Marc Cruts

List of Publications by Year
in descending order

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142 papers	17,421 citations	14655 66 h-index	14208 128 g-index
155 all docs	155 docs citations	155 times ranked	15239 citing authors

#	Article	IF	Citations
1	Null mutations in progranulin cause ubiquitin-positive frontotemporal dementia linked to chromosome 17q21. Nature, 2006, 442, 920-924.	27.8	1,386
2	The C9orf72 GGGGCC Repeat Is Translated into Aggregating Dipeptide-Repeat Proteins in FTLD/ALS. Science, 2013, 339, 1335-1338.	12.6	1,095
3	Presenile dementia and cerebral haemorrhage linked to a mutation at codon 692 of the Î²â€“amyloid precursor protein gene. Nature Genetics, 1992, 1, 218-221.	21.4	715
4	A C9orf72 promoter repeat expansion in a Flanders-Belgian cohort with disorders of the frontotemporal lobar degeneration-amyotrophic lateral sclerosis spectrum: a gene identification study. Lancet Neurology, The, 2012, 11, 54-65.	10.2	565
5	Apolipoprotein E4 allele in a populationâ€“based study of earlyâ€“onset Alzheimer's disease. Nature Genetics, 1994, 7, 74-78.	21.4	460
6	Genetic etiology of Parkinson disease associated with mutations in the SNCA, PARK2, PINK1, PARK7, and LRRK2 genes: a mutation update. Human Mutation, 2010, 31, 763-780.	2.5	428
7	Bidirectional transcripts of the expanded C9orf72 hexanucleotide repeat are translated into aggregating dipeptide repeat proteins. Acta Neuropathologica, 2013, 126, 881-893.	7.7	427
8	Genetic Association of Apolipoprotein E with Age-Related Macular Degeneration. American Journal of Human Genetics, 1998, 63, 200-206.	6.2	425
9	Locusâ€specific mutation databases for neurodegenerative brain diseases. Human Mutation, 2012, 33, 1340-1344.	2.5	414
10	Estimation of the Genetic Contribution of Presenilin-1 and -2 Mutations in a Population-Based Study of Presenile Alzheimer Disease. Human Molecular Genetics, 1998, 7, 43-51.	2.9	396
11	Risk Estimates of Dementia by Apolipoprotein E Genotypes From a Population-Based Incidence Study: The Rotterdam Study. Archives of Neurology, 1998, 55, 964.	4.5	378
12	APP duplication is sufficient to cause early onset Alzheimer's dementia with cerebral amyloid angiopathy. Brain, 2006, 129, 2977-2983.	7.6	337
13	The role of tau (MAPT) in frontotemporal dementia and related tauopathies. Human Mutation, 2004, 24, 277-295.	2.5	323
14	Mapping of a gene predisposing to earlyâ€“onset Alzheimer's disease to chromosome 14q24.3. Nature Genetics, 1992, 2, 335-339.	21.4	321
15	Mean age-of-onset of familial alzheimer disease caused by presenilin mutations correlates with both increased AÎ²42 and decreased AÎ²40. Human Mutation, 2006, 27, 686-695.	2.5	306
16	hnRNP A3 binds to GGGGCC repeats and is a constituent of p62-positive/TDP43-negative inclusions in the hippocampus of patients with C9orf72 mutations. Acta Neuropathologica, 2013, 125, 413-423.	7.7	302
17	Frontotemporal dementia and its subtypes: a genome-wide association study. Lancet Neurology, The, 2014, 13, 686-699.	10.2	302
18	novoSNP, a novel computational tool for sequence variation discovery. Genome Research, 2005, 15, 436-442.	5.5	254

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19	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
20	The genetics and neuropathology of frontotemporal lobar degeneration. Acta Neuropathologica, 2012, 124, 353-372.	7.7	242
21	Drosophila screen connects nuclear transport genes to DPR pathology in c9ALS/FTD. Scientific Reports, 2016, 6, 20877.	3.3	239
22	A novel presenilin 1 mutation associated with Pick's disease but not Î²â€amyloid plaques. Annals of Neurology, 2004, 55, 617-626.	5.3	210
23	Serum biomarker for progranulinâ€associated frontotemporal lobar degeneration. Annals of Neurology, 2009, 65, 603-609.	5.3	195
24	Alzheimer risk associated with a copy number variation in the complement receptor 1 increasing C3b/C4b binding sites. Molecular Psychiatry, 2012, 17, 223-233.	7.9	179
25	Molecular genetic analysis of familial early-onset Alzheimer's disease linked to chromosome 14q24.3. Human Molecular Genetics, 1995, 4, 2363-2371.	2.9	171
26	The presenilin genes: a new gene family involved in Alzheimer disease pathology. Human Molecular Genetics, 1996, 5, 1449-1455.	2.9	169
27	Presenilin mutations in Alzheimer's disease. Human Mutation, 1998, 11, 183-190.	2.5	169
28	Progranulin null mutations in both sporadic and familial frontotemporal dementia. Human Mutation, 2007, 28, 846-855.	2.5	162
29	Investigating the role of rare heterozygous TREM2 variants in Alzheimer's disease and frontotemporal dementia. Neurobiology of Aging, 2014, 35, 726.e11-726.e19.	3.1	158
30	High-density SNP haplotyping suggests altered regulation of tau gene expression in progressive supranuclear palsy. Human Molecular Genetics, 2005, 14, 3281-3292.	2.9	156
31	Sequencing an Ashkenazi reference panel supports population-targeted personal genomics and illuminates Jewish and European origins. Nature Communications, 2014, 5, 4835.	12.8	156
32	Current insights into the C9orf72 repeat expansion diseases of the FTLD/ALS spectrum. Trends in Neurosciences, 2013, 36, 450-459.	8.6	151
33	Loss of TBK1 is a frequent cause of frontotemporal dementia in a Belgian cohort. Neurology, 2015, 85, 2116-2125.	1.1	151
34	Alzheimer and Parkinson Diagnoses in Progranulin Null Mutation Carriers in an Extended Founder Family. Archives of Neurology, 2007, 64, 1436.	4.5	143
35	Nonfibrillar diffuse amyloid deposition due to a gamma42-secretase site mutation points to an essential role for N-truncated Abeta42 in Alzheimer's disease. Human Molecular Genetics, 2000, 9, 2589-2598.	2.9	135
36	Reduced functional brain activity response in cognitively intact apolipoprotein E Îµ4 carriers. Brain, 2006, 129, 1240-1248.	7.6	133

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37	Molecular genetics of Alzheimer's disease. Annals of Medicine, 1998, 30, 560-565.	3.8	131
38	APOE genotype does not modulate age of onset in families with chromosome 14 encoded Alzheimer's disease. Neuroscience Letters, 1994, 169, 179-180.	2.1	130
39	Granulin mutations associated with frontotemporal lobar degeneration and related disorders: An update. Human Mutation, 2008, 29, 1373-1386.	2.5	126
40	Loss of ALS-associated TDP-43 in zebrafish causes muscle degeneration, vascular dysfunction, and reduced motor neuron axon outgrowth. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 4986-4991.	7.1	126
41	TMEM106B is a genetic modifier of frontotemporal lobar degeneration with C9orf72 hexanucleotide repeat expansions. Acta Neuropathologica, 2014, 127, 407-418.	7.7	123
42	The pursuit of susceptibility genes for Alzheimer's disease: progress and prospects. Trends in Genetics, 2010, 26, 84-93.	6.7	122
43	Rescue of Progranulin Deficiency Associated with Frontotemporal Lobar Degeneration by Alkalizing Reagents and Inhibition of Vacuolar ATPase. Journal of Neuroscience, 2011, 31, 1885-1894.	3.6	121
44	Mutations other than null mutations producing a pathogenic loss of progranulin in frontotemporal dementia. Human Mutation, 2007, 28, 416-416.	2.5	116
45	Relationship between C9orf72 repeat size and clinical phenotype. Current Opinion in Genetics and Development, 2017, 44, 117-124.	3.3	114
46	The influence of apoe status on episodic and semantic memory: Data from a population-based study.. Neuropsychology, 2006, 20, 645-657.	1.3	112
47	Reduced hippocampal volume in non-demented carriers of the apolipoprotein E É›4: Relation to chronological age and recognition memory. Neuroscience Letters, 2006, 396, 23-27.	2.1	112
48	Amyloid precursor protein gene mutation in early-onset Alzheimer's disease. Lancet, The, 1991, 337, 978.	13.7	110
49	Loss of progranulin function in frontotemporal lobar degeneration. Trends in Genetics, 2008, 24, 186-194.	6.7	110
50	TMEM106B is associated with frontotemporal lobar degeneration in a clinically diagnosed patient cohort. Brain, 2011, 134, 808-815.	7.6	110
51	Genetic and Clinical Features of Progranulin-Associated Frontotemporal Lobar Degeneration. Archives of Neurology, 2011, 68, 488.	4.5	108
52	Tau negative frontal lobe dementia at 17q21: significant finemapping of the candidate region to a 4.8 cM interval. Molecular Psychiatry, 2002, 7, 1064-1074.	7.9	103
53	PRNP Val129 homozygosity increases risk for earlyâ€onset Alzheimer's disease. Annals of Neurology, 2003, 53, 409-412.	5.3	103
54	Rare mutations in SQSTM1 modify susceptibility to frontotemporal lobar degeneration. Acta Neuropathologica, 2014, 128, 397-410.	7.7	93

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55	A Belgian ancestral haplotype harbours a highly prevalent mutation for 17q21-linked tau-negative FTLD. Brain, 2006, 129, 841-852.	7.6	88
56	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
57	Clinical features ofTBK1carriers compared withC9orf72,GRNand non-mutation carriers in a Belgian cohort. Brain, 2016, 139, 452-467.	7.6	86
58	Progranulin locus deletion in frontotemporal dementia. Human Mutation, 2008, 29, 53-58.	2.5	85
59	Distinct Clinical Characteristics of C9orf72 Expansion Carriers Compared With GRN, MAPT, and Nonmutation Carriers in a Flanders-Belgian FTLD Cohort. JAMA Neurology, 2013, 70, 365.	9.0	85
60	Genomic architecture of human 17q21 linked to frontotemporal dementia uncovers a highly homologous family of low-copy repeats in the tau region. Human Molecular Genetics, 2005, 14, 1753-1762.	2.9	82
61	Genetic variability in the regulatory region of presenilin 1 associated with risk for Alzheimer's disease and variable expression. Human Molecular Genetics, 2000, 9, 325-331.	2.9	77
62	Apolipoprotein E and Carotid Artery Atherosclerosis. Stroke, 2001, 32, 1947-1952.	2.0	75
63	A Presenilin-1 Truncating Mutation Is Present in Two Cases with Autopsy-Confirmed Early-Onset Alzheimer Disease. American Journal of Human Genetics, 1998, 62, 70-76.	6.2	74
64	Tau (MAPT) mutation Arg406Trp presenting clinically with Alzheimer disease does not share a common founder in Western Europe. Human Mutation, 2003, 22, 409-411.	2.5	72
65	Genetic association of CR1 with Alzheimer's disease: A tentative disease mechanism. Neurobiology of Aging, 2012, 33, 2949.e5-2949.e12.	3.1	72
66	Cerebral amyloid angiopathy is a pathogenic lesion in Alzheimer's disease due to a novel presenilin 1 mutation. Brain, 2001, 124, 2383-2392.	7.6	70
67	Neuronal inclusion protein TDP-43 has no primary genetic role in FTD and ALS. Neurobiology of Aging, 2009, 30, 1329-1331.	3.1	67
68	C9orf72 G4C2 repeat expansions in Alzheimer's disease and mild cognitive impairment. Neurobiology of Aging, 2013, 34, 1712.e1-1712.e7.	3.1	65
69	Dose dependent effect of APOE É›4 on behavioral symptoms in frontal lobe dementia. Neurobiology of Aging, 2006, 27, 285-292.	3.1	64
70	Alzheimer dementia caused by a novel mutation located in the APP C-terminal intracytosolic fragment. Human Mutation, 2006, 27, 888-896.	2.5	62
71	Familial clustering and genetic risk for dementia in a genetically isolated Dutch population. Brain, 2004, 127, 1641-1649.	7.6	60
72	Progranulin Mutations in Ubiquitin-Positive Frontotemporal Dementia Linked to Chromosome 17q21. Current Alzheimer Research, 2006, 3, 485-491.	1.4	60

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73	Serum apolipoprotein E level is not increased in Alzheimer's disease: the Rotterdam study. Neuroscience Letters, 1998, 248, 21-24.	2.1	58
74	Cholesterol and Triglycerides Moderate the Effect of Apolipoprotein E on Memory Functioning in Older Adults. Journals of Gerontology - Series B Psychological Sciences and Social Sciences, 2007, 62, P112-P118.	3.9	57
75	Increased risk of dementia following mild head injury for carriers but not for non-carriers of the APOE Îµ4 allele. International Psychogeriatrics, 2007, 19, 159.	1.0	57
76	Global investigation and meta-analysis of the C9orf72 (G ₄ C ₂) _n repeat in Parkinson disease. Neurology, 2014, 83, 1906-1913.	1.1	56
77	Clinical Evidence of Disease Anticipation in Families Segregating a C9orf72 Repeat Expansion. JAMA Neurology, 2017, 74, 445.	9.0	56
78	Tau is central in the genetic Alzheimerâ€“frontotemporal dementia spectrum. Trends in Genetics, 2005, 21, 664-672.	6.7	55
79	Systematic genetic study of Alzheimer disease in Latin America: Mutation frequencies of the amyloid ? precursor protein and presenilin genes in Colombia. American Journal of Medical Genetics Part A, 2001, 103, 138-143.	2.4	54
80	The microtubule associated protein Tau gene and Alzheimer's disease â€“ an association study and meta-analysis. Neuroscience Letters, 2001, 314, 92-96.	2.1	52
81	Diagnostic accuracy of the Preclinical AD Scale (PAS) in cognitively mildly impaired subjects. Journal of Neurology, 2002, 249, 312-319.	3.6	51
82	Apolipoprotein E genotype and progression of Alzheimer's disease: the Rotterdam Study. Journal of Neurology, 1999, 246, 304-308.	3.6	50
83	Genetic association of the presenilin-1 regulatory region with early-onset Alzheimer's disease in a population-based sample. European Journal of Human Genetics, 1999, 7, 801-806.	2.8	49
84	Evidence That AÎ²42 Plasma Levels in Presenilin-1 Mutation Carriers Do not Allow for Prediction of Their Clinical Phenotype. Neurobiology of Disease, 1999, 6, 280-287.	4.4	48
85	Linkage and Association Studies Identify a Novel Locus for Alzheimer Disease at 7q36 in a Dutch Population-Based Sample. American Journal of Human Genetics, 2005, 77, 643-652.	6.2	48
86	The Glu318Gly Substitution in Presenilin 1 Is Not Causally Related to Alzheimer Disease. American Journal of Human Genetics, 1999, 64, 290-292.	6.2	47
87	Amyloid Î² secretase gene (BACE) is neither mutated in nor associated with early-onset Alzheimer's disease. Neuroscience Letters, 2001, 313, 105-107.	2.1	47
88	Fatigue before and after mild traumatic brain injury: Preâ€“post-injury comparisons in relation toApolipoproteinE. Brain Injury, 2007, 21, 1049-1054.	1.2	47
89	Identification of 2 Loci at Chromosomes 9 and 14 in a Multiplex Family With Frontotemporal Lobar Degeneration and Amyotrophic Lateral Sclerosis. Archives of Neurology, 2010, 67, 606-16.	4.5	47
90	Reduced secreted clusterin as a mechanism for Alzheimer-associated CLU mutations. Molecular Neurodegeneration, 2015, 10, 30.	10.8	46

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91	Course of objective memory impairment in non-demented subjects attending a memory clinic and predictors of outcome. International Journal of Geriatric Psychiatry, 2000, 15, 363-372.	2.7	45
92	The Gene Encoding Nicastrin, a Major Î³-Secretase Component, Modifies Risk for Familial Early-Onset Alzheimer Disease in a Dutch Population-Based Sample. American Journal of Human Genetics, 2002, 70, 1568-1574.	6.2	45
93	Alzheimer-associated C allele of the promoter polymorphism -22C>T causes a critical neuron-specific decrease of presenilin 1 expression. Human Molecular Genetics, 2003, 12, 869-877.	2.9	45
94	Characterization of Ubiquitinated Intraneuronal Inclusions in a Novel Belgian Frontotemporal Lobar Degeneration Family. Journal of Neuropathology and Experimental Neurology, 2006, 65, 289-301.	1.7	45
95	Influence of the prion protein and the apolipoprotein E genotype on the Creutzfeldtâ€“Jakob Disease phenotype. Neuroscience Letters, 2001, 313, 69-72.	2.1	44
96	Mutation screening of the tau gene in patients with early-onset Alzheimer's disease. Neuroscience Letters, 1999, 277, 137-139.	2.1	43
97	Early cognitive decline is associated with prion protein codon 129 polymorphism. Annals of Neurology, 2003, 54, 275-276.	5.3	43
98	Promoter DNA methylation regulates progranulin expression and is altered in FTLD. Acta Neuropathologica Communications, 2013, 1, 16.	5.2	43
99	Genetics of Early-Onset Alzheimer Dementia. Scientific World Journal, The, 2003, 3, 497-519.	2.1	40
100	Molecular Pathways of Frontotemporal Lobar Degeneration. Annual Review of Neuroscience, 2010, 33, 71-88.	10.7	39
101	The â^'491 A/T polymorphism in the regulatory region of the Apolipoprotein E gene and early-onset Alzheimer's disease. Neuroscience Letters, 1998, 258, 65-68.	2.1	38
102	Explorative genetic study of UBQLN2 and PFN1 in an extended Flanders-Belgian cohort of frontotemporal lobar degeneration patients. Neurobiology of Aging, 2013, 34, 1711.e1-1711.e5.	3.1	36
103	Familial Creutzfeldt-Jakob disease in a patient carrying both a presenilin 1 missense substitution and a prion protein gene insertion. Journal of Neurology, 2000, 247, 364-368.	3.6	32
104	Ataxin-2 polyQ expansions in FTLD-ALS spectrum disorders in Flanders-Belgian cohorts. Neurobiology of Aging, 2012, 33, 1004.e17-1004.e20.	3.1	32
105	APOE epsilon4 and Alzheimer's disease: positive association in a Colombian clinical series and review of the Latin-American studies. Arquivos De Neuro-Psiquiatria, 2001, 59, 11-17.	0.8	29
106	Genetic and physical characterization of the early-onset Alzheimer's disease AD3 locus on chromosome 14q24.3. Human Molecular Genetics, 1995, 4, 1355-1364.	2.9	27
107	Novel APP mutation V715A associated with presenile Alzheimer?s disease in a German family. Journal of Neurology, 2003, 250, 1374-1375.	3.6	26
108	Modifiers of GRN -Associated Frontotemporal Lobar Degeneration. Trends in Molecular Medicine, 2017, 23, 962-979.	6.7	26

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109	Islands of euchromatin-like sequence and expressed polymorphic sequences within the short arm of human chromosome 21. Genome Research, 2007, 17, 1690-1696.	5.5	25
110	APOLIPOPROTEIN E AND LONGEVITY: THE ROTTERDAM STUDY. Journal of the American Geriatrics Society, 2001, 49, 1258-1259.	2.6	24
111	A High-Resolution Physical Map of Human Chromosome 21p Using Yeast Artificial Chromosomes. Genome Research, 1999, 9, 1059-1073.	5.5	23
112	Familial primary lateral sclerosis or dementia associated with Arg573GlyTBK1mutation. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, 996-997.	1.9	23
113	Frontotemporal Lobar Degeneration with Ubiquitin-Positive Inclusions: A Molecular Genetic Update. Neurodegenerative Diseases, 2007, 4, 227-235.	1.4	21
114	A yeast artificial chromosome contig from human chromosome 14q24 spanning the Alzheimer's disease locus AD3. Human Molecular Genetics, 1995, 4, 1347-1354.	2.9	20
115	Missense mutation in exon 11 (codon 378) of the presenilin-1 gene in a French family with early-onset Alzheimer's disease and transmission study by mismatch enhanced allele specific amplification. Human Mutation, 1998, 11, 481-481.	2.5	19
116	Contribution of TARDBP to Alzheimer's Disease Genetic Etiology. Journal of Alzheimer's Disease, 2010, 21, 423-430.	2.6	19
117	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
118	The Genetics ofC9orf72Expansions. Cold Spring Harbor Perspectives in Medicine, 2018, 8, a026757.	6.2	19
119	Genetic analysis of the cellular oncogene fos in patients with chromosome 14 encoded Alzheimer's disease. Neuroscience Letters, 1994, 174, 97-100.	2.1	18
120	Novel PSEN1 Mutation in a Bulgarian Patient With Very Early-Onset Alzheimerâ€™s Disease, Spastic Paraparesis, and Extrapyramidal Signs. American Journal of Alzheimer's Disease and Other Dementias, 2009, 24, 404-407.	1.9	18
121	Clinical variability and onset age modifiers in an extended Belgian GRN founder family. Neurobiology of Aging, 2018, 67, 84-94.	3.1	17
122	Presenilin mutations in Alzheimer's disease. Human Mutation, 1998, 11, 183-190.	2.5	17
123	Genetic Testing Has No Place as a Routine Diagnostic Test in Sporadic and Familial Cases of Alzheimer's Disease. Journal of the American Geriatrics Society, 2004, 52, 2110-2113.	2.6	16
124	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
125	Visualization ofMAPT inversion on stretched chromosomes of tau-negative frontotemporal dementia patients. Human Mutation, 2006, 27, 1057-1059.	2.5	14
126	Investigating the role of filamin C in Belgian patients with frontotemporal dementia linked to GRN deficiency in FTLD-TDP brains. Acta Neuropathologica Communications, 2015, 3, 68.	5.2	13

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127	Reduced secretion and altered proteolytic processing caused by missense mutations in progranulin. Neurobiology of Aging, 2016, 39, 220.e17-220.e26.	3.1	11
128	Extended FTLD pedigree segregating a Belgian GRN-null mutation: neuropathological heterogeneity in one family. Alzheimer's Research and Therapy, 2018, 10, 7.	6.2	10
129	Mutation analysis of the chromosome 14q24.3 dihydrolipoyl succinyltransferase (DLST) gene in patients with early-onset Alzheimer disease. Neuroscience Letters, 1995, 199, 73-77.	2.1	9
130	Determination of the genomic organization of human presenilin 1 by fiber-FISH analysis and restriction mapping of cloned DNA. Mammalian Genome, 1999, 10, 410-414.	2.2	9
131	TheTNFRSF6 gene is not implicated in familial early-onset Alzheimer's disease. Human Genetics, 2001, 108, 552-553.	3.8	6
132	Data Mining: Applying the AD&FTD Mutation Database to Progranulin. Methods in Molecular Biology, 2018, 1806, 81-92.	0.9	6
133	Call for participation in the neurogenetics consortium within the Human Variome Project. Neurogenetics, 2011, 12, 169-173.	1.4	5
134	Subregional localization of the chromosome 21 loci D21S24 and D21S26 using physical mapping techniques. Human Genetics, 1991, 87, 109-111.	3.8	2
135	Molecular Pathogenesis of Frontotemporal Lobar Degeneration. Archives of Neurology, 2008, 65, 700-4.	4.5	2
136	A Major Genetic Factor at Chromosome 9p Implicated in Amyotrophic Lateral Sclerosis (ALS) and Frontotemporal Lobar Degeneration (FTLD). , 2012, , .		2
137	Genetics of frontotemporal dementia and related disorders. , 0, , 185-196.		1
138	Frontotemporal dementia. , 2017, , 199-249.		1
139	O3-02-07 Linkage analysis for AD using amyloid beta 42 levels shows evidence for a novel AD gene on chromosome 19. Neurobiology of Aging, 2004, 25, S56.	3.1	0
140	O4-06-03 A novel NR4A2 promoter variation associated with Parkinson's disease alters gene expression. Neurobiology of Aging, 2004, 25, S85.	3.1	0
141	Chromosome 17-linked Frontotemporal dementia with Ubiquitin-Positive, Tau-Negative Inclusions. Research and Perspectives in Alzheimer's Disease, 2005, , 117-137.	0.1	0
142	O3-13-03: Massive parallel gene panel sequencingÂin a belgian ftld cohort of causal genes associated with diverse neurodegenerative brain diseases. , 2015, 11, P251-P251.		0