## Raquel Sanchez-Valle

List of Publications by Year in descending order

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277	12,461	<sup>31976</sup> 53	<sup>37204</sup>
papers	citations	h-index	g-index
313	313	313	13279
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	New insights into the genetic etiology of Alzheimer's disease and related dementias. Nature Genetics, 2022, 54, 412-436.	21.4	700
2	Neurodegenerative disease status and post-mortem pathology in idiopathic rapid-eye-movement sleep behaviour disorder: an observational cohort study. Lancet Neurology, The, 2013, 12, 443-453.	10.2	602
3	TREM2 mutations implicated in neurodegeneration impair cell surface transport and phagocytosis. Science Translational Medicine, 2014, 6, 243ra86.	12.4	600
4	Neurodegenerative Disorder Risk in Idiopathic REM Sleep Behavior Disorder: Study in 174 Patients. PLoS ONE, 2014, 9, e89741.	2.5	407
5	<scp>sTREM</scp> 2 cerebrospinal fluid levels are a potential biomarker for microglia activity in earlyâ€stage Alzheimer's disease and associate with neuronal injury markers. EMBO Molecular Medicine, 2016, 8, 466-476.	6.9	392
6	Anti-Tr antibodies as markers of paraneoplastic cerebellar degeneration and Hodgkin's disease. Neurology, 2003, 60, 230-234.	1.1	297
7	A Panâ€ <scp>E</scp> uropean Study of the <i>C9orf72</i> Repeat Associated with <scp>FTLD</scp> : Geographic Prevalence, Genomic Instability, and Intermediate Repeats. Human Mutation, 2013, 34, 363-373.	2.5	247
8	Neurofilament light chain: a biomarker for genetic frontotemporal dementia. Annals of Clinical and Translational Neurology, 2016, 3, 623-636.	3.7	207
9	Genetic screening of Alzheimer's disease genes in Iberian and African samples yields novel mutations in presenilins and APP. Neurobiology of Aging, 2010, 31, 725-731.	3.1	196
10	A trial of gantenerumab or solanezumab in dominantly inherited Alzheimer's disease. Nature Medicine, 2021, 27, 1187-1196.	30.7	182
11	Age at symptom onset and death and disease duration in genetic frontotemporal dementia: an international retrospective cohort study. Lancet Neurology, The, 2020, 19, 145-156.	10.2	175
12	Detection of α-synuclein in CSF by RT-QuIC in patients with isolated rapid-eye-movement sleep behaviour disorder: a longitudinal observational study. Lancet Neurology, The, 2021, 20, 203-212.	10.2	174
13	Low cerebrospinal fluid concentration of mitochondrial DNA in preclinical Alzheimer disease. Annals of Neurology, 2013, 74, 655-668.	5.3	171
14	Dopamine transporter imaging deficit predicts early transition to synucleinopathy in idiopathic rapid eye movement sleep behavior disorder. Annals of Neurology, 2017, 82, 419-428.	5.3	161
15	YKL-40 in the brain and cerebrospinal fluid of neurodegenerative dementias. Molecular Neurodegeneration, 2017, 12, 83.	10.8	140
16	Common variants in Alzheimer's disease and risk stratification by polygenic risk scores. Nature Communications, 2021, 12, 3417.	12.8	140
17	Increased Cortical Thickness and Caudate Volume Precede Atrophy in PSEN1 Mutation Carriers. Journal of Alzheimer's Disease, 2010, 22, 909-922.	2.6	136
18	Cognitive reserve modulates task-induced activations and deactivations in healthy elders, amnestic mild cognitive impairment and mild Alzheimer's disease. Cortex, 2010, 46, 451-461.	2.4	136

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19	Cerebrospinal fluid biomarker supported diagnosis of Creutzfeldt–Jakob disease and rapid dementias: a longitudinal multicentre study over 10 years. Brain, 2012, 135, 3051-3061.	7.6	135
20	Prevalence of amyloidâ€Î² pathology in distinct variants of primary progressive aphasia. Annals of Neurology, 2018, 84, 729-740.	5.3	132
21	Serum neurofilament light chain in genetic frontotemporal dementia: a longitudinal, multicentre cohort study. Lancet Neurology, The, 2019, 18, 1103-1111.	10.2	128
22	Screening for amnestic mild cognitive impairment and early Alzheimer's disease with M@T (Memory) Tj ETQq0 C 294-304.	) 0 rgBT /0 2.7	verlock 10 Tf 124
23	Clinical features and <i>APOE</i> genotype of pathologically proven early-onset Alzheimer disease. Neurology, 2011, 76, 1720-1725.	1.1	123
24	TMEM106B is a genetic modifier of frontotemporal lobar degeneration with C9orf72 hexanucleotide repeat expansions. Acta Neuropathologica, 2014, 127, 407-418.	7.7	123
25	Plasma miR-34a-5p and miR-545-3p as Early Biomarkers of Alzheimer's Disease: Potential and Limitations. Molecular Neurobiology, 2017, 54, 5550-5562.	4.0	119
26	Changes in Synaptic Proteins Precede Neurodegeneration Markers in Preclinical Alzheimer's Disease Cerebrospinal Fluid. Molecular and Cellular Proteomics, 2019, 18, 546-560.	3.8	115
27	LifeTime and improving European healthcare through cell-based interceptive medicine. Nature, 2020, 587, 377-386.	27.8	108
28	Plasma glial fibrillary acidic protein is raised in progranulin-associated frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 263-270.	1.9	106
29	Distinct patterns of APP processing in the CNS in autosomal-dominant and sporadic Alzheimer disease. Acta Neuropathologica, 2013, 125, 201-213.	7.7	103
30	Cerebrospinal Fluid Level of YKL-40 Protein in Preclinical and Prodromal Alzheimer's Disease. Journal of Alzheimer's Disease, 2014, 42, 901-908.	2.6	102
31	Active Aβ immunotherapy CAD106 in Alzheimer's disease: A phase 2b study. Alzheimer's and Dementia: Translational Research and Clinical Interventions, 2017, 3, 10-22.	3.7	102
32	CSF sAPPβ, YKL-40, and neurofilament light in frontotemporal lobar degeneration. Neurology, 2017, 89, 178-188.	1.1	100
33	Characterization of the repeat expansion size in C9orf72 in amyotrophic lateral sclerosis and frontotemporal dementia. Human Molecular Genetics, 2014, 23, 749-754.	2.9	98
34	Potential genetic modifiers of disease risk and age at onset in patients with frontotemporal lobar degeneration and GRN mutations: a genome-wide association study. Lancet Neurology, The, 2018, 17, 548-558.	10.2	97
35	Cognitively Preserved Subjects with Transitional Cerebrospinal Fluid ß-Amyloid 1-42 Values Have Thicker Cortex in Alzheimer's Disease Vulnerable Areas. Biological Psychiatry, 2011, 70, 183-190. 	1.3	93
36	Rare mutations in SQSTM1 modify susceptibility to frontotemporal lobar degeneration. Acta Neuropathologica, 2014, 128, 397-410.	7.7	93

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37	Assessing the role of the TREM2 p.R47H variant as a risk factor for Alzheimer's disease and frontotemporal dementia. Neurobiology of Aging, 2014, 35, 444.e1-444.e4.	3.1	92
38	A preliminary study of the whole-genome expression profile of sporadic and monogenic early-onset Alzheimer's disease. Neurobiology of Aging, 2013, 34, 1772-1778.	3.1	87
39	<i>TBK1</i> Mutation Spectrum in an Extended European Patient Cohort with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. Human Mutation, 2017, 38, 297-309.	2.5	87
40	Cerebrospinal fluid sTREM2 levels are associated with gray matter volume increases and reduced diffusivity in early Alzheimer's disease. Alzheimer's and Dementia, 2016, 12, 1259-1272.	0.8	86
41	Comparison of Pittsburgh compound B and florbetapir in crossâ€sectional and longitudinal studies. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2019, 11, 180-190.	2.4	84
42	A comprehensive study of the genetic impact of rare variants in SORL1 in European early-onset Alzheimer's disease. Acta Neuropathologica, 2016, 132, 213-224.	7.7	83
43	Interactions of cognitive reserve with regional brain anatomy and brain function during a working memory task in healthy elders. Biological Psychology, 2009, 80, 256-259.	2.2	81
44	Validation of 14-3-3 Protein as a Marker in Sporadic Creutzfeldt-Jakob Disease Diagnostic. Molecular Neurobiology, 2016, 53, 2189-2199.	4.0	80
45	Cerebrospinal fluid biomarkers in human genetic transmissible spongiform encephalopathies. Journal of Neurology, 2009, 256, 1620-1628.	3.6	77
46	Influence of timing on CSF tests value for Creutzfeldt-Jakob disease diagnosis. Journal of Neurology, 2007, 254, 901-906.	3.6	72
47	14-3-3 protein in the CSF as prognostic marker in early multiple sclerosis. Neurology, 2001, 57, 722-724.	1.1	68
48	Cerebrospinal Fluid Biomarkers and Memory Present Distinct Associations along the Continuum from Healthy Subjects to AD Patients. Journal of Alzheimer's Disease, 2011, 23, 319-326.	2.6	66
49	CSF analysis in patients with sporadic CJD and other transmissible spongiform encephalopathies. European Journal of Neurology, 2007, 14, 121-124.	3.3	61
50	Cerebrospinal fluid neurofilament light levels in neurodegenerative dementia: Evaluation of diagnostic accuracy in the differential diagnosis of prion diseases. Alzheimer's and Dementia, 2018, 14, 751-763.	0.8	61
51	Effects of <scp>lgLON5</scp> Antibodies on Neuronal Cytoskeleton: A Link between Autoimmunity and Neurodegeneration. Annals of Neurology, 2020, 88, 1023-1027.	5.3	61
52	Distinct Functional Activity of the Precuneus and Posterior Cingulate Cortex During Encoding in the Preclinical Stage of Alzheimer's Disease. Journal of Alzheimer's Disease, 2012, 31, 517-526.	2.6	59
53	Determination of Neuronal Antibodies in Suspected and Definite Creutzfeldt-Jakob Disease. JAMA Neurology, 2014, 71, 74.	9.0	59
54	CSF Biomarkers in COVID-19 Associated Encephalopathy and Encephalitis Predict Long-Term Outcome. Frontiers in Immunology, 2022, 13, 866153.	4.8	57

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55	Analysis of the <i>CHCHD10</i> gene in patients with frontotemporal dementia and amyotrophic lateral sclerosis from Spain. Brain, 2015, 138, e400-e400.	7.6	56
56	Plasma phosphorylated TDP-43 levels are elevated in patients with frontotemporal dementia carrying a C9orf72 repeat expansion or a GRN mutation. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 684-691.	1.9	55
57	Neuronal pentraxin 2: a synapse-derived CSF biomarker in genetic frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 612-621.	1.9	55
58	CSF YKL-40 and pTau181 are related to different cerebral morphometric patterns in early AD. Neurobiology of Aging, 2016, 38, 47-55.	3.1	54
59	Serum neurofilament light levels correlate with severity measures and neurodegeneration markers in autosomal dominant Alzheimer's disease. Alzheimer's Research and Therapy, 2018, 10, 113.	6.2	54
60	Neuroimaging and Biochemical Markers in the Three Variants of Primary Progressive Aphasia. Dementia and Geriatric Cognitive Disorders, 2013, 35, 106-117.	1.5	53
61	Deleterious ABCA7 mutations and transcript rescue mechanisms in early onset Alzheimer's disease. Acta Neuropathologica, 2017, 134, 475-487.	7.7	53
62	Clinical, Neuropathologic, and Biochemical Profile of the Amyloid Precursor Protein I716F Mutation. Journal of Neuropathology and Experimental Neurology, 2010, 69, 53-59.	1.7	52
63	Presentations and mechanisms of CNS disorders related to COVID-19. Neurology: Neuroimmunology and NeuroInflammation, 2021, 8, .	6.0	52
64	Plasma Neurofilament Light for Prediction of Disease Progression in Familial Frontotemporal Lobar Degeneration. Neurology, 2021, 96, e2296-e2312.	1.1	52
65	Different profiles of Alzheimer's disease cerebrospinal fluid biomarkers in controls and subjects with subjective memory complaints. Journal of Neural Transmission, 2011, 118, 259-262.	2.8	49
66	Poly(GP), neurofilament and grey matter deficits in <i>C9orf72</i> expansion carriers. Annals of Clinical and Translational Neurology, 2018, 5, 583-597.	3.7	48
67	<scp><i>TARDBP</i></scp> mutation p. <scp>I</scp> le383 <scp>V</scp> al associated with semantic dementia and complex proteinopathy. Neuropathology and Applied Neurobiology, 2014, 40, 225-230.	3.2	48
68	Cerebral amyloid angiopathy in Down syndrome and sporadic and autosomalâ€dominant Alzheimer's disease. Alzheimer's and Dementia, 2017, 13, 1251-1260.	0.8	47
69	Synaptic, axonal damage and inflammatory cerebrospinal fluid biomarkers in neurodegenerative dementias. Alzheimer's and Dementia, 2020, 16, 262-272.	0.8	47
70	CSF microRNA Profiling in Alzheimer's Disease: a Screening and Validation Study. Molecular Neurobiology, 2017, 54, 6647-6654.	4.0	45
71	Cortical microstructure in the behavioural variant of frontotemporal dementia: looking beyond atrophy. Brain, 2019, 142, 1121-1133.	7.6	45
72	Neuronal intranuclear (hyaline) inclusion disease and fragile X-associated tremor/ataxia syndrome: a morphological and molecular dilemma. Brain, 2017, 140, e51-e51.	7.6	43

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73	Longitudinal Study of Amnesic Patients at High Risk for Alzheimer's Disease: Clinical, Neuropsychological and Magnetic Resonance Spectroscopy Features. Dementia and Geriatric Cognitive Disorders, 2007, 24, 402-410.	1.5	42
74	Novel <scp>CSF</scp> biomarkers in genetic frontotemporal dementia identified by proteomics. Annals of Clinical and Translational Neurology, 2019, 6, 698-707.	3.7	42
75	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. JAMA Network Open, 2021, 4, e2030194.	5.9	42
76	Cortical Brain Metabolism as Measured by Proton Spectroscopy Is Related to Memory Performance in Patients with Amnestic Mild Cognitive Impairment and Alzheimer's Disease. Dementia and Geriatric Cognitive Disorders, 2007, 24, 274-279.	1.5	41
77	Clinicopathological and genetic correlates of frontotemporal lobar degeneration and corticobasal degeneration. Journal of Neurology, 2008, 255, 488-494.	3.6	40
78	Rapidly Progressive Dementia. Alzheimer Disease and Associated Disorders, 2012, 26, 267-271.	1.3	40
79	Cognitive Reserve Proxies Relate to Gray Matter Loss in Cognitively Healthy Elderly with Abnormal Cerebrospinal Fluid Amyloid-β Levels. Journal of Alzheimer's Disease, 2013, 35, 715-726.	2.6	40
80	Donepezil Treatment Stabilizes Functional Connectivity During Resting State and Brain Activity During Memory Encoding in Alzheimer's Disease. Journal of Clinical Psychopharmacology, 2013, 33, 199-205.	1.4	40
81	The memory alteration test (M@T) discriminates between subjective memory complaints, mild cognitive impairment and Alzheimer's disease. Archives of Gerontology and Geriatrics, 2010, 50, 171-174.	3.0	38
82	Applying the new research diagnostic criteria: MRI findings and neuropsychological correlations of prodromal AD. International Journal of Geriatric Psychiatry, 2012, 27, 127-134.	2.7	38
83	Phosphorylated tau in cerebrospinal fluid as a marker for Creutzfeldt-Jakob disease. Journal of Neurology, Neurosurgery and Psychiatry, 2002, 73, 79-81.	1.9	37
84	Challenges associated with biomarkerâ€based classification systems for Alzheimer's disease. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2018, 10, 346-357.	2.4	37
85	Cerebrospinal Fluid Biomarkers Predict Clinical Evolution in Patients with Subjective Cognitive Decline and Mild Cognitive Impairment. Neurodegenerative Diseases, 2016, 16, 69-76.	1.4	36
86	The <i>APOE</i> ε4 genotype modulates CSF YKLâ€40 levels and their structural brain correlates in the continuum of Alzheimer's disease but not those of sTREM2. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2017, 6, 50-59.	2.4	36
87	Brain functional network integrity sustains cognitive function despite atrophy in presymptomatic genetic frontotemporal dementia. Alzheimer's and Dementia, 2021, 17, 500-514.	0.8	36
88	Clinicopathological Correlations and Concomitant Pathologies in Rapidly Progressive Dementia: A Brain Bank Series. Neurodegenerative Diseases, 2015, 15, 350-360.	1.4	35
89	Role for ATXN1, ATXN2, and HTT intermediate repeats in frontotemporal dementia and Alzheimer's disease. Neurobiology of Aging, 2020, 87, 139.e1-139.e7.	3.1	35
90	Investigation of the role of rare TREM2 variants in frontotemporal dementia subtypes. Neurobiology of Aging, 2014, 35, 2657.e13-2657.e19.	3.1	34

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91	Genetic variability in SQSTM1 and risk of early-onset Alzheimer dementia: a European early-onset dementia consortium study. Neurobiology of Aging, 2015, 36, 2005.e15-2005.e22.	3.1	34
92	Validation of α-Synuclein as a CSF Biomarker for Sporadic Creutzfeldt-Jakob Disease. Molecular Neurobiology, 2018, 55, 2249-2257.	4.0	34
93	Sex differences in the behavioral variant of frontotemporal dementia: A new window to executive and behavioral reserve. Alzheimer's and Dementia, 2021, 17, 1329-1341.	0.8	34
94	The inner fluctuations of the brain in presymptomatic Frontotemporal Dementia: The chronnectome fingerprint. NeuroImage, 2019, 189, 645-654.	4.2	33
95	Usefulness of Biomarkers in the Diagnosis and Prognosis of Early-Onset Cognitive Impairment. Journal of Alzheimer's Disease, 2014, 40, 919-927.	2.6	32
96	Serum Progranulin Levels in Patients with Frontotemporal Lobar Degeneration and Alzheimer's Disease: Detection of GRN Mutations in a Spanish Cohort. Journal of Alzheimer's Disease, 2012, 31, 581-591.	2.6	31
97	Frontotemporal Dementia Caused by the P301L Mutation in <b> </b> the <b><i> MAPT</i></b> Gene: Clinicopathological Features of 13 Cases from the Same Geographical Origin in Barcelona, Spain. Dementia and Geriatric Cognitive Disorders, 2017, 44, 213-221.	1.5	31
98	Apathy in presymptomatic genetic frontotemporal dementia predicts cognitive decline and is driven by structural brain changes. Alzheimer's and Dementia, 2021, 17, 969-983.	0.8	31
99	Association of Rare <i>APOE</i> Missense Variants V236E and R251G With Risk of Alzheimer Disease. JAMA Neurology, 2022, 79, 652.	9.0	31
100	14-3-3 protein isoforms and atypical patterns of the 14-3-3 assay in the diagnosis of Creutzfeldt–Jakob disease. Neuroscience Letters, 2002, 320, 69-72.	2.1	30
101	Neuropsychological profile of prodromal Alzheimer's disease (Prd-AD) and their radiological correlates. Archives of Gerontology and Geriatrics, 2011, 52, 190-196.	3.0	30
102	Evolving brain structural changes in PSEN1 mutation carriers. Neurobiology of Aging, 2015, 36, 1261-1270.	3.1	30
103	Clinical and video-polysomnographic analysis of rapid eye movement sleep behavior disorder and other sleep disturbances in dementia with Lewy bodies. Sleep, 2019, 42, .	1.1	30
104	Network structure and transcriptomic vulnerability shape atrophy in frontotemporal dementia. Brain, 2023, 146, 321-336.	7.6	30
105	Clinical and genetic features of human prion diseases in Catalonia: 1993-2002. European Journal of Neurology, 2004, 11, 649-655.	3.3	29
106	Hippocampal atrophy has limited usefulness as a diagnostic biomarker on the early onset Alzheimer's disease patients: A comparison between visual and quantitative assessment. NeuroImage: Clinical, 2019, 23, 101927.	2.7	29
107	Mechanisms of functional compensation, delineated by eigenvector centrality mapping, across the pathophysiological continuum of Alzheimer's disease. NeuroImage: Clinical, 2019, 22, 101777.	2.7	29
108	Multicenter Alzheimer's and Parkinson's disease immune biomarker verification study. Alzheimer's and Dementia, 2020, 16, 292-304.	0.8	29

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109	Telemedicine assessment of long-term cognitive and functional status in anti-leucine-rich, glioma-inactivated 1 encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	29
110	Characterizing the Clinical Features and Atrophy Patterns of <i>MAPT</i> -Related Frontotemporal Dementia With Disease Progression Modeling. Neurology, 2021, 97, e941-e952.	1.1	29
111	CSF 14-3-3 protein assay and MRI as prognostic markers in patients with a clinically isolated syndrome suggestive of MS. Journal of Neurology, 2004, 251, 1278-1279.	3.6	28
112	PICOGEN: experiencia de 5 años de un programa de asesoramiento genético en demencia. NeurologÃa, 2011, 26, 143-149.	0.7	28
113	C-terminal fragments of the amyloid precursor protein in cerebrospinal fluid as potential biomarkers for Alzheimer disease. Scientific Reports, 2017, 7, 2477.	3.3	28
114	Digital biomarkerâ€based individualized prognosis for people at risk of dementia. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2020, 12, e12073.	2.4	28
115	Differential early subcortical involvement in genetic FTD within the GENFI cohort. NeuroImage: Clinical, 2021, 30, 102646.	2.7	28
116	Longitudinal brain atrophy and CSF biomarkers in early-onset Alzheimer's disease. NeuroImage: Clinical, 2021, 32, 102804.	2.7	28
117	White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal GENFI study. NeuroImage: Clinical, 2019, 24, 102077.	2.7	27
118	Biphasic cortical macro―and microstructural changes in autosomal dominant Alzheimer's disease. Alzheimer's and Dementia, 2021, 17, 618-628.	0.8	27
119	Modelling the cascade of biomarker changes in <i>GRN</i> -related frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 494-501.	1.9	27
120	A data-driven disease progression model of fluid biomarkers in genetic frontotemporal dementia. Brain, 2022, 145, 1805-1817.	7.6	27
121	Rare Variants in Calcium Homeostasis Modulator 1 (CALHM1) Found in Early Onset Alzheimer's Disease Patients Alter Calcium Homeostasis. PLoS ONE, 2013, 8, e74203.	2.5	26
122	Diagnostic accuracy of behavioral variant frontotemporal dementia consortium criteria (FTDC) in a clinicopathological cohort. Neuropathology and Applied Neurobiology, 2015, 41, 882-892.	3.2	26
123	CSF glial biomarkers YKL40 and sTREM2 are associated with longitudinal volume and diffusivity changes in cognitively unimpaired individuals. NeuroImage: Clinical, 2019, 23, 101801.	2.7	26
124	Social cognition impairment in genetic frontotemporal dementia within the GENFI cohort. Cortex, 2020, 133, 384-398.	2.4	26
125	From Progressive Nonfluent Aphasia to Corticobasal Syndrome: A Case Report of Corticobasal Degeneration. Neurocase, 2006, 12, 355-359.	0.6	25
126	Structural Connectivity Alterations Along the Alzheimer's Disease Continuum: Reproducibility Across Two Independent Samples and Correlation with Cerebrospinal Fluid Amyloid-β and Tau. Journal of Alzheimer's Disease, 2018, 61, 1575-1587.	2.6	25

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127	Early symptoms in symptomatic and preclinical genetic frontotemporal lobar degeneration. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 975-984.	1.9	25
128	Cerebrospinal Fluid Biomarkers in Alzheimer's Disease Families with <i>PSEN1</i> Mutations. Neurodegenerative Diseases, 2011, 8, 202-207.	1.4	24
129	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. Alzheimer's and Dementia, 2022, 18, 1408-1423.	0.8	24
130	A novel mutation in the <i>PSEN1</i> gene (L286P) associated with familial earlyâ€onset dementia of Alzheimer type and lobar haematomas. European Journal of Neurology, 2007, 14, 1409-1412.	3.3	23
131	Voxel based morphometry features and followâ€up of amnestic patients at high risk for Alzheimer's disease conversion. International Journal of Geriatric Psychiatry, 2009, 24, 875-884.	2.7	23
132	Rare Variants in <i>PLD3</i> Do Not Affect Risk for Early-Onset Alzheimer Disease in a European Consortium Cohort. Human Mutation, 2015, 36, 1226-1235.	2.5	23
133	Education modulates brain maintenance in presymptomatic frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 1124-1130.	1.9	23
134	Contribution of CSF biomarkers to earlyâ€onset Alzheimer's disease and frontotemporal dementia neuroimaging signatures. Human Brain Mapping, 2020, 41, 2004-2013.	3.6	22
135	A novel PSEN1 mutation (K239N) associated with Alzheimer's disease with wide range age of onset and slow progression. European Journal of Neurology, 2010, 17, 994-996.	3.3	21
136	Interrupted CAG expansions in ATXN2 gene expand the genetic spectrum of frontotemporal dementias. Acta Neuropathologica Communications, 2018, 6, 41.	5.2	21
137	The hippocampal longitudinal axis—relevance for underlying tau and TDP-43 pathology. Neurobiology of Aging, 2018, 70, 1-9.	3.1	21
138	Neuroanatomical and cognitive correlates of visual hallucinations in Parkinson's disease and dementia with Lewy bodies: Voxel-based morphometry and neuropsychological meta-analysis. Neuroscience and Biobehavioral Reviews, 2021, 128, 367-382.	6.1	21
139	Stratifying the Presymptomatic Phase of Genetic Frontotemporal Dementia by Serum <scp>NfL</scp> and <scp>pNfH</scp> : A Longitudinal Multicentre Study. Annals of Neurology, 2022, 91, 33-47.	5.3	21
140	Cerebrospinal Fluid Total Prion Protein in the Spectrum of Prion Diseases. Molecular Neurobiology, 2019, 56, 2811-2821.	4.0	20
141	Clinical applicability of diagnostic biomarkers in earlyâ€onset cognitive impairment. European Journal of Neurology, 2019, 26, 1098-1104.	3.3	20
142	Plasma levels of soluble TREM2 and neurofilament light chain in TREM2 rare variant carriers. Alzheimer's Research and Therapy, 2019, 11, 94.	6.2	20
143	Analysis of brain atrophy and local gene expression in genetic frontotemporal dementia. Brain Communications, 2020, 2, .	3.3	20
144	Association between cerebrospinal fluid tau and brain atrophy is not related to clinical severity in the Alzheimer's disease continuum. Psychiatry Research - Neuroimaging, 2011, 192, 140-146.	1.8	19

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145	Rare nonsynonymous variants in SORT1 are associated with increased risk for frontotemporal dementia. Neurobiology of Aging, 2018, 66, 181.e3-181.e10.	3.1	19
146	Clinical value of cerebrospinal fluid neurofilament light chain in semantic dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 997-1004.	1.9	19
147	Faster Cortical Thinning and Surface Area Loss in Presymptomatic and Symptomatic <i>C9orf72</i> Repeat Expansion Adult Carriers. Annals of Neurology, 2020, 88, 113-122.	5.3	19
148	Multiâ€cohort profiling reveals elevated CSF levels of brainâ€enriched proteins in Alzheimer's disease. Annals of Clinical and Translational Neurology, 2021, 8, 1456-1470.	3.7	19
149	Evolving Brain Functional Abnormalities in PSEN1 Mutation Carriers: A Resting and Visual Encoding fMRI Study. Journal of Alzheimer's Disease, 2013, 36, 165-175.	2.6	19
150	Distinctive age-related temporal cortical thinning in asymptomatic granulin gene mutation carriers. Neurobiology of Aging, 2013, 34, 1462-1468.	3.1	18
151	Alpha-synuclein Aggregates in Labial Salivary Glands of Idiopathic Rapid Eye Movement Sleep Behavior Disorder. Sleep, 2018, 41, .	1.1	18
152	Systematic Screening of Ubiquitin/p62 Aggregates in Cerebellar Cortex Expands the Neuropathological Phenotype of the C9orf72 Expansion Mutation. Journal of Neuropathology and Experimental Neurology, 2018, 77, 703-709.	1.7	18
153	A modified Camel and Cactus Test detects presymptomatic semantic impairment in genetic frontotemporal dementia within the GENFI cohort. Applied Neuropsychology Adult, 2022, 29, 112-119.	1.2	18
154	TREM2 expression in the brain and biological fluids in prion diseases. Acta Neuropathologica, 2021, 141, 841-859.	7.7	18
155	The amyloid-β isoform pattern in cerebrospinal fluid in familial PSEN1 M139T- and L286P-associated Alzheimer's disease. Molecular Medicine Reports, 2012, 5, 1111-1115.	2.4	17
156	"Preclinical―MSA in definite Creutzfeldtâ€Jakob disease. Neuropathology, 2012, 32, 158-163.	1.2	17
157	Quantitative Magnetic Resonance Abnormalities in Creutzfeldt-Jakob Disease and Fatal Insomnia. Journal of Alzheimer's Disease, 2016, 55, 431-443.	2.6	17
158	White Matter Abnormalities Track Disease Progression in PSEN1 Autosomal Dominant Alzheimer's Disease. Journal of Alzheimer's Disease, 2016, 51, 827-835.	2.6	17
159	A novel <i>MAPT</i> mutation (P301T) associated with familial frontotemporal dementia. European Journal of Neurology, 2007, 14, e9-10.	3.3	16
160	Molecular evidence of founder effects of fatal familial insomnia through SNP haplotypes around the D178N mutation. Neurogenetics, 2008, 9, 109-118.	1.4	16
161	Common and rare TBK1 variants in early-onset Alzheimer disease in a European cohort. Neurobiology of Aging, 2018, 62, 245.e1-245.e7.	3.1	16
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