## Christopher M Morris

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	High levels of mitochondrial DNA deletions in substantia nigra neurons in aging and Parkinson disease. Nature Genetics, 2006, 38, 515-517.	9.4	1,363
2	Mutation in the gene encoding ferritin light polypeptide causes dominant adult-onset basal ganglia disease. Nature Genetics, 2001, 28, 350-354.	9.4	533
3	D2 dopamine receptor gene (DRD2) Taql A polymorphism: reduced dopamine D2 receptor binding in the human striatum associated with the A1 allele. Pharmacogenetics and Genomics, 1997, 7, 479-484.	5.7	518
4	Identification of common variants influencing risk of the tauopathy progressive supranuclear palsy. Nature Genetics, 2011, 43, 699-705.	9.4	502
5	Alteration in nicotine binding sites in Parkinson's disease, Lewy body dementia and Alzheimer's disease: Possible index of early neuropathology. Neuroscience, 1995, 64, 385-395.	1.1	396
6	A Multicenter Study of Glucocerebrosidase Mutations in Dementia With Lewy Bodies. JAMA Neurology, 2013, 70, 727.	4.5	374
7	Frontotemporal dementia and its subtypes: a genome-wide association study. Lancet Neurology, The, 2014, 13, 686-699.	4.9	302
8	Increase in Interleukin-1β in Late-Life Depression. American Journal of Psychiatry, 2005, 162, 175-177.	4.0	269
9	Common genetic variation within the Low-Density Lipoprotein Receptor-Related Protein 6 and late-onset Alzheimer's disease. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 9434-9439.	3.3	252
10	The MAPT H1c risk haplotype is associated with increased expression of tau and especially of 4 repeat containing transcripts. Neurobiology of Disease, 2007, 25, 561-570.	2.1	231
11	Linkage disequilibrium fine mapping and haplotype association analysis of the tau gene in progressive supranuclear palsy and corticobasal degeneration. Journal of Medical Genetics, 2005, 42, 837-846.	1.5	225
12	Cholinergic Transmitter and Neurotrophic Activities in Lewy Body Dementia. Alzheimer Disease and Associated Disorders, 1993, 7, 69-79.	0.6	219
13	The H1c haplotype at the MAPT locus is associated with Alzheimer's disease. Human Molecular Genetics, 2005, 14, 2399-2404.	1.4	205
14	Dementia with Lewy bodies: an update and outlook. Molecular Neurodegeneration, 2019, 14, 5.	4.4	203
15	Genome sequencing analysis identifies new loci associated with Lewy body dementia and provides insights into its genetic architecture. Nature Genetics, 2021, 53, 294-303.	9.4	198
16	Histochemical Distribution of Non-Haem Iron in the Human Brain. Cells Tissues Organs, 1992, 144, 235-257.	1.3	193
17	Mitochondrial <scp>DNA</scp> Depletion in Respiratory Chain–Deficient <scp>P</scp> arkinson Disease Neurons. Annals of Neurology, 2016, 79, 366-378.	2.8	189
18	Single-cell sequencing of human midbrain reveals glial activation and a Parkinson-specific neuronal state. Brain, 2022, 145, 964-978.	3.7	177

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19	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.	2.6	157
20	White matter connections of the supplementary motor area in humans. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 1377-1385.	0.9	151
21	Up-regulation of the inflammatory cytokines IFN-Î <sup>3</sup> and IL-12 and down-regulation of IL-4 in cerebral cortex regions of APPSWE transgenic mice. Journal of Neuroimmunology, 2002, 126, 50-57.	1.1	150
22	Genetic modifiers of risk and age at onset in GBA associated Parkinson's disease and Lewy body dementia. Brain, 2020, 143, 234-248.	3.7	149
23	Peripheral inflammation in prodromal Alzheimer's and Lewy body dementias. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 339-345.	0.9	141
24	SIRT1 ameliorates oxidative stress induced neural cell death and is down-regulated in Parkinson's disease. BMC Neuroscience, 2017, 18, 46.	0.8	140
25	Quantification of Alzheimer pathology in ageing and dementia: age-related accumulation of amyloid-beta(42) peptide in vascular dementia. Neuropathology and Applied Neurobiology, 2006, 32, 103-118.	1.8	131
26	Distribution of Nicotinic Receptors in the Human Hippocampus and Thalamus. European Journal of Neuroscience, 1994, 6, 1596-1604.	1.2	130
27	Uptake and Distribution of Iron and Transferrin in the Adult Rat Brain. Journal of Neurochemistry, 1992, 59, 300-306.	2.1	129
28	Dopamine and nicotinic receptor binding and the levels of dopamine and homovanillic acid in human brain related to tobacco use. Neuroscience, 1998, 87, 63-78.	1.1	127
29	Expression analysis of dopaminergic neurons in Parkinson's disease and aging links transcriptional dysregulation of energy metabolism to cell death. Acta Neuropathologica, 2011, 122, 75-86.	3.9	127
30	Dementia with Lewy bodies. Seminars in Clinical Neuropsychiatry, 2003, 8, 46-57.	1.9	124
31	Nature of Mitochondrial DNA Deletions in Substantia Nigra Neurons. American Journal of Human Genetics, 2008, 82, 228-235.	2.6	123
32	Selective loss of glucocerebrosidase activity in sporadic Parkinson's disease and dementia with Lewy bodies. Molecular Neurodegeneration, 2015, 10, 15.	4.4	120
33	Towards a definition of neurodisability: a Delphi survey. Developmental Medicine and Child Neurology, 2013, 55, 1103-1108.	1.1	113
34	The progression of cognitive impairment in dementia with Lewy bodies, vascular dementia and Alzheimer's disease. International Journal of Geriatric Psychiatry, 2001, 16, 499-503.	1.3	106
35	Does the mitochondrial genome play a role in the etiology of Alzheimer's disease?. Human Genetics, 2006, 119, 241-254	1.8	102
36	Dementia with Lewy Bodies. A Distinct Nonâ€Alzheimer Dementia Syndrome?. Brain Pathology, 1998, 8, 299-324.	2.1	96

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37	Cortical serotonin-S2 receptor binding in Lewy body dementia, Alzheimer's and Parkinson's diseases. Journal of the Neurological Sciences, 1991, 106, 50-55.	0.3	95
38	Mitochondrial dysfunction within the synapses of substantia nigra neurons in Parkinson's disease. Npj Parkinson's Disease, 2018, 4, 9.	2.5	92
39	A nonsynonymous mutation in PLCG2 reduces the risk of Alzheimer's disease, dementia with Lewy bodies and frontotemporal dementia, and increases the likelihood of longevity. Acta Neuropathologica, 2019, 138, 237-250.	3.9	87
40	Regulation of attention and response to therapy in dementia by butyrylcholinesterase. Pharmacogenetics and Genomics, 2003, 13, 231-239.	5.7	85
41	Brain iron homeostasis. Journal of Inorganic Biochemistry, 1992, 47, 257-265.	1.5	84
42	Aberrant Splicing in the Presenilin-1 Intron 4 Mutation Causes Presenile Alzheimer's Disease by Increased AÂ42 Secretion. Human Molecular Genetics, 1999, 8, 1529-1540.	1.4	84
43	Apolipoprotein E genes in Lewy body and Parkinson's disease. Lancet, The, 1994, 343, 1565.	6.3	83
44	Mitochondrial DNA point mutations and relative copy number in 1363 disease and control human brains. Acta Neuropathologica Communications, 2017, 5, 13.	2.4	83
45	Apolipoprotein E ϵ4 Allele, Temporal Lobe Atrophy, and White Matter Lesions in Late-Life Dementias. Archives of Neurology, 1999, 56, 961.	4.9	82
46	Apolipoprotein Â3 allele is associated with persistent hepatitis C virus infection. Gut, 2006, 55, 715-718.	6.1	81
47	Brain matrix metalloproteinase 1 levels are elevated in Alzheimer's disease. Neuroscience Letters, 2000, 291, 201-203.	1.0	80
48	Protective effect of apoE ∈2 in Alzheimer's disease. Lancet, The, 1994, 344, 473-474.	6.3	76
49	Decreased Fractalkine and Increased IP-10 Expression in Aged Brain of APPswe Transgenic Mice. Neurochemical Research, 2008, 33, 1085-1089.	1.6	74
50	FUS and TDP43 genetic variability in FTD and CBS. Neurobiology of Aging, 2012, 33, 1016.e9-1016.e17.	1.5	69
51	APOA1 polymorphism influences risk for early-onset nonfamiliar AD. Annals of Neurology, 2005, 58, 436-441.	2.8	68
52	Interventions utilising contact with people with disabilities to improve children's attitudes towards disability: A systematic review and meta-analysis. Disability and Health Journal, 2017, 10, 11-22.	1.6	68
53	Neuroferritinopathy: A Window on the Role of Iron in Neurodegeneration. Blood Cells, Molecules, and Diseases, 2002, 29, 522-531.	0.6	67
54	Comparison of the regional distribution of transferrin receptors and aluminium in the forebrain of chronic renal dialysis patients. Journal of the Neurological Sciences, 1989, 94, 295-306.	0.3	66

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55	Lack of association between the dopamine D2 receptor gene allele DRD2*A1 and cigarette smoking in a United Kingdom population. Pharmacogenetics and Genomics, 1998, 8, 125-128.	5.7	66
56	Parkinson's disease is not associated with the combined ?-synuclein/apolipoprotein E susceptibility genotype. Annals of Neurology, 2001, 49, 665-668.	2.8	66
57	Preliminary observation of elevated levels of nanocrystalline iron oxide in the basal ganglia of neuroferritinopathy patients. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2007, 1772, 21-25.	1.8	66
58	The role of the cholinergic system in the development of the human cerebellum. Developmental Brain Research, 1995, 90, 159-167.	2.1	64
59	Selective Nicotinic Receptor Consequences in APPSWE Transgenic Mice. Molecular and Cellular Neurosciences, 2002, 20, 354-365.	1.0	60
60	Morphometric analysis of neuronal and glial cell pathology in the dorsolateral prefrontal cortex in late-life depression. British Journal of Psychiatry, 2009, 195, 163-169.	1.7	59
61	No association between the K variant of the butyrylcholinesterase gene and pathologically confirmed Alzheimer's disease. Human Molecular Genetics, 1998, 7, 937-939.	1.4	58
62	Glucocerebrosidase Mutations alter the endoplasmic reticulum and lysosomes in Lewy body disease. Journal of Neurochemistry, 2012, 123, 298-309.	2.1	58
63	Analysis of primary visual cortex in dementia with Lewy bodies indicates GABAergic involvement associated with recurrent complex visual hallucinations. Acta Neuropathologica Communications, 2016, 4, 66.	2.4	58
64	Autoradiographic comparison of cholinergic and other transmitter receptors in the normal human hippocampus. Hippocampus, 1993, 3, 307-315.	0.9	57
65	Clinical and Neuropathological Correlates of Apolipoprotein E Genotype in Dementia with Lewy Bodies. Dementia and Geriatric Cognitive Disorders, 2002, 14, 167-175.	0.7	57
66	HLA-DR antigens associated with major genetic risk for late-onset Alzheimer's disease. NeuroReport, 1997, 8, 1467-1469.	0.6	56
67	A Systematic Review of Generic Multidimensional Patient-Reported Outcome Measures for Children, Part I: Descriptive Characteristics. Value in Health, 2015, 18, 315-333.	0.1	56
68	Comparative proteomic analysis using samples obtained with laser microdissection and saturation dye labelling. Proteomics, 2005, 5, 3851-3858.	1.3	55
69	Gallium-67 as a Potential Marker for Aluminium Transport in Rat Brain: Implications for Alzheimer's Disease. Journal of Neurochemistry, 1990, 55, 251-259.	2.1	54
70	Intralobar fibres of the occipital lobe: A post mortem dissection study. Cortex, 2014, 56, 145-156.	1.1	54
71	High prevalence of focal and multi-focal somatic genetic variants in the human brain. Nature Communications, 2018, 9, 4257.	5.8	54
72	Chronic glial activation, neurodegeneration, and APP immunoreactive deposits following acute administration of double-stranded RNA. Glia, 2003, 44, 1-12.	2.5	53

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73	Mitochondrial Abnormality Associates with Type-Specific Neuronal Loss and Cell Morphology Changes in the Pedunculopontine Nucleus in Parkinson Disease. American Journal of Pathology, 2013, 183, 1826-1840.	1.9	53
74	Pyroglutamylated amyloid-β is associated with hyperphosphorylated tau and severity of Alzheimer's disease. Acta Neuropathologica, 2014, 128, 67-79.	3.9	53
75	Hereditary multi-infarct dementia of the Swedish type is a novel disorder different from NOTCH3 causing CADASIL. Brain, 2007, 130, 357-367.	3.7	51
76	Transferrin and HFE genes interact in Alzheimer's disease risk: the Epistasis Project. Neurobiology of Aging, 2012, 33, 202.e1-202.e13.	1.5	51
77	Somatic variants in autosomal dominant genes are a rare cause of sporadic Alzheimer's disease. Alzheimer's and Dementia, 2018, 14, 1632-1639.	0.4	51
78	Distribution of neuronal nicotinic receptor subunits in human brain. Neurochemistry International, 1994, 25, 69-71.	1.9	50
79	C9orf72 intermediate repeats are associated with corticobasal degeneration, increased C9orf72 expression and disruption of autophagy. Acta Neuropathologica, 2019, 138, 795-811.	3.9	50
80	Evidence for the localization of haemopexin immunoreactivity in neurones in the human brain. Neuroscience Letters, 1993, 149, 141-144.	1.0	49
81	Transferrin receptors in the Parkinsonian midbrain. Neuropathology and Applied Neurobiology, 1994, 20, 468-472.	1.8	49
82	Singleâ€cell expression profiling of dopaminergic neurons combined with association analysis identifies pyridoxal kinase as Parkinson's disease gene. Annals of Neurology, 2009, 66, 792-798.	2.8	49
83	Variation in tau isoform expression in different brain regions and disease states. Neurobiology of Aging, 2013, 34, 1922.e7-1922.e12.	1.5	49
84	Genetic Variability in <i>CHMP2B</i> and Frontotemporal Dementia. Neurodegenerative Diseases, 2006, 3, 129-133.	0.8	47
85	A morphometric examination of neuronal and glial cell pathology in the orbitofrontal cortex in late-life depression. International Psychogeriatrics, 2011, 23, 132-140.	0.6	45
86	Mitochondrial DNA changes in pedunculopontine cholinergic neurons in Parkinson disease. Annals of Neurology, 2017, 82, 1016-1021.	2.8	45
87	Neuroblastoma and Alzheimer's Disease Brain Cells Contain Aromatase Activity. Steroids, 1998, 63, 263-267.	0.8	44
88	Genetic compendium of 1511 human brains available through the UK Medical Research Council Brain Banks Network Resource. Genome Research, 2017, 27, 165-173.	2.4	44
89	Relationship Between Mitochondria and α-Synuclein. Archives of Neurology, 2012, 69, 385.	4.9	43
90	Development of passive CLARITY and immunofluorescent labelling of multiple proteins in human cerebellum: understanding mechanisms of neurodegeneration in mitochondrial disease. Scientific Reports, 2016, 6, 26013.	1.6	43

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91	Novel presenilin 1 mutation with profound neurofibrillary pathology in an indigenous Southern African family with early-onset Alzheimer's disease. Brain, 2004, 127, 133-142.	3.7	42
92	Diagnosis of Mitochondrial Disease: Assessment of Mitochondrial DNA Heteroplasmy in Blood. Biochemical and Biophysical Research Communications, 1998, 251, 883-887.	1.0	41
93	Brain oestradiol and testosterone levels in Alzheimer's disease. Neuroscience Letters, 2000, 286, 1-4.	1.0	41
94	Low-level repeated exposure to diazinon and chlorpyrifos decrease anxiety-like behaviour in adult male rats as assessed by marble burying behaviour. NeuroToxicology, 2015, 50, 149-156.	1.4	41
95	The tau locus is not significantly associated with pathologically confirmed sporadic Parkinson's disease. Neuroscience Letters, 2002, 330, 201-203.	1.0	39
96	BuChEâ€K and APOE Ϊμ4 allele frequencies in Lewy body dementias, and influence of genotype and hyperhomocysteinemia on cognitive decline. Movement Disorders, 2009, 24, 392-400.	2.2	39
97	A C6orf10/LOC101929163 locus is associated with age of onset in C9orf72 carriers. Brain, 2018, 141, 2895-2907.	3.7	39
98	Effect of apolipoprotein E genotype on Alzheimer's disease neuropathology in a cohort of elderly Norwegians. Neuroscience Letters, 1995, 201, 45-48.	1.0	38
99	Nitric oxide synthase gene polymorphisms in Alzheimer's disease and dementia with Lewy bodies. Neuroscience Letters, 2001, 303, 33-36.	1.0	38
100	Ubiquilin 1 polymorphisms are not associated with late-onset Alzheimer's disease. Annals of Neurology, 2006, 59, 21-26.	2.8	37
101	Immunocytochemical Localisation of Transferrin in the Human Brain. Cells Tissues Organs, 1992, 143, 14-18.	1.3	36
102	Morphometric Analysis of Neuronal and Glial Cell Pathology in the Caudate Nucleus in Late-Life Depression. American Journal of Geriatric Psychiatry, 2011, 19, 132-141.	0.6	36
103	Association studies between risk for late-onset Alzheimer's disease and variants in insulin degrading enzyme. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2005, 136B, 62-68.	1.1	35
104	Exome sequencing in dementia with Lewy bodies. Translational Psychiatry, 2016, 6, e728-e728.	2.4	35
105	Effects of Apolipoprotein E Genotype on Cortical Neuropathology in Senile Dementia of the Lewy Body and Alzheimer's Disease. Experimental Neurology, 1995, 4, 443-448.	1.7	34
106	Polymorphism in the human DJ-1 gene is not associated with sporadic dementia with Lewy bodies or Parkinson's disease. Neuroscience Letters, 2003, 352, 151-153.	1.0	34
107	Anatomic Connections of the Subgenual Cingulate Region. Neurosurgery, 2016, 79, 465-472.	0.6	34
108	Sirtuin-2 Protects Neural Cells from Oxidative Stress and Is Elevated in Neurodegeneration. Parkinson's Disease, 2017, 2017, 1-17.	0.6	34

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109	Degeneration of dopaminergic circuitry influences depressive symptoms in Lewy body disorders. Brain Pathology, 2019, 29, 544-557.	2.1	33
110	The immunhistochemical examination of GABAergic interneuron markers in the dorsolateral prefrontal cortex of patients with late-life depression. International Psychogeriatrics, 2011, 23, 644-653.	0.6	32
111	Mechanism for the acute effects of organophosphate pesticides on the adult 5-HT system. Chemico-Biological Interactions, 2016, 245, 82-89.	1.7	32
112	Specific patterns of neuronal loss in the pulvinar nucleus in dementia with lewy bodies. Movement Disorders, 2017, 32, 414-422.	2.2	32
113	Gene expression analysis reveals chronic low level exposure to the pesticide diazinon affects psychological disorders gene sets in the adult rat. Toxicology, 2018, 393, 90-101.	2.0	32
114	Distribution of transferrin receptors in relation to cytochrome oxidase activity in the human spinal cord, lower brainstem and cerebellum. Journal of the Neurological Sciences, 1992, 111, 158-172.	0.3	31
115	Transferrin gene polymorphism in Alzheimer's disease and dementia with Lewy bodies in humans. Neuroscience Letters, 2002, 317, 13-16.	1.0	31
116	The human brainome: network analysis identifies HSPA2 as a novel Alzheimer's disease target. Brain, 2018, 141, 2721-2739.	3.7	31
117	Inflammation in mild cognitive impairment due to Parkinson's disease, Lewy body disease, and Alzheimer's disease. International Journal of Geriatric Psychiatry, 2019, 34, 1244-1250.	1.3	31
118	Altered ceramide metabolism is a feature in the extracellular vesicle-mediated spread of alpha-synuclein in Lewy body disorders. Acta Neuropathologica, 2021, 142, 961-984.	3.9	31
119	Diquat causes caspase-independent cell death in SH-SY5Y cells by production of ROS independently of mitochondria. Archives of Toxicology, 2015, 89, 1811-1825.	1.9	30
120	Telomerase Activity is Downregulated Early During Human Brain Development. Genes, 2016, 7, 27.	1.0	30
121	Regional levels of physiological α-synuclein are directly associated with Lewy body pathology. Acta Neuropathologica, 2018, 135, 153-154.	3.9	30
122	The imaging and quantification of aluminium in the human brain using dynamic secondary ion mass spectrometry (SIMS). Biology of the Cell, 1992, 74, 109-118.	0.7	29
123	α2-Macroglobulin polymorphisms in Alzheimer's disease and dementia with Lewy bodies. NeuroReport, 1999, 10, 1507-1510.	0.6	29
124	The CCTTT polymorphism in the NOS2A gene is associated with dementia with Lewy bodies. NeuroReport, 2000, 11, 297-299.	0.6	29
125	Impact of Hypertension and Apolipoprotein E4 on Poststroke Cognition in Subjects >75 Years of Age. Stroke, 2005, 36, 1864-1868.	1.0	29
126	Neuronal Loss and Î <sup>r</sup> -Synuclein Pathology in the Superior Colliculus and Its Relationship to Visual Hallucinations in Dementia with Lewy Bodies. American Journal of Geriatric Psychiatry, 2017, 25, 595-604.	0.6	29

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127	Transferrin receptors in the normal human hippocampus and in Alzheimer's disease. Neuropathology and Applied Neurobiology, 1994, 20, 473-477.	1.8	28
128	Hereditary Vascular Dementia Linked to Notch 3 Mutations: CADASIL in British Families. Annals of the New York Academy of Sciences, 2000, 903, 293-298.	1.8	28
129	Distribution of Amyloid beta42 in Relation to the Cerebral Microvasculature in an Elderly Cohort with Alzheimer's Disease. Annals of the New York Academy of Sciences, 2000, 903, 83-88.	1.8	27
130	Synaptic Protein Alterations in Parkinson's Disease. Molecular Neurobiology, 2012, 45, 126-143.	1.9	27
131	Heterogeneity in αâ€synuclein subtypes and their expression in cortical brain tissue lysates from Lewy body diseases and Alzheimer's disease. Neuropathology and Applied Neurobiology, 2019, 45, 597-608.	1.8	27
132	Feasibility of a randomised controlled trial to evaluate home-based virtual reality therapy in children with cerebral palsy. Disability and Rehabilitation, 2021, 43, 85-97.	0.9	27
133	Hippocampal nerve growth factor receptor immunoreactivity in patients with Alzheimer's and Parkinson's disease. Neuroscience Letters, 1992, 143, 101-104.	1.0	26
134	The low abundance of clonally expanded mitochondrial DNA point mutations in aged substantia nigra neurons. Aging Cell, 2009, 8, 496-498.	3.0	26
135	The role of transferrin in the uptake of aluminium and manganese by the IMR 32 neuroblastoma cell line. Biochemical Society Transactions, 1987, 15, 498-498.	1.6	25
136	Iron uptake in the brain of the myelin-deficient rat. Neuroscience Letters, 1993, 154, 187-190.	1.0	25
137	Risk for Alzheimer's disease in older late-onset cases is associated with HLA-DRB1*03. Neuroscience Letters, 1999, 275, 137-140.	1.0	25
138	Distribution of nerve growth factor receptor immunoreactivity in the human hippocampus. Neuroscience Letters, 1991, 121, 178-182.	1.0	24
139	Neuroferritinopathy. International Review of Neurobiology, 2013, 110, 91-123.	0.9	24
140	Molecular changes in the absence of severe pathology in the pulvinar in dementia with Lewy bodies. Movement Disorders, 2018, 33, 982-991.	2.2	24
141	Apolipoprotein E genotyping in Alzheimer's disease. Lancet, The, 1996, 347, 1775-1776.	6.3	23
142	LRP gene and late-onset Alzheimer's disease. Lancet, The, 1998, 352, 239-240.	6.3	23
143	Cellular pathology within the anterior cingulate cortex of patients with late-life depression: A morphometric study. Psychiatry Research - Neuroimaging, 2011, 194, 184-189.	0.9	23
144	A Low Mortality, High Morbidity Reduced Intensity Status Epilepticus (RISE) Model of Epilepsy and Epileptogenesis in the Rat. PLoS ONE, 2016, 11, e0147265.	1.1	23

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145	Familial neurocardiogenic (vasovagal) syncope. American Journal of Medical Genetics, Part A, 2005, 133A, 176-179.	0.7	22
146	The K Variant of the Butyrylcholinesterase Gene Is Associated with Reduced Phosphorylation of Tau in Dementia Patients. Dementia and Geriatric Cognitive Disorders, 2005, 19, 357-360.	0.7	22
147	Changes to the lateral geniculate nucleus in A lzheimer's disease but not dementia with L ewy bodies. Neuropathology and Applied Neurobiology, 2016, 42, 366-376.	1.8	22
148	Effect of aluminium on expression and processing of amyloid precursor protein. , 1996, 46, 395-403.		21
149	Assessment of APOE in atypical parkinsonism syndromes. Neurobiology of Disease, 2019, 127, 142-146.	2.1	21
150	Health outcomes for children with neurodisability: what do professionals regard as primary targets?. Archives of Disease in Childhood, 2014, 99, 927-932.	1.0	20
151	Frequency and signature of somatic variants in 1461 human brain exomes. Genetics in Medicine, 2019, 21, 904-912.	1.1	20
152	Frequency of HLA-A and B alleles in early and late-onset Alzheimer's disease. Neuroscience Letters, 1999, 262, 140-142.	1.0	19
153	Soluble cell adhesion molecules in late-life depression. International Psychogeriatrics, 2007, 19, 914-920.	0.6	19
154	Trichloroethylene and its metabolite TaClo lead to degeneration of substantia nigra dopaminergic neurones: Effects in wild type and human A30P mutant α-synuclein mice. Neuroscience Letters, 2019, 711, 134437.	1.0	19
155	No pathogenic mutations in the synphilin-1 gene in Parkinson's disease. Neuroscience Letters, 2001, 307, 125-127.	1.0	18
156	Extended post-mortem delay times should not be viewed as a deterrent to the scientific investigation of human brain tissue: a study from the Brains for Dementia Research Network Neuropathology Study Group, UK. Acta Neuropathologica, 2016, 132, 753-755.	3.9	18
157	Transferrin and transferrin receptors in normal brain and in Alzheimer's disease. Biochemical Society Transactions, 1987, 15, 891-892.	1.6	17
158	No association between an intronic polymorphism in the presenilin-1 gene and Alzheimer's disease. Neuroscience Letters, 1997, 234, 19-22.	1.0	17
159	Sex differences in effects of low level domoic acid exposure. NeuroToxicology, 2013, 34, 1-8.	1.4	17
160	Oligogenic genetic variation of neurodegenerative disease genes in 980 postmortem human brains. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 813-816.	0.9	17
161	Genetic evaluation of dementia with Lewy bodies implicates distinct disease subgroups. Brain, 2022, 145, 1757-1762.	3.7	17
162	Butyrylcholinesterase K: an association with dementia with Lewy bodies. Lancet, The, 1998, 351, 1818.	6.3	15

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163	Screening of the regulatory and coding regions of vascular endothelial growth factor in amyotrophic lateral sclerosis. Neurogenetics, 2005, 6, 101-104.	0.7	15
164	Angiotensin converting enzyme insertion/deletion polymorphisms in vasovagal syncope. Europace, 2005, 7, 396-399.	0.7	15
165	NOS3 gene rs1799983 polymorphism and incident dementia in elderly stroke survivors. Neurobiology of Aging, 2011, 32, 554.e1-554.e6.	1.5	15
166	Neuropathological Changes in Dementia With Lewy Bodies and the Cingulate Island Sign. Journal of Neuropathology and Experimental Neurology, 2019, 78, 717-724.	0.9	15
167	Pathological Changes to the Subcortical Visual System and its Relationship to Visual Hallucinations in Dementia with Lewy Bodies. Neuroscience Bulletin, 2019, 35, 295-300.	1.5	15
168	Investigating the presence of doubly phosphorylated αâ€synuclein at tyrosine 125 and serine 129 in idiopathic Lewy body diseases. Brain Pathology, 2020, 30, 831-843.	2.1	15
169	Peripheral inflammation in mild cognitive impairment with possible and probable Lewy body disease and Alzheimer's disease. International Psychogeriatrics, 2019, 31, 551-560.	0.6	14
170	Prospective longitudinal evaluation of cytokines in mild cognitive impairment due to <scp>AD</scp> and Lewy body disease. International Journal of Geriatric Psychiatry, 2020, 35, 1250-1259.	1.3	14
171	RT-QuIC Using C-Terminally Truncated α-Synuclein Forms Detects Differences in Seeding Propensity of Different Brain Regions from Synucleinopathies. Biomolecules, 2021, 11, 820.	1.8	14
172	High throughput approaches in neuroscience. International Journal of Developmental Neuroscience, 2004, 22, 515-522.	0.7	13
173	Core Health Outcomes In Childhood Epilepsy (CHOICE): protocol for the selection of a core outcome set. Trials, 2017, 18, 572.	0.7	13
174	Labelâ€Free Nanoimaging of Neuromelanin in the Brain by Soft Xâ€ray Spectromicroscopy. Angewandte Chemie - International Edition, 2020, 59, 11984-11991.	7.2	13
175	Hippocampal p75 Nerve Growth Factor Receptor Immunoreactivity in Development, Normal Aging and Senescence. Cells Tissues Organs, 1993, 147, 216-222.	1.3	12
176	Presenilin polymorphisms in Alzheimer's disease. Lancet, The, 1997, 350, 958-959.	6.3	12
177	Parent-to-parent support interventions for parents of babies cared for in a neonatal unit—protocol of a systematic review of qualitative and quantitative evidence. Systematic Reviews, 2018, 7, 179.	2.5	12
178	Brain transferrin receptors and the distribution of cytochrome oxidase. Biochemical Society Transactions, 1990, 18, 647-648.	1.6	11
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