# Piero Parchi

## List of Publications by Year in Descending Order



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| 241 papers | 13,600 citations | 62 h-index | 111 g-index |
| 260 ext. papers | 15,647 ext. citations | 7.4 avg, IF | 5.81 L-index |

| # | Paper | IF | Citations |
|---|---|---|---|
| 241 | Prevalence Estimates of Amyloid Abnormality Across the Alzheimer Disease Clinical Spectrum.. *JAMA Neurology*, **2022**, | 17.2 | 9 |
| 240 | Validation of Revised International Creutzfeldt-Jakob Disease Surveillance Network Diagnostic Criteria for Sporadic Creutzfeldt-Jakob Disease.. *JAMA Network Open*, **2022**, 5, e2146319 | 10.4 | 3 |
| 239 | Identification of recurrent genetic patterns from targeted sequencing panels with advanced data science: a case-study on sporadic and genetic neurodegenerative diseases.. *BMC Medical Genomics*, **2022**, 15, 26 | 3.7 | 1 |
| 238 | Phenotypic Heterogeneity of Variably Protease-Sensitive Prionopathy: A Report of Three Cases Carrying Different Genotypes at Codon 129.. *Viruses*, **2022**, 14, | 6.2 | 2 |
| 237 | More Than Meets the Eye: "In Vivo" Diagnosis of Heidenhain Variant of Sporadic Creutzfeldt-Jakob Disease.. *Journal of Neuro-Ophthalmology*, **2022**, 42, e423-e426 | 2.6 | |
| 236 | PMCA-Based Detection of Prions in the Olfactory Mucosa of Patients With Sporadic Creutzfeldt-Jakob Disease.. *Frontiers in Aging Neuroscience*, **2022**, 14, 848991 | 5.3 | 0 |
| 235 | The Use of Real-Time Quaking-Induced Conversion for the Diagnosis of Human Prion Diseases.. *Frontiers in Aging Neuroscience*, **2022**, 14, 874734 | 5.3 | 0 |
| 234 | Homozygous R136S mutation in PRNP gene causes inherited early onset prion disease. *Alzheimer's Research and Therapy*, **2021**, 13, 176 | 9 | 2 |
| 233 | Homozygous R136S mutation in PRNP gene causes inherited early onset prion disease. *Alzheimer's Research and Therapy*, **2021**, 13, 176 | 9 | 1 |
| 232 | Plasma and CSF Neurofilament Light Chain in Amyotrophic Lateral Sclerosis: A Cross-Sectional and Longitudinal Study. *Frontiers in Aging Neuroscience*, **2021**, 13, 753242 | 5.3 | 4 |
| 231 | Neurofilament light chain and □synuclein RT-QuIC as differential diagnostic biomarkers in parkinsonisms and related syndromes. *Npj Parkinson's Disease*, **2021**, 7, 93 | 9.7 | 4 |
| 230 | The First Sporadic Creutzfeldt-Jakob Disease Case with a Rare Molecular Subtype VV1 and 1-Octapeptide Repeat Deletion in. *Viruses*, **2021**, 13, | 6.2 | 1 |
| 229 | Association between CSF alpha-synuclein seeding activity and genetic status in Parkinson's disease and dementia with Lewy bodies. *Acta Neuropathologica Communications*, **2021**, 9, 175 | 7.3 | 8 |
| 228 | Targeted sequencing panels in Italian ALS patients support different etiologies in the ALS/FTD continuum. *Journal of Neurology*, **2021**, 268, 3766-3776 | 5.5 | 1 |
| 227 | Early sensory disturbances and seizures are common manifestations of familial Creutzfeldt-Jakob disease due to E200K PRNP mutation: Case report from two Peruvian families. *Clinical Neurology and Neurosurgery*, **2021**, 202, 106490 | 2 | |
| 226 | Biomarkers and diagnostic guidelines for sporadic Creutzfeldt-Jakob disease. *Lancet Neurology, The* , **2021**, 20, 235-246 | 24.1 | 47 |
| 225 | LRP10 interacts with SORL1 in the intracellular vesicle trafficking pathway in non-neuronal brain cells and localises to Lewy bodies in Parkinson's disease and dementia with Lewy bodies. *Acta Neuropathologica*, **2021**, 142, 117-137 | 14.3 | 0 |

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| 224 | RT-QuIC Detection of Pathological α-Synuclein in Skin Punches of Patients with Lewy Body Disease. *Movement Disorders*, **2021**, 36, 2173-2177 | 7 | 14 |
| 223 | Diagnostic Value of the CSF α-Synuclein Real-Time Quaking-Induced Conversion Assay at the Prodromal MCI Stage of Dementia With Lewy Bodies. *Neurology*, **2021**, 97, e930-e940 | 6.5 | 9 |
| 222 | Phenotypic diversity of genetic Creutzfeldt-Jakob disease: a histo-molecular-based classification. *Acta Neuropathologica*, **2021**, 142, 707-728 | 14.3 | 5 |
| 221 | Characterization of novel progranulin gene variants in Italian patients with neurodegenerative diseases. *Neurobiology of Aging*, **2021**, 97, 145.e7-145.e15 | 5.6 | 1 |
| 220 | Diagnostic and prognostic performance of CSF α-synuclein in prion disease in the context of rapidly progressive dementia. *Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring*, **2021**, 13, e12214 | 5.2 | 0 |
| 219 | Cerebrospinal Fluid and Blood Neurofilament Light Chain Protein in Prion Disease and Other Rapidly Progressive Dementias: Current State of the Art. *Frontiers in Neuroscience*, **2021**, 15, 648743 | 5.1 | 4 |
| 218 | The In Vivo Diagnosis of Concomitant Alzheimer and Lewy Body Pathology: A Case Report. *Journal of Neuropathology and Experimental Neurology*, **2021**, | 3.1 | 1 |
| 217 | The clinical spectrum of multisystem proteinopathy: Data from a neurodegenerative cohort. *Journal of the Neurological Sciences*, **2021**, 426, 117478 | 3.2 | 2 |
| 216 | Cerebrospinal fluid biomarkers of neurodegeneration in narcolepsy type 1. *Sleep*, **2020**, 43, | 1.1 | 2 |
| 215 | Anterior Callosal Angle: A New Marker of Idiopathic Normal Pressure Hydrocephalus?. *World Neurosurgery*, **2020**, 139, e548-e552 | 2.1 | 6 |
| 214 | Genetic Creutzfeldt-Jakob disease in Sardinia: a case series linked to the PRNP R208H mutation due to a single founder effect. *Neurogenetics*, **2020**, 21, 251-257 | 3 | 2 |
| 213 | Detection of prions in skin punch biopsies of Creutzfeldt-Jakob disease patients. *Annals of Clinical and Translational Neurology*, **2020**, 7, 559-564 | 5.3 | 25 |
| 212 | Diagnostic-prognostic value and electrophysiological correlates of CSF biomarkers of neurodegeneration and neuroinflammation in amyotrophic lateral sclerosis. *Journal of Neurology*, **2020**, 267, 1699-1708 | 5.5 | 22 |
| 211 | CSF SerpinA1 in Creutzfeldt-Jakob disease and frontotemporal lobar degeneration. *Annals of Clinical and Translational Neurology*, **2020**, 7, 191-199 | 5.3 | 8 |
| 210 | CSF Ubiquitin Levels Are Higher in Alzheimer's Disease than in Frontotemporal Dementia and Reflect the Molecular Subtype in Prion Disease. *Biomolecules*, **2020**, 10, | 5.9 | 5 |
| 209 | Ultrasensitive RT-QuIC assay with high sensitivity and specificity for Lewy body-associated synucleinopathies. *Acta Neuropathologica*, **2020**, 140, 49-62 | 14.3 | 77 |
| 208 | Neurofilaments in motor neuron disorders: towards promising diagnostic and prognostic biomarkers. *Molecular Neurodegeneration*, **2020**, 15, 58 | 19 | 25 |
| 207 | Towards an improved early diagnosis of neurodegenerative diseases: the emerging role of in vitro conversion assays for protein amyloids. *Acta Neuropathologica Communications*, **2020**, 8, 117 | 7.3 | 22 |

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|---|---|---|---|
| 206 | Gerstmann-Str□ssler-Scheinker disease ( p.D202N) presenting with atypical parkinsonism. *Neurology: Genetics*, **2020**, 6, e400 | 3.8 | 3 |
| 205 | Comparison between plasma and cerebrospinal fluid biomarkers for the early diagnosis and association with survival in prion disease. *Journal of Neurology, Neurosurgery and Psychiatry*, **2020**, 91, 1181-1188 | 5.5 | 16 |
| 204 | Identification of novel risk loci and causal insights for sporadic Creutzfeldt-Jakob disease: a genome-wide association study. *Lancet Neurology, The*, **2020**, 19, 840-848 | 24.1 | 15 |
| 203 | Spatial Epidemiology of Sporadic Creutzfeldt-Jakob Disease in Apulia, Italy. *Neuroepidemiology*, **2020**, 54, 83-90 | 5.4 | 3 |
| 202 | 4-Repeat tau seeds and templating subtypes as brain and CSF biomarkers of frontotemporal lobar degeneration. *Acta Neuropathologica*, **2020**, 139, 63-77 | 14.3 | 49 |
| 201 | First case of an gene mutation causing frontotemporal dementia preceded by adult onset psychiatric symptoms. *Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration*, **2020**, 21, 467-469 | 3.6 | |
| 200 | □Synuclein RT-QuIC assay in cerebrospinal fluid of patients with dementia with Lewy bodies. *Annals of Clinical and Translational Neurology*, **2019**, 6, 2120-2126 | 5.3 | 42 |
| 199 | Diagnostic value of surrogate CSF biomarkers for Creutzfeldt-Jakob disease in the era of RT-QuIC. *Journal of Neurology*, **2019**, 266, 3136-3143 | 5.5 | 21 |
| 198 | Molecular Characterization of the Danish Prion Diseases Cohort With Special Emphasis on Rare and Unique Cases. *Journal of Neuropathology and Experimental Neurology*, **2019**, 78, 980-992 | 3.1 | 4 |
| 197 | Sudden cardiac death in a patient with LGI1 antibody-associated encephalitis. *Seizure: the Journal of the British Epilepsy Association*, **2019**, 65, 148-150 | 3.2 | 1 |
| 196 | Age at onset in genetic prion disease and the design of preventive clinical trials. *Neurology*, **2019**, 93, e125-e134 | 6.5 | 34 |
| 195 | A Novel Eight Octapeptide Repeat Insertion in PRNP Causing Prion Disease in a Danish Family. *Journal of Neuropathology and Experimental Neurology*, **2019**, 78, 595-604 | 3.1 | 5 |
| 194 | Predicting conversion from mild cognitive impairment to Alzheimer's disease using brain H-MRS and volumetric changes: A two- year retrospective follow-up study. *NeuroImage: Clinical*, **2019**, 23, 101843 | 5.3 | 19 |
| 193 | Clinicopathological features of the rare form of Creutzfeldt-Jakob disease in R208H-V129V PRNP carrier. *Acta Neuropathologica Communications*, **2019**, 7, 47 | 7.3 | 3 |
| 192 | Revisiting the Cerebrospinal Fluid Biomarker Profile in Idiopathic Normal Pressure Hydrocephalus: The Bologna Pro-Hydro Study. *Journal of Alzheimer's Disease*, **2019**, 68, 723-733 | 4.3 | 11 |
| 191 | The characterization of AD/PART co-pathology in CJD suggests independent pathogenic mechanisms and no cross-seeding between misfolded A□and prion proteins. *Acta Neuropathologica Communications*, **2019**, 7, 53 | 7.3 | 10 |
| 190 | Prion protein quantification in human cerebrospinal fluid as a tool for prion disease drug development. *Proceedings of the National Academy of Sciences of the United States of America*, **2019**, 116, 7793-7798 | 11.5 | 29 |
| 189 | Understanding Prion Strains: Evidence from Studies of the Disease Forms Affecting Humans. *Viruses*, **2019**, 11, | 6.2 | 27 |

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| 188 | A prospective evaluation of clinical and instrumental features before and after ventriculo-peritoneal shunt in patients with idiopathic Normal pressure hydrocephalus: The Bologna PRO-Hydro study. *Parkinsonism and Related Disorders*, **2019**, 66, 117-124 | 3.6 | 14 |
| 187 | CSF biomarkers of neuroinflammation in distinct forms and subtypes of neurodegenerative dementia. *Alzheimer's Research and Therapy*, **2019**, 12, 2 | 9 | 44 |
| 186 | Two distinct prions in fatal familial insomnia and its sporadic form. *Brain Communications*, **2019**, 1, fcz045 | 4.5 | 8 |
| 185 | Large expert-curated database for benchmarking document similarity detection in biomedical literature search. *Database: the Journal of Biological Databases and Curation*, **2019**, 2019, | 5 | 4 |
| 184 | Domain-specific Quantification of Prion Protein in Cerebrospinal Fluid by Targeted Mass Spectrometry. *Molecular and Cellular Proteomics*, **2019**, 18, 2388-2400 | 7.6 | 17 |
| 183 | Analysis of RNA Expression Profiles Identifies Dysregulated Vesicle Trafficking Pathways in Creutzfeldt-Jakob Disease. *Molecular Neurobiology*, **2019**, 56, 5009-5024 | 6.2 | 8 |
| 182 | Variable Protease-Sensitive Prionopathy Transmission to Bank Voles. *Emerging Infectious Diseases*, **2019**, 25, 73-81 | 10.2 | 13 |
| 181 | Recent advances in the histo-molecular pathology of human prion disease. *Brain Pathology*, **2019**, 29, 278-300 | 6 | 35 |
| 180 | Prion-related peripheral neuropathy in sporadic Creutzfeldt-Jakob disease. *Journal of Neurology, Neurosurgery and Psychiatry*, **2019**, 90, 424-427 | 5.5 | 20 |
| 179 | Antemortem CSF A42/A40 ratio predicts Alzheimer's disease pathology better than A42 in rapidly progressive dementias. *Annals of Clinical and Translational Neurology*, **2019**, 6, 263-273 | 5.3 | 19 |
| 178 | The First Historically Reported Italian Family with FTD/ALS Teaches a Lesson on C9orf72 RE: Clinical Heterogeneity and Oligogenic Inheritance. *Journal of Alzheimer's Disease*, **2018**, 62, 687-697 | 4.3 | 4 |
| 177 | Identification of rare genetic variants in Italian patients with dementia by targeted gene sequencing. *Neurobiology of Aging*, **2018**, 66, 180.e23-180.e31 | 5.6 | 15 |
| 176 | An in vivo C-PK PET study of microglia activation in Fatal Familial Insomnia. *Annals of Clinical and Translational Neurology*, **2018**, 5, 11-18 | 5.3 | 8 |
| 175 | Cerebral Mitochondrial Microangiopathy Leads to Leukoencephalopathy in Mitochondrial Neurogastrointestinal Encephalopathy. *American Journal of Neuroradiology*, **2018**, 39, 427-434 | 4.4 | 13 |
| 174 | Regional pattern of microgliosis in sporadic Creutzfeldt-Jakob disease in relation to phenotypic variants and disease progression. *Neuropathology and Applied Neurobiology*, **2018**, 44, 574-589 | 5.2 | 11 |
| 173 | Rapidly Progressive Alzheimer's Disease: Contributions to Clinical-Pathological Definition and Diagnosis. *Journal of Alzheimer's Disease*, **2018**, 63, 887-897 | 4.3 | 11 |
| 172 | Two novel truncating mutations broaden the spectrum of prion amyloidosis. *Annals of Clinical and Translational Neurology*, **2018**, 5, 777-783 | 5.3 | 11 |
| 171 | Age at onset of genetic (E200K) and sporadic Creutzfeldt-Jakob diseases is modulated by the gene. *Journal of Neurology, Neurosurgery and Psychiatry*, **2018**, 89, 1243-1249 | 5.5 | 9 |

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| 170 | Sporadic Fatal Insomnia in Europe: Phenotypic Features and Diagnostic Challenges. *Annals of Neurology*, **2018**, 84, 347-360 | 9.4 | 18 |
| 169 | A novel prion protein gene-truncating mutation causing autonomic neuropathy and diarrhea. *European Journal of Neurology*, **2018**, 25, e91-e92 | 6 | 8 |
| 168 | Unusual Clinical Presentations Challenging the Early Clinical Diagnosis of Creutzfeldt-Jakob Disease. *Journal of Alzheimer's Disease*, **2018**, 64, 1051-1065 | 4.3 | 24 |
| 167 | The CSF neurofilament light signature in rapidly progressive neurodegenerative dementias. *Alzheimer's Research and Therapy*, **2018**, 10, 3 | 9 | 57 |
| 166 | Iatrogenic Creutzfeldt-Jakob disease with Amyloid-β pathology: an international study. *Acta Neuropathologica Communications*, **2018**, 6, 5 | 7.3 | 55 |
| 165 | Familial human prion diseases associated with prion protein mutations Y226X and G131V are transmissible to transgenic mice expressing human prion protein. *Acta Neuropathologica Communications*, **2018**, 6, 13 | 7.3 | 13 |
| 164 | LRP10 genetic variants in familial Parkinson's disease and dementia with Lewy bodies: a genome-wide linkage and sequencing study. *Lancet Neurology, The*, **2018**, 17, 597-608 | 24.1 | 68 |
| 163 | Sporadic Creutzfeldt-Jakob disease. *Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn*, **2018**, 153, 155-174 | 3 | 49 |
| 162 | Clinical Reasoning: Rapidly progressive dementia in a patient with HIV after an exotic journey. *Neurology*, **2018**, 91, e1360-e1364 | 6.5 | 1 |
| 161 | Variably protease-sensitive prionopathy presenting within ALS/FTD spectrum. *Annals of Clinical and Translational Neurology*, **2018**, 5, 1297-1302 | 5.3 | 8 |
| 160 | Cerebrospinal Fluid Biomarkers in Patients with Frontotemporal Dementia Spectrum: A Single-Center Study. *Journal of Alzheimer's Disease*, **2018**, 66, 551-563 | 4.3 | 29 |
| 159 | Prion-specific and surrogate CSF biomarkers in Creutzfeldt-Jakob disease: diagnostic accuracy in relation to molecular subtypes and analysis of neuropathological correlates of p-tau and Aβ42 levels. *Acta Neuropathologica*, **2017**, 133, 559-578 | 14.3 | 95 |
| 158 | Multiple variants in families with amyotrophic lateral sclerosis and frontotemporal dementia related to C9orf72 repeat expansion: further observations on their oligogenic nature. *Journal of Neurology*, **2017**, 264, 1426-1433 | 5.5 | 22 |
| 157 | Muscle ceroid lipofuscin-like deposits in a patient with corticobasal syndrome due to a progranulin mutation. *Movement Disorders*, **2017**, 32, 1259-1260 | 7 | 2 |
| 156 | Atypical Creutzfeldt-Jakob disease with PrP-amyloid plaques in white matter: molecular characterization and transmission to bank voles show the M1 strain signature. *Acta Neuropathologica Communications*, **2017**, 5, 87 | 7.3 | 17 |
| 155 | Diagnostic Accuracy of a Combined Analysis of Cerebrospinal Fluid t-PrP, t-tau, p-tau, and Aβ42 in the Differential Diagnosis of Creutzfeldt-Jakob Disease from Alzheimer's Disease with Emphasis on Atypical Disease Variants. *Journal of Alzheimer's Disease*, **2017**, 55, 1471-1480 | 4.3 | 30 |
| 154 | High diagnostic value of second generation CSF RT-QuIC across the wide spectrum of CJD prions. *Scientific Reports*, **2017**, 7, 10655 | 4.9 | 105 |
| 153 | Towards an early clinical diagnosis of sporadic CJD VV2 (ataxic type). *Journal of Neurology, Neurosurgery and Psychiatry*, **2017**, 88, 764-772 | 5.5 | 22 |

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| 152 | Patient with rapidly evolving neurological disease with neuropathological lesions of Creutzfeldt-Jakob disease, Lewy body dementia, chronic subcortical vascular encephalopathy and meningothelial meningioma. *Neuropathology*, **2017**, 37, 110-115 | 2 | 4 |
| 151 | Iodine-123-meta-iodobenzylguanidine Myocardial Scintigraphy in Isolated Autonomic Failure: Potential Red Flag for Future Multiple System Atrophy. *Frontiers in Neurology*, **2017**, 8, 225 | 4.1 | 6 |
| 150 | Quantifying prion disease penetrance using large population control cohorts. *Science Translational Medicine*, **2016**, 8, 322ra9 | 17.5 | 205 |
| 149 | Neuropathological and biochemical criteria to identify acquired Creutzfeldt-Jakob disease among presumed sporadic cases. *Neuropathology*, **2016**, 36, 305-10 | 2 | 17 |
| 148 | Distinctive properties of plaque-type dura mater graft-associated Creutzfeldt-Jakob disease in cell-protein misfolding cyclic amplification. *Laboratory Investigation*, **2016**, 96, 581-7 | 5.9 | 13 |
| 147 | Revisiting the Heidenhain Variant of Creutzfeldt-Jakob Disease: Evidence for Prion Type Variability Influencing Clinical Course and Laboratory Findings. *Journal of Alzheimerʼs Disease*, **2016**, 50, 465-76 | 4.3 | 42 |
| 146 | Analysis of Conformational Stability of Abnormal Prion Protein Aggregates across the Spectrum of Creutzfeldt-Jakob Disease Prions. *Journal of Virology*, **2016**, 90, 6244-6254 | 6.6 | 27 |
| 145 | Cerebrospinal fluid real-time quaking-induced conversion is a robust and reliable test for sporadic creutzfeldt-jakob disease: An international study. *Annals of Neurology*, **2016**, 80, 160-5 | 9.4 | 89 |
| 144 | Messenger RNA processing is altered in autosomal dominant leukodystrophy. *Human Molecular Genetics*, **2015**, 24, 2746-56 | 5.6 | 21 |
| 143 | Transmission properties of atypical Creutzfeldt-Jakob disease: a clue to disease etiology?. *Journal of Virology*, **2015**, 89, 3939-46 | 6.6 | 22 |
| 142 | Creutzfeldt-Jakob disease masked by head trauma and features of Wilson's disease. *International Journal of Neuroscience*, **2015**, 125, 312-4 | 2 | |
| 141 | Neuropathological assessments of the pathology in frontotemporal lobar degeneration with TDP43-positive inclusions: an inter-laboratory study by the BrainNet Europe consortium. *Journal of Neural Transmission*, **2015**, 122, 957-72 | 4.3 | 19 |
| 140 | Creutzfeldt-Jakob disease manifesting as stroke mimic in a 78-year-old patient: pitfalls and tips in the diagnosis. *Journal of the Neurological Sciences*, **2014**, 346, 343-4 | 3.2 | 7 |
| 139 | Mutant PrPCJD prevails over wild-type PrPCJD in the brain of V210I and R208H genetic Creutzfeldt-Jakob disease patients. *Biochemical and Biophysical Research Communications*, **2014**, 454, 289-94 | 3.4 | 6 |
| 138 | A CTNNA3 compound heterozygous deletion implicates a role for α-catenin in susceptibility to autism spectrum disorder. *Journal of Neurodevelopmental Disorders*, **2014**, 6, 17 | 4.6 | 32 |
| 137 | Wait and see: a 5 year history of 'recurrent dementia'. *BMJ Case Reports*, **2014**, 2014, | 0.9 | |
| 136 | Primary progressive narcolepsy type 1: the other side of the coin. *Neurology*, **2014**, 83, 2189-90 | 6.5 | 40 |
| 135 | Narcolepsy is a common phenotype in HSAN IE and ADCA-DN. *Brain*, **2014**, 137, 1643-55 | 11.2 | 36 |

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| 134 | Pearls & Oy-sters: rapidly progressive dementia: prions or immunomediated?. *Neurology*, **2014**, 82, e149-52 | 8.3 | 3 |
| 133 | Gait disorders in fatal familial insomnia. *Movement Disorders*, **2014**, 29, 420-4 | 7 | 18 |
| 132 | Maternally inherited genetic variants of CADPS2 are present in autism spectrum disorders and intellectual disability patients. *EMBO Molecular Medicine*, **2014**, 6, 795-809 | 12 | 28 |
| 131 | R208H-129VV haplotype in the prion protein gene: phenotype and neuroimaging of a patient with genetic Creutzfeldt-Jakob disease. *Journal of Neurology*, **2013**, 260, 2650-2 | 5.5 | 9 |
| 130 | Atypical neuropathological sCJD-MM phenotype with abundant white matter Kuru-type plaques sparing the cerebellar cortex. *Neuropathology*, **2013**, 33, 204-8 | 2 | 6 |
| 129 | CSF biomarker variability in the Alzheimer's Association quality control program. *Alzheimer's and Dementia*, **2013**, 9, 251-61 | 1.2 | 289 |
| 128 | Prion protein misfolding, strains, and neurotoxicity: an update from studies on Mammalian prions. *International Journal of Cell Biology*, **2013**, 2013, 910314 | 2.6 | 37 |
| 127 | Analyses of protease resistance and aggregation state of abnormal prion protein across the spectrum of human prions. *Journal of Biological Chemistry*, **2013**, 288, 27972-85 | 5.4 | 30 |
| 126 | Divergent clinical and neuropathological phenotype in a Gerstmann-Strüssler-Scheinker P102L family. *Acta Neurologica Scandinavica*, **2012**, 126, 315-23 | 3.8 | 11 |
| 125 | Selection of novel reference genes for use in the human central nervous system: a BrainNet Europe Study. *Acta Neuropathologica*, **2012**, 124, 893-903 | 14.3 | 79 |
| 124 | Consensus classification of human prion disease histotypes allows reliable identification of molecular subtypes: an inter-rater study among surveillance centres in Europe and USA. *Acta Neuropathologica*, **2012**, 124, 517-29 | 14.3 | 148 |
| 123 | The need to unify neuropathological assessments of vascular alterations in the ageing brain: multicentre survey by the BrainNet Europe consortium. *Experimental Gerontology*, **2012**, 47, 825-33 | 4.5 | 50 |
| 122 | Creutzfeldt-Jakob disease: an under-recognized cause of dementia. *Journal of the American Geriatrics Society*, **2012**, 60, 156-7 | 5.6 | 1 |
| 121 | Human prion diseases in the Netherlands (1998-2009): clinical, genetic and molecular aspects. *PLoS ONE*, **2012**, 7, e36333 | 3.7 | 37 |
| 120 | Molecular pathology, classification, and diagnosis of sporadic human prion disease variants **2012**, 50, 20-45 | | 32 |
| 119 | A second case of Gerstmann-Strüssler-Scheinker disease linked to the G131V mutation in the prion protein gene in a Dutch patient. *Journal of Neuropathology and Experimental Neurology*, **2011**, 70, 698-702 | 3.1 | 18 |
| 118 | The first case of fatal familial insomnia (FFI) in the Netherlands: a patient from Egyptian descent with concurrent four repeat tau deposits. *Neuropathology and Applied Neurobiology*, **2011**, 37, 549-53 | 5.2 | 10 |
| 117 | New lexicon and criteria for the diagnosis of Alzheimer's disease. *Lancet Neurology, The*, **2011**, 10, 298-9; author reply 300-1 | 24.1 | 19 |

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| 116 | A novel seven-octapeptide repeat insertion in the prion protein gene (PRNP) in a Dutch pedigree with Gerstmann-Str□ssler-Scheinker disease phenotype: comparison with similar cases from the literature. *Acta Neuropathologica*, **2011**, 121, 59-68 | 14.3 | 24 |
| 115 | Genetic Creutzfeldt-Jakob disease and fatal familial insomnia: insights into phenotypic variability and disease pathogenesis. *Acta Neuropathologica*, **2011**, 121, 21-37 | 14.3 | 99 |
| 114 | Phenotypic variability of sporadic human prion disease and its molecular basis: past, present, and future. *Acta Neuropathologica*, **2011**, 121, 91-112 | 14.3 | 106 |
| 113 | Multiorgan detection and characterization of protease-resistant prion protein in a case of variant CJD examined in the United States. *PLoS ONE*, **2010**, 5, e8765 | 3.7 | 46 |
| 112 | PrP conformational transitions alter species preference of a PrP-specific antibody. *Journal of Biological Chemistry*, **2010**, 285, 13874-84 | 5.4 | 49 |
| 111 | Reply to Kascsak: Definition of the PrP 3F4 Epitope Revisited. *Journal of Biological Chemistry*, **2010**, 285, le6 | 5.4 | 78 |
| 110 | Agent strain variation in human prion disease: insights from a molecular and pathological review of the National Institutes of Health series of experimentally transmitted disease. *Brain*, **2010**, 133, 3030-42 | 11.2 | 61 |
| 109 | Cathepsin D (C224T) polymorphism in sporadic and genetic Creutzfeldt-Jakob disease. *Alzheimer Disease and Associated Disorders*, **2010**, 24, 104-7 | 2.5 | 9 |
| 108 | Prion protein amyloidosis with divergent phenotype associated with two novel nonsense mutations in PRNP. *Acta Neuropathologica*, **2010**, 119, 189-97 | 14.3 | 101 |
| 107 | An atypical phenotype of CJD associated with the E200K mutation in the prion protein gene. *Neurological Sciences*, **2010**, 31, 837-9 | 3.5 | 4 |
| 106 | Variably protease-sensitive prionopathy: a new sporadic disease of the prion protein. *Annals of Neurology*, **2010**, 68, 162-72 | 9.4 | 168 |
| 105 | 'Agrypnia excitata' in a case of sporadic Creutzfeldt-Jakob disease VV2. *Journal of Neurology, Neurosurgery and Psychiatry*, **2009**, 80, 244-6 | 5.5 | 15 |
| 104 | Pathologic correlates of diffusion MRI changes in Creutzfeldt-Jakob disease. *Neurology*, **2009**, 72, 1425-31 | 8.15 | 67 |
| 103 | Co-existence of scrapie prion protein types 1 and 2 in sporadic Creutzfeldt-Jakob disease: its effect on the phenotype and prion-type characteristics. *Brain*, **2009**, 132, 2643-58 | 11.2 | 111 |
| 102 | Inherited Creutzfeldt-Jakob disease in a Dutch patient with a novel five octapeptide repeat insertion and unusual cerebellar morphology. *Journal of Neurology, Neurosurgery and Psychiatry*, **2009**, 80, 1386-9 | 5.5 | 12 |
| 101 | Magnetic resonance diagnostic markers in clinically sporadic prion disease: a combined brain magnetic resonance imaging and spectroscopy study. *Brain*, **2009**, 132, 2669-79 | 11.2 | 37 |
| 100 | Creutzfeldt-Jakob disease with E200K PRNP mutation: a case report and revision of the literature. *Neurological Sciences*, **2009**, 30, 417-20 | 3.5 | 13 |
| 99 | Assessment of beta-amyloid deposits in human brain: a study of the BrainNet Europe Consortium. *Acta Neuropathologica*, **2009**, 117, 309-20 | 14.3 | 119 |

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| 98 | Staging/typing of Lewy body related alpha-synuclein pathology: a study of the BrainNet Europe Consortium. *Acta Neuropathologica*, **2009**, 117, 635-52 | 14.3 | 189 |
| 97 | Incidence and spectrum of sporadic Creutzfeldt-Jakob disease variants with mixed phenotype and co-occurrence of PrPSc types: an updated classification. *Acta Neuropathologica*, **2009**, 118, 659-71 | 14.3 | 184 |
| 96 | A case of fatal familial insomnia in Africa. *Journal of Neurology*, **2009**, 256, 1778-9 | 5.5 | 10 |
| 95 | Striatal [123I] FP-CIT SPECT demonstrates dopaminergic deficit in a sporadic case of Creutzfeldt-Jakob disease. *Acta Neurologica Scandinavica*, **2009**, 119, 131-4 | 3.8 | 19 |
| 94 | Inter-laboratory assessment of PrPSc typing in creutzfeldt-jakob disease: a Western blot study within the NeuroPrion Consortium. *Brain Pathology*, **2009**, 19, 384-91 | 6 | 49 |
| 93 | Staging of neurofibrillary pathology in Alzheimer's disease: a study of the BrainNet Europe Consortium. *Brain Pathology*, **2008**, 18, 484-96 | 6 | 270 |
| 92 | Characterization of truncated forms of abnormal prion protein in Creutzfeldt-Jakob disease. *Journal of Biological Chemistry*, **2008**, 283, 30557-65 | 5.4 | 67 |
| 91 | Diffusion-weighted brain imaging study of patients with clinical diagnosis of corticobasal degeneration, progressive supranuclear palsy and Parkinson's disease. *Brain*, **2008**, 131, 2690-700 | 11.2 | 114 |
| 90 | Mixed brain pathologies in dementia: the BrainNet Europe consortium experience. *Dementia and Geriatric Cognitive Disorders*, **2008**, 26, 343-50 | 2.6 | 119 |
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