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List of Publications by Year in descending order

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54 papers	10,283 citations	331670 21 h-index	189892 50 g-index
64 all docs	64 docs citations	64 times ranked	10774 citing authors

#	Article	IF	Citations
1	Network structure and transcriptomic vulnerability shape atrophy in frontotemporal dementia. Brain, 2023, 146, 321-336.	7.6	30
2	Practice effects in genetic frontotemporal dementia and at-risk individuals: a GENFI study. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 336-339.	1.9	1
3	A data-driven disease progression model of fluid biomarkers in genetic frontotemporal dementia. Brain, 2022, 145, 1805-1817.	7.6	27
4	Cognitive composites for genetic frontotemporal dementia: GENFI-Cog. Alzheimer's Research and Therapy, 2022, 14, 10.	6.2	4
5	The behavioral variant of Alzheimerâ€™s disease does not show a selective loss of Von Economo and phylogenetically related neurons in the anterior cingulate cortex. Alzheimer's Research and Therapy, 2022, 14, 11.	6.2	3
6	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. Alzheimer's and Dementia, 2022, 18, 1408-1423.	0.8	24
7	Structural brain splitting is a hallmark of Granulin-related frontotemporal dementia. Neurobiology of Aging, 2022, , .	3.1	1
8	Anomia is present pre-symptomatically in frontotemporal dementia due to MAPT mutations. Journal of Neurology, 2022, 269, 4322-4332.	3.6	1
9	Development of a sensitive trial-ready poly(GP) CSF biomarker assay forC9orf72-associated frontotemporal dementia and amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 761-771.	1.9	12
10	Differential linguistic features of verbal fluency in behavioral variant frontotemporal dementia and primary progressive aphasia. Applied Neuropsychology Adult, 2022, , 1-9.	1.2	4
11	Longitudinal Cognitive Changes in Genetic Frontotemporal Dementia Within the GENFI Cohort. Neurology, 2022, 99, .	1.1	5
12	Underlying genetic variation in familial frontotemporal dementia: sequencing of 198 patients. Neurobiology of Aging, 2021, 97, 148.e9-148.e16.	3.1	17
13	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
14	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. JAMA Network Open, 2021, 4, e2030194.	5.9	42
15	Distinctive pattern of temporal atrophy in patients with frontotemporal dementia and the I383V variant in TARDBP. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 787-789.	1.9	5
16	CSF sTREM2 is elevated in a subset in GRN-related frontotemporal dementia. Neurobiology of Aging, 2021, 103, 158.e1-158.e5.	3.1	8
17	Novel TUBA4A Variant Associated With Familial Frontotemporal Dementia. Neurology: Genetics, 2021, 7, e596.	1.9	18
18	Characterizing the Clinical Features and Atrophy Patterns of MAPT -Related Frontotemporal Dementia With Disease Progression Modeling. Neurology, 2021, 97, e941-e952.	1.1	29

#	Article	IF	Citations
19	[¹⁸ F]Flortaucipir PET Across Various MAPT Mutations in Presymptomatic and Symptomatic Carriers. Neurology, 2021, 97, e1017-e1030.	1.1	16
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.	6.2	12
21	Genome-wide association study of frontotemporal dementia identifies a C9ORF72 haplotype with a median of 12-G4C2 repeats that predisposes to pathological repeat expansions. Translational Psychiatry, 2021, 11, 451.	4.8	6
22	Differential early subcortical involvement in genetic FTD within the GENFI cohort. NeuroImage: Clinical, 2021, 30, 102646.	2.7	28
23	Unravelling the clinical spectrum and the role of repeat length in C9ORF72 repeat expansions. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 502-509.	1.9	28
24	In vivo PET imaging of neuroinflammation in familial frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 231-231.	1.9	3
25	A panel of CSF proteins separates genetic frontotemporal dementia from presymptomatic mutation carriers: a GENFI study. Molecular Neurodegeneration, 2021, 16, 79.	10.8	9
26	Neuroanatomy of FTD: Wholeâ€brain correlations between symptoms and pathologies. Alzheimer's and Dementia, 2021, 17, e056016.	0.8	0
27	Proteomics of the dentate gyrus reveals semantic-dementia-specific biology.. Alzheimer's and Dementia, 2021, 17 Suppl 3, e052092.	0.8	0
28	Mapping tau burden and neuronal loss in MAPT-associated frontotemporal lobar degeneration.. Alzheimer's and Dementia, 2021, 17 Suppl 3, e054141.	0.8	0
29	Genetic screening in early-onset Alzheimer's disease identified three novel presenilin mutations. Neurobiology of Aging, 2020, 86, 201.e9-201.e14.	3.1	16
30	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
31	SomaticTARDBPvariants as a cause of semantic dementia. Brain, 2020, 143, 3827-3841.	7.6	12
32	Dissecting frontotemporal dementia: Correlations between neuropsychiatric symptoms and neuropathology. Alzheimer's and Dementia, 2020, 16, e038926.	0.8	0
33	Recommendations to distinguish behavioural variant frontotemporal dementia from psychiatric disorders. Brain, 2020, 143, 1632-1650.	7.6	158
34	Cognitive profiles discriminate between genetic variants of behavioral frontotemporal dementia. Journal of Neurology, 2020, 267, 1603-1612.	3.6	17
35	Neuronal pentraxin 2: a synapse-derived CSF biomarker in genetic frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 612-621.	1.9	55
36	Frontotemporal Dementia: Correlations Between Psychiatric Symptoms and Pathology. Annals of Neurology, 2020, 87, 950-961.	5.3	30

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37	EIF2AK3 variants in Dutch patients with Alzheimer's disease. Neurobiology of Aging, 2019, 73, 229.e11-229.e18.	3.1	25
38	Clinical value of cerebrospinal fluid neurofilament light chain in semantic dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 997-1004.	1.9	19
39	Refining the Spectrum of Neuronal Intranuclear Inclusion Disease: A Case Report. Journal of Neuropathology and Experimental Neurology, 2019, 78, 665-670.	1.7	21
40	Genome-wide analyses as part of the international FTLD-TDP whole-genome sequencing consortium reveals novel disease risk factors and increases support for immune dysfunction in FTLD. Acta Neuropathologica, 2019, 137, 879-899.	7.7	90
41	Potential genetic modifiers of disease risk and age at onset in patients with frontotemporal lobar degeneration and GRN mutations: a genome-wide association study. Lancet Neurology, The, 2018, 17, 548-558.	10.2	97
42	P2â€291: THE DIAGNOSTIC CHALLENGE OF NEUROPSYCHIATRIC SYMPTOMS IN ALZHEIMER'S DISEASE: A CASE REPORT. Alzheimer's and Dementia, 2018, 14, P792.	0.8	0
43	Neuropsychiatric Symptoms Complicating the Diagnosis of Alzheimerâ€™s Disease: A Case Report. Journal of Alzheimer's Disease, 2018, 66, 1363-1369.	2.6	5
44	Prevalence of amyloidâ€Î²2 pathology in distinct variants of primary progressive aphasia. Annals of Neurology, 2018, 84, 729-740.	5.3	132
45	Three VCP Mutations in Patients with Frontotemporal Dementia. Journal of Alzheimer's Disease, 2018, 65, 1139-1146.	2.6	19
46	Cerebral blood flow in presymptomatic MAPT and GRN mutation carriers: A longitudinal arterial spin labeling study. NeuroImage: Clinical, 2016, 12, 460-465.	2.7	46
47	Structural and functional brain connectivity in presymptomatic familial frontotemporal dementia. Neurology, 2014, 83, e19-26.	1.1	127
48	Frontotemporal dementia and its subtypes: a genome-wide association study. Lancet Neurology, The, 2014, 13, 686-699.	10.2	302
49	Clinical, genetic and pathological heterogeneity of frontotemporal dementia: a review. Journal of Neurology, Neurosurgery and Psychiatry, 2011, 82, 476-486.	1.9	508
50	Sensitivity of revised diagnostic criteria for the behavioural variant of frontotemporal dementia. Brain, 2011, 134, 2456-2477.	7.6	3,913
51	A Hexanucleotide Repeat Expansion in C9ORF72 Is the Cause of Chromosome 9p21-Linked ALS-FTD. Neuron, 2011, 72, 257-268.	8.1	3,833
52	Brain perfusion patterns in familial frontotemporal lobar degeneration. Neurology, 2011, 77, 384-392.	1.1	17
53	Frequency of ubiquitin and FUS-positive, TDP-43-negative frontotemporal lobar degeneration. Journal of Neurology, 2010, 257, 747-753.	3.6	131
54	Distinct genetic forms of frontotemporal dementia. Neurology, 2008, 71, 1220-1226.	1.1	184