Benedetta Nacmias

List of Publications by Year in Descending Order

336 papers	21,774 citations	63 h-index	140 g-index
364 ext. papers	26,408 ext. citations	6.9 avg, IF	5.46 L-index

#	Paper	IF	Citations
336	Cognitive composites for genetic frontotemporal dementia: GENFI-Cog.. *Alzheimerls Research and Therapy*, **2022**, 14, 10	9	0
335	Examining empathy deficits across familial forms of frontotemporal dementia within the GENFI cohort.. *Cortex*, **2022**, 150, 12-28	3.8	
334	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	4.1	1
333	Huntingtin gene intermediate alleles influence the progression from subjective cognitive decline to mild cognitive impairment: A 14-year follow-up study.. *European Journal of Neurology*, **2022**,	6	1
332	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	5	0
331	New insights into the genetic etiology of Alzheimer's disease and related dementias.. *Nature Genetics*, **2022**,	36.3	27
330	Unravelling neural correlates of empathy deficits in Subjective Cognitive Decline, Mild Cognitive Impairment and Alzheimer's Disease.. *Behavioural Brain Research*, **2022**, 113893	3.4	1
329	Intermediate alleles of HTT: A new pathway in longevity.. *Journal of the Neurological Sciences*, **2022**, 438, 120274	3.2	
328	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. *Alzheimerls and Dementia*, **2021**,	1.2	2
327	Stratifying the Presymptomatic Phase of Genetic Frontotemporal Dementia by Serum NfL and pNfH: A Longitudinal Multicentre Study. *Annals of Neurology*, **2021**,	9.4	2
326	Cerebral amyloid load determination in a clinical setting: interpretation of amyloid biomarker discordances aided by tau and neurodegeneration measurements. *Neurological Sciences*, **2021**, 1	3.5	
325	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*, **2021**, 1-10	6.9	3
324	A panel of CSF proteins separates genetic frontotemporal dementia from presymptomatic mutation carriers: a GENFI study. *Molecular Neurodegeneration*, **2021**, 16, 79	19	0
323	Gender differences in cognitive reserve: implication for subjective cognitive decline in women. *Neurological Sciences*, **2021**, 1	3.5	4
322	Effect of BDNF Val66Met polymorphism on hippocampal subfields in multiple sclerosis patients. *Molecular Psychiatry*, **2021**,	15.1	2
321	A data-driven disease progression model of fluid biomarkers in genetic frontotemporal dementia. *Brain*, **2021**,	11.2	3
320	Alpha-synuclein seeds in olfactory mucosa and cerebrospinal fluid of patients with dementia with Lewy bodies. *Brain Communications*, **2021**, 3, fcab045	4.5	11

319	Late-onset Huntington disease: An Italian cohort. *Journal of Clinical Neuroscience*, **2021**, 86, 58-63	2.2	
318	Predictors of Function, Activity, and Participation of Stroke Patients Undergoing Intensive Rehabilitation: A Multicenter Prospective Observational Study Protocol. *Frontiers in Neurology*, **2021**, 12, 632672	4.1	4
317	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	7.9	3
316	Whole-genome sequencing analysis of semi-supercentenarians. *ELife*, **2021**, 10,	8.9	11
315	Common variants in Alzheimer's disease and risk stratification by polygenic risk scores. *Nature Communications*, **2021**, 12, 3417	17.4	23
314	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,	3.8	3
313	The Revised Self-Monitoring Scale detects early impairment of social cognition in genetic frontotemporal dementia within the GENFI cohort. *Alzheimerls Research and Therapy*, **2021**, 13, 127	9	2
312	Shared genetic risk between eating disorder- and substance-use-related phenotypes: Evidence from genome-wide association studies. *Addiction Biology*, **2021**, 26, e12880	4.6	12
311	Genetic variation in APOE, GRN, and TP53 are phenotype modifiers in frontotemporal dementia. *Neurobiology of Aging*, **2021**, 99, 99.e15-99.e22	5.6	3
310	PER2 C111G polymorphism, cognitive reserve and cognition in subjective cognitive decline and mild cognitive impairment: a 10-year follow-up study. *European Journal of Neurology*, **2021**, 28, 56-65	6	1
309	Brain functional network integrity sustains cognitive function despite atrophy in presymptomatic genetic frontotemporal dementia. *Alzheimerls and Dementia*, **2021**, 17, 500-514	1.2	8
308	Impairment of episodic memory in genetic frontotemporal dementia: A GENFI study. *Alzheimerls and Dementia: Diagnosis, Assessment and Disease Monitoring*, **2021**, 13, e12185	5.2	1
307	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. *JAMA Network Open*, **2021**, 4, e2030194	10.4	14
306	The Brain-Derived Neurotrophic Factor Val66Met Polymorphism Can Protect Against Cognitive Impairment in Multiple Sclerosis. *Frontiers in Neurology*, **2021**, 12, 645220	4.1	3
305	Neurofilament Light Chain and Intermediate HTT Alleles as Combined Biomarkers in Italian ALS Patients. *Frontiers in Neuroscience*, **2021**, 15, 695049	5.1	
304	Differential early subcortical involvement in genetic FTD within the GENFI cohort. *NeuroImage: Clinical*, **2021**, 30, 102646	5.3	6
303	Matching Clinical Diagnosis and Amyloid Biomarkers in Alzheimer's Disease and Frontotemporal Dementia. *Journal of Personalized Medicine*, **2021**, 11,	3.6	1
302	SLITRK2, an X-linked modifier of the age at onset in C9orf72 frontotemporal lobar degeneration. *Brain*, **2021**, 144, 2798-2811	11.2	2

301	Influence of Genotype and T3111C Interaction with Cardiovascular Risk Factors on the Progression to Alzheimer's Disease in Subjective Cognitive Decline and Mild Cognitive Impairment Patients. *Journal of Personalized Medicine*, **2020**, 10,	3.6	6
300	SIRT1 accelerates the progression of activity-based anorexia. *Nature Communications*, **2020**, 11, 2814	17.4	9
299	Early functional MRI changes in a prodromal semantic variant of primary progressive aphasia: a longitudinal case report. *Journal of Neurology*, **2020**, 267, 3100-3104	5.5	1
298	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	8	23
297	Incomplete penetrance in familial Alzheimer's disease with PSEN1 Ala260Gly mutation. *Neurological Sciences*, **2020**, 41, 2263-2266	3.5	3
296	Role for ATXN1, ATXN2, and HTT intermediate repeats in frontotemporal dementia and Alzheimer's disease. *Neurobiology of Aging*, **2020**, 87, 139.e1-139.e7	5.6	13
295	Cerebrospinal fluid biomarkers for dementia: A case of post-lumbar puncture epidural hematoma. *Clinical Neurology and Neurosurgery*, **2020**, 190, 105638	2	
294	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	24.1	90
293	A case of limbic encephalitis evolving into a frontotemporal dementia-like picture. *Psychogeriatrics*, **2020**, 20, 355-357	1.8	
292	Linguistic profiles, brain metabolic patterns and rates of amyloid-□biomarker positivity in patients with mixed primary progressive aphasia. *Neurobiology of Aging*, **2020**, 96, 155-164	5.6	1
291	Assessing the effectiveness of subjective cognitive decline plus criteria in predicting the progression to Alzheimer's disease: an 11-year follow-up study. *European Journal of Neurology*, **2020**, 27, 894-899	6	7
290	Mendelian randomization implies no direct causal association between leukocyte telomere length and amyotrophic lateral sclerosis. *Scientific Reports*, **2020**, 10, 12184	4.9	1
289	Challenges in Alzheimer's Disease Diagnostic Work-Up: Amyloid Biomarker Incongruences. *Journal of Alzheimerls Disease*, **2020**, 77, 203-217	4.3	1
288	, age at onset, and ancestry help discriminate behavioral from language variants in FTLD cohorts. *Neurology*, **2020**, 95, e3288-e3302	6.5	5
287	Tumor Necrosis Factor □nfluences Phenotypic Plasticity and Promotes Epigenetic Changes in Human Basal Forebrain Cholinergic Neuroblasts. *International Journal of Molecular Sciences*, **2020**, 21,	6.3	9
286	The implication of BDNF Val66Met polymorphism in progression from subjective cognitive decline to mild cognitive impairment and Alzheimer's disease: a 9-year follow-up study. *European Archives of Psychiatry and Clinical Neuroscience*, **2020**, 270, 471-482	5.1	11
285	Faster Cortical Thinning and Surface Area Loss in Presymptomatic and Symptomatic C9orf72 Repeat Expansion Adult Carriers. *Annals of Neurology*, **2020**, 88, 113-122	9.4	11
284	Clinical and neuroimaging profiles to identify C9orf72 -FTD patients and serum Neurofilament to monitor the progression and the severity of the disease. *Neurology and Clinical Neuroscience*, **2019**, 7, 326-333	0.3	0

283	Transethnic meta-analysis of rare coding variants in PLCG2, ABI3, and TREM2 supports their general contribution to Alzheimer's disease. *Translational Psychiatry*, **2019**, 9, 55	8.6	19
282	The inner fluctuations of the brain in presymptomatic Frontotemporal Dementia: The chronnectome fingerprint. *NeuroImage*, **2019**, 189, 645-654	7.9	18
281	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	7.9	24
280	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	5.5	0
279	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	3.5	2
278	Primary Progressive Aphasia: Natural History in an Italian Cohort. *Alzheimer Disease and Associated Disorders*, **2019**, 33, 42-46	2.5	6
277	Genome-wide association study identifies eight risk loci and implicates metabo-psychiatric origins for anorexia nervosa. *Nature Genetics*, **2019**, 51, 1207-1214	36.3	303
276	Serum neurofilament light chain in genetic frontotemporal dementia: a longitudinal, multicentre cohort study. *Lancet Neurology, The*, **2019**, 18, 1103-1111	24.1	68
275	High Frequency of Crossed Aphasia in Dextral in an Italian Cohort of Patients with Logopenic Primary Progressive Aphasia. *Journal of Alzheimerls Disease*, **2019**, 72, 1089-1096	4.3	2
274	Genetic meta-analysis of diagnosed Alzheimer's disease identifies new risk loci and implicates A tau, immunity and lipid processing. *Nature Genetics*, **2019**, 51, 414-430	36.3	917
273	White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal GENFI study. *NeuroImage: Clinical*, **2019**, 24, 102077	5.3	13
272	Genomic Relationships, Novel Loci, and Pleiotropic Mechanisms across Eight Psychiatric Disorders. *Cell*, **2019**, 179, 1469-1482.e11	56.2	402
271	Spatiotemporal analysis for detection of pre-symptomatic shape changes in neurodegenerative diseases: Initial application to the GENFI cohort. *NeuroImage*, **2019**, 188, 282-290	7.9	10
270	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	5.5	33
269	Functional network resilience to pathology in presymptomatic genetic frontotemporal dementia. *Neurobiology of Aging*, **2019**, 77, 169-177	5.6	24
268	Novel GRN Mutations in Alzheimer's Disease and Frontotemporal Lobar Degeneration. *Journal of Alzheimerls Disease*, **2018**, 62, 1683-1689	4.3	7
267	Biomarkers study in atypical dementia: proof of a diagnostic work-up. *Neurological Sciences*, **2018**, 39, 1203-1210	3.5	3
266	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	24.1	60

265	Rare Dementias **2018**, 313-336		
264	Rare nonsynonymous variants in SORT1 are associated with increased risk for frontotemporal dementia. *Neurobiology of Aging*, **2018**, 66, 181.e3-181.e10	5.6	12
263	Comparison of arterial spin labeling registration strategies in the multi-center GENetic frontotemporal dementia initiative (GENFI). *Journal of Magnetic Resonance Imaging*, **2018**, 47, 131-140	5.6	32
262	Patterns of gray matter atrophy in genetic frontotemporal dementia: results from the GENFI study. *Neurobiology of Aging*, **2018**, 62, 191-196	5.6	104
261	Presymptomatic white matter integrity loss in familial frontotemporal dementia in the GENFI cohort: A cross-sectional diffusion tensor imaging study. *Annals of Clinical and Translational Neurology*, **2018**, 5, 1025-1036	5.3	29
260	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	5.6	11
259	Association of the New Variant Tyr424Asp at TBK1 Gene with Amyotrophic Lateral Sclerosis and Cognitive Decline. *Journal of Alzheimerls Disease*, **2018**, 61, 41-46	4.3	2
258	Alzheimer's Disease Progression: Factors Influencing Cognitive Decline. *Journal of Alzheimerls Disease*, **2018**, 61, 785-791	4.3	26
257	Impact of demography and population dynamics on the genetic architecture of human longevity. *Aging*, **2018**, 10, 1947-1963	5.6	13
256	The diagnosis of dementias: a practical tool not to miss rare causes. *Neurological Sciences*, **2018**, 39, 615-627	1.7	11
255	Genetic Heterogeneity of Alzheimer's Disease: Embracing Research Partnerships. *Journal of Alzheimerls Disease*, **2018**, 62, 903-911	4.3	6
254	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	5.6	30
253	Common and rare TBK1 variants in early-onset Alzheimer disease in a European cohort. *Neurobiology of Aging*, **2018**, 62, 245.e1-245.e7	5.6	12
252	Contribution of Bilingualism to Cognitive Reserve of an Italian Literature Professor at High Risk for Alzheimer's Disease. *Journal of Alzheimerls Disease*, **2018**, 66, 1389-1395	4.3	7
251	A C6orf10/LOC101929163 locus is associated with age of onset in C9orf72 carriers. *Brain*, **2018**, 141, 2895-2907	11.2	25
250	Genetically elevated high-density lipoprotein cholesterol through the cholesteryl ester transfer protein gene does not associate with risk of Alzheimer's disease. *Alzheimerls and Dementia: Diagnosis, Assessment and Disease Monitoring*, **2018**, 10, 595-598	5.2	
249	Quantitative Genetics Validates Previous Genetic Variants and Identifies Novel Genetic Players Influencing Alzheimer's Disease Cerebrospinal Fluid Biomarkers. *Journal of Alzheimerls Disease*, **2018**, 66, 639-652	4.3	8
248	From Subjective Cognitive Decline to Alzheimer's Disease: The Predictive Role of Neuropsychological Assessment, Personality Traits, and Cognitive Reserve. A 7-Year Follow-Up Study. *Journal of Alzheimerls Disease*, **2018**, 63, 1523-1535	4.3	48

247	Crossed aphasia in nonfluent variant of primary progressive aphasia carrying a GRN mutation. *Journal of the Neurological Sciences*, **2018**, 392, 34-37	3.2	4
246	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	5.5	76
245	Effects of Multiple Genetic Loci on Age at Onset in Frontotemporal Dementia. *Journal of Alzheimerls Disease*, **2017**, 56, 1271-1278	4.3	3
244	The Methylenetetrahydrofolate Reductase C677T Polymorphism and Risk for Late-Onset Alzheimer's disease: Further Evidence in an Italian Multicenter Study. *Journal of Alzheimerls Disease*, **2017**, 56, 1451-1457	4.3	17
243	Genetics of vascular dementia - review from the ICVD working group. *BMC Medicine*, **2017**, 15, 48	11.4	39
242	Significant Locus and Metabolic Genetic Correlations Revealed in Genome-Wide Association Study of Anorexia Nervosa. *American Journal of Psychiatry*, **2017**, 174, 850-858	11.9	276
241	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	5.6	12
240	Cognitive reserve and TMEM106B genotype modulate brain damage in presymptomatic frontotemporal dementia: a GENFI study. *Brain*, **2017**, 140, 1784-1791	11.2	31
239	Deleterious ABCA7 mutations and transcript rescue mechanisms in early onset Alzheimer's disease. *Acta Neuropathologica*, **2017**, 134, 475-487	14.3	34
238	White matter hyperintensities are seen only in mutation carriers in the GENFI cohort. *NeuroImage: Clinical*, **2017**, 15, 171-180	5.3	43
237	TBK1 Mutation Spectrum in an Extended European Patient Cohort with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. *Human Mutation*, **2017**, 38, 297-309	4.7	66
236	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. *Nature Genetics*, **2017**, 49, 1373-1384	36.3	508
235	Low Florbetapir PET Uptake and Normal Aβ1-42 Cerebrospinal Fluid in an APP Ala713Thr Mutation Carrier. *Journal of Alzheimerls Disease*, **2017**, 57, 697-703	4.3	5
234	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	3.7	38
233	A novel network analysis approach reveals DNA damage, oxidative stress and calcium/cAMP homeostasis-associated biomarkers in frontotemporal dementia. *PLoS ONE*, **2017**, 12, e0185797	3.7	14
232	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, 156-60	9.9	6
231	Novel presenilin 1 mutation (Ile408Thr) in an Italian family with late-onset Alzheimer's disease. *Neuroscience Letters*, **2016**, 610, 150-3	3.3	2
230	PRNP P39L Variant is a Rare Cause of Frontotemporal Dementia in Italian Population. *Journal of Alzheimerls Disease*, **2016**, 50, 353-7	4.3	12

229	Genetic Counseling and Testing for Alzheimer's Disease and Frontotemporal Lobar Degeneration: An Italian Consensus Protocol. *Journal of Alzheimerls Disease*, **2016**, 51, 277-91	4.3	13
228	Shared genetic contribution to Ischaemic Stroke and Alzheimer's Disease. *Annals of Neurology*, **2016**, 79, 739-747	9.4	42
227	The C9orf72 repeat expansion itself is methylated in ALS and FTLD patients. *Acta Neuropathologica*, **2015**, 129, 715-27	14.3	101
226	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.e13-26	5.6	34
225	Genetic variability in SQSTM1 and risk of early-onset Alzheimer dementia: a European early-onset dementia consortium study. *Neurobiology of Aging*, **2015**, 36, 2005.e15-22	5.6	29
224	Csf p-tau181/tau ratio as biomarker for TDP pathology in frontotemporal dementia. *Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration*, **2015**, 16, 86-91	3.6	47
223	Mutation analysis of CHCHD10 in different neurodegenerative diseases. *Brain*, **2015**, 138, e380	11.2	67
222	A new social-family model for eating disorders: A European multicentre project using a case-control design. *Appetite*, **2015**, 95, 544-53	4.5	10
221	Convergent genetic and expression data implicate immunity in Alzheimer's disease. *Alzheimerls and Dementia*, **2015**, 11, 658-71	1.2	146
220	Mutation analysis of patients with neurodegenerative disorders using NeuroX array. *Neurobiology of Aging*, **2015**, 36, 545.e9-14	5.6	24
219	Long-term effect of HCV eradication in patients with mixed cryoglobulinemia: a prospective, controlled, open-label, cohort study. *Hepatology*, **2015**, 61, 1145-53	11.2	98
218	Epigenetic modifications in Alzheimer's disease: cause or effect?. *Journal of Alzheimerls Disease*, **2015**, 43, 1169-73	4.3	23
217	Rare Variants in PLD3 Do Not Affect Risk for Early-Onset Alzheimer Disease in a European Consortium Cohort. *Human Mutation*, **2015**, 36, 1226-35	4.7	20
216	Left hippocampus-amygdala complex macro- and microstructural variation is associated with BDNF plasma levels in healthy elderly individuals. *Brain and Behavior*, **2015**, 5, e00334	3.4	8
215	Donor-Specific Anti-HLA Antibodies in Huntington's Disease Recipients of Human Fetal Striatal Grafts. *Cell Transplantation*, **2015**, 24, 811-7	4	10
214	Monomeric □amyloid interacts with type-1 insulin-like growth factor receptors to provide energy supply to neurons. *Frontiers in Cellular Neuroscience*, **2015**, 9, 297	6.1	29
213	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, 253-62	24.1	328
212	Italian Frontotemporal Dementia Network (FTD Group-SINDEM): sharing clinical and diagnostic procedures in Frontotemporal Dementia in Italy. *Neurological Sciences*, **2015**, 36, 751-7	3.5	8

211	Effects of a novel schizophrenia risk variant rs7914558 at CNNM2 on brain structure and attributional style. *British Journal of Psychiatry*, **2014**, 204, 115-21	5.4	25
210	Rare mutations in SQSTM1 modify susceptibility to frontotemporal lobar degeneration. *Acta Neuropathologica*, **2014**, 128, 397-410	14.3	83
209	Frontotemporal dementia and its subtypes: a genome-wide association study. *Lancet Neurology, The*, **2014**, 13, 686-99	24.1	207
208	Heterozygous TREM2 mutations in frontotemporal dementia. *Neurobiology of Aging*, **2014**, 35, 934.e7-10	5.6	107
207	Plasma membrane injury depends on bilayer lipid composition in Alzheimer's disease. *Journal of Alzheimerls Disease*, **2014**, 41, 289-300	4.3	15
206	Association of the variant Cys139Arg at GRN gene to the clinical spectrum of frontotemporal lobar degeneration. *Journal of Alzheimerls Disease*, **2014**, 40, 679-85	4.3	8
205	P4-074: ITALIAN NETWORK FOR AUTOSOMAL DOMINANT ALZHEIMER'S DISEASE AND FRONTOTEMPORAL LOBAR DEGENERATION (ITALIANDIAFN) **2014**, 10, P810-P810		
204	Hypermethylation of the CpG-island near the C9orf72 G□□repeat expansion in FTLD patients. *Human Molecular Genetics*, **2014**, 23, 5630-7	5.6	68
203	A pilot study evaluating the contribution of SLC19A1 (RFC-1) 80G>a polymorphism to Alzheimer's disease in Italian Caucasians. *BioMed Research International*, **2014**, 2014, 608104	3	1
202	Increased levels of serum IL-18 are associated with the long-term outcome of severe traumatic brain injury. *NeuroImmunoModulation*, **2014**, 21, 8-12	2.5	15
201	A psychosocial risk factor model for female eating disorders: a European multicentre project. *Journal of Eating Disorders*, **2014**, 2,	4.1	78
200	Advances in imaging-genetic relationships for Alzheimer's disease: clinical implications. *Neurodegenerative Disease Management*, **2014**, 4, 73-81	2.8	4
199	Imaging and cognitive reserve studies predict dementia in presymptomatic Alzheimer's disease subjects. *Neurodegenerative Diseases*, **2014**, 13, 157-9	2.3	18
198	Gene-wide analysis detects two new susceptibility genes for Alzheimer's disease. *PLoS ONE*, **2014**, 9, e94661	3.7	90
197	Genetics of Alzheimer's Disease and Frontotemporal Dementia. *Current Molecular Medicine*, **2014**, 14, 993-1000	2.5	15
196	Uncommon Dementias **2014**, 193-214		1
195	FDG PET and the genetics of dementia. *Clinical and Translational Imaging*, **2013**, 1, 235-246	2	2
194	Meta-analysis of 74,046 individuals identifies 11 new susceptibility loci for Alzheimer's disease. *Nature Genetics*, **2013**, 45, 1452-8	36.3	2714

193	The stimulation of dendritic cells by amyloid beta 1-42 reduces BDNF production in Alzheimer's disease patients. *Brain, Behavior, and Immunity*, **2013**, 32, 29-32	16.6	24
192	A pan-European study of the C9orf72 repeat associated with FTLD: geographic prevalence, genomic instability, and intermediate repeats. *Human Mutation*, **2013**, 34, 363-73	4.7	208
191	Autosomal dominant frontotemporal lobar degeneration due to the C9ORF72 hexanucleotide repeat expansion: late-onset psychotic clinical presentation. *Biological Psychiatry*, **2013**, 74, 384-91	7.9	94
190	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	3.1	59
189	TOMM40 polymorphisms in Italian Alzheimer's disease and frontotemporal dementia patients. *Neurological Sciences*, **2013**, 34, 995-8	3.5	22
188	The SIRT2 polymorphism rs10410544 and risk of Alzheimer's disease in two Caucasian case-control cohorts. *Alzheimerls and Dementia*, **2013**, 9, 392-9	1.2	36
187	Ataxia-telangiectasia mutated (ATM) genetic variant in Italian centenarians. *Neurological Sciences*, **2013**, 34, 573-5	3.5	14
186	A case of atypical early-onset Alzheimer's disease carrying the missense mutation Thr354Ile in exon 10 of the PSEN1 gene. *Neurological Sciences*, **2013**, 34, 1691-2	3.5	2
185	Neuropsychiatric symptoms and interleukin-6 serum levels in acute stroke. *Journal of Neuropsychiatry and Clinical Neurosciences*, **2013**, 25, 255-63	2.7	34
184	Genome-wide haplotype association study identifies the FRMD4A gene as a risk locus for Alzheimer's disease. *Molecular Psychiatry*, **2013**, 18, 461-70	15.1	77
183	Plasma fatty acid lipidomics in amnestic mild cognitive impairment and Alzheimer's disease. *Journal of Alzheimerls Disease*, **2013**, 36, 545-53	4.3	46
182	Blood dendritic cell frequency declines in idiopathic Parkinson's disease and is associated with motor symptom severity. *PLoS ONE*, **2013**, 8, e65352	3.7	28
181	Lipid rafts mediate amyloid-induced calcium dyshomeostasis and oxidative stress in Alzheimer's disease. *Current Alzheimer Research*, **2013**, 10, 143-53	3	34
180	Genetics of familial and sporadic Alzheimer's disease. *Frontiers in Bioscience - Elite*, **2013**, 5, 167-77	1.6	122
179	Clinical heterogeneity in Italian patients with amyotrophic lateral sclerosis. *Clinical Genetics*, **2012**, 82, 83-7	4	9
178	Progranulin genetic screening in frontotemporal lobar degeneration patients from central Italy. *Cellular and Molecular Neurobiology*, **2012**, 32, 13-6	4.6	9
177	Position paper of the Italian Society for the study of Dementias (SINDEM) on the proposal of a new lexicon on Alzheimer disease. *Neurological Sciences*, **2012**, 33, 201-8	3.5	3
176	EFNS-ENS Guidelines on the diagnosis and management of disorders associated with dementia. *European Journal of Neurology*, **2012**, 19, 1159-79	6	191

175	Increased levels of serum MAP-2 at 6-months correlate with improved outcome in survivors of severe traumatic brain injury. *Brain Injury*, **2012**, 26, 1629-35	2.1	40
174	Elevated levels of circulating IL-18BP and perturbed regulation of IL-18 in schizophrenia. *Journal of Neuroinflammation*, **2012**, 9, 206	10.1	24
173	High serum levels of transforming growth factor 1 are associated with increased cortical thickness in cingulate and right frontal areas in healthy subjects. *Journal of Neuroinflammation*, **2012**, 9, 42	10.1	6
172	Suitability of neuropsychological tests in patients with vascular dementia (VaD). *Journal of the Neurological Sciences*, **2012**, 322, 41-5	3.2	9
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2	New insights on the genetic etiology of Alzheimer and related dementia		25
1	Identifying tissues implicated in Anorexia Nervosa using Transcriptomic Imputation		1