Raquel Sanchez-Valle

List of Publications by Citations

269 papers	7,819 citations	47 h-index	79 g-index
313 ext. papers	10,208 ext. citations	6.2 avg, IF	5.44 L-index

#	Paper	IF	Citations
269	Neurodegenerative disease status and post-mortem pathology in idiopathic rapid-eye-movement sleep behaviour disorder: an observational cohort study. *Lancet Neurology, The*, **2013**, 12, 443-53	24.1	472
268	TREM2 mutations implicated in neurodegeneration impair cell surface transport and phagocytosis. *Science Translational Medicine*, **2014**, 6, 243ra86	17.5	436
267	Neurodegenerative disorder risk in idiopathic REM sleep behavior disorder: study in 174 patients. *PLoS ONE*, **2014**, 9, e89741	3.7	308
266	sTREM2 cerebrospinal fluid levels are a potential biomarker for microglia activity in early-stage Alzheimer's disease and associate with neuronal injury markers. *EMBO Molecular Medicine*, **2016**, 8, 466-76	12	256
265	Anti-Tr antibodies as markers of paraneoplastic cerebellar degeneration and Hodgkin's disease. *Neurology*, **2003**, 60, 230-4	6.5	246
264	A pan-European study of the C9orf72 repeat associated with FTLD: geographic prevalence, genomic instability, and intermediate repeats. *Human Mutation*, **2013**, 34, 363-73	4.7	208
263	Neurofilament light chain: a biomarker for genetic frontotemporal dementia. *Annals of Clinical and Translational Neurology*, **2016**, 3, 623-36	5.3	163
262	Genetic screening of Alzheimer's disease genes in Iberian and African samples yields novel mutations in presenilins and APP. *Neurobiology of Aging*, **2010**, 31, 725-31	5.6	162
261	Low cerebrospinal fluid concentration of mitochondrial DNA in preclinical Alzheimer disease. *Annals of Neurology*, **2013**, 74, 655-68	9.4	129
260	Dopamine transporter imaging deficit predicts early transition to synucleinopathy in idiopathic rapid eye movement sleep behavior disorder. *Annals of Neurology*, **2017**, 82, 419-428	9.4	108
259	Cognitive reserve modulates task-induced activations and deactivations in healthy elders, amnestic mild cognitive impairment and mild Alzheimer's disease. *Cortex*, **2010**, 46, 451-61	3.8	104
258	Increased cortical thickness and caudate volume precede atrophy in PSEN1 mutation carriers. *Journal of Alzheimer's Disease*, **2010**, 22, 909-22	4.3	103
257	Cerebrospinal fluid biomarker supported diagnosis of Creutzfeldt-Jakob disease and rapid dementias: a longitudinal multicentre study over 10 years. *Brain*, **2012**, 135, 3051-61	11.2	102
256	TMEM106B is a genetic modifier of frontotemporal lobar degeneration with C9orf72 hexanucleotide repeat expansions. *Acta Neuropathologica*, **2014**, 127, 407-18	14.3	97
255	Clinical features and APOE genotype of pathologically proven early-onset Alzheimer disease. *Neurology*, **2011**, 76, 1720-5	6.5	91
254	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	24.1	90
253	YKL-40 in the brain and cerebrospinal fluid of neurodegenerative dementias. *Molecular Neurodegeneration*, **2017**, 12, 83	19	86

252	Characterization of the repeat expansion size in C9orf72 in amyotrophic lateral sclerosis and frontotemporal dementia. *Human Molecular Genetics*, **2014**, 23, 749-54	5.6	84
251	Rare mutations in SQSTM1 modify susceptibility to frontotemporal lobar degeneration. *Acta Neuropathologica*, **2014**, 128, 397-410	14.3	83
250	Screening for amnestic mild cognitive impairment and early Alzheimer's disease with M@T (Memory Alteration Test) in the primary care population. *International Journal of Geriatric Psychiatry*, **2007**, 22, 294-304	3.9	82
249	Distinct patterns of APP processing in the CNS in autosomal-dominant and sporadic Alzheimer disease. *Acta Neuropathologica*, **2013**, 125, 201-13	14.3	81
248	Plasma miR-34a-5p and miR-545-3p as Early Biomarkers of Alzheimer's Disease: Potential and Limitations. *Molecular Neurobiology*, **2017**, 54, 5550-5562	6.2	81
247	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.e1-4	5.6	81
246	Cerebrospinal fluid level of YKL-40 protein in preclinical and prodromal Alzheimer's disease. *Journal of Alzheimers Disease*, **2014**, 42, 901-8	4.3	79
245	CSF sAPP, YKL-40, and neurofilament light in frontotemporal lobar degeneration. *Neurology*, **2017**, 89, 178-188	6.5	75
244	Cognitively preserved subjects with transitional cerebrospinal fluid amyloid 1-42 values have thicker cortex in Alzheimer's disease vulnerable areas. *Biological Psychiatry*, **2011**, 70, 183-90	7.9	75
243	Prevalence of amyloid- pathology in distinct variants of primary progressive aphasia. *Annals of Neurology*, **2018**, 84, 729-740	9.4	74
242	Serum neurofilament light chain in genetic frontotemporal dementia: a longitudinal, multicentre cohort study. *Lancet Neurology, The*, **2019**, 18, 1103-1111	24.1	68
241	Cerebrospinal fluid biomarkers in human genetic transmissible spongiform encephalopathies. *Journal of Neurology*, **2009**, 256, 1620-8	5.5	68
240	Cerebrospinal fluid sTREM2 levels are associated with gray matter volume increases and reduced diffusivity in early Alzheimer's disease. *Alzheimers and Dementia*, **2016**, 12, 1259-1272	1.2	67
239	TBK1 Mutation Spectrum in an Extended European Patient Cohort with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. *Human Mutation*, **2017**, 38, 297-309	4.7	66
238	Influence of timing on CSF tests value for Creutzfeldt-Jakob disease diagnosis. *Journal of Neurology*, **2007**, 254, 901-6	5.5	66
237	Changes in Synaptic Proteins Precede Neurodegeneration Markers in Preclinical Alzheimer's Disease Cerebrospinal Fluid. *Molecular and Cellular Proteomics*, **2019**, 18, 546-560	7.6	66
236	Validation of 14-3-3 Protein as a Marker in Sporadic Creutzfeldt-Jakob Disease Diagnostic. *Molecular Neurobiology*, **2016**, 53, 2189-99	6.2	65
235	Active Immunotherapy CAD106 in Alzheimer's disease: A phase 2b study. *Alzheimers and Dementia: Translational Research and Clinical Interventions*, **2017**, 3, 10-22	6	65

234	Interactions of cognitive reserve with regional brain anatomy and brain function during a working memory task in healthy elders. *Biological Psychology*, **2009**, 80, 256-9	3.2	65
233	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	14.3	62
232	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	24.1	60
231	A preliminary study of the whole-genome expression profile of sporadic and monogenic early-onset Alzheimer's disease. *Neurobiology of Aging*, **2013**, 34, 1772-8	5.6	59
230	14-3-3 protein in the CSF as prognostic marker in early multiple sclerosis. *Neurology*, **2001**, 57, 722-4	6.5	59
229	Cerebrospinal fluid biomarkers and memory present distinct associations along the continuum from healthy subjects to AD patients. *Journal of Alzheimer's Disease*, **2011**, 23, 319-26	4.3	56
228	LifeTime and improving European healthcare through cell-based interceptive medicine. *Nature*, **2020**, 587, 377-386	50.4	56
227	CSF analysis in patients with sporadic CJD and other transmissible spongiform encephalopathies. *European Journal of Neurology*, **2007**, 14, 121-4	6	53
226	Detection of □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	24.1	52
225	A trial of gantenerumab or solanezumab in dominantly inherited Alzheimer's disease. *Nature Medicine*, **2021**, 27, 1187-1196	50.5	51
224	Analysis of the CHCHD10 gene in patients with frontotemporal dementia and amyotrophic lateral sclerosis from Spain. *Brain*, **2015**, 138, e400	11.2	47
223	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, 684-91	5.5	47
222	Comparison of Pittsburgh compound B and florbetapir in cross-sectional and longitudinal studies. *Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring*, **2019**, 11, 180-190	5.2	46
221	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	1.2	45
220	Determination of neuronal antibodies in suspected and definite Creutzfeldt-Jakob disease. *JAMA Neurology*, **2014**, 71, 74-8	17.2	43
219	Neuroimaging and biochemical markers in the three variants of primary progressive aphasia. *Dementia and Geriatric Cognitive Disorders*, **2013**, 35, 106-17	2.6	43
218	CSF YKL-40 and pTau181 are related to different cerebral morphometric patterns in early AD. *Neurobiology of Aging*, **2016**, 38, 47-55	5.6	42
217	Different profiles of Alzheimer's disease cerebrospinal fluid biomarkers in controls and subjects with subjective memory complaints. *Journal of Neural Transmission*, **2011**, 118, 259-62	4.3	41

216	Plasma glial fibrillary acidic protein is raised in progranulin-associated frontotemporal dementia. *Journal of Neurology, Neurosurgery and Psychiatry*, **2020**, 91, 263-270	5.5	40
215	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, 517-26	4.3	39
214	Cerebral amyloid angiopathy in Down syndrome and sporadic and autosomal-dominant Alzheimer's disease. *Alzheimer's and Dementia*, **2017**, 13, 1251-1260	1.2	38
213	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	9	38
212	Clinical, neuropathologic, and biochemical profile of the amyloid precursor protein I716F mutation. *Journal of Neuropathology and Experimental Neurology*, **2010**, 69, 53-9	3.1	37
211	Clinicopathological and genetic correlates of frontotemporal lobar degeneration and corticobasal degeneration. *Journal of Neurology*, **2008**, 255, 488-94	5.5	36
210	TARDBP mutation p.Ile383Val associated with semantic dementia and complex proteinopathy. *Neuropathology and Applied Neurobiology*, **2014**, 40, 225-30	5.2	36
209	Rapidly progressive dementia: experience in a tertiary care medical center. *Alzheimer Disease and Associated Disorders*, **2012**, 26, 267-71	2.5	35
208	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, 402-10	2.6	35
207	Deleterious ABCA7 mutations and transcript rescue mechanisms in early onset Alzheimer's disease. *Acta Neuropathologica*, **2017**, 134, 475-487	14.3	34
206	Cerebrospinal Fluid Biomarkers Predict Clinical Evolution in Patients with Subjective Cognitive Decline and Mild Cognitive Impairment. *Neurodegenerative Diseases*, **2016**, 16, 69-76	2.3	34
205	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, 274-9	2.6	34
204	Applying the new research diagnostic criteria: MRI findings and neuropsychological correlations of prodromal AD. *International Journal of Geriatric Psychiatry*, **2012**, 27, 127-34	3.9	33
203	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, 715-26	4.3	33
202	Cortical microstructure in the behavioural variant of frontotemporal dementia: looking beyond atrophy. *Brain*, **2019**, 142, 1121-1133	11.2	32
201	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	1.7	32
200	Investigation of the role of rare TREM2 variants in frontotemporal dementia subtypes. *Neurobiology of Aging*, **2014**, 35, 2657.e13-2657.e19	5.6	31
199	Validation of β-Synuclein as a CSF Biomarker for Sporadic Creutzfeldt-Jakob Disease. *Molecular Neurobiology*, **2018**, 55, 2249-2257	6.2	30

198	Phosphorylated tau in cerebrospinal fluid as a marker for Creutzfeldt-Jakob disease. *Journal of Neurology, Neurosurgery and Psychiatry*, **2002**, 73, 79-81	5.5	30
197	The ε4 genotype modulates CSF YKL-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	5.2	29
196	Genetic variability in SQSTM1 and risk of early-onset Alzheimer dementia: a European early-onset dementia consortium study. *Neurobiology of Aging*, **2015**, 36, 2005.e15-22	5.6	29
195	Poly(GP), neurofilament and grey matter deficits in expansion carriers. *Annals of Clinical and Translational Neurology*, **2018**, 5, 583-597	5.3	29
194	CSF microRNA Profiling in Alzheimer's Disease: a Screening and Validation Study. *Molecular Neurobiology*, **2017**, 54, 6647-6654	6.2	29
193	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, 581-91	4.3	29
192	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, 171-4	4	29
191	Synaptic, axonal damage and inflammatory cerebrospinal fluid biomarkers in neurodegenerative dementias. *Alzheimer's and Dementia*, **2020**, 16, 262-272	1.2	29
190	Presentations and mechanisms of CNS disorders related to COVID-19. *Neurology: Neuroimmunology and NeuroInflammation*, **2021**, 8,	9.1	28
189	Neuronal intranuclear (hyaline) inclusion disease and fragile X-associated tremor/ataxia syndrome: a morphological and molecular dilemma. *Brain*, **2017**, 140, e51	11.2	27
188	Clinicopathological Correlations and Concomitant Pathologies in Rapidly Progressive Dementia: A Brain Bank Series. *Neurodegenerative Diseases*, **2015**, 15, 350-60	2.3	27
187	Evolving brain structural changes in PSEN1 mutation carriers. *Neurobiology of Aging*, **2015**, 36, 1261-70	5.6	27
186	Usefulness of biomarkers in the diagnosis and prognosis of early-onset cognitive impairment. *Journal of Alzheimer's Disease*, **2014**, 40, 919-27	4.3	27
185	New insights into the genetic etiology of Alzheimer's disease and related dementias.. *Nature Genetics*, **2022**,	36.3	27
184	Novel CSF biomarkers in genetic frontotemporal dementia identified by proteomics. *Annals of Clinical and Translational Neurology*, **2019**, 6, 698-707	5.3	25
183	Challenges associated with biomarker-based classification systems for Alzheimer's disease. *Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring*, **2018**, 10, 346-357	5.2	25
182	Effects of IgLON5 Antibodies on Neuronal Cytoskeleton: A Link between Autoimmunity and Neurodegeneration. *Annals of Neurology*, **2020**, 88, 1023-1027	9.4	25
181	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, 1278-9	5.5	24

180	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	3.3	24
179	Frontotemporal Dementia Caused by the P301L Mutation in the MAPT Gene: Clinicopathological Features of 13 Cases from the Same Geographical Origin in Barcelona, Spain. *Dementia and Geriatric Cognitive Disorders*, **2017**, 44, 213-221	2.6	23
178	Cerebrospinal fluid biomarkers in Alzheimer's disease families with PSEN1 mutations. *Neurodegenerative Diseases*, **2011**, 8, 202-7	2.3	23
177	Common variants in Alzheimer's disease and risk stratification by polygenic risk scores. *Nature Communications*, **2021**, 12, 3417	17.4	23
176	Neuronal pentraxin 2: a synapse-derived CSF biomarker in genetic frontotemporal dementia. *Journal of Neurology, Neurosurgery and Psychiatry*, **2020**, 91, 612-621	5.5	22
175	Rare variants in calcium homeostasis modulator 1 (CALHM1) found in early onset Alzheimer's disease patients alter calcium homeostasis. *PLoS ONE*, **2013**, 8, e74203	3.7	22
174	From progressive nonfluent aphasia to corticobasal syndrome: a case report of corticobasal degeneration. *Neurocase*, **2006**, 12, 355-9	0.8	22
173	Clinical and genetic features of human prion diseases in Catalonia: 1993-2002. *European Journal of Neurology*, **2004**, 11, 649-55	6	22
172	C-terminal fragments of the amyloid precursor protein in cerebrospinal fluid as potential biomarkers for Alzheimer disease. *Scientific Reports*, **2017**, 7, 2477	4.9	21
171	Diagnostic accuracy of behavioral variant frontotemporal dementia consortium criteria (FTDC) in a clinicopathological cohort. *Neuropathology and Applied Neurobiology*, **2015**, 41, 882-92	5.2	21
170	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	4.3	20
169	Rare Variants in PLD3 Do Not Affect Risk for Early-Onset Alzheimer Disease in a European Consortium Cohort. *Human Mutation*, **2015**, 36, 1226-35	4.7	20
168	Neuropsychological profile of prodromal Alzheimer's disease (Prd-AD) and their radiological correlates. *Archives of Gerontology and Geriatrics*, **2011**, 52, 190-6	4	20
167	PICOGEN: five years experience with a genetic counselling program for dementia. *Neurologia*, **2011**, 26, 143-9	1.4	19
166	The inner fluctuations of the brain in presymptomatic Frontotemporal Dementia: The chronnectome fingerprint. *NeuroImage*, **2019**, 189, 645-654	7.9	18
165	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	5.3	18
164	Mechanisms of functional compensation, delineated by eigenvector centrality mapping, across the pathophysiological continuum of Alzheimer's disease. *NeuroImage: Clinical*, **2019**, 22, 101777	5.3	17
163	CSF glial biomarkers YKL40 and sTREM2 are associated with longitudinal volume and diffusivity changes in cognitively unimpaired individuals. *NeuroImage: Clinical*, **2019**, 23, 101801	5.3	17

162	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, 994-6	6	17
161	A novel mutation in the PSEN1 gene (L286P) associated with familial early-onset dementia of Alzheimer type and lobar haematomas. *European Journal of Neurology*, **2007**, 14, 1409-12	6	17
160	Cerebrospinal Fluid Total Prion Protein in the Spectrum of Prion Diseases. *Molecular Neurobiology*, **2019**, 56, 2811-2821	6.2	16
159	Multicenter Alzheimer's and Parkinson's disease immune biomarker verification study. *Alzheimer's and Dementia*, **2020**, 16, 292-304	1.2	16
158	The hippocampal longitudinal axis-relevance for underlying tau and TDP-43 pathology. *Neurobiology of Aging*, **2018**, 70, 1-9	5.6	15
157	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, 140-6	2.9	15
156	Voxel based morphometry features and follow-up of amnestic patients at high risk for Alzheimer's disease conversion. *International Journal of Geriatric Psychiatry*, **2009**, 24, 875-84	3.9	15
155	Evolving brain functional abnormalities in PSEN1 mutation carriers: a resting and visual encoding fMRI study. *Journal of Alzheimer's Disease*, **2013**, 36, 165-75	4.3	15
154	Early symptoms in symptomatic and preclinical genetic frontotemporal lobar degeneration. *Journal of Neurology, Neurosurgery and Psychiatry*, **2020**, 91, 975-984	5.5	15
153	Plasma levels of soluble TREM2 and neurofilament light chain in TREM2 rare variant carriers. *Alzheimer's Research and Therapy*, **2019**, 11, 94	9	15
152	"Preclinical" MSA in definite Creutzfeldt-Jakob disease. *Neuropathology*, **2012**, 32, 158-63	2	14
151	Distinctive age-related temporal cortical thinning in asymptomatic granulin gene mutation carriers. *Neurobiology of Aging*, **2013**, 34, 1462-8	5.6	14
150	The amyloid-isoform pattern in cerebrospinal fluid in familial PSEN1 M139T- and L286P-associated Alzheimer's disease. *Molecular Medicine Reports*, **2012**, 5, 1111-5	2.9	14
149	MAPT gene duplications are not a cause of frontotemporal lobar degeneration. *Neuroscience Letters*, **2007**, 424, 61-5	3.3	14
148	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. *JAMA Network Open*, **2021**, 4, e2030194	10.4	14
147	Clinical value of cerebrospinal fluid neurofilament light chain in semantic dementia. *Journal of Neurology, Neurosurgery and Psychiatry*, **2019**, 90, 997-1004	5.5	13
146	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	13
145	Telemedicine assessment of long-term cognitive and functional status in anti-leucine-rich, glioma-inactivated 1 encephalitis. *Neurology: Neuroimmunology and NeuroInflammation*, **2020**, 7,	9.1	13

144	A novel MAPT mutation (P301T) associated with familial frontotemporal dementia. *European Journal of Neurology*, **2007**, 14, e9-10	6	13
143	Late-onset frontotemporal dementia associated with a novel PGRN mutation. *Journal of Neural Transmission*, **2007**, 114, 1051-4	4.3	13
142	Role for ATXN1, ATXN2, and HTT intermediate repeats in frontotemporal dementia and Alzheimer's disease. *Neurobiology of Aging*, **2020**, 87, 139.e1-139.e7	5.6	13
141	Quantitative Magnetic Resonance Abnormalities in Creutzfeldt-Jakob Disease and Fatal Insomnia. *Journal of Alzheimers Disease*, **2017**, 55, 431-443	4.3	13
140	White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal GENFI study. *NeuroImage: Clinical*, **2019**, 24, 102077	5.3	13
139	Rare nonsynonymous variants in SORT1 are associated with increased risk for frontotemporal dementia. *Neurobiology of Aging*, **2018**, 66, 181.e3-181.e10	5.6	12
138	Identifying earlier Alzheimer's disease: insights from the preclinical and prodromal phases. *Neurodegenerative Diseases*, **2012**, 10, 158-60	2.3	12
137	Molecular evidence of founder effects of fatal familial insomnia through SNP haplotypes around the D178N mutation. *Neurogenetics*, **2008**, 9, 109-18	3	12
136	Plasma Neurofilament Light for Prediction of Disease Progression in Familial Frontotemporal Lobar Degeneration. *Neurology*, **2021**, 96, e2296-e2312	6.5	12
135	White Matter Abnormalities Track Disease Progression in PSEN1 Autosomal Dominant Alzheimer's Disease. *Journal of Alzheimers Disease*, **2016**, 51, 827-35	4.3	12
134	Common and rare TBK1 variants in early-onset Alzheimer disease in a European cohort. *Neurobiology of Aging*, **2018**, 62, 245.e1-245.e7	5.6	12
133	Synuclein aggregates in labial salivary glands of idiopathic rapid eye movement sleep behavior disorder. *Sleep*, **2018**, 41,	1.1	12
132	Clinical applicability of diagnostic biomarkers in early-onset cognitive impairment. *European Journal of Neurology*, **2019**, 26, 1098-1104	6	11
131	No supportive evidence for TIA1 gene mutations in a European cohort of ALS-FTD spectrum patients. *Neurobiology of Aging*, **2018**, 69, 293.e9-293.e11	5.6	11
130	A novel PSEN1 gene mutation (L235R) associated with familial early-onset Alzheimer's disease. *Neuroscience Letters*, **2011**, 496, 40-2	3.3	11
129	Faster Cortical Thinning and Surface Area Loss in Presymptomatic and Symptomatic C9orf72 Repeat Expansion Adult Carriers. *Annals of Neurology*, **2020**, 88, 113-122	9.4	11
128	Biphasic cortical macro- and microstructural changes in autosomal dominant Alzheimer's disease. *Alzheimers and Dementia*, **2021**, 17, 618-628	1.2	11
127	Modelling the cascade of biomarker changes in -related frontotemporal dementia. *Journal of Neurology, Neurosurgery and Psychiatry*, **2021**, 92, 494-501	5.5	11

126	Education modulates brain maintenance in presymptomatic frontotemporal dementia. *Journal of Neurology, Neurosurgery and Psychiatry*, **2019**, 90, 1124-1130	5.5	10
125	Interrupted CAG expansions in ATXN2 gene expand the genetic spectrum of frontotemporal dementias. *Acta Neuropathologica Communications*, **2018**, 6, 41	7.3	10
124	First demonstrated de novo insertion in the prion protein gene in a young patient with dementia. *Journal of Neurology, Neurosurgery and Psychiatry*, **2008**, 79, 845-6	5.5	10
123	APP-derived peptides reflect neurodegeneration in frontotemporal dementia. *Annals of Clinical and Translational Neurology*, **2019**, 6, 2518-2530	5.3	10
122	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	1.2	10
121	Diagnostic Accuracy of MRI Visual Rating Scales in the Diagnosis of Early Onset Cognitive Impairment. *Journal of Alzheimer's Disease*, **2020**, 73, 1575-1583	4.3	9
120	Altered Blood Gene Expression of Tumor-Related Genes (PRKCB, BECN1, and CDKN2A) in Alzheimer's Disease. *Molecular Neurobiology*, **2016**, 53, 5902-5911	6.2	9
119	Naming is associated with left temporal pole metabolite levels in neurodegenerative diseases. *Dementia and Geriatric Cognitive Disorders*, **2008**, 25, 212-7	2.6	9
118	Common variants in Alzheimer's disease: Novel association of six genetic variants with AD and risk stratification by polygenic risk scores		9
117	Digital biomarker-based individualized prognosis for people at risk of dementia. *Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring*, **2020**, 12, e12073	5.2	9
116	Sporadic MM2-thalamic + cortical Creutzfeldt-Jakob disease: Utility of diffusion tensor imaging in the detection of cortical involvement in vivo. *Neuropathology*, **2016**, 36, 199-204	2	9
115	Apathy in presymptomatic genetic frontotemporal dementia predicts cognitive decline and is driven by structural brain changes. *Alzheimer's and Dementia*, **2021**, 17, 969-983	1.2	9
114	A modified Camel and Cactus Test detects presymptomatic semantic impairment in genetic frontotemporal dementia within the GENFI cohort. *Applied Neuropsychology Adult*, **2020**, 1-8	1.9	8
113	Contribution of CSF biomarkers to early-onset Alzheimer's disease and frontotemporal dementia neuroimaging signatures. *Human Brain Mapping*, **2020**, 41, 2004-2013	5.9	8
112	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	3.1	8
111	Breakpoint sequence analysis of an APP locus duplication associated with autosomal dominant Alzheimer's disease and severe cerebral amyloid angiopathy. *Journal of Alzheimer's Disease*, **2012**, 28, 303-8	4.3	8
110	Brain functional network integrity sustains cognitive function despite atrophy in presymptomatic genetic frontotemporal dementia. *Alzheimer's and Dementia*, **2021**, 17, 500-514	1.2	8
109	Variably protease-sensitive prionopathy presenting within ALS/FTD spectrum. *Annals of Clinical and Translational Neurology*, **2018**, 5, 1297-1302	5.3	8

108	Improved Cerebrospinal Fluid-Based Discrimination between Alzheimer's Disease Patients and Controls after Correction for Ventricular Volumes. *Journal of Alzheimer's Disease*, **2017**, 56, 543-555	4.3	7
107	Assessing the role of TUBA4A gene in frontotemporal degeneration. *Neurobiology of Aging*, **2016**, 38, 215.e13-215.e14	5.6	7
106	Large APP locus duplication in a sporadic case of cerebral haemorrhage. *Neurogenetics*, **2014**, 15, 145-9	3	7
105	Inherited prion disease with 4-octapeptide repeat insertion linked to valine at codon 129. *Brain*, **2012**, 135, e212	11.2	7
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