.	3.1	49
71	Functional and genetic analysis of haplotypic sequence variation at the nicastrin genomic locus. Neurobiology of Aging, 2012, 33, 1848.e1-1848.e13.	3.1	5
72	Genetic variants influencing human aging from late-onset Alzheimer's disease (LOAD) genome-wide association studies (GWAS). Neurobiology of Aging, 2012, 33, 1849.e5-1849.e18.	3.1	43

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73	C9ORF72 repeat expansion in a large Italian ALS cohort: evidence of a founder effect. Neurobiology of Aging, 2012, 33, 2528.e7-2528.e14.	3.1	74
74	Complement activation as a biomarker for Alzheimer's disease. Immunobiology, 2012, 217, 204-215.	1.9	59
75	Missense substitutions associated with behavioural disturbances in Alzheimer's disease (AD). Brain Research Bulletin, 2012, 88, 394-405.	3.0	6
76	Confirmation that the AKT1 (rs2494732) Genotype Influences the Risk of Psychosis in Cannabis Users. Biological Psychiatry, 2012, 72, 811-816.	1.3	212
77	Identification of <i>cis-</i> regulatory variation influencing protein abundance levels in human plasma. Human Molecular Genetics, 2012, 21, 3719-3726.	2.9	94
78	ALSoD: A user-friendly online bioinformatics tool for amyotrophic lateral sclerosis genetics. Human Mutation, 2012, 33, 1345-1351.	2.5	262
79	The impact of the Val ¹⁵⁸ Met catechol- <i>O</i> -methyltransferase genotype on neural correlates of sad facial affect processing in patients with bipolar disorder and their relatives. Psychological Medicine, 2011, 41, 779-788.	4.5	58
80	A Multiple Indicators Multiple Causes (MIMIC) model of Behavioural and Psychological Symptoms in Dementia (BPSD). Neurobiology of Aging, 2011, 32, 434-442.	3.1	64
81	No association of DPP6 with amyotrophic lateral sclerosis in an Italian population. Neurobiology of Aging, 2011, 32, 966-967.	3.1	28
82	Common variants at ABCA7, MS4A6A/MS4A4E, EPHA1, CD33 and CD2AP are associated with Alzheimer's disease. Nature Genetics, 2011, 43, 429-435.	21.4	1,708
83	Genome-wide association with MRI atrophy measures as a quantitative trait locus for Alzheimer's disease. Molecular Psychiatry, 2011, 16, 1130-1138.	7.9	133
84	No evidence that extended tracts of homozygosity are associated with Alzheimer's disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 764-771.	1.7	17
85	Does intravenous î"9-tetrahydrocannabinol increase dopamine release? A SPET study. Journal of Psychopharmacology, 2011, 25, 1462-1468.	4.0	84
86	Do COMT, BDNF and NRG1 polymorphisms influence P50 sensory gating in psychosis?. Psychological Medicine, 2011, 41, 263-276.	4.5	34
87	Deep Sequencing of the Nicastrin Gene in Pooled DNA, the Identification of Genetic Variants That Affect Risk of Alzheimer's Disease. PLoS ONE, 2011, 6, e17298.	2.5	21
88	Education, occupation and retirement age effects on the age of onset of Alzheimer's disease. International Journal of Geriatric Psychiatry, 2010, 25, 30-36.	2.7	34
89	Relapse to smoking during unaided cessation: clinical, cognitive and motivational predictors. Psychopharmacology, 2010, 212, 537-549.	3.1	146
90	Chromosome 9p21 in sporadic amyotrophic lateral sclerosis in the UK and seven other countries: a genome-wide association study. Lancet Neurology, The, 2010, 9, 986-994.	10.2	205

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91	Effect of APOE ε4 Allele on Cortical Thicknesses and Volumes: The AddNeuroMed Study. Journal of Alzheimer's Disease, 2010, 21, 947-966.	2.6	82
92	Genetic Evidence Implicates the Immune System and Cholesterol Metabolism in the Aetiology of Alzheimer's Disease. PLoS ONE, 2010, 5, e13950.	2.5	347
93	Evidence for Varied Aetiologies Regulating the Transmission of Prion Disease: Implications for Understanding the Heritable Basis of Prion Incubation Times. PLoS ONE, 2010, 5, e14186.	2.5	8
94	APOE ε2 Allele Is Associated with Larger Regional Cortical Thicknesses and Volumes. Dementia and Geriatric Cognitive Disorders, 2010, 30, 229-237.	1.5	40
95	PONM19 The ALS Online Genetics Database. Journal of Neurology, Neurosurgery and Psychiatry, 2010, 81, e65-e65.	1.9	0
96	Association of Plasma Clusterin Concentration With Severity, Pathology, and Progression in Alzheimer Disease. Archives of General Psychiatry, 2010, 67, 739.	12.3	353
97	CANNABIS USE AND PSYCHOTIC EXPERIENCES IN A HEALTHY POPULATION SAMPLE. Schizophrenia Research, 2010, 117, 315-316.	2.0	0
98	Variation in DRD2 dopamine gene predicts Extraverted personality. Neuroscience Letters, 2010, 468, 234-237.	2.1	54
99	Genes of the serotonergic and dopaminergic pathways and their interaction affect the expression of Behavioural and Psychological Symptoms in Dementia (BPSD) Nature Precedings, 2009, , .	0.1	0
100	Variants of the elongator protein 3 (ELP3) gene are associated with motor neuron degeneration. Human Molecular Genetics, 2009, 18, 472-481.	2.9	512
101	The acute effects of synthetic intravenous Δ9-tetrahydrocannabinol on psychosis, mood and cognitive functioning. Psychological Medicine, 2009, 39, 1607.	4.5	259
102	Vulnerability to depression: what is the role of stress genes in gene × environment interaction?. Psychological Medicine, 2009, 39, 1407-1411.	4.5	50
103	Reduced expression of the <i>Kinesin-Associated Protein 3</i> (<i>KIFAP3</i>) gene increases survival in sporadic amyotrophic lateral sclerosis. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 9004-9009.	7.1	177
104	A functional polymorphism of the brain derived neurotrophic factor gene and cortical anatomy in autism spectrum disorder. Journal of Neurodevelopmental Disorders, 2009, 1, 215-223.	3.1	37
105	Patterns of change in withdrawal symptoms, desire to smoke, reward motivation and response inhibition across 3 months of smoking abstinence. Addiction, 2009, 104, 850-858.	3.3	66
106	Genome-wide association study identifies variants at CLU and PICALM associated with Alzheimer's disease. Nature Genetics, 2009, 41, 1088-1093.	21.4	2,697
107	High-potency cannabis and the risk of psychosis. British Journal of Psychiatry, 2009, 195, 488-491.	2.8	465
108	Meta-analysis of linkage studies for Alzheimer's disease—A web resource. Neurobiology of Aging, 2009, 30, 1037-1047.	3.1	58

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109	Evidence that variation in the oligodendrocyte lineage transcription factor 2 (OLIG2) gene is associated with psychosis in Alzheimer's disease. Neuroscience Letters, 2009, 461, 54-59.	2.1	30
110	Serotonin transporter genotype and neuroanatomy in autism spectrum disorders. Psychiatric Genetics, 2009, 19, 147-150.	1.1	19
111	The effects of gender and COMT Val158Met polymorphism on fearful facial affect recognition: a fMRI study. International Journal of Neuropsychopharmacology, 2009, 12, 371.	2.1	77
112	What is the mechanism whereby cannabis use increases risk of psychosis?. Neurotoxicity Research, 2008, 14, 105-112.	2.7	53
113	Association analysis of 528 intraâ€genic SNPs in a region of chromosome 10 linked to late onset Alzheimer's disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 727-731.	1.7	40
114	Glycogen synthase kinaseâ€3β and tau genes interact in Alzheimer's disease. Annals of Neurology, 2008, 64, 446-454.	5.3	65
115	Positional Pathway Screen of wnt Signaling Genes in Schizophrenia: Association with DKK4. Biological Psychiatry, 2008, 63, 13-16.	1.3	37
116	ALSOD: The Amyotrophic Lateral Sclerosis Online Database. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2008, 9, 249-250.	2.1	128
117	Association study on glutathione Sâ€ŧransferase omega 1 and 2 and familial ALS. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2008, 9, 81-84.	2.1	19
118	Catechol-O-Methyltransferase (COMT) Val158Met Genotype is Associated with BOLD Response as a Function of Task Characteristic. Neuropsychopharmacology, 2008, 33, 3046-3057.	5.4	51
119	Interaction between theADAM12 andSH3MD1 genes may confer susceptibility to late-onset Alzheimer's disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 448-452.	1.7	27
120	Candidate gene association study of insulin signaling genes and Alzheimer's disease: Evidence forSOS2,PCK1, andPPARγas susceptibility loci. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 508-516.	1.7	54
121	Increased familial risk and genomewide significant linkage for Alzheimer's disease with psychosis. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 841-848.	1.7	45
122	A double-blind placebo-controlled experimental study of nicotine: II—Effects on response inhibition and executive functioning. Psychopharmacology, 2007, 190, 457-467.	3.1	73
123	Age at onset in sod1-mediated amyotrophic lateral sclerosis shows familiality. Neurogenetics, 2007, 8, 235-236.	1.4	14
124	Complement Factor H Y402H Polymorphism is not Associated with Late-onset Alzheimer's Disease. NeuroMolecular Medicine, 2007, 9, 331-334.	3.4	22
125	A Scan of Chromosome 10 Identifies a Novel Locus Showing Strong Association with Late-Onset Alzheimer Disease. American Journal of Human Genetics, 2006, 78, 78-88.	6.2	157
126	Proteome-based plasma biomarkers for Alzheimer's disease. Brain, 2006, 129, 3042-3050.	7.6	427

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127	Polymorphisms in the phosphate and tensin homolog gene are not associated with late-onset Alzheimer's disease. Neuroscience Letters, 2006, 401, 77-80.	2.1	7
128	An association analysis of candidate genes on chromosome 15 q11–13 and autism spectrum disorder. Molecular Psychiatry, 2006, 11, 709-713.	7.9	19
129	A double-blind placebo controlled experimental study of nicotine: l—effects on incentive motivation. Psychopharmacology, 2006, 189, 355-367.	3.1	88
130	The BDNF val66met polymorphism is not associated with late onset Alzheimer's disease in three case–control samples. Molecular Psychiatry, 2005, 10, 809-810.	7.9	33
131	Genetic association of the APP binding protein 2 gene (APBB2) with late onset Alzheimer disease. Human Mutation, 2005, 25, 270-277.	2.5	36
132	Molecular and phenotypic characterization of ring chromosome 22. American Journal of Medical Genetics, Part A, 2005, 137A, 139-147.	1.2	86
133	Candidate gene association studies of genes involved in neuronal cholinergic transmission in Alzheimer's disease suggests choline acetyltransferase as a candidate deserving further study. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2005, 132B, 5-8.	1.7	37
134	A central resource for accurate allele frequency estimation from pooled DNA genotyped on DNA microarrays. Nucleic Acids Research, 2005, 33, e25-e25.	14.5	39
135	Nicastrin gene polymorphisms, cognitive ability level and cognitive ageing. Neuroscience Letters, 2005, 373, 110-114.	2.1	24
136	MaGIC: a program to generate targeted marker sets for genome-wide association studies. BioTechniques, 2004, 37, 996-999.	1.8	10
137	Association of late-onset Alzheimer's disease with genetic variation in multiple members of the <i>GAPD</i> gene family. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 15688-15693.	7.1	134
138	α-T-Catenin Is Expressed in Human Brain and Interacts With the Wnt Signaling Pathway But Is Not Responsible for Linkage to Chromosome 10 in Alzheimer's Disease. NeuroMolecular Medicine, 2004, 5, 133-146.	3.4	41
139	P4-122 Genetic association of an APP binding protein gene with late onset Alzheimer's disease. Neurobiology of Aging, 2004, 25, S510.	3.1	0
140	ACE genotype and cognitive decline in an African-Caribbean population. Neurobiology of Aging, 2004, 25, 1369-1375.	3.1	18
141	Candidate gene association studies of the α4 (CHRNA4) and β2 (CHRNB2) neuronal nicotinic acetylcholine receptor subunit genes in Alzheimer's disease. Neuroscience Letters, 2004, 358, 142-146.	2.1	40
142	Association of ABCA1 with late-onset Alzheimer's disease is not observed in a case-control study. Neuroscience Letters, 2004, 366, 268-271.	2.1	58
143	Glycogen synthase kinase-3 is increased in white cells early in Alzheimer's disease. Neuroscience Letters, 2004, 373, 1-4.	2.1	112
144	Cognitive and psychological correlates of smoking abstinence, and predictors of successful cessation. Addictive Behaviors, 2004, 29, 1407-1426.	3.0	68

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145	Variants in the ALS2 gene are not associated with sporadic amyotrophic lateral sclerosis. Neurogenetics, 2003, 4, 221-222.	1.4	18
146	Sequence variation in the CHAT locus shows no association with late-onset Alzheimer's disease. Human Genetics, 2003, 113, 258-267.	3.8	33
147	Depression in Alzheimer's disease: The effect of serotonin receptor gene variation. American Journal of Medical Genetics Part A, 2003, 119B, 40-43.	2.4	58
148	β-1,3-Glucuronyltransferase-1 gene implicated as a candidate for a schizophrenia-like psychosis through molecular analysis of a balanced translocation. Molecular Psychiatry, 2003, 8, 654-663.	7.9	34
149	Two Families with Familial Amyotrophic Lateral Sclerosis Are Linked to a Novel Locus on Chromosome 16q. American Journal of Human Genetics, 2003, 73, 390-396.	6.2	76
150	Genetics, molecular biology, neuropathology and phenotype of frontal lobe dementia. British Journal of Psychiatry, 2002, 180, 455-460.	2.8	3
151	Genetic variability in the insulin signalling pathway may contribute to the risk of late onset Alzheimer's disease. Journal of Neurology, Neurosurgery and Psychiatry, 2002, 73, 261-266.	1.9	38
152	Identification of genomic organisation, sequence variants and analysis of the role of the human dishevelled 1 gene in late onset Alzheimer's disease. Molecular Psychiatry, 2002, 7, 104-109.	7.9	9
153	The extended haplotype of the microtubule associated protein tau gene is not associated with Pick's disease. Neuroscience Letters, 2001, 299, 156-158.	2.1	35
154	The microtubule associated protein Tau gene and Alzheimer's disease – an association study and meta-analysis. Neuroscience Letters, 2001, 314, 92-96.	2.1	52
155	Intron 7 retention and exon 9 skipping EAAT2 mRNA variants are not associated with amyotrophic lateral sclerosis. Annals of Neurology, 2001, 49, 643-649.	5.3	68
156	Association study of the 5-HT2A receptor gene polymorphism, T102C and essential hypertension. Journal of Human Hypertension, 2001, 15, 335-339.	2.2	33
157	Identification of sequence variants and analysis of the role of the glycogen synthase kinase 3 β gene and promoter in late onset Alzheimer's disease. Molecular Psychiatry, 2001, 6, 320-324.	7.9	61
158	Psychosis and aggression in Alzheimer's disease: the effect of dopamine receptor gene variation. Journal of Neurology, Neurosurgery and Psychiatry, 2001, 71, 777-779.	1.9	98
159	Systematic screening of the 14-3-3 eta (?) chain gene for polymorphic variants and case-control analysis in schizophrenia. American Journal of Medical Genetics Part A, 2000, 96, 736-743.	2.4	38
160	Variation in DCP1, encoding ACE, is associated with susceptibility to Alzheimer disease. Nature Genetics, 1999, 21, 71-72.	21.4	260
161	Autosome search for schizophrenia susceptibility genes in multiply affected families. Molecular Psychiatry, 1999, 4, 353-359.	7.9	20
162	Mutations in the gene encoding human persyn are not associated with amyotrophic lateral sclerosis or familial Parkinson's disease. Neuroscience Letters, 1999, 274, 21-24.	2.1	16

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163	Circadian regulation of prion protein messenger RNA in the rat forebrain: a widespread and synchronous rhythm. Neuroscience, 1999, 91, 1201-1204.	2.3	28
164	Deletions of the heavy neurofilament subunit tail in amyotrophic lateral sclerosis. Human Molecular Genetics, 1999, 8, 157-164.	2.9	303
165	Mutations in all five exons of <i>SODâ€l </i> may cause ALS. Annals of Neurology, 1998, 43, 390-394.	5.3	153
166	Variation in the expression of the mRNA for protein kinase C isoforms in the rat suprachiasmatic nuclei, caudate putamen and cerebral cortex. Molecular Brain Research, 1998, 53, 277-284.	2.3	24
167	Apolipoprotein E: Depressive illness, depressive symptoms, and Alzheimer's disease. Biological Psychiatry, 1998, 43, 159-164.	1.3	44
168	5-HT2A and 5-HT2C receptor polymorphisms and psychopathology in late onset Alzheimer's disease. Human Molecular Genetics, 1998, 7, 1507-1509.	2.9	175
169	Recessive amyotrophic lateral sclerosis families with the D90A SOD1 mutation share a common founder: evidence for a linked protective factor. Human Molecular Genetics, 1998, 7, 2045-2050.	2.9	132
170	Failure to exclude a possible schizophrenia susceptibility locus on chromosome 13q14.1-q32 in Southern African Bantu-speaking families. Psychiatric Genetics, 1998, 8, 155-162.	1.1	15
171	Endothelial Nitric Oxide Synthase Exon 7 Polymorphism, Ischemic Cerebrovascular Disease, and Carotid Atheroma. Stroke, 1998, 29, 1908-1911.	2.0	140
172	Autism and multiple exostoses associated with an X;8 translocation occurring within the GRPR gene and 3' to the SDC2 gene. Human Molecular Genetics, 1997, 6, 1241-1250.	2.9	100
173	Allelic functional variation of serotonin transporter expression is a susceptibility factor for late onset Alzheimer's disease. NeuroReport, 1997, 8, 683-686.	1.2	103
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