## Paola Caroppo

List of Publications by Year in descending order

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79 2,481 26 45
papers citations h-index g-index

81 81 81 3982 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Age at symptom onset and death and disease duration in genetic frontotemporal dementia: an international retrospective cohort study. Lancet Neurology, The, 2020, 19, 145-156.	10.2	175
2	Behavioral and Psychological Effects of Coronavirus Disease-19 Quarantine in Patients With Dementia. Frontiers in Psychiatry, 2020, 11, 578015.	2.6	157
3	Early Cognitive, Structural, and Microstructural Changes in Presymptomatic <i>C9orf72</i> Carriers Younger Than 40 Years. JAMA Neurology, 2018, 75, 236.	9.0	108
4	Plasma glial fibrillary acidic protein is raised in progranulin-associated frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 263-270.	1.9	106
5	Brain volumetric abnormalities in patients with anorexia and bulimia nervosa: A Voxel-based morphometry study. Psychiatry Research - Neuroimaging, 2013, 213, 210-216.	1.8	91
6	The Recognition of Facial Emotions in Spinocerebellar Ataxia Patients. Cerebellum, 2011, 10, 600-610.	2.5	87
7	The Impact of COVID-19 Quarantine on Patients With Dementia and Family Caregivers: A Nation-Wide Survey. Frontiers in Aging Neuroscience, 2020, 12, 625781.	3.4	85
8	Homozygous TREM2 mutation in a family with atypical frontotemporal dementia. Neurobiology of Aging, 2014, 35, 2419.e23-2419.e25.	3.1	84
9	The CSF neurofilament light signature in rapidly progressive neurodegenerative dementias. Alzheimer's Research and Therapy, 2018, 10, 3.	6.2	76
10	The effect of gender on planning: An fMRI study using the Tower of London task. NeuroImage, 2006, 33, 999-1010.	4.2	71
11	Extensive White Matter Involvement in Patients With Frontotemporal Lobar Degeneration. JAMA Neurology, 2014, 71, 1562.	9.0	68
12	TBK1 mutation frequencies in French frontotemporal dementia and amyotrophic lateral sclerosis cohorts. Neurobiology of Aging, 2015, 36, 3116.e5-3116.e8.	3.1	63
13	17q21.31 duplication causes prominent tau-related dementia with increased MAPT expression. Molecular Psychiatry, 2017, 22, 1119-1125.	7.9	57
14	Defining the spectrum of frontotemporal dementias associated with <i>TARDBP</i> mutations. Neurology: Genetics, 2016, 2, e80.	1.9	56
15	Pathogenic Huntingtin Repeat Expansions in Patients with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. Neuron, 2021, 109, 448-460.e4.	8.1	56
16	TDP-43 real-time quaking induced conversion reaction optimization and detection of seeding activity in CSF of amyotrophic lateral sclerosis and frontotemporal dementia patients. Brain Communications, 2020, 2, fcaa142.	3.3	55
17	Intrinsic Connectivity Networks Within Cerebellum and Beyond in Eating Disorders. Cerebellum, 2013, 12, 623-631.	2.5	53
18	<i>DCTN1</i> Mutation Analysis in Families With Progressive Supranuclear Palsy–Like Phenotypes. JAMA Neurology, 2014, 71, 208.	9.0	48

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19	Functional network resilience to pathology in presymptomatic genetic frontotemporal dementia. Neurobiology of Aging, 2019, 77, 169-177.	3.1	47
20	Linking coordinative and executive dysfunctions to atrophy in spinocerebellar ataxia 2 patients. Brain Structure and Function, 2011, 216, 275-288.	2.3	42
21	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. JAMA Network Open, 2021, 4, e2030194.	5.9	42
22	Understanding the Pathophysiology of Cerebral Amyloid Angiopathy. International Journal of Molecular Sciences, 2020, 21, 3435.	4.1	39
23	Brain functional network integrity sustains cognitive function despite atrophy in presymptomatic genetic frontotemporal dementia. Alzheimer's and Dementia, 2021, 17, 500-514.	0.8	36
24	Role for ATXN1, ATXN2, and HTT intermediate repeats in frontotemporal dementia and Alzheimer's disease. Neurobiology of Aging, 2020, 87, 139.e1-139.e7.	3.1	35
25	Lateral Temporal Lobe: An Early Imaging Marker of the Presymptomatic GRN Disease?. Journal of Alzheimer's Disease, 2015, 47, 751-759.	2.6	34
26	The inner fluctuations of the brain in presymptomatic Frontotemporal Dementia: The chronnectome fingerprint. Neurolmage, 2019, 189, 645-654.	4.2	33
27	Neuropsychological picture of 33 spinocerebellar ataxia cases. Journal of Clinical and Experimental Neuropsychology, 2011, 33, 315-325.	1.3	32
28	Prion-related peripheral neuropathy in sporadic Creutzfeldt-Jakob disease. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 424-427.	1.9	31
29	Differential early subcortical involvement in genetic FTD within the GENFI cohort. NeuroImage: Clinical, 2021, 30, 102646.	2.7	28
30	White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal GENFI study. NeuroImage: Clinical, 2019, 24, 102077.	2.7	27
31	A data-driven disease progression model of fluid biomarkers in genetic frontotemporal dementia. Brain, 2022, 145, 1805-1817.	7.6	27
32	Semantic and nonfluent aphasic variants, secondarily associated with amyotrophic lateral sclerosis, are predominant frontotemporal lobar degeneration phenotypes in <i>TBK1</i> carriers. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2015, 1, 481-486.	2.4	26
33	latrogenic early onset cerebral amyloid angiopathy 30 years after cerebral trauma with neurosurgery: vascular amyloid deposits are made up of both Al²40 and Al²42. Acta Neuropathologica Communications, 2019, 7, 70.	5.2	26
34	Social cognition impairment in genetic frontotemporal dementia within the GENFI cohort. Cortex, 2020, 133, 384-398.	2.4	26
35	Brain correlates of alexithymia in eating disorders: A voxelâ€based morphometry study. Psychiatry and Clinical Neurosciences, 2015, 69, 708-716.	1.8	24
36	Cognitive and Neurophysiological Effects of Non-invasive Brain Stimulation in Stroke Patients after Motor Rehabilitation. Frontiers in Behavioral Neuroscience, 2016, 10, 135.	2.0	24

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37	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. Alzheimer's and Dementia, 2022, 18, 1408-1423.	0.8	24
38	Neurologic and cognitive outcomes after aortic arch operation with hypothermic circulatory arrest. Surgery, 2016, 160, 796-804.	1.9	22
39	Posterior Cortical Atrophy as an Extreme Phenotype of <i>GRN </i> Mutations. JAMA Neurology, 2015, 72, 224.	9.0	21
40	Stratifying the Presymptomatic Phase of Genetic Frontotemporal Dementia by Serum <scp>NfL</scp> and <scp>pNfH</scp> : A Longitudinal Multicentre Study. Annals of Neurology, 2022, 91, 33-47.	5.3	21
41	White matter lesions in FTLD: distinct phenotypes characterize <i>GRN</i> and <i>C9ORF72</i> mutations. Neurology: Genetics, 2016, 2, e47.	1.9	20
42	Faster Cortical Thinning and Surface Area Loss in Presymptomatic and Symptomatic <i>C9orf72</i> Repeat Expansion Adult Carriers. Annals of Neurology, 2020, 88, 113-122.	5.3	19
43	Structural MRI Signatures in Genetic Presentations of the Frontotemporal Dementia/Motor Neuron Disease Spectrum. Neurology, 2021, 97, e1594-e1607.	1.1	19
44	Alzheimer neuropathology without frontotemporal lobar degeneration hallmarks ( <scp>TAR) Tj ETQq0 0 0 rgBT <scp>C</scp>ys139<scp>A</scp>rg. Brain Pathology, 2018, 28, 72-76.</scp>	Overlock 4.1	10 Tf 50 467 16
45	Neurofunctional Signature of Hyperfamiliarity for Unknown Faces. PLoS ONE, 2015, 10, e0129970.	2.5	15
46	Altered Expression of Circulating Cdc42 in Frontotemporal Lobar Degeneration. Journal of Alzheimer's Disease, 2018, 61, 1477-1483.	2.6	15
47	Machine Learning Profiling of Alzheimer's Disease Patients Based on Current Cerebrospinal Fluid Markers and Iron Content in Biofluids. Frontiers in Aging Neuroscience, 2021, 13, 607858.	3.4	15
48	Multivariate analysis of brain metabolism reveals chemotherapy effects on prefrontal cerebellar system when related to dorsal attention network. EJNMMI Research, 2013, 3, 22.	2.5	14
49	Cerebral amyloid angiopathy in a 51-year-old patient with embolization by dura mater extract and surgery for nasopharyngeal angiofibroma at age 17. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, 28, 142-143.	3.0	14
50	Partial deletions of the GRN gene are a cause of frontotemporal lobar degeneration. Neurogenetics, 2014, 15, 95-100.	1.4	11
51	Impairment of episodic memory in genetic frontotemporal dementia: A GENFI study. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2021, 13, e12185.	2.4	11
52	Double-Cone Coil TMS Stimulation of the Medial Cortex Inhibits Central Pain Habituation. PLoS ONE, 2015, 10, e0128765.	2.5	11
53	A previously undiagnosed case of Gerstmannâ€6trässlerâ€6cheinker disease revealed by <i>PRNP</i> gene analysis in patients with adultâ€onset ataxia. Movement Disorders, 2008, 23, 1468-1471.	3.9	10
54	A new NOTCH3 mutation presenting as primary intracerebral haemorrhage. Journal of the Neurological Sciences, 2012, 315, 143-145.	0.6	10

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55	A panel of CSF proteins separates genetic frontotemporal dementia from presymptomatic mutation carriers: a GENFI study. Molecular Neurodegeneration, 2021, 16, 79.	10.8	9
56	Mutations in the POLG1 gene are not a relevant cause of cerebellar ataxia in Italy. Journal of Neurology, 2008, 255, 1079-1080.	3.6	8
57	Neuroimaging Correlates of Frontotemporal Dementia Associated with SQSTM1 Mutations. Journal of Alzheimer's Disease, 2016, 53, 303-313.	2.6	8
58	Missense mutation in GRN gene affecting RNA splicing and plasma progranulin level in a family affected by frontotemporal lobar degeneration. Neurobiology of Aging, 2017, 54, 214.e1-214.e6.	3.1	8
59	Factors influencing the age at onset in familial frontotemporal lobar dementia. Neurology: Genetics, 2017, 3, e203.	1.9	8
60	Genetic variation in APOE, GRN, and TP53 are phenotype modifiers in frontotemporal dementia. Neurobiology of Aging, 2021, 99, 99.e15-99.e22.	3.1	8
61	Disease-related cortical thinning in presymptomatic granulin mutation carriers. Neurolmage: Clinical, 2021, 29, 102540.	2.7	8
62	A unique common ancestor introduced P301L mutation in MAPT gene in frontotemporal dementia patients from Barcelona (Baix Llobregat, Spain). Neurobiology of Aging, 2019, 84, 236.e9-236.e15.	3.1	7
63	Neuropsychological and functional study in a case of partial cerebellar agenesis. Neurocase, 2009, 15, 373-383.	0.6	6
64	Brain metabolism changes after therapy with chenodeoxycholic acid in a case of cerebrotendinous xanthomatosis. Neurological Sciences, 2013, 34, 1693-1696.	1.9	6
65	Chemotherapy-Induced Neurotoxicity: Evidence of a Protective Role of CC Homozygosis in the Interleukin-11 <sup>2</sup> Gene-511 C>T Polymorphism. Neurotoxicity Research, 2016, 30, 521-529.	2.7	6
66	The Neurobiological Basis of the Distress Thermometer: A PET Study in Cancer Patients. Stress and Health, 2015, 31, 197-203.	2.6	5
67	A cluster of progranulin C157KfsX97 mutations in Southern Italy: clinical characterization and genetic correlations. Neurobiology of Aging, 2017, 49, 219.e5-219.e13.	3.1	4
68	Neuropathological Alzheimer's Disease Lesions in Nasu-Hakola Disease with TREM2 Mutation: Atypical Distribution of Neurofibrillary Changes. Journal of Alzheimer's Disease, 2021, 79, 25-30.	2.6	4
69	Cognitive composites for genetic frontotemporal dementia: GENFI-Cog. Alzheimer's Research and Therapy, 2022, 14, 10.	6.2	4
70	Resting state functional brain networks associated with emotion processing in frontotemporal lobar degeneration. Molecular Psychiatry, 2022, 27, 4809-4821.	7.9	4
71	Discovering the Italian phenotype of cerebral amyloid angiopathy (CAA): the SENECA project. Neurological Sciences, 2020, 41, 2193-2200.	1.9	3
72	Sporadic MM-1 Type Creutzfeldt-Jakob Disease With Hemiballic Presentation and No Cognitive Impairment Until Death: How New NCJDRSU Diagnostic Criteria May Allow Early Diagnosis. Frontiers in Neurology, 2018, 9, 739.	2.4	2

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73	New MAPT variant in a FTD patient with Alzheimer's disease phenotype at onset. Neurological Sciences, 2021, 42, 2111-2114.	1.9	2
74	Emotional imagination of negative situations: Functional neuroimaging in anorexia and bulimia. PLoS ONE, 2021, 16, e0231684.	2.5	2
75	Examining empathy deficits across familial forms of frontotemporal dementia within the GENFI cohort. Cortex, 2022, 150, 12-28.	2.4	2
76	The Rise of the GRN C157KfsX97 Mutation in Southern Italy: Going Back to the Fall of the Western Roman Empire. Journal of Alzheimer's Disease, 2020, 78, 387-394.	2.6	1
77	MAPT Q336H mutation: Intrafamilial phenotypic heterogeneity in a new Italian family. European Journal of Neurology, 2022, , .	3.3	1
78	Structural brain splitting is a hallmark of Granulin-related frontotemporal dementia. Neurobiology of Aging, 2022, , .	3.1	1
79	18F-FDG in the differential diagnosis of neurodegenerative dementias. Clinical and Translational Imaging, 2019, 7, 437-445.	2.1	0