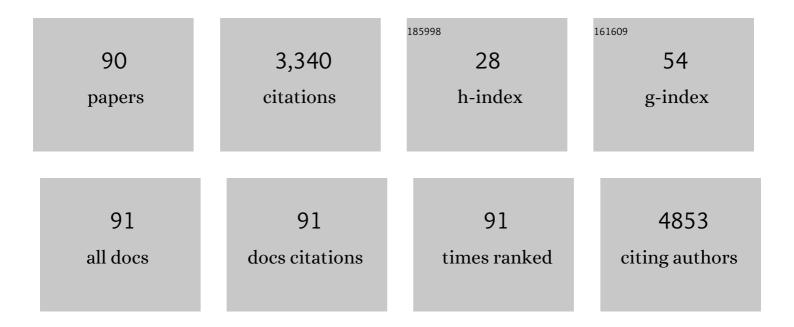
## Giorgio G Fumagalli

List of Publications by Year in descending order

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| #  | Article   | IF  | CITATIONS |
|----|---|-----|-----------|
| 1  | A modified Camel and Cactus Test detects presymptomatic semantic impairment in genetic<br>frontotemporal dementia within the GENFI cohort. Applied Neuropsychology Adult, 2022, 29, 112-119.  | 0.7 | 18        |
| 2  | A data-driven disease progression model of fluid biomarkers in genetic frontotemporal dementia.<br>Brain, 2022, 145, 1805-1817.   | 3.7 | 27        |
| 3  | Stratifying the Presymptomatic Phase of Genetic Frontotemporal Dementia by Serum <scp>NfL</scp><br>and <scp>pNfH</scp> : A Longitudinal Multicentre Study. Annals of Neurology, 2022, 91, 33-47.  | 2.8 | 21        |
| 4  | Caregiver Tele-Assistance for Reduction of Emotional Distress During the COVID-19 Pandemic.<br>Psychological Support to Caregivers of People with Dementia: The Italian Experience. Journal of<br>Alzheimer's Disease, 2022, 85, 1045-1052. | 1.2 | 7         |
| 5  | Association of Superficial White Matter Alterations with Cerebrospinal Fluid Biomarkers and<br>Cognitive Decline in Neurodegenerative Dementia. Journal of Alzheimer's Disease, 2022, 85, 431-442.  | 1.2 | 2         |
| 6  | Cognitive composites for genetic frontotemporal dementia: GENFI-Cog. Alzheimer's Research and Therapy, 2022, 14, 10.  | 3.0 | 4         |
| 7  | Unravelling the Association Between Amyloid-PET and Cerebrospinal Fluid Biomarkers in the<br>Alzheimer's Disease Spectrum: Who Really Deserves an A+?. Journal of Alzheimer's Disease, 2022, 85,<br>1009-1020.                              | 1.2 | 5         |
| 8  | Examining empathy deficits across familial forms of frontotemporal dementia within the GENFI cohort. Cortex, 2022, 150, 12-28.  | 1.1 | 2         |
| 9  | Amyloid PET imaging and dementias: potential applications in detecting and quantifying early white matter damage. Alzheimer's Research and Therapy, 2022, 14, 33.   | 3.0 | 9         |
| 10 | Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia.<br>Alzheimer's and Dementia, 2022, 18, 1408-1423.   | 0.4 | 24        |
| 11 | Structural brain splitting is a hallmark of Granulin-related frontotemporal dementia. Neurobiology of Aging, 2022, , .  | 1.5 | 1         |
| 12 | The <scp>CBIâ€R</scp> detects early behavioural impairment in genetic frontotemporal dementia. Annals of Clinical and Translational Neurology, 2022, 9, 644-658.  | 1.7 | 1         |
| 13 | Brain functional network integrity sustains cognitive function despite atrophy in presymptomatic genetic frontotemporal dementia. Alzheimer's and Dementia, 2021, 17, 500-514.  | 0.4 | 36        |
| 14 | Impairment of episodic memory in genetic frontotemporal dementia: A GENFI study. Alzheimer's and<br>Dementia: Diagnosis, Assessment and Disease Monitoring, 2021, 13, e12185.   | 1.2 | 11        |
| 15 | Facing the digital divide into a dementia clinic during COVID-19 pandemic: caregiver age matters.<br>Neurological Sciences, 2021, 42, 1247-1251.  | 0.9 | 47        |
| 16 | Detection of the SQSTM1 Mutation in a Patient with Early-Onset Hippocampal Amnestic Syndrome.<br>Journal of Alzheimer's Disease, 2021, 79, 477-481.   | 1.2 | 2         |
| 17 | Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic<br>Frontotemporal Dementia. JAMA Network Open, 2021, 4, e2030194.   | 2.8 | 42        |
| 18 | Diogenes syndrome in dementia: a case report. BJPsych Open, 2021, 7, e43.   | 0.3 | 0         |

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|----|--|-----|-----------|
| 19 | Analysis of C9orf72 Intermediate Alleles in a Retrospective Cohort of Neurological Patients: Risk<br>Factors for Alzheimer's Disease?. Journal of Alzheimer's Disease, 2021, 81, 1445-1451.      | 1.2 | 6         |
| 20 | The Revised Self-Monitoring Scale detects early impairment of social cognition in genetic frontotemporal dementia within the GENFI cohort. Alzheimer's Research and Therapy, 2021, 13, 127.      | 3.0 | 12        |
| 21 | Niemann-Pick Type C 1 (NPC1) and NPC2 Gene Variability in Demented Patients with Evidence of Brain<br>Amyloid Deposition. Journal of Alzheimer's Disease, 2021, 83, 1313-1323.                   | 1.2 | 5         |
| 22 | Differential early subcortical involvement in genetic FTD within the GENFI cohort. NeuroImage: Clinical, 2021, 30, 102646.   | 1.4 | 28        |
| 23 | Disease-related cortical thinning in presymptomatic granulin mutation carriers. NeuroImage: Clinical, 2021, 29, 102540.  | 1.4 | 8         |
| 24 | Unravelling the association between amyloid-pet and CSF biomarkers: Who really deserves an A +?.<br>Journal of the Neurological Sciences, 2021, 429, 117853.                                     | 0.3 | 0         |
| 25 | A panel of CSF proteins separates genetic frontotemporal dementia from presymptomatic mutation carriers: a GENFI study. Molecular Neurodegeneration, 2021, 16, 79.                               | 4.4 | 9         |
| 26 | Low CSF β-amyloid levels predict early regional grey matter atrophy in multiple sclerosis. Multiple<br>Sclerosis and Related Disorders, 2020, 39, 101899.  | 0.9 | 5         |
| 27 | 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.                   | 4.9 | 175       |
| 28 | Parieto-occipital sulcus widening differentiates posterior cortical atrophy from typical Alzheimer<br>disease. NeuroImage: Clinical, 2020, 28, 102453.   | 1.4 | 11        |
| 29 | Phenotypic heterogeneity of the rare R377W PSEN1 mutation: Lateâ€onset presentation with mixed<br>Alzheimer's and frontotemporal dementia features. Alzheimer's and Dementia, 2020, 16, e042581. | 0.4 | Ο         |
| 30 | Alemtuzumab in multiple sclerosis during the COVID-19 pandemic: A mild uncomplicated infection despite intense immunosuppression. Multiple Sclerosis Journal, 2020, 26, 1268-1269.               | 1.4 | 35        |
| 31 | MiRNA Profiling in Plasma Neural-Derived Small Extracellular Vesicles from Patients with Alzheimer's<br>Disease. Cells, 2020, 9, 1443.   | 1.8 | 60        |
| 32 | Evidence of retinal anterograde neurodegeneration in the very early stages of multiple sclerosis: a<br>longitudinal OCT study. Neurological Sciences, 2020, 41, 3175-3183.                       | 0.9 | 16        |
| 33 | Crossing Borders Between Frontotemporal Dementia and Psychiatric Disorders: An Updated Overview.<br>Journal of Alzheimer's Disease, 2020, 75, 661-673.   | 1.2 | 3         |
| 34 | Faster Cortical Thinning and Surface Area Loss in Presymptomatic and Symptomatic <i>C9orf72</i> Repeat Expansion Adult Carriers. Annals of Neurology, 2020, 88, 113-122.                         | 2.8 | 19        |
| 35 | Social cognition impairment in genetic frontotemporal dementia within the GENFI cohort. Cortex, 2020, 133, 384-398.  | 1.1 | 26        |
| 36 | CSF β-amyloid predicts prognosis in patients with multiple sclerosis. Multiple Sclerosis Journal, 2019, 25, 1223-1231.   | 1.4 | 19        |

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|----|--|-----|-----------|
| 37 | Testing the 2018 NIA-AA research framework in a retrospective large cohort of patients with cognitive impairment: from biological biomarkers to clinical syndromes. Alzheimer's Research and Therapy, 2019, 11, 84.                        | 3.0 | 28        |
| 38 | Serum neurofilament light chain in genetic frontotemporal dementia: a longitudinal, multicentre<br>cohort study. Lancet Neurology, The, 2019, 18, 1103-1111.   | 4.9 | 128       |
| 39 | Monozygotic Twins with Frontotemporal Dementia Due To Thr272fs GRN Mutation Discordant for Age<br>At Onset. Journal of Alzheimer's Disease, 2019, 67, 1173-1179.   | 1.2 | 4         |
| 40 | Inflammatory expression profile in peripheral blood mononuclear cells from patients with<br>Nasu-Hakola Disease. Cytokine, 2019, 116, 115-119.   | 1.4 | 6         |
| 41 | The inner fluctuations of the brain in presymptomatic Frontotemporal Dementia: The chronnectome fingerprint. Neurolmage, 2019, 189, 645-654.   | 2.1 | 33        |
| 42 | Cerebrospinal Fluid Level of Aquaporin4: A New Window on Glymphatic System Involvement in Neurodegenerative Disease?. Journal of Alzheimer's Disease, 2019, 69, 663-669.   | 1.2 | 21        |
| 43 | Cerebral perfusion changes in presymptomatic genetic frontotemporal dementia: a GENFI study. Brain, 2019, 142, 1108-1120.  | 3.7 | 41        |
| 44 | White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal<br>GENFI study. NeuroImage: Clinical, 2019, 24, 102077.  | 1.4 | 27        |
| 45 | Spatiotemporal analysis for detection of pre-symptomatic shape changes in neurodegenerative diseases: Initial application to the GENFI cohort. NeuroImage, 2019, 188, 282-290.   | 2.1 | 16        |
| 46 | Functional network resilience to pathology in presymptomatic genetic frontotemporal dementia.<br>Neurobiology of Aging, 2019, 77, 169-177.   | 1.5 | 47        |
| 47 | Amyloid PET as a marker of normal-appearing white matter early damage in multiple sclerosis:<br>correlation with CSF β-amyloid levels and brain volumes. European Journal of Nuclear Medicine and<br>Molecular Imaging, 2019, 46, 280-287. | 3.3 | 28        |
| 48 | The loss of macular ganglion cells begins from the early stages of disease and correlates with brain atrophy in multiple sclerosis patients. Multiple Sclerosis Journal, 2019, 25, 31-38.  | 1.4 | 39        |
| 49 | Drug Prescription and Delirium in Older Inpatients. Journal of Clinical Psychiatry, 2019, 80, .  | 1.1 | 16        |
| 50 | Profiling of Specific Gene Expression Pathways in Peripheral Cells from Prodromal Alzheimer's<br>Disease Patients. Journal of Alzheimer's Disease, 2018, 61, 1289-1294.  | 1.2 | 2         |
| 51 | Potential genetic modifiers of disease risk and age at onset in patients with frontotemporal lobar<br>degeneration and GRN mutations: a genome-wide association study. Lancet Neurology, The, 2018, 17,<br>548-558.                        | 4.9 | 97        |
| 52 | Diagnosis of Frontotemporal Dementia. , 2018, , 113-121.   |     | 0         |
| 53 | Comparison of arterial spin labeling registration strategies in the multiâ€center GENetic<br>frontotemporal dementia initiative (GENFI). Journal of Magnetic Resonance Imaging, 2018, 47, 131-140.   | 1.9 | 41        |
| 54 | CSF β-amyloid and white matter damage: a new perspective on Alzheimer's disease. Journal of Neurology,<br>Neurosurgery and Psychiatry, 2018, 89, 352-357.  | 0.9 | 36        |

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|----|---|-----|-----------|
| 55 | Patterns of gray matter atrophy in genetic frontotemporal dementia: results from the GENFI study.<br>Neurobiology of Aging, 2018, 62, 191-196.  | 1.5 | 151       |
| 56 | Progranulin plasma levels predict the presence of GRN mutations in asymptomatic subjects and do not correlate with brain atrophy: results from the GENFI study. Neurobiology of Aging, 2018, 62, 245.e9-245.e12.  | 1.5 | 40        |
| 57 | Behavioral and Neurophysiological Effects of Transcranial Direct Current Stimulation (tDCS) in<br>Fronto-Temporal Dementia. Frontiers in Behavioral Neuroscience, 2018, 12, 235.  | 1.0 | 19        |
| 58 | Uncovering the heterogeneity and temporal complexity of neurodegenerative diseases with Subtype and Stage Inference. Nature Communications, 2018, 9, 4273.  | 5.8 | 263       |
| 59 | Distinct patterns of brain atrophy in Genetic Frontotemporal Dementia Initiative (GENFI) cohort revealed by visual rating scales. Alzheimer's Research and Therapy, 2018, 10, 46.   | 3.0 | 34        |
| 60 | Presymptomatic white matter integrity loss in familial frontotemporal dementia in the<br><scp>GENFI</scp> cohort: A crossâ€sectional diffusion tensor imaging study. Annals of Clinical and<br>Translational Neurology, 2018, 5, 1025-1036.             | 1.7 | 39        |
| 61 | Cognitive reserve and TMEM106B genotype modulate brain damage in presymptomatic frontotemporal dementia: a GENFI study. Brain, 2017, 140, 1784-1791.  | 3.7 | 55        |
| 62 | White matter hyperintensities are seen only in GRN mutation carriers in the GENFI cohort.<br>NeuroImage: Clinical, 2017, 15, 171-180.   | 1.4 | 63        |
| 63 | Evidence of CNS β-amyloid deposition in Nasu-Hakola disease due to the <i>TREM2</i> Q33X mutation.<br>Neurology, 2017, 89, 2503-2505.   | 1.5 | 26        |
| 64 | Word and Picture Version of the Free and Cued Selective Reminding Test (FCSRT): Is There Any Difference?. Journal of Alzheimer's Disease, 2017, 61, 47-52.  | 1.2 | 8         |
| 65 | CSF β-amyloid as a putative biomarker of disease progression in multiple sclerosis. Multiple Sclerosis<br>Journal, 2017, 23, 1085-1091.   | 1.4 | 33        |
| 66 | The Italian dementia with Lewy bodies study group (DLB-SINdem): toward a standardization of clinical procedures and multicenter cohort studies design. Neurological Sciences, 2017, 38, 83-91.  | 0.9 | 11        |
| 67 | Alzheimer's Disease Diagnosis: Discrepancy between Clinical, Neuroimaging, and Cerebrospinal Fluid<br>Biomarkers Criteria in an Italian Cohort of Geriatric Outpatients: A Retrospective Cross-sectional<br>Study. Frontiers in Medicine, 2017, 4, 203. | 1.2 | 8         |
| 68 | PRNP P39L Variant is a Rare Cause ofÂFrontotemporal Dementia in Italian Population. Journal of<br>Alzheimer's Disease, 2016, 50, 353-357.   | 1.2 | 15        |
| 69 | P1â€025: Cerebral Perfusion as an Imaging Biomarker of Presymptomatic Genetic Frontotemporal<br>Dementia: Preliminary Results from the Genetic Frontotemporal Dementia Initiative (GENFI).<br>Alzheimer's and Dementia, 2016, 12, P409.                 | 0.4 | 0         |
| 70 | MRI visual rating scales in the diagnosis of dementia: evaluation in 184 post-mortem confirmed cases.<br>Brain, 2016, 139, 1211-1225.   | 3.7 | 174       |
| 71 | O4-08-06: Visual assessment in postmortem-proven dementias: Clinical expertise versus machine<br>learning. , 2015, 11, P289-P289.   |     | 0         |
| 72 | Presymptomatic cognitive and neuroanatomical changes in genetic frontotemporal dementia in the<br>Genetic Frontotemporal dementia Initiative (GENFI) study: a cross-sectional analysis. Lancet<br>Neurology, The, 2015, 14, 253-262.                    | 4.9 | 432       |

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|----|--|-----|-----------|
| 73 | Inflammatory molecules in Frontotemporal Dementia: Cerebrospinal fluid signature of progranulin mutation carriers. Brain, Behavior, and Immunity, 2015, 49, 182-187.   | 2.0 | 51        |
| 74 | Profiling of Ubiquitination Pathway Genes in Peripheral Cells from Patients with Frontotemporal<br>Dementia due to C9ORF72 and GRN Mutations. International Journal of Molecular Sciences, 2015, 16,<br>1385-1394.   | 1.8 | 14        |
| 75 | Usefulness of Multi-Parametric MRI for the Investigation of Posterior Cortical Atrophy. PLoS ONE, 2015, 10, e0140639.  | 1.1 | 4         |
| 76 | Partial recovery after severe immune reconstitution inflammatory syndrome in a multiple sclerosis patient with progressive multifocal leukoencephalopathy. Immunotherapy, 2014, 6, 23-28.                            | 1.0 | 3         |
| 77 | Circulating miRNAs as Potential Biomarkers in Alzheimer's Disease. Journal of Alzheimer's Disease, 2014, 42, 1261-1267.  | 1.2 | 188       |
| 78 | Autosomal Dominant Frontotemporal Lobar Degeneration Due to the C9ORF72 Hexanucleotide Repeat<br>Expansion: Late-Onset Psychotic Clinical Presentation. Biological Psychiatry, 2013, 74, 384-391.                    | 0.7 | 105       |
| 79 | A 66-year-old patient with vanishing white matter disease due to the p.Ala87Val <i>EIF2B3</i> mutation.<br>Neurology, 2012, 79, 2077-2078.   | 1.5 | 16        |
| 80 | Early Onset Behavioral Variant Frontotemporal Dementia due to the C9ORF72 Hexanucleotide Repeat<br>Expansion: Psychiatric Clinical Presentations. Journal of Alzheimer's Disease, 2012, 31, 447-452.                 | 1.2 | 60        |
| 81 | Sciatic endometriosis presenting as periodic (catamenial) sciatic radiculopathy. Journal of Neurology, 2012, 259, 1470-1471.   | 1.8 | 12        |
| 82 | BAG1 is a Protective Factor for Sporadic Frontotemporal Lobar Degeneration but not for Alzheimer's<br>Disease. Journal of Alzheimer's Disease, 2011, 23, 701-707.  | 1.2 | 12        |
| 83 | Phenotypic Heterogeneity of the GRN Asp22fs Mutation in a Large Italian Kindred. Journal of Alzheimer's Disease, 2011, 24, 253-259.  | 1.2 | 62        |
| 84 | A Novel MAPT Mutation Associated with the Clinical Phenotype of Progressive Nonfluent Aphasia.<br>Journal of Alzheimer's Disease, 2011, 26, 19-26.   | 1.2 | 28        |
| 85 | Cell-dependent kinase inhibitor 2A and 2B genetic variability in patients with Alzheimer's disease.<br>Journal of Neurology, 2011, 258, 704-705.   | 1.8 | 1         |
| 86 | Role of <i>hnRNP-A1</i> and miR-590-3p in Neuronal Death: Genetics and Expression Analysis in Patients<br>with Alzheimer Disease and Frontotemporal Lobar Degeneration. Rejuvenation Research, 2011, 14,<br>275-281. | 0.9 | 57        |
| 87 | Progranulin gene variability increases the risk for primary progressive multiple sclerosis in males.<br>Genes and Immunity, 2010, 11, 497-503.   | 2.2 | 17        |
| 88 | Cerebrospinal fluid progranulin levels in patients with different multiple sclerosis subtypes.<br>Neuroscience Letters, 2010, 469, 234-236.  | 1.0 | 24        |
| 89 | ls KIF24 a genetic risk factor for Frontotemporal Lobar Degeneration?. Neuroscience Letters, 2010, 482, 240-244.   | 1.0 | 9         |
| 90 | Teaching Neuroimage: Crowned Dens Syndrome, an Acute Attack of Calcium Pyrophosphate Deposition<br>Disease Mimicking Acute Meningitis. Neurology, 0, , 10.1212/WNL.0000000000200949.                                 | 1.5 | 0         |