

Raquel Sanchez-Valle

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/1317273/raquel-sanchez-valle-publications-by-year.pdf>

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

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	Cognitive composites for genetic frontotemporal dementia: GENFI-Cog.. <i>Alzheimer's Research and Therapy</i> , 2022 , 14, 10	9	0
268	Cortical microstructure in primary progressive aphasia: a multicenter study.. <i>Alzheimer's Research and Therapy</i> , 2022 , 14, 27	9	0
267	Examining empathy deficits across familial forms of frontotemporal dementia within the GENFI cohort.. <i>Cortex</i> , 2022 , 150, 12-28	3.8	
266	Data-driven staging of genetic frontotemporal dementia using multi-modal MRI.. <i>Human Brain Mapping</i> , 2022 ,	5.9	1
265	Diagnostic accuracy of cerebrospinal fluid biomarkers in genetic prion diseases.. <i>Brain</i> , 2022 ,	11.2	1
264	Anomia is present pre-symptomatically in frontotemporal dementia due to MAPT mutations.. <i>Journal of Neurology</i> , 2022 , 1	5.5	
263	A neurodegenerative disease landscape of rare mutations in Colombia due to founder effects.. <i>Genome Medicine</i> , 2022 , 14, 27	14.4	1
262	Development of a sensitive trial-ready poly(GP) CSF biomarker assay for -associated frontotemporal dementia and amyotrophic lateral sclerosis.. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2022 ,	5.5	2
261	New insights into the genetic etiology of Alzheimer's disease and related dementias.. <i>Nature Genetics</i> , 2022 ,	36.3	27
260	CSF Biomarkers in COVID-19 Associated Encephalopathy and Encephalitis Predict Long-Term Outcome.. <i>Frontiers in Immunology</i> , 2022 , 13, 866153	8.4	2
259	Homozygous R136S mutation in PRNP gene causes inherited early onset prion disease. <i>Alzheimer's Research and Therapy</i> , 2021 , 13, 176	9	2
258	Sex differences in the behavioral variant of frontotemporal dementia: A new window to executive and behavioral reserve. <i>Alzheimer's and Dementia</i> , 2021 , 17,	1.2	3
257	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. <i>Alzheimer's and Dementia</i> , 2021 ,	1.2	2
256	Stratifying the Presymptomatic Phase of Genetic Frontotemporal Dementia by Serum NfL and pNfH: A Longitudinal Multicentre Study. <i>Annals of Neurology</i> , 2021 ,	9.4	2
255	Homozygous R136S mutation in PRNP gene causes inherited early onset prion disease. <i>Alzheimer's Research and Therapy</i> , 2021 , 13, 176	9	1
254	A panel of CSF proteins separates genetic frontotemporal dementia from presymptomatic mutation carriers: a GENFI study. <i>Molecular Neurodegeneration</i> , 2021 , 16, 79	19	0
253	Baseline MRI atrophy predicts 2-year cognitive outcomes in early-onset Alzheimer's disease. <i>Journal of Neurology</i> , 2021 , 1	5.5	0

252	A data-driven disease progression model of fluid biomarkers in genetic frontotemporal dementia. <i>Brain</i> , 2021 ,	11.2	3
251	Prion diseases. <i>European Neuropsychopharmacology</i> , 2021 , 55, 1-3	1.2	
250	Detection of β -synuclein in CSF by RT-QuIC in patients with isolated rapid-eye-movement sleep behaviour disorder: a longitudinal observational study. <i>Lancet Neurology</i> , 2021 , 20, 203-212	24.1	52
249	MRI data-driven algorithm for the diagnosis of behavioural variant frontotemporal dementia. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2021 ,	5.5	3
248	Diagnostic Utility of Measuring Cerebral Atrophy in the Behavioral Variant of Frontotemporal Dementia and Association With Clinical Deterioration. <i>JAMA Network Open</i> , 2021 , 4, e211290	10.4	3
247	CSF sTREM2 is elevated in a subset in GRN-related frontotemporal dementia. <i>Neurobiology of Aging</i> , 2021 , 103, 158.e1-158.e5	5.6	2
246	TREM2 expression in the brain and biological fluids in prion diseases. <i>Acta Neuropathologica</i> , 2021 , 141, 841-859	14.3	5
245	Accelerated long-term forgetting in individuals with subjective cognitive decline and amyloid- β positivity. <i>International Journal of Geriatric Psychiatry</i> , 2021 , 36, 1037-1049	3.9	2
244	Plasma Neurofilament Light for Prediction of Disease Progression in Familial Frontotemporal Lobar Degeneration. <i>Neurology</i> , 2021 , 96, e2296-e2312	6.5	12
243	Characterizing the Clinical Features and Atrophy Patterns of β -Related Frontotemporal Dementia With Disease Progression Modeling. <i>Neurology</i> , 2021 , 97, e941-e952	6.5	3
242	Common variants in Alzheimer's disease and risk stratification by polygenic risk scores. <i>Nature Communications</i> , 2021 , 12, 3417	17.4	23
241	Multi-cohort profiling reveals elevated CSF levels of brain-enriched proteins in Alzheimer's disease. <i>Annals of Clinical and Translational Neurology</i> , 2021 , 8, 1456-1470	5.3	3
240	A trial of gantenerumab or solanezumab in dominantly inherited Alzheimer's disease. <i>Nature Medicine</i> , 2021 , 27, 1187-1196	50.5	51
239	The Revised Self-Monitoring Scale detects early impairment of social cognition in genetic frontotemporal dementia within the GENFI cohort. <i>Alzheimer's Research and Therapy</i> , 2021 , 13, 127	9	2
238	Genetic variation in APOE, GRN, and TP53 are phenotype modifiers in frontotemporal dementia. <i>Neurobiology of Aging</i> , 2021 , 99, 99.e15-99.e22	5.6	3
237	Biphasic cortical macro- and microstructural changes in autosomal dominant Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2021 , 17, 618-628	1.2	11
236	Brain functional network integrity sustains cognitive function despite atrophy in presymptomatic genetic frontotemporal dementia. <i>Alzheimer's and Dementia</i> , 2021 , 17, 500-514	1.2	8
235	Presentations and mechanisms of CNS disorders related to COVID-19. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2021 , 8,	9.1	28

234	Errorless Learning Therapy in Semantic Variant of Primary Progressive Aphasia. <i>Journal of Alzheimer's Disease</i> , 2021 , 79, 415-422	4.3	2
233	Apathy in presymptomatic genetic frontotemporal dementia predicts cognitive decline and is driven by structural brain changes. <i>Alzheimer's and Dementia</i> , 2021 , 17, 969-983	1.2	9
232	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. <i>JAMA Network Open</i> , 2021 , 4, e2030194	10.4	14
231	Modelling the cascade of biomarker changes in -related frontotemporal dementia. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2021 , 92, 494-501	5.5	11
230	Cognitive decline in amyotrophic lateral sclerosis: Neuropathological substrate and genetic determinants. <i>Brain Pathology</i> , 2021 , 31, e12942	6	1
229	Sex differences in the behavioral variant of frontotemporal dementia: A new window to executive and behavioral reserve. <i>Alzheimer's and Dementia</i> , 2021 , 17, 1329-1341	1.2	10
228	Pioglitazone for prevention of cognitive impairment: results and lessons. <i>Lancet Neurology</i> , 2021 , 20, 500-502	24.1	0
227	Practice effects in genetic frontotemporal dementia and at-risk individuals: a GENFI study. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2021 ,	5.5	
226	Dissemination in time and space in presymptomatic granulin mutation carriers: a GENFI spatial chronnectome study. <i>Neurobiology of Aging</i> , 2021 , 108, 155-167	5.6	0
225	Neuroanatomical and cognitive correlates of visual hallucinations in Parkinson's disease and dementia with Lewy bodies: Voxel-based morphometry and neuropsychological meta-analysis. <i>Neuroscience and Biobehavioral Reviews</i> , 2021 , 128, 367-382	9	3
224	Differential early subcortical involvement in genetic FTD within the GENFI cohort. <i>NeuroImage: Clinical</i> , 2021 , 30, 102646	5.3	6
223	Disease-related cortical thinning in presymptomatic granulin mutation carriers. <i>NeuroImage: Clinical</i> , 2021 , 29, 102540	5.3	2
222	Longitudinal brain atrophy and CSF biomarkers in early-onset Alzheimer's disease. <i>NeuroImage: Clinical</i> , 2021 , 32, 102804	5.3	3
221	SLITRK2, an X-linked modifier of the age at onset in C9orf72 frontotemporal lobar degeneration. <i>Brain</i> , 2021 , 144, 2798-2811	11.2	2
220	Accelerated long-term forgetting over three months in asymptomatic APOE e4 carriers. <i>Annals of Clinical and Translational Neurology</i> , 2021 , 8, 477-484	5.3	5
219	ALTOIDA-iADL for the diagnosis of Mild Cognitive Impairment and early Alzheimer's disease.. <i>Alzheimer's and Dementia</i> , 2021 , 17 Suppl 12, e057982	1.2	
218	Impact of COVID-19 pandemic in an early-onset dementia clinic in Barcelona.. <i>Alzheimer's and Dementia</i> , 2021 , 17 Suppl 7, e052114	1.2	
217	Functional network alterations in early-onset Alzheimer's disease studied with resting-state fMRI. <i>Alzheimer's and Dementia</i> , 2020 , 16, e043307	1.2	

216	Telemedicine assessment of long-term cognitive and functional status in anti-leucine-rich, glioma-inactivated 1 encephalitis. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2020 , 7,	9.1	13
215	Autoantibodies against the prion protein in individuals with mutations. <i>Neurology</i> , 2020 , 95, e2028-e2037.	5.5	5
214	A modified Camel and Cactus Test detects presymptomatic semantic impairment in genetic frontotemporal dementia within the GENFI cohort. <i>Applied Neuropsychology Adult</i> , 2020 , 1-8	1.9	8
213	Plasma glial fibrillary acidic protein is raised in progranulin-associated frontotemporal dementia. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020 , 91, 263-270	5.5	40
212	Diagnostic Accuracy of MRI Visual Rating Scales in the Diagnosis of Early Onset Cognitive Impairment. <i>Journal of Alzheimer's Disease</i> , 2020 , 73, 1575-1583	4.3	9
211	Contribution of CSF biomarkers to early-onset Alzheimer's disease and frontotemporal dementia neuroimaging signatures. <i>Human Brain Mapping</i> , 2020 , 41, 2004-2013	5.9	8
210	Neuronal pentraxin 2: a synapse-derived CSF biomarker in genetic frontotemporal dementia. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020 , 91, 612-621	5.5	22
209	Screening of dementia genes by whole-exome sequencing in Spanish patients with early-onset dementia: likely pathogenic, uncertain significance and risk variants. <i>Neurobiology of Aging</i> , 2020 , 93, e1-e9	5.6	4
208	Social cognition impairment in genetic frontotemporal dementia within the GENFI cohort. <i>Cortex</i> , 2020 , 133, 384-398	3.8	7
207	Synaptic, axonal damage and inflammatory cerebrospinal fluid biomarkers in neurodegenerative dementias. <i>Alzheimer's and Dementia</i> , 2020 , 16, 262-272	1.2	29
206	Role for ATXN1, ATXN2, and HTT intermediate repeats in frontotemporal dementia and Alzheimer's disease. <i>Neurobiology of Aging</i> , 2020 , 87, 139.e1-139.e7	5.6	13
205	Multicenter Alzheimer's and Parkinson's disease immune biomarker verification study. <i>Alzheimer's and Dementia</i> , 2020 , 16, 292-304	1.2	16
204	Age at symptom onset and death and disease duration in genetic frontotemporal dementia: an international retrospective cohort study. <i>Lancet Neurology, The</i> , 2020 , 19, 145-156	24.1	90
203	Sex Differences of Longitudinal Brain Changes in Cognitively Unimpaired Adults. <i>Journal of Alzheimer's Disease</i> , 2020 , 76, 1413-1422	4.3	1
202	Use of Antipsychotics in Patients with Behavioral and Psychological Symptoms of Dementia: Results of a Spanish Delphi Consensus. <i>Dementia and Geriatric Cognitive Disorders</i> , 2020 , 49, 573-582	2.6	1
201	LifeTime and improving European healthcare through cell-based interceptive medicine. <i>Nature</i> , 2020 , 587, 377-386	50.4	56
200	Early symptoms in symptomatic and preclinical genetic frontotemporal lobar degeneration. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020 , 91, 975-984	5.5	15
199	Abnormal pain perception is associated with thalamo-cortico-striatal atrophy in expansion carriers in the GENFI cohort. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020 , 91, 1325-1328	5.5	5

198	Effects of IgLON5 Antibodies on Neuronal Cytoskeleton: A Link between Autoimmunity and Neurodegeneration. <i>Annals of Neurology</i> , 2020 , 88, 1023-1027	9.4	25
197	, age at onset, and ancestry help discriminate behavioral from language variants in FTL D cohorts. <i>Neurology</i> , 2020 , 95, e3288-e3302	6.5	5
196	Digital biomarker-based individualized prognosis for people at risk of dementia. <i>Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2020 , 12, e12073	5.2	9
195	Analysis of brain atrophy and local gene expression in genetic frontotemporal dementia. <i>Brain Communications</i> , 2020 , 2,	4.5	6
194	Faster Cortical Thinning and Surface Area Loss in Presymptomatic and Symptomatic C9orf72 Repeat Expansion Adult Carriers. <i>Annals of Neurology</i> , 2020 , 88, 113-122	9.4	11
193	Microglial Hyperreactivity Evolved to Immunosuppression in the Hippocampus of a Mouse Model of Accelerated Aging and Alzheimer's Disease Traits. <i>Frontiers in Aging Neuroscience</i> , 2020 , 12, 622360	5.3	1
192	Evolution of Clinical-Pathological Correlations in Early-Onset Alzheimer's Disease Over a 25-Year Period in an Academic Brain Bank. <i>Journal of Alzheimer's Disease</i> , 2020 , 1-11	4.3	0
191	Early detection of subtle motor dysfunction in cognitively normal subjects with amyloid- β positivity. <i>Cortex</i> , 2019 , 121, 117-124	3.8	3
190	The inner fluctuations of the brain in presymptomatic Frontotemporal Dementia: The chronnectome fingerprint. <i>NeuroImage</i> , 2019 , 189, 645-654	7.9	18
189	Tau Protein is Associated with Longitudinal Memory Decline in Cognitively Healthy Subjects with Normal Alzheimer's Disease Cerebrospinal Fluid Biomarker Levels. <i>Journal of Alzheimer's Disease</i> , 2019 , 70, 211-225	4.3	6
188	Clinical value of cerebrospinal fluid neurofilament light chain in semantic dementia. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019 , 90, 997-1004	5.5	13
187	Education modulates brain maintenance in presymptomatic frontotemporal dementia. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019 , 90, 1124-1130	5.5	10
186	Regional patterns of 18F-florbetaben uptake in presenilin 1 mutation carriers. <i>Neurobiology of Aging</i> , 2019 , 81, 1-8	5.6	4
185	Novel CSF biomarkers in genetic frontotemporal dementia identified by proteomics. <i>Annals of Clinical and Translational Neurology</i> , 2019 , 6, 698-707	5.3	25
184	Cortical microstructure in the behavioural variant of frontotemporal dementia: looking beyond atrophy. <i>Brain</i> , 2019 , 142, 1121-1133	11.2	32
183	Mechanisms of functional compensation, delineated by eigenvector centrality mapping, across the pathophysiological continuum of Alzheimer's disease. <i>NeuroImage: Clinical</i> , 2019 , 22, 101777	5.3	17
182	Comparison of Pittsburgh compound B and florbetapir in cross-sectional and longitudinal studies. <i>Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2019 , 11, 180-190	5.2	46
181	CSF glial biomarkers YKL40 and sTREM2 are associated with longitudinal volume and diffusivity changes in cognitively unimpaired individuals. <i>NeuroImage: Clinical</i> , 2019 , 23, 101801	5.3	17

180	Clinical and video-polysomnographic analysis of rapid eye movement sleep behavior disorder and other sleep disturbances in dementia with Lewy bodies. <i>Sleep</i> , 2019 , 42,	1.1	13
179	Clinical applicability of diagnostic biomarkers in early-onset cognitive impairment. <i>European Journal of Neurology</i> , 2019 , 26, 1098-1104	6	11
178	Cerebrospinal Fluid Total Prion Protein in the Spectrum of Prion Diseases. <i>Molecular Neurobiology</i> , 2019 , 56, 2811-2821	6.2	16
177	Hippocampal atrophy has limited usefulness as a diagnostic biomarker on the early onset Alzheimer's disease patients: A comparison between visual and quantitative assessment. <i>NeuroImage: Clinical</i> , 2019 , 23, 101927	5.3	18
176	Novel P397S MAPT variant associated with late onset and slow progressive frontotemporal dementia. <i>Annals of Clinical and Translational Neurology</i> , 2019 , 6, 1559-1565	5.3	3
175	A unique common ancestor introduced P301L mutation in MAPT gene in frontotemporal dementia patients from Barcelona (Baix Llobregat, Spain). <i>Neurobiology of Aging</i> , 2019 , 84, 236.e9-236.e15	5.6	6
174	Serum neurofilament light chain in genetic frontotemporal dementia: a longitudinal, multicentre cohort study. <i>Lancet Neurology</i> , 2019 , 18, 1103-1111	24.1	68
173	IC-02-05: GENDER DIFFERENCES IN THE ASSOCIATION BETWEEN LONGITUDINAL BRAIN CHANGES AND BASELINE LEVELS OF CSF ALZHEIMER'S DISEASE AND GLIAL BIOMARKERS IN HEALTHY ELDERLY 2019 , 15, P4-P4		
172	Plasma levels of soluble TREM2 and neurofilament light chain in TREM2 rare variant carriers. <i>Alzheimer's Research and Therapy</i> , 2019 , 11, 94	9	15
171	APP-derived peptides reflect neurodegeneration in frontotemporal dementia. <i>Annals of Clinical and Translational Neurology</i> , 2019 , 6, 2518-2530	5.3	10
170	White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal GENFI study. <i>NeuroImage: Clinical</i> , 2019 , 24, 102077	5.3	13
169	Evaluation of Human Cerebrospinal Fluid Malate Dehydrogenase 1 as a Marker in Genetic Prion Disease Patients. <i>Biomolecules</i> , 2019 , 9,	5.9	4
168	CSF-ApoER2 fragments as a read-out of reelin signaling: Distinct patterns in sporadic and autosomal-dominant Alzheimer disease. <i>Clinica Chimica Acta</i> , 2019 , 490, 6-11	6.2	1
167	Changes in Synaptic Proteins Precede Neurodegeneration Markers in Preclinical Alzheimer's Disease Cerebrospinal Fluid. <i>Molecular and Cellular Proteomics</i> , 2019 , 18, 546-560	7.6	66
166	Does ALS-FUS without FUS mutation represent ALS-FET? Report of three cases. <i>Neuropathology and Applied Neurobiology</i> , 2019 , 45, 421-426	5.2	3
165	Challenges associated with biomarker-based classification systems for Alzheimer's disease. <i>Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2018 , 10, 346-357	5.2	25
164	Poly(GP), neurofilament and grey matter deficits in expansion carriers. <i>Annals of Clinical and Translational Neurology</i> , 2018 , 5, 583-597	5.3	29
163	Cerebrospinal fluid neurofilament light levels in neurodegenerative dementia: Evaluation of diagnostic accuracy in the differential diagnosis of prion diseases. <i>Alzheimer's and Dementia</i> , 2018 , 14, 751-763	1.2	45

162	Structural Connectivity Alterations Along the Alzheimer's Disease Continuum: Reproducibility Across Two Independent Samples and Correlation with Cerebrospinal Fluid Amyloid- β and Tau. <i>Journal of Alzheimer's Disease</i> , 2018 , 61, 1575-1587	4.3	20
161	Tauopathy with Hippocampal 4-Repeat Tau Immunoreactive Spherical Inclusions in a Patient with PSP. <i>Brain Pathology</i> , 2018 , 28, 284-286	6	2
160	Potential genetic modifiers of disease risk and age at onset in patients with frontotemporal lobar degeneration and GRN mutations: a genome-wide association study. <i>Lancet Neurology</i> , 2018 , 17, 548-558	24.1	60
159	Rare nonsynonymous variants in SORT1 are associated with increased risk for frontotemporal dementia. <i>Neurobiology of Aging</i> , 2018 , 66, 181.e3-181.e10	5.6	12
158	Validation of β -Synuclein as a CSF Biomarker for Sporadic Creutzfeldt-Jakob Disease. <i>Molecular Neurobiology</i> , 2018 , 55, 2249-2257	6.2	30
157	Interrupted CAG expansions in ATXN2 gene expand the genetic spectrum of frontotemporal dementias. <i>Acta Neuropathologica Communications</i> , 2018 , 6, 41	7.3	10
156	The hippocampal longitudinal axis-relevance for underlying tau and TDP-43 pathology. <i>Neurobiology of Aging</i> , 2018 , 70, 1-9	5.6	15
155	Systematic Screening of Ubiquitin/p62 Aggregates in Cerebellar Cortex Expands the Neuropathological Phenotype of the C9orf72 Expansion Mutation. <i>Journal of Neuropathology and Experimental Neurology</i> , 2018 , 77, 703-709	3.1	8
154	No supportive evidence for TIA1 gene mutations in a European cohort of ALS-FTD spectrum patients. <i>Neurobiology of Aging</i> , 2018 , 69, 293.e9-293.e11	5.6	11
153	Common and rare TBK1 variants in early-onset Alzheimer disease in a European cohort. <i>Neurobiology of Aging</i> , 2018 , 62, 245.e1-245.e7	5.6	12
152	P2-347: THE HIPPOCAMPAL LONGITUDINAL AXIS: RELEVANCE FOR UNDERLYING TAU AND TDP-43 PATHOLOGY 2018 , 14, P819-P820		
151	P1-146: WHOLE EXOME SEQUENCING IN PATIENTS WITH EARLY-ONSET ALZHEIMER'S DISEASE AND FRONTOTEMPORAL DEMENTIA: MUTATION DETECTION IN CAUSAL AND RISK GENES FOR DEMENTIA 2018 , 14, P332-P332		
150	P2-262: A CEREBROSPINAL FLUID PANEL OF SYNAPTIC PROTEINS ACROSS THE ENTIRE ALZHEIMER'S DISEASE CONTINUUM 2018 , 14, P777-P777		
149	P3-394: CORTICAL MEAN DIFFUSIVITY MAY BE MORE SENSITIVE IN DETECTING STRUCTURAL CHANGES IN FRONTOTEMPORAL DEMENTIA THAN CORTICAL THICKNESS 2018 , 14, P1248-P1249		
148	P1-432: REGIONAL PATTERNS OF 18F-FLORBETABEN UPTAKE IN PRESENILIN 1 MUTATION CARRIERS 2018 , 14, P475-P475		
147	O3-09-03: SERUM NEUROFILAMENT LIGHT LEVELS CORRELATE WITH SEVERITY MEASURES AND NEURODEGENERATION MARKERS IN AUTOSOMAL DOMINANT ALZHEIMER'S DISEASE 2018 , 14, P1037-P1037		
146	Serum neurofilament light levels correlate with severity measures and neurodegeneration markers in autosomal dominant Alzheimer's disease. <i>Alzheimer's Research and Therapy</i> , 2018 , 10, 113	9	38
145	Prevalence of amyloid- β pathology in distinct variants of primary progressive aphasia. <i>Annals of Neurology</i> , 2018 , 84, 729-740	9.4	74

144	Variably protease-sensitive prionopathy presenting within ALS/FTD spectrum. <i>Annals of Clinical and Translational Neurology</i> , 2018 , 5, 1297-1302	5.3	8
143	P2-230: CHALLENGES ASSOCIATED WITH BIOMARKER-BASED CLASSIFICATIONS SYSTEMS FOR ALZHEIMER'S DISEASE 2018 , 14, P756-P757		
142	βSynuclein aggregates in labial salivary glands of idiopathic rapid eye movement sleep behavior disorder. <i>Sleep</i> , 2018 , 41,	1.1	12
141	Adaptation and validation of a Spanish-language version of the Frontotemporal Dementia Rating Scale (FTD-FRS). <i>Neurología</i> , 2017 , 32, 290-299	1.4	3
140	Active Aβ immunotherapy CAD106 in Alzheimer's disease: A phase 2b study. <i>Alzheimer's and Dementia: Translational Research and Clinical Interventions</i> , 2017 , 3, 10-22	6	65
139	The ε genotype modulates CSF YKL-40 levels and their structural brain correlates in the continuum of Alzheimer's disease but not those of sTREM2. <i>Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2017 , 6, 50-59	5.2	29
138	A Common Variant in the MC1R Gene (p.V92M) is associated with Alzheimer's Disease Risk. <i>Journal of Alzheimer's Disease</i> , 2017 , 56, 1065-1074	4.3	1
137	Improved Cerebrospinal Fluid-Based Discrimination between Alzheimer's Disease Patients and Controls after Correction for Ventricular Volumes. <i>Journal of Alzheimer's Disease</i> , 2017 , 56, 543-555	4.3	7
136	Cerebral amyloid angiopathy in Down syndrome and sporadic and autosomal-dominant Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2017 , 13, 1251-1260	1.2	38
135	Deleterious ABCA7 mutations and transcript rescue mechanisms in early onset Alzheimer's disease. <i>Acta Neuropathologica</i> , 2017 , 134, 475-487	14.3	34
134	C-terminal fragments of the amyloid precursor protein in cerebrospinal fluid as potential biomarkers for Alzheimer disease. <i>Scientific Reports</i> , 2017 , 7, 2477	4.9	21
133	CSF sAPPβ, YKL-40, and neurofilament light in frontotemporal lobar degeneration. <i>Neurology</i> , 2017 , 89, 178-188	6.5	75
132	TBK1 Mutation Spectrum in an Extended European Patient Cohort with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. <i>Human Mutation</i> , 2017 , 38, 297-309	4.7	66
131	YKL-40 in the brain and cerebrospinal fluid of neurodegenerative dementias. <i>Molecular Neurodegeneration</i> , 2017 , 12, 83	19	86
130	Dopamine transporter imaging deficit predicts early transition to synucleinopathy in idiopathic rapid eye movement sleep behavior disorder. <i>Annals of Neurology</i> , 2017 , 82, 419-428	9.4	108
129	Frontotemporal Dementia Caused by the P301L Mutation in the MAPT Gene: Clinicopathological Features of 13 Cases from the Same Geographical Origin in Barcelona, Spain. <i>Dementia and Geriatric Cognitive Disorders</i> , 2017 , 44, 213-221	2.6	23
128	Adaptation and validation of a Spanish-language version of the Frontotemporal Dementia Rating Scale (FTD-FRS). <i>Neurología (English Edition)</i> , 2017 , 32, 290-299	0.4	0
127	[P4007]: A PRESCREENING STUDY IN SPAIN TO IDENTIFY INDIVIDUALS WHO MIGHT BE SUITABLE FOR PARTICIPATION IN CLINICAL STUDIES IN EARLY ALZHEIMER'S DISEASE UTILIZING REFERRALS FROM GENERAL PRACTITIONER NETWORKS 2017 , 13, P1345-P1346		

126	[P4189]: SYMPTOM ONSET IN GENETIC FRONTOTEMPORAL DEMENTIA 2017 , 13, P1337-P1337		1
125	Neuronal intranuclear (hyaline) inclusion disease and fragile X-associated tremor/ataxia syndrome: a morphological and molecular dilemma. <i>Brain</i> , 2017 , 140, e51	11.2	27
124	CSF microRNA Profiling in Alzheimer's Disease: a Screening and Validation Study. <i>Molecular Neurobiology</i> , 2017 , 54, 6647-6654	6.2	29
123	Plasma miR-34a-5p and miR-545-3p as Early Biomarkers of Alzheimer's Disease: Potential and Limitations. <i>Molecular Neurobiology</i> , 2017 , 54, 5550-5562	6.2	81
122	[P2110]: VISUAL AND QUANTITATIVE ASSESSMENT OF HIPPOCAMPAL ATROPHY IN EARLY ONSET ALZHEIMER'S DISEASE PATIENTS 2017 , 13, P789-P789		
121	[P4114]: TOWARD A FUNCTIONAL NEUROMARKER FOR PRECLINICAL AD: EIGENVECTOR CENTRALITY REVEALS PRECLINICAL DIFFERENCES OF FUNCTIONAL INFORMATION FLOW IN THE HIPPOCAMPUS, PRECUNEUS, CEREBELLUM AND INFERIOR PARIETAL LOBULE 2017 , 13, P1348-P1349		
120	[P1113]: VISUAL PATTERNS OF FLORBETABEN UPTAKE IN PRESENILIN 1 MUTATION CARRIERS 2017 , 13, P435-P435		
119	Clinical Neuropathology image 4-2017: High-resolution 7 Tesla MRI of postmortem brain specimens: improving neuroimaging-neuropathology correlations 2017 , 36, 162-163		1
118	Quantitative Magnetic Resonance Abnormalities in Creutzfeldt-Jakob Disease and Fatal Insomnia. <i>Journal of Alzheimer's Disease</i> , 2017 , 55, 431-443	4.3	13
117	Validation of 14-3-3 Protein as a Marker in Sporadic Creutzfeldt-Jakob Disease Diagnostic. <i>Molecular Neurobiology</i> , 2016 , 53, 2189-99	6.2	65
116	Neurofilament light chain: a biomarker for genetic frontotemporal dementia. <i>Annals of Clinical and Translational Neurology</i> , 2016 , 3, 623-36	5.3	163
115	Preservation of cell-survival mechanisms by the presenilin-1 K239N mutation may cause its milder clinical phenotype. <i>Neurobiology of Aging</i> , 2016 , 46, 169-79	5.6	3
114	Cerebrospinal fluid sTREM2 levels are associated with gray matter volume increases and reduced diffusivity in early Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2016 , 12, 1259-1272	1.2	67
113	Cerebrospinal Fluid Biomarkers Predict Clinical Evolution in Patients with Subjective Cognitive Decline and Mild Cognitive Impairment. <i>Neurodegenerative Diseases</i> , 2016 , 16, 69-76	2.3	34
112	Assessing the role of TUBA4A gene in frontotemporal degeneration. <i>Neurobiology of Aging</i> , 2016 , 38, 215.e13-215.e14	5.6	7
111	CSF YKL-40 and pTau181 are related to different cerebral morphometric patterns in early AD. <i>Neurobiology of Aging</i> , 2016 , 38, 47-55	5.6	42
110	Altered Blood Gene Expression of Tumor-Related Genes (PRKCB, BECN1, and CDKN2A) in Alzheimer's Disease. <i>Molecular Neurobiology</i> , 2016 , 53, 5902-5911	6.2	9
109	Sporadic MM2-thalamic + cortical Creutzfeldt-Jakob disease: Utility of diffusion tensor imaging in the detection of cortical involvement in vivo. <i>Neuropathology</i> , 2016 , 36, 199-204	2	9

108	stREM2 cerebrospinal fluid levels are a potential biomarker for microglia activity in early-stage Alzheimer's disease and associate with neuronal injury markers. <i>EMBO Molecular Medicine</i> , 2016 , 8, 466-76	12	256
107	White Matter Abnormalities Track Disease Progression in PSEN1 Autosomal Dominant Alzheimer's Disease. <i>Journal of Alzheimer's Disease</i> , 2016 , 51, 827-35	4.3	12
106	IC-02-05: Cerebrospinal Fluid Strem2 Levels are Associated with Gray Matter Volume Increases and Reduced Diffusivity in Early Alzheimer's Disease 2016 , 12, P8-P8		
105	A comprehensive study of the genetic impact of rare variants in SORL1 in European early-onset Alzheimer's disease. <i>Acta Neuropathologica</i> , 2016 , 132, 213-224	14.3	62
104	Cerebrospinal fluid Presenilin-1 increases at asymptomatic stage in genetically determined Alzheimer's disease. <i>Molecular Neurodegeneration</i> , 2016 , 11, 66	19	6
103	Gerstmann-Strüssler-Scheinker Disease Presenting with Atypical Parkinsonism, but Typical Magnetic Resonance Imaging Findings of Prion Disease. <i>Movement Disorders Clinical Practice</i> , 2016 , 3, 93-95	2.2	4
102	Analysis of the CHCHD10 gene in patients with frontotemporal dementia and amyotrophic lateral sclerosis from Spain. <i>Brain</i> , 2015 , 138, e400	11.2	47
101	Genetic variability in SQSTM1 and risk of early-onset Alzheimer dementia: a European early-onset dementia consortium study. <i>Neurobiology of Aging</i> , 2015 , 36, 2005.e15-22	5.6	29
100	O3-13-06: Targeted re-sequencing of sorl1 in early-onset Alzheimer's dementia: The european early onset dementia consortium 2015 , 11, P253-P253		
99	Rare Variants in PLD3 Do Not Affect Risk for Early-Onset Alzheimer Disease in a European Consortium Cohort. <i>Human Mutation</i> , 2015 , 36, 1226-35	4.7	20
98	Clinicopathological Correlations and Concomitant Pathologies in Rapidly Progressive Dementia: A Brain Bank Series. <i>Neurodegenerative Diseases</i> , 2015 , 15, 350-60	2.3	27
97	Diagnostic accuracy of behavioral variant frontotemporal dementia consortium criteria (FTDC) in a clinicopathological cohort. <i>Neuropathology and Applied Neurobiology</i> , 2015 , 41, 882-92	5.2	21
96	Evolving brain structural changes in PSEN1 mutation carriers. <i>Neurobiology of Aging</i> , 2015 , 36, 1261-70	5.6	27
95	Rapidly progressive dementia with psychotic onset in a patient with the C9ORF72 mutation 2015 , 34, 294-7		6
94	TMEM106B is a genetic modifier of frontotemporal lobar degeneration with C9orf72 hexanucleotide repeat expansions. <i>Acta Neuropathologica</i> , 2014 , 127, 407-18	14.3	97
93	Large APP locus duplication in a sporadic case of cerebral haemorrhage. <i>Neurogenetics</i> , 2014 , 15, 145-9	3	7
92	Characterization of the repeat expansion size in C9orf72 in amyotrophic lateral sclerosis and frontotemporal dementia. <i>Human Molecular Genetics</i> , 2014 , 23, 749-54	5.6	84
91	Plasma phosphorylated TDP-43 levels are elevated in patients with frontotemporal dementia carrying a C9orf72 repeat expansion or a GRN mutation. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014 , 85, 684-91	5.5	47

90	Cerebrospinal fluid level of YKL-40 protein in preclinical and prodromal Alzheimer's disease. <i>Journal of Alzheimer's Disease</i> , 2014 , 42, 901-8	4.3	79
89	Investigation of the role of rare TREM2 variants in frontotemporal dementia subtypes. <i>Neurobiology of Aging</i> , 2014 , 35, 2657.e13-2657.e19	5.6	31
88	Rare mutations in SQSTM1 modify susceptibility to frontotemporal lobar degeneration. <i>Acta Neuropathologica</i> , 2014 , 128, 397-410	14.3	83
87	O4-11-04: ACTIVE ADJUVANT IMMUNOTHERAPY CAD106 PHASE II DOSE-ADJUVANT FINDING STUDY: SAFETY AND CNS BIOMARKERS 2014 , 10, P274-P274		2
86	Usefulness of biomarkers in the diagnosis and prognosis of early-onset cognitive impairment. <i>Journal of Alzheimer's Disease</i> , 2014 , 40, 919-27	4.3	27
85	Determination of neuronal antibodies in suspected and definite Creutzfeldt-Jakob disease. <i>JAMA Neurology</i> , 2014 , 71, 74-8	17.2	43
84	In vivo decreased dopamine transporter uptake in corticobasal degeneration presenting with primary progressive aphasia without parkinsonism. <i>European Journal of Neurology</i> , 2014 , 21, e56-7	6	1
83	Reply: To PMID 23794434. <i>Annals of Neurology</i> , 2014 , 75, 460-1	9.4	
82	TREM2 mutations implicated in neurodegeneration impair cell surface transport and phagocytosis. <i>Science Translational Medicine</i> , 2014 , 6, 243ra86	17.5	436
81	Assessing the role of the TREM2 p.R47H variant as a risk factor for Alzheimer's disease and frontotemporal dementia. <i>Neurobiology of Aging</i> , 2014 , 35, 444.e1-4	5.6	81
80	TARDBP mutation p.Ile383Val associated with semantic dementia and complex proteinopathy. <i>Neuropathology and Applied Neurobiology</i> , 2014 , 40, 225-30	5.2	36
79	Neurodegenerative disorder risk in idiopathic REM sleep behavior disorder: study in 174 patients. <i>PLoS ONE</i> , 2014 , 9, e89741	3.7	308
78	Low cerebrospinal fluid concentration of mitochondrial DNA in preclinical Alzheimer disease. <i>Annals of Neurology</i> , 2013 , 74, 655-68	9.4	129
77	A pan-European study of the C9orf72 repeat associated with FTLD: geographic prevalence, genomic instability, and intermediate repeats. <i>Human Mutation</i> , 2013 , 34, 363-73	4.7	208
76	Atypical neuropathological sCJD-MM phenotype with abundant white matter Kuru-type plaques sparing the cerebellar cortex. <i>Neuropathology</i> , 2013 , 33, 204-8	2	6
75	Distinct patterns of APP processing in the CNS in autosomal-dominant and sporadic Alzheimer disease. <i>Acta Neuropathologica</i> , 2013 , 125, 201-13	14.3	81
74	A preliminary study of the whole-genome expression profile of sporadic and monogenic early-onset Alzheimer's disease. <i>Neurobiology of Aging</i> , 2013 , 34, 1772-8	5.6	59
73	Distinctive age-related temporal cortical thinning in asymptomatic granulin gene mutation carriers. <i>Neurobiology of Aging</i> , 2013 , 34, 1462-8	5.6	14

72	Neurodegenerative disease status and post-mortem pathology in idiopathic rapid-eye-movement sleep behaviour disorder: an observational cohort study. <i>Lancet Neurology, The</i> , 2013 , 12, 443-53	24.1	472
71	Neuroimaging and biochemical markers in the three variants of primary progressive aphasia. <i>Dementia and Geriatric Cognitive Disorders</i> , 2013 , 35, 106-17	2.6	43
70	Cognitive reserve proxies relate to gray matter loss in cognitively healthy elderly with abnormal cerebrospinal fluid amyloid- β levels. <i>Journal of Alzheimer's Disease</i> , 2013 , 35, 715-26	4.3	33
69	Donepezil treatment stabilizes functional connectivity during resting state and brain activity during memory encoding in Alzheimer's disease. <i>Journal of Clinical Psychopharmacology</i> , 2013 , 33, 199-205	1.7	32
68	Decreased striatal dopamine transporter uptake in the non-fluent/agrammatic variant of primary progressive aphasia. <i>European Journal of Neurology</i> , 2013 , 20, 1459-e126	6	6
67	Rare variants in calcium homeostasis modulator 1 (CALHM1) found in early onset Alzheimer's disease patients alter calcium homeostasis. <i>PLoS ONE</i> , 2013 , 8, e74203	3.7	22
66	Evolving brain functional abnormalities in PSEN1 mutation carriers: a resting and visual encoding fMRI study. <i>Journal of Alzheimer's Disease</i> , 2013 , 36, 165-75	4.3	15
65	"Preclinical" MSA in definite Creutzfeldt-Jakob disease. <i>Neuropathology</i> , 2012 , 32, 158-63	2	14
64	Applying the new research diagnostic criteria: MRI findings and neuropsychological correlations of prodromal AD. <i>International Journal of Geriatric Psychiatry</i> , 2012 , 27, 127-34	3.9	33
63	Serum progranulin levels in patients with frontotemporal lobar degeneration and Alzheimer's disease: detection of GRN mutations in a Spanish cohort. <i>Journal of Alzheimer's Disease</i> , 2012 , 31, 581-94	4.3	29
62	Distinct functional activity of the precuneus and posterior cingulate cortex during encoding in the preclinical stage of Alzheimer's disease. <i>Journal of Alzheimer's Disease</i> , 2012 , 31, 517-26	4.3	39
61	Hirano body-rich subtypes of Creutzfeldt-Jakob disease. <i>Neuropathology and Applied Neurobiology</i> , 2012 , 38, 153-61	5.2	6
60	Rapidly progressive dementia: experience in a tertiary care medical center. <i>Alzheimer Disease and Associated Disorders</i> , 2012 , 26, 267-71	2.5	35
59	Inherited prion disease with 4-octapeptide repeat insertion linked to valine at codon 129. <i>Brain</i> , 2012 , 135, e212	11.2	7
58	Breakpoint sequence analysis of an A β P locus duplication associated with autosomal dominant Alzheimer's disease and severe cerebral amyloid angiopathy. <i>Journal of Alzheimer's Disease</i> , 2012 , 28, 303-8	4.3	8
57	Cerebrospinal fluid biomarker supported diagnosis of Creutzfeldt-Jakob disease and rapid dementias: a longitudinal multicentre study over 10 years. <i>Brain</i> , 2012 , 135, 3051-61	11.2	102
56	Identifying earlier Alzheimer's disease: insights from the preclinical and prodromal phases. <i>Neurodegenerative Diseases</i> , 2012 , 10, 158-60	2.3	12
55	PSEN1 mutation carriers present lower cerebrospinal fluid amyloid- β 2 levels than sporadic early-onset Alzheimer's disease patients but no differences in neuronal injury biomarkers. <i>Journal of Alzheimer's Disease</i> , 2012 , 30, 605-16	4.3	6

54	The amyloid- β isoform pattern in cerebrospinal fluid in familial PSEN1 M139T- and L286P-associated Alzheimer's disease. <i>Molecular Medicine Reports</i> , 2012 , 5, 1111-5	2.9	14
53	PICOGEN: Five years experience with a genetic counselling program for dementia. <i>Neurología (English Edition)</i> , 2011 , 26, 143-149	0.4	
52	Cognitively preserved subjects with transitional cerebrospinal fluid β amyloid 1-42 values have thicker cortex in Alzheimer's disease vulnerable areas. <i>Biological Psychiatry</i> , 2011 , 70, 183-90	7.9	75
51	A novel PSEN1 gene mutation (L235R) associated with familial early-onset Alzheimer's disease. <i>Neuroscience Letters</i> , 2011 , 496, 40-2	3.3	11
50	Focusing on atypical symptoms for improved diagnosis of early-onset Alzheimer's disease. <i>Future Neurology</i> , 2011 , 6, 575-578	1.5	3
49	Cerebrospinal fluid biomarkers and memory present distinct associations along the continuum from healthy subjects to AD patients. <i>Journal of Alzheimer's Disease</i> , 2011 , 23, 319-26	4.3	56
48	PICOGEN: five years experience with a genetic counselling program for dementia. <i>Neurología</i> , 2011 , 26, 143-9	1.4	19
47	Association between cerebrospinal fluid tau and brain atrophy is not related to clinical severity in the Alzheimer's disease continuum. <i>Psychiatry Research - Neuroimaging</i> , 2011 , 192, 140-6	2.9	15
46	Different profiles of Alzheimer's disease cerebrospinal fluid biomarkers in controls and subjects with subjective memory complaints. <i>Journal of Neural Transmission</i> , 2011 , 118, 259-62	4.3	41
45	Neuropsychological profile of prodromal Alzheimer's disease (Prd-AD) and their radiological correlates. <i>Archives of Gerontology and Geriatrics</i> , 2011 , 52, 190-6	4	20
44	Cerebrospinal fluid biomarkers in Alzheimer's disease families with PSEN1 mutations. <i>Neurodegenerative Diseases</i> , 2011 , 8, 202-7	2.3	23
43	Clinical features and APOE genotype of pathologically proven early-onset Alzheimer disease. <i>Neurology</i> , 2011 , 76, 1720-5	6.5	91
42	New mutation in the PSEN1 (E120G) gene associated with early onset Alzheimer's disease. <i>Neurología (English Edition)</i> , 2010 , 25, 13-16	0.4	
41	Cognitive reserve modulates task-induced activations and deactivations in healthy elders, amnesic mild cognitive impairment and mild Alzheimer's disease. <i>Cortex</i> , 2010 , 46, 451-61	3.8	104
40	Genetic screening of Alzheimer's disease genes in Iberian and African samples yields novel mutations in presenilins and APP. <i>Neurobiology of Aging</i> , 2010 , 31, 725-31	5.6	162
39	Clinical, neuropathologic, and biochemical profile of the amyloid precursor protein I716F mutation. <i>Journal of Neuropathology and Experimental Neurology</i> , 2010 , 69, 53-9	3.1	37
38	Increased cortical thickness and caudate volume precede atrophy in PSEN1 mutation carriers. <i>Journal of Alzheimer's Disease</i> , 2010 , 22, 909-22	4.3	103
37	Nueva mutaci3n en el gen PSEN1 (E120G) asociada a enfermedad de Alzheimer de inicio precoz. <i>Neurología</i> , 2010 , 25, 13-16	1.4	7

36	The memory alteration test (M@T) discriminates between subjective memory complaints, mild cognitive impairment and Alzheimer's disease. <i>Archives of Gerontology and Geriatrics</i> , 2010 , 50, 171-4	4	29
35	A novel PSEN1 mutation (K239N) associated with Alzheimer's disease with wide range age of onset and slow progression. <i>European Journal of Neurology</i> , 2010 , 17, 994-6	6	17
34	Voxel based morphometry features and follow-up of amnesic patients at high risk for Alzheimer's disease conversion. <i>International Journal of Geriatric Psychiatry</i> , 2009 , 24, 875-84	3.9	15
33	Cerebrospinal fluid biomarkers in human genetic transmissible spongiform encephalopathies. <i>Journal of Neurology</i> , 2009 , 256, 1620-8	5.5	68
32	Interactions of cognitive reserve with regional brain anatomy and brain function during a working memory task in healthy elders. <i>Biological Psychology</i> , 2009 , 80, 256-9	3.2	65
31	Atrofia cortical posterior. Perfil neuropsicológico y diferencias con la enfermedad de Alzheimer típica. <i>Revista De Neurologia</i> , 2009 , 48, 178	24	3
30	Validez discriminativa y asociación del test minimal (MMSE) y del test de alteración de memoria (T@M) con una batería neuropsicológica en pacientes con deterioro cognitivo leve amnésico y enfermedad de Alzheimer. <i>Revista De Neurologia</i> , 2009 , 49, 169	24	4
29	Naming is associated with left temporal pole metabolite levels in neurodegenerative diseases. <i>Dementia and Geriatric Cognitive Disorders</i> , 2008 , 25, 212-7	2.6	9
28	First demonstrated de novo insertion in the prion protein gene in a young patient with dementia. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2008 , 79, 845-6	5.5	10
27	Molecular evidence of founder effects of fatal familial insomnia through SNP haplotypes around the D178N mutation. <i>Neurogenetics</i> , 2008 , 9, 109-18	3	12
26	Clinicopathological and genetic correlates of frontotemporal lobar degeneration and corticobasal degeneration. <i>Journal of Neurology</i> , 2008 , 255, 488-94	5.5	36
25	Screening for amnesic mild cognitive impairment and early Alzheimer's disease with M@T (Memory Alteration Test) in the primary care population. <i>International Journal of Geriatric Psychiatry</i> , 2007 , 22, 294-304	3.9	82
24	CSF analysis in patients with sporadic CJD and other transmissible spongiform encephalopathies. <i>European Journal of Neurology</i> , 2007 , 14, 121-4	6	53
23	A novel MAPT mutation (P301T) associated with familial frontotemporal dementia. <i>European Journal of Neurology</i> , 2007 , 14, e9-10	6	13
22	A novel mutation in the PSEN1 gene (L286P) associated with familial early-onset dementia of Alzheimer type and lobar haematomas. <i>European Journal of Neurology</i> , 2007 , 14, 1409-12	6	17
21	Late-onset frontotemporal dementia associated with a novel PGRN mutation. <i>Journal of Neural Transmission</i> , 2007 , 114, 1051-4	4.3	13
20	Brain tau expression and correlation with the H1/H1 tau genotype in frontotemporal lobar degeneration patients. <i>Journal of Neural Transmission</i> , 2007 , 114, 1585-8	4.3	6
19	Influence of timing on CSF tests value for Creutzfeldt-Jakob disease diagnosis. <i>Journal of Neurology</i> , 2007 , 254, 901-6	5.5	66

18	Cortical brain metabolism as measured by proton spectroscopy is related to memory performance in patients with amnesic mild cognitive impairment and Alzheimer's disease. <i>Dementia and Geriatric Cognitive Disorders</i> , 2007 , 24, 274-9	2.6	34
17	Longitudinal study of amnesic patients at high risk for Alzheimer's disease: clinical, neuropsychological and magnetic resonance spectroscopy features. <i>Dementia and Geriatric Cognitive Disorders</i> , 2007 , 24, 402-10	2.6	35
16	MAPT gene duplications are not a cause of frontotemporal lobar degeneration. <i>Neuroscience Letters</i> , 2007 , 424, 61-5	3.3	14
15	From progressive nonfluent aphasia to corticobasal syndrome: a case report of corticobasal degeneration. <i>Neurocase</i> , 2006 , 12, 355-9	0.8	22
14	Ancestral origins of the prion protein gene D178N mutation in the Basque Country. <i>Human Genetics</i> , 2005 , 117, 61-9	6.3	7
13	Discrepancies in the clinical utility of the 14-3-3 protein for the diagnosis of sporadic Creutzfeldt-Jakob disease. <i>Archives of Neurology</i> , 2004 , 61, 604		2
12	Clinical and genetic features of human prion diseases in Catalonia: 1993-2002. <i>European Journal of Neurology</i> , 2004 , 11, 649-55	6	22
11	CSF 14-3-3 protein assay and MRI as prognostic markers in patients with a clinically isolated syndrome suggestive of MS. <i>Journal of Neurology</i> , 2004 , 251, 1278-9	5.5	24
10	Anti-Tr antibodies as markers of paraneoplastic cerebellar degeneration and Hodgkin's disease. <i>Neurology</i> , 2003 , 60, 230-4	6.5	246
9	Analysis of the exon 1 polymorphism in the Tau gene in transmissible spongiform encephalopathies. <i>Journal of Neurology</i> , 2002 , 249, 938-9	5.5	4
8	Phosphorylated tau in cerebrospinal fluid as a marker for Creutzfeldt-Jakob disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2002 , 73, 79-81	5.5	30
7	14-3-3 Protein isoforms and atypical patterns of the 14-3-3 assay in the diagnosis of Creutzfeldt-Jakob disease. <i>Neuroscience Letters</i> , 2002 , 320, 69-72	3.3	24
6	14-3-3 protein in the CSF as prognostic marker in early multiple sclerosis. <i>Neurology</i> , 2001 , 57, 722-4	6.5	59
5	Autoantibodies against the prion protein in individuals with PRNP mutations		1
4	Brain functional network integrity sustains cognitive function despite atrophy in presymptomatic genetic frontotemporal dementia		2
3	AETIONOMY, a Cross-Sectional Study Aimed at validating a new taxonomy of Neurodegenerative Diseases: Study design and subject characteristics		2
2	Common variants in Alzheimer's disease: Novel association of six genetic variants with AD and risk stratification by polygenic risk scores		9
1	Analysis of brain atrophy and local gene expression in genetic frontotemporal dementia		2

