Benedetta Nacmias

List of Publications by Year
in descending order

350 papers	30,019 citations	12330 69 h-index	6471 157 g-index
364 all docs	364 docs citations	364 times ranked	30666 citing authors

#	Article	IF	Citations
1	A proof-of-concept study applying machine learning methods to putative risk factors for eating disorders: results from the multi-centre European project on healthy eating. Psychological Medicine, 2023, 53, 2913-2922.	4.5	9
2	Gender differences in cognitive reserve: implication for subjective cognitive decline in women. Neurological Sciences, 2022, 43, 2499-2508.	1.9	12
3	Effect of BDNF Val66Met polymorphism on hippocampal subfields in multiple sclerosis patients. Molecular Psychiatry, 2022, 27, 1010-1019.	7.9	10
4	A data-driven disease progression model of fluid biomarkers in genetic frontotemporal dementia. Brain, 2022, 145, 1805-1817.	7.6	27
5	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
6	Cognitive composites for genetic frontotemporal dementia: GENFI-Cog. Alzheimer's Research and Therapy, 2022, 14, 10.	6.2	4
7	Examining empathy deficits across familial forms of frontotemporal dementia within the GENFI cohort. Cortex, 2022, 150, 12-28.	2.4	2
8	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. Alzheimer's and Dementia, 2022, 18, 1408-1423.	0.8	24
9	Clinical, Neurophysiological, and Genetic Predictors of Recovery in Patients With Severe Acquired Brain Injuries (PRABI): A Study Protocol for a Longitudinal Observational Study. Frontiers in Neurology, 2022, 13, 711312.	2.4	11
10	Structural brain splitting is a hallmark of Granulin-related frontotemporal dementia. Neurobiology of Aging, 2022, , .	3.1	1
11	Huntingtin gene intermediate alleles influence the progression from subjective cognitive decline to mild cognitive impairment: A 14â€year followâ€up study. European Journal of Neurology, 2022, 29, 1600-1609.	3.3	4
12	CAG Repeats Within the Non-pathological Range in the HTT Gene Influence Personality Traits in Patients With Subjective Cognitive Decline: A 13-Year Follow-Up Study. Frontiers in Psychiatry, 2022, 13, 826135.	2.6	2
13	The <scp>CBIâ€R</scp> detects early behavioural impairment in genetic frontotemporal dementia. Annals of Clinical and Translational Neurology, 2022, 9, 644-658.	3.7	1
14	Plasma neurofilament light chain as a biomarker of Alzheimerâ€™s disease in Subjective Cognitive Decline and Mild Cognitive Impairment. Journal of Neurology, 2022, 269, 4270-4280.	3.6	30
15	New insights into the genetic etiology of Alzheimerâ€™s disease and related dementias. Nature Genetics, 2022, 54, 412-436.	21.4	700
16	Unravelling neural correlates of empathy deficits in Subjective Cognitive Decline, Mild Cognitive Impairment and Alzheimerâ€™s Disease. Behavioural Brain Research, 2022, 428, 113893.	2.2	6
17	Neurofilaments as Decay Rate Biomarker in Spinocerebellar Ataxia Type 1. Neurology, 2022, 98, 821-822.	1.1	2
18	Intermediate alleles of HTT: A new pathway in longevity. Journal of the Neurological Sciences, 2022, 438, 120274.	0.6	3

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19	Long-term use of pharmacological treatment in Alzheimerâ€™s disease: a retrospective cohort study in real-world clinical practice. European Journal of Clinical Pharmacology, 2022, 78, 1155-1163.	1.9	2
20	Loss of speech and functional impairment in Alzheimer's disease-related primary progressive aphasia: predictive factors of decline. Neurobiology of Aging, 2022, 117, 59-70.	3.1	6
21	Association of Rare APOE Missense Variants V236E and R251G With Risk of Alzheimer Disease. JAMA Neurology, 2022, 79, 652.	9.0	31
22	Alzheimerâ€™s Disease CSF Biomarker Profiles in Idiopathic Normal Pressure Hydrocephalus. Journal of Personalized Medicine, 2022, 12, 935.	2.5	4
23	Shared genetic risk between eating disorderâ€•and substanceâ€•useâ€•related phenotypes: Evidence from genomeâ€•wide association studies. Addiction Biology, 2021, 26, e12880.	2.6	28
24	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
25	PER2 C111G polymorphism, cognitive reserve and cognition in subjective cognitive decline and mild cognitive impairment: a 10â€•year followâ€•up study. European Journal of Neurology, 2021, 28, 56-65.	3.3	7
26	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
27	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
28	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. JAMA Network Open, 2021, 4, e2030194.	5.9	42
29	The Brain-Derived Neurotrophic Factor Val66Met Polymorphism Can Protect Against Cognitive Impairment in Multiple Sclerosis. Frontiers in Neurology, 2021, 12, 645220.	2.4	13
30	Alpha-synuclein seeds in olfactory mucosa and cerebrospinal fluid of patients with dementia with Lewy bodies. Brain Communications, 2021, 3, fcab045.	3.3	37
31	Late-onset Huntington disease: An Italian cohort. Journal of Clinical Neuroscience, 2021, 86, 58-63.	1.5	3
32	Predictors of Function, Activity, and Participation of Stroke Patients Undergoing Intensive Rehabilitation: A Multicenter Prospective Observational Study Protocol. Frontiers in Neurology, 2021, 12, 632672.	2.4	15
33	Gene Expression Imputation Across Multiple Tissue Types Provides Insight Into the Genetic Architecture of Frontotemporal Dementia and Its Clinical Subtypes. Biological Psychiatry, 2021, 89, 825-835.	1.3	10
34	Dual Effect of PER2 C111G Polymorphism on Cognitive Functions across Progression from Subjective Cognitive Decline to Mild Cognitive Impairment. Diagnostics, 2021, 11, 718.	2.6	3
35	Whole-genome sequencing analysis of semi-supercentenarians. ELife, 2021, 10, .	6.0	37
36	Common variants in Alzheimerâ€™s disease and risk stratification by polygenic risk scores. Nature Communications, 2021, 12, 3417.	12.8	140

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37	The Effect of CAG Repeats within the Non-Pathological Range in the HTT Gene on Cognitive Functions in Patients with Subjective Cognitive Decline and Mild Cognitive Impairment. Diagnostics, 2021, 11, 1051.	2.6	7
38	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
39	Neurofilament Light Chain and Intermediate HTT Alleles as Combined Biomarkers in Italian ALS Patients. Frontiers in Neuroscience, 2021, 15, 695049.	2.8	1
40	Differential early subcortical involvement in genetic FTD within the GENFI cohort. NeuroImage: Clinical, 2021, 30, 102646.	2.7	28
41	Matching Clinical Diagnosis and Amyloid Biomarkers in Alzheimerâ€™s Disease and Frontotemporal Dementia. Journal of Personalized Medicine, 2021, 11, 47.	2.5	9
42	SLITRK2, an X-linked modifier of the age at onset in C9orf72 frontotemporal lobar degeneration. Brain, 2021, 144, 2798-2811.	7.6	7
43	Plasma neurofilament light chain as a useful biomarker in prodromal phases of Alzheimer's disease. Journal of the Neurological Sciences, 2021, 429, 117826.	0.6	0
44	Predictive factors of progression to total loss of language and functional autonomy in primary progressive aphasia related to Alzheimer's disease. Journal of the Neurological Sciences, 2021, 429, 117851.	0.6	0
45	Gender differences in cognitive reserve: Implication for subjective cognitive decline in women. Journal of the Neurological Sciences, 2021, 429, 117827.	0.6	1
46	THE effect of trinucleotide repeats in the HTT gene on the progression from subjective cognitive decline to mild cognitive impairment: A 12 years follow-up study. Journal of the Neurological Sciences, 2021, 429, 119020.	0.6	0
47	Cerebral amyloid load determination in a clinical setting: interpretation of amyloid biomarker discordances aided by tau and neurodegeneration measurements. Neurological Sciences, 2021, , 1.	1.9	0
48	A panel of CSF proteins separates genetic frontotemporal dementia from presymptomatic mutation carriers: a GENFI study. Molecular Neurodegeneration, 2021, 16, 79.	10.8	9
49	The implication of BDNF Val66Met polymorphism in progression from subjective cognitive decline to mild cognitive impairment and Alzheimerâ€™s disease: a 9-year follow-up study. European Archives of Psychiatry and Clinical Neuroscience, 2020, 270, 471-482.	3.2	25
50	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
51	Cerebrospinal fluid biomarkers for dementia: A case of post-lumbar puncture epidural hematoma. Clinical Neurology and Neurosurgery, 2020, 190, 105638.	1.4	0
52	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
53	A case of limbic encephalitis evolving into a frontotemporal dementiaâ€like picture. Psychogeriatrics, 2020, 20, 355-357.	1.2	0
54	Linguistic profiles, brain metabolic patterns and rates of amyloid-Î² biomarker positivity in patients with mixed primary progressive aphasia. Neurobiology of Aging, 2020, 96, 155-164.	3.1	9

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55	Assessing the effectiveness of subjective cognitive decline plus criteria in predicting the progression to Alzheimerâ€™s disease: an 11â€year followâ€up study. European Journal of Neurology, 2020, 27, 894-899.	3.3	14
56	Mendelian randomization implies no direct causal association between leukocyte telomere length and amyotrophic lateral sclerosis. Scientific Reports, 2020, 10, 12184.	3.3	4
57	Challenges in Alzheimerâ€™s Disease Diagnostic Work-Up: Amyloid Biomarker Incongruences. Journal of Alzheimer's Disease, 2020, 77, 203-217.	2.6	3
58	C9orf72, age at onset, and ancestry help discriminate behavioral from language variants in FTLD cohorts. Neurology, 2020, 95, e3288-e3302.	1.1	7
59	Tumor Necrosis Factor Î± Influences Phenotypic Plasticity and Promotes Epigenetic Changes in Human Basal Forebrain Cholinergic Neuroblasts. International Journal of Molecular Sciences, 2020, 21, 6128.	4.1	17
60	Influence of ApoE Genotype and Clock T3111C Interaction with Cardiovascular Risk Factors on the Progression to Alzheimerâ€™s Disease in Subjective Cognitive Decline and Mild Cognitive Impairment Patients. Journal of Personalized Medicine, 2020, 10, 45.	2.5	15
61	SIRT1 accelerates the progression of activity-based anorexia. Nature Communications, 2020, 11, 2814.	12.8	16
62	Early functional MRI changes in a prodromal semantic variant of primary progressive aphasia: a longitudinal case report. Journal of Neurology, 2020, 267, 3100-3104.	3.6	4
63	Human iPSC-Derived Hippocampal Spheroids: An Innovative Tool for Stratifying Alzheimer Disease Patient-Specific Cellular Phenotypes and Developing Therapies. Stem Cell Reports, 2020, 15, 256-273.	4.8	49
64	Incomplete penetrance in familial Alzheimerâ€™s disease with PSEN1 Ala260Gly mutation. Neurological Sciences, 2020, 41, 2263-2266.	1.9	3
65	Faster Cortical Thinning and Surface Area Loss in Presymptomatic and Symptomatic C9orf72 Repeat Expansion Adult Carriers. Annals of Neurology, 2020, 88, 113-122.	5.3	19
66	Genome-wide association study identifies eight risk loci and implicates metabo-psychiatric origins for anorexia nervosa. Nature Genetics, 2019, 51, 1207-1214.	21.4	641
67	Serum neurofilament light chain in genetic frontotemporal dementia: a longitudinal, multicentre cohort study. Lancet Neurology, The, 2019, 18, 1103-1111.	10.2	128
68	High Frequency of Crossed Aphasia in Dextral in an Italian Cohort of Patients with Logopenic Primary Progressive Aphasia. Journal of Alzheimer's Disease, 2019, 72, 1089-1096.	2.6	2
69	Clinical and neuroimaging profiles to identify C9orf72 â€FTD patients and serum Neurofilament to monitor the progression and the severity of the disease. Neurology and Clinical Neuroscience, 2019, 7, 326-333.	0.4	1
70	Transethnic meta-analysis of rare coding variants in PLCG2, ABI3, and TREM2 supports their general contribution to Alzheimerâ€™s disease. Translational Psychiatry, 2019, 9, 55.	4.8	32
71	The inner fluctuations of the brain in presymptomatic Frontotemporal Dementia: The chronnectome fingerprint. NeuroImage, 2019, 189, 645-654.	4.2	33
72	Associations Between Attention-Deficit/Hyperactivity Disorder and Various Eating Disorders: A Swedish Nationwide Population Study Using Multiple Genetically Informative Approaches. Biological Psychiatry, 2019, 86, 577-586.	1.3	43

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73	Crossed aphasia confirmed by fMRI in a case with nonfluent variant of primary progressive aphasia carrying a GRN mutation. Journal of Neurology, 2019, 266, 1274-1279.	3.6	3
74	KIBRA T allele influences memory performance and progression of cognitive decline: a 7-year follow-up study in subjective cognitive decline and mild cognitive impairment. Neurological Sciences, 2019, 40, 1559-1566.	1.9	10
75	Primary Progressive Aphasia. Alzheimer Disease and Associated Disorders, 2019, 33, 42-46.	1.3	12
76	Genetic meta-analysis of diagnosed Alzheimerâ€™s disease identifies new risk loci and implicates AÎ², tau, immunity and lipid processing. Nature Genetics, 2019, 51, 414-430.	21.4	1,962
77	White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal GENFI study. NeuroImage: Clinical, 2019, 24, 102077.	2.7	27
78	Genomic Relationships, Novel Loci, and Pleiotropic Mechanisms across Eight Psychiatric Disorders. Cell, 2019, 179, 1469-1482.e11.	28.9	935
79	Spatiotemporal analysis for detection of pre-symptomatic shape changes in neurodegenerative diseases: Initial application to the GENFI cohort. NeuroImage, 2019, 188, 282-290.	4.2	16
80	The dual role of cognitive reserve in subjective cognitive decline and mild cognitive impairment: a 7-year follow-up study. Journal of Neurology, 2019, 266, 487-497.	3.6	51
81	Functional network resilience to pathology in presymptomatic genetic frontotemporal dementia. Neurobiology of Aging, 2019, 77, 169-177.	3.1	47
82	Novel GRN Mutations in Alzheimerâ€™s Disease and Frontotemporal Lobar Degeneration. Journal of Alzheimer's Disease, 2018, 62, 1683-1689.	2.6	12
83	Biomarkers study in atypical dementia: proof of a diagnostic work-up. Neurological Sciences, 2018, 39, 1203-1210.	1.9	3
84	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
85	Rare Dementias. , 2018, , 313-336.		0
86	Rare nonsynonymous variants in SORT1 are associated with increased risk for frontotemporal dementia. Neurobiology of Aging, 2018, 66, 181.e3-181.e10.	3.1	19
87	Comparison of arterial spin labeling registration strategies in the multiâ€center GENetic frontotemporal dementia initiative (GENFI). Journal of Magnetic Resonance Imaging, 2018, 47, 131-140.	3.4	41
88	Patterns of gray matter atrophy in genetic frontotemporal dementia: results from the GENFI study. Neurobiology of Aging, 2018, 62, 191-196.	3.1	151
89	The diagnosis of dementias: a practical tool not to miss rare causes. Neurological Sciences, 2018, 39, 615-627.	1.9	14
90	Genetic Heterogeneity of Alzheimerâ€™s Disease: Embracing Research Partnerships. Journal of Alzheimer's Disease, 2018, 62, 903-911.	2.6	10

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91	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.	3.1	40
92	Common and rare TBK1 variants in early-onset Alzheimer disease in a European cohort. Neurobiology of Aging, 2018, 62, 245.e1-245.e7.	3.1	16
93	G02â€...Decision-making in predictive testing in huntingtonâ€™s disease. , 2018, , .		0
94	Contribution of Bilingualism to Cognitive Reserve of an Italian Literature Professor at High Risk for Alzheimerâ€™s Disease. Journal of Alzheimer's Disease, 2018, 66, 1389-1395.	2.6	8
95	A C6orf10/LOC101929163 locus is associated with age of onset in C9orf72 carriers. Brain, 2018, 141, 2895-2907.	7.6	39
96	Genetically elevated highâ€‐density lipoprotein cholesterol through the cholesteryl ester transfer protein gene does not associate with risk of Alzheimer's disease. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2018, 10, 595-598.	2.4	2
97	Quantitative Genetics Validates Previous Genetic Variants and Identifies Novel Genetic Players Influencing Alzheimerâ€™s Disease Cerebrospinal Fluid Biomarkers. Journal of Alzheimer's Disease, 2018, 66, 639-652.	2.6	12
98	From Subjective Cognitive Decline to Alzheimerâ€™s Disease: The Predictive Role of Neuropsychological Assessment, Personality Traits, and Cognitive Reserve. A 7-Year Follow-Up Study. Journal of Alzheimer's Disease, 2018, 63, 1523-1535.	2.6	68
99	Crossed aphasia in nonfluent variant of primary progressive aphasia carrying a GRN mutation. Journal of the Neurological Sciences, 2018, 392, 34-37.	0.6	7
100	Presymptomatic white matter integrity loss in familial frontotemporal dementia in the <scp>GENFI</scp> cohort: A crossâ€‐sectional diffusion tensor imaging study. Annals of Clinical and Translational Neurology, 2018, 5, 1025-1036.	3.7	39
101	No supportive evidence for TIA1 gene mutations in a European cohort of ALS-FTD spectrum patients. Neurobiology of Aging, 2018, 69, 293.e9-293.e11.	3.1	15
102	Impact of demography and population dynamics on the genetic architecture of human longevity. Aging, 2018, 10, 1947-1963.	3.1	16
103	Genetic architecture of sporadic frontotemporal dementia and overlap with Alzheimer's and Parkinson's diseases. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, 152-164.	1.9	107
104	Effects of Multiple Genetic Loci on Age atÂOnset in Frontotemporal Dementia. Journal of Alzheimer's Disease, 2017, 56, 1271-1278.	2.6	4
105	The Methylenetetrahydrofolate Reductase C677T Polymorphism and Risk for Late-Onset Alzheimerâ€™s disease: Further Evidence in an Italian Multicenter Study. Journal of Alzheimer's Disease, 2017, 56, 1451-1457.	2.6	20
106	Genetics of vascular dementia â€“ review from the ICVD working group. BMC Medicine, 2017, 15, 48.	5.5	59
107	Significant Locus and Metabolic Genetic Correlations Revealed in Genome-Wide Association Study of Anorexia Nervosa. American Journal of Psychiatry, 2017, 174, 850-858.	7.2	410
108	The novel PSEN1 M84V mutation associated to frontal dysexecutive syndrome, spastic paraparesis, and cerebellar atrophy in a dominant Alzheimer's disease family. Neurobiology of Aging, 2017, 56, 213.e7-213.e12.	3.1	19

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109	Cognitive reserve and TMEM106B genotype modulate brain damage in presymptomatic frontotemporal dementia: a GENFI study. Brain, 2017, 140, 1784-1791.	7.6	55
110	Deleterious ABCA7 mutations and transcript rescue mechanisms in early onset Alzheimerâ€™s disease. Acta Neuropathologica, 2017, 134, 475-487.	7.7	53
111	White matter hyperintensities are seen only in GRN mutation carriers in the GENFI cohort. NeuroImage: Clinical, 2017, 15, 171-180.	2.7	63
112	TBK1 Mutation Spectrum in an Extended European Patient Cohort with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. Human Mutation, 2017, 38, 297-309.	2.5	87
113	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. Nature Genetics, 2017, 49, 1373-1384.	21.4	783
114	Association of the New Variant Tyr424Asp at TBK1 Gene with Amyotrophic Lateral Sclerosis and Cognitive Decline. Journal of Alzheimer's Disease, 2017, 61, 41-46.	2.6	3
115	Low Florbetapir PET Uptake and Normal AÎ²1-42 Cerebrospinal Fluid in an APP Ala713Thr Mutation Carrier. Journal of Alzheimer's Disease, 2017, 57, 697-703.	2.6	5
116	Alzheimerâ€™s Disease Progression: Factors Influencing Cognitive Decline. Journal of Alzheimer's Disease, 2017, 61, 785-791.	2.6	37
117	Fat mass and obesity-associated gene (FTO) is associated to eating disorders susceptibility and moderates the expression of psychopathological traits. PLoS ONE, 2017, 12, e0173560.	2.5	45
118	A novel network analysis approach reveals DNA damage, oxidative stress and calcium/cAMP homeostasis-associated biomarkers in frontotemporal dementia. PLoS ONE, 2017, 12, e0185797.	2.5	21
119	PRNP P39L Variant is a Rare Cause ofÂFrontotemporal Dementia in Italian Population. Journal of Alzheimer's Disease, 2016, 50, 353-357.	2.6	15
120	Genetic Counseling and Testing for Alzheimerâ€™s Disease and Frontotemporal Lobar Degeneration: An Italian Consensus Protocol. Journal of Alzheimer's Disease, 2016, 51, 277-291.	2.6	18
121	Shared genetic contribution to ischemic stroke and Alzheimer's disease. Annals of Neurology, 2016, 79, 739-747.	5.3	56
122	Analyses of the role of the glucocorticoid receptor gene polymorphism (rs41423247) as a potential moderator in the association between childhood overweight, psychopathology, and clinical outcomes in Eating Disorders patients: A 6 years follow up study. Psychiatry Research, 2016, 243, 156-160.	3.3	7
123	Novel presenilin 1 mutation (Ile408Thr) in an Italian family with late-onset Alzheimerâ€™s disease. Neuroscience Letters, 2016, 610, 150-153.	2.1	4
124	Rare Variants inPLD3Do Not Affect Risk for Early-Onset Alzheimer Disease in a European Consortium Cohort. Human Mutation, 2015, 36, 1226-1235.	2.5	23
125	Left hippocampusâ€“amygdala complex macroâ€•and microstructural variation is associated with <scp>BDNF</scp> plasma levels in healthy elderly individuals. Brain and Behavior, 2015, 5, e00334.	2.2	10
126	Donor-Specific Anti-HLA Antibodies in Huntington's Disease Recipients of Human Fetal Striatal Grafts. Cell Transplantation, 2015, 24, 811-817.	2.5	12

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127	Monomeric ÃŸ-amyloid interacts with type-1 insulin-like growth factor receptors to provide energy supply to neurons. Frontiers in Cellular Neuroscience, 2015, 9, 297.	3.7	44
128	Presymptomatic cognitive and neuroanatomical changes in genetic frontotemporal dementia in the Genetic Frontotemporal dementia Initiative (GENFI) study: a cross-sectional analysis. Lancet Neurology, The, 2015, 14, 253-262.	10.2	432
129	Italian Frontotemporal Dementia Network (FTD Group-SINDEM): sharing clinical and diagnostic procedures in Frontotemporal Dementia in Italy. Neurological Sciences, 2015, 36, 751-757.	1.9	9
130	The C9orf72 repeat expansion itself is methylated in ALS and FTLD patients. Acta Neuropathologica, 2015, 129, 715-727.	7.7	114
131	A genome-wide screening and SNPs-to-genes approach to identify novel genetic risk factors associated with frontotemporal dementia. Neurobiology of Aging, 2015, 36, 2904.e13-2904.e26.	3.1	48
132	Genetic variability in SQSTM1 and risk of early-onset Alzheimer dementia: a European early-onset dementia consortium study. Neurobiology of Aging, 2015, 36, 2005.e15-2005.e22.	3.1	34
133	Csf p-tau₁₈₁/tau ratio as biomarker for TDP pathology in frontotemporal dementia. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2015, 16, 86-91.	1.7	65
134	Mutation analysis ofCHCHD10in different neurodegenerative diseases. Brain, 2015, 138, e380-e380.	7.6	86
135	A new social-family model for eating disorders: A European multicentre project using a caseâ€“control design. Appetite, 2015, 95, 544-553.	3.7	14
136	Convergent genetic and expression data implicate immunity in Alzheimer's disease. Alzheimer's and Dementia, 2015, 11, 658-671.	0.8	173
137	Mutation analysis of patients with neurodegenerative disorders using NeuroX array. Neurobiology of Aging, 2015, 36, 545.e9-545.e14.	3.1	36
138	Longâ€term effect of HCV eradication in patients with mixed cryoglobulinemia: A prospective, controlled, openâ€label, cohort study. Hepatology, 2015, 61, 1145-1153.	7.3	107
139	Hypermethylation of the CpG-island near the C9orf72 G4C2-repeat expansion in FTLD patients. Human Molecular Genetics, 2014, 23, 5630-5637.	2.9	74
140	A Pilot Study Evaluating the Contribution ofSLC19A1(RFC-1) 80G>A Polymorphism to Alzheimerâ€™s Disease in Italian Caucasians. BioMed Research International, 2014, 2014, 1-6.	1.9	1
141	Increased Levels of Serum IL-18 Are Associated with the Long-Term Outcome of Severe Traumatic Brain Injury. NeuroImmunoModulation, 2014, 21, 8-12.	1.8	25
142	A psychosocial risk factor model for female eating disorders: a European multicentre project. Journal of Eating Disorders, 2014, 2, .	2.7	0
143	Advances in imagingâ€“genetic relationships for Alzheimerâ€™s disease: clinical implications. Neurodegenerative Disease Management, 2014, 4, 73-81.	2.2	5
144	Imaging and Cognitive Reserve Studies Predict Dementia in Presymptomatic Alzheimer's Disease Subjects. Neurodegenerative Diseases, 2014, 13, 157-159.	1.4	18

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145	Effects of a novel schizophrenia risk variant rs7914558 atCNNM2on brain structure and attributional style. British Journal of Psychiatry, 2014, 204, 115-121.	2.8	30
146	Rare mutations in SQSTM1 modify susceptibility to frontotemporal lobar degeneration. Acta Neuropathologica, 2014, 128, 397-410.	7.7	93
147	Frontotemporal dementia and its subtypes: a genome-wide association study. Lancet Neurology, The, 2014, 13, 686-699.	10.2	302
148	Heterozygous TREM2 mutations in frontotemporal dementia. Neurobiology of Aging, 2014, 35, 934.e7-934.e10.	3.1	134
149	Epigenetic Modifications in Alzheimer's Disease: Cause or Effect?. Journal of Alzheimer's Disease, 2014, 43, 1169-1173.	2.6	27
150	Plasma Membrane Injury Depends on Bilayer Lipid Composition in Alzheimer's Disease. Journal of Alzheimer's Disease, 2014, 41, 289-300.	2.6	23
151	Association of the Variant Cys139Arg at GRN Gene to the Clinical Spectrum of Frontotemporal Lobar Degeneration. Journal of Alzheimer's Disease, 2014, 40, 679-685.	2.6	11
152	P4-074: ITALIAN NETWORK FOR AUTOSOMAL DOMINANT ALZHEIMER'S DISEASE AND FRONTOTEMPORAL LOBAR DEGENERATION (ITALIANDIAFN). , 2014, 10, P810-P810.		0
153	Uncommon Dementias. , 2014, , 193-214.		1
154	Gene-Wide Analysis Detects Two New Susceptibility Genes for Alzheimer's Disease. PLoS ONE, 2014, 9, e94661.	2.5	155
155	Genetics of Alzheimerâ€™s Disease and Frontotemporal Dementia. Current Molecular Medicine, 2014, 14, 993-1000.	1.3	16
156	FDG PET and the genetics of dementia. Clinical and Translational Imaging, 2013, 1, 235-246.	2.1	2
157	Meta-analysis of 74,046 individuals identifies 11 new susceptibility loci for Alzheimer's disease. Nature Genetics, 2013, 45, 1452-1458.	21.4	3,741
158	The stimulation of dendritic cells by amyloid beta 1â€“42 reduces BDNF production in Alzheimerâ€™s disease patients. Brain, Behavior, and Immunity, 2013, 32, 29-32.	4.1	29
159	A Panâ€<scp>E</scp>uropean Study of theC9orf72Repeat Associated with<scp>FTLD</scp>: Geographic Prevalence, Genomic Instability, and Intermediate Repeats. Human Mutation, 2013, 34, 363-373.	2.5	247
160	Autosomal Dominant Frontotemporal Lobar Degeneration Due to the C9ORF72 Hexanucleotide Repeat Expansion: Late-Onset Psychotic Clinical Presentation. Biological Psychiatry, 2013, 74, 384-391.	1.3	105
161	Low social interactions in eating disorder patients in childhood and adulthood: A multi-centre European case control study. Journal of Health Psychology, 2013, 18, 26-37.	2.3	70
162	Tomm40 polymorphisms in Italian Alzheimerâ€™s disease and frontotemporal dementia patients. Neurological Sciences, 2013, 34, 995-998.	1.9	28

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