Debby W Tsuang

List of Publications by Year
in descending order

170	24,242	15504 65	8630 146
papers	citations	h-index	g-index

175	175	175	28298
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	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
2	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
3	Common variants at MS4A4/MS4A6E, CD2AP, CD33 and EPHA1 are associated with late-onset Alzheimer's disease. Nature Genetics, 2011, 43, 436-441.	21.4	1,676
4	Two Phase 3 Trials of Bapineuzumab in Mild-to-Moderate Alzheimer's Disease. New England Journal of Medicine, 2014, 370, 322-333.	27.0	1,613
5	Mapping genomic loci implicates genes and synaptic biology in schizophrenia. Nature, 2022, 604, 502-508.	27.8	929
6	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
7	Review Article: Genetics of Alzheimer Disease. Journal of Geriatric Psychiatry and Neurology, 2010, 23, 213-227.	2.3	748
8	Spatial and Temporal Mapping of De Novo Mutations in Schizophrenia to a Fetal Prefrontal Cortical Network. Cell, 2013, 154, 518-529.	28.9	507
9	Neuropathological and genetic correlates of survival and dementia onset in synucleinopathies: a retrospective analysis. Lancet Neurology, The, 2017, 16, 55-65.	10.2	394
10	Co-morbidity of TDP-43 proteinopathy in Lewy body related diseases. Acta Neuropathologica, 2007, 114, 221-229.	7.7	378
11	Meta-analysis Confirms CR1, CLU, and PICALM as Alzheimer Disease Risk Loci and Reveals Interactions With APOE Genotypes. Archives of Neurology, 2010, 67, 1473.	4.5	376
12	Variants in the ATP-Binding Cassette Transporter (ABCA7), Apolipoprotein E Ïµ4, and the Risk of Late-Onset Alzheimer Disease in African Americans. JAMA - Journal of the American Medical Association, 2013, 309, 1483.	7.4	360
13	Clinicoâ€Neuropathological Correlation of Alzheimer's Disease in a Communityâ€Based Case Series. Journal of the American Geriatrics Society, 1999, 47, 564-569.	2.6	354
14	Initial Heritability Analyses of Endophenotypic Measures for Schizophrenia. Archives of General Psychiatry, 2007, 64, 1242.	12.3	351
15	Genome-Wide Association Meta-analysis of Neuropathologic Features of Alzheimer's Disease and Related Dementias. PLoS Genetics, 2014, 10, e1004606.	3.5	305
16	Common genetic variants in the CLDN2 and PRSS1-PRSS2 loci alter risk for alcohol-related and sporadic pancreatitis. Nature Genetics, 2012, 44, 1349-1354.	21.4	303
17	APOE Ïµ4 Increases Risk for Dementia in Pure Synucleinopathies. JAMA Neurology, 2013, 70, 223.	9.0	302
18	A novel Alzheimer disease locus located near the gene encoding tau protein. Molecular Psychiatry, 2016, 21, 108-117.	7.9	260

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19	Alzheimerâ€™s Disease: Analyzing the Missing Heritability. PLoS ONE, 2013, 8, e79771.	2.5	257
20	Analysis of 94 Candidate Genes and 12 Endophenotypes for Schizophrenia From the Consortium on the Genetics of Schizophrenia. American Journal of Psychiatry, 2011, 168, 930-946.	7.2	241
21	Cognitive differences in dementia patients with autopsy-verified AD, Lewy body pathology, or both. Neurology, 2005, 64, 2069-2073.	1.1	238
22	Brain Expression Genome-Wide Association Study (eGWAS) Identifies Human Disease-Associated Variants. PLoS Genetics, 2012, 8, e1002707.	3.5	225
23	Proteomic Identification of Novel Proteins in Cortical Lewy Bodies. Brain Pathology, 2007, 17, 139-145.	4.1	194
24	Glucocerebrosidase Gene Mutations. Archives of Neurology, 2008, 65, 379-82.	4.5	188
25	Assessment of the genetic variance of late-onset Alzheimer's disease. Neurobiology of Aging, 2016, 41, 200.e13-200.e20.	3.1	174
26	Convergent genetic and expression data implicate immunity in Alzheimer's disease. Alzheimer's and Dementia, 2015, 11, 658-671.	0.8	173
27	Effects of Multiple Genetic Loci on Age at Onset in Late-Onset Alzheimer Disease. JAMA Neurology, 2014, 71, 1394.	9.0	166
28	Transethnic genomeâ€wide scan identifies novel Alzheimer's disease loci. Alzheimer's and Dementia, 2017, 13, 727-738.	0.8	166
29	Modeling Deficits From Early Auditory Information Processing to Psychosocial Functioning in Schizophrenia. JAMA Psychiatry, 2017, 74, 37.	11.0	163
30	GBA Variants are associated with a distinct pattern of cognitive deficits in <scp>P</scp>arkinson's disease. Movement Disorders, 2016, 31, 95-102.	3.9	158
31	SNCA Variant Associated With Parkinson Disease and Plasma Î±-Synuclein Level. Archives of Neurology, 2010, 67, 1350-6.	4.5	157
32	Gene-Wide Analysis Detects Two New Susceptibility Genes for Alzheimer's Disease. PLoS ONE, 2014, 9, e94661.	2.5	155
33	Validation of mismatch negativity and P3a for use in multi-site studies of schizophrenia: Characterization of demographic, clinical, cognitive, and functional correlates in COGS-2. Schizophrenia Research, 2015, 163, 63-72.	2.0	154
34	Associations between Potentially Modifiable Risk Factors and Alzheimer Disease: A Mendelian Randomization Study. PLoS Medicine, 2015, 12, e1001841.	8.4	153
35	Comprehensive analysis of APOE and selected proximate markers for late-onset Alzheimer's disease: Patterns of linkage disequilibrium and disease/marker association. Genomics, 2007, 89, 655-665.	2.9	149
36	SORL1 Is Genetically Associated with Late-Onset Alzheimerâ€™s Disease in Japanese, Koreans and Caucasians. PLoS ONE, 2013, 8, e58618.	2.5	149

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37	Novel late-onset Alzheimer disease loci variants associate with brain gene expression. Neurology, 2012, 79, 221-228.	1.1	144
38	Novel Alzheimer Disease Risk Loci and Pathways in African American Individuals Using the African Genome Resources Panel. JAMA Neurology, 2021, 78, 102.	9.0	144
39	Serum cholesterol and risk of Alzheimer disease. Neurology, 2005, 65, 1045-1050.	1.1	140
40	GBA mutations increase risk for Lewy body disease with and without Alzheimer disease pathology. Neurology, 2012, 79, 1944-1950.	1.1	138
41	Clinical and Neuropathological Characteristics of Hippocampal Sclerosis. Archives of Neurology, 2002, 59, 1099.	4.5	136
42	The Consortium on the Genetics of Endophenotypes in Schizophrenia: Model Recruitment, Assessment, and Endophenotyping Methods for a Multisite Collaboration. Schizophrenia Bulletin, 2006, 33, 33-48.	4.3	134
43	Lewy Body Pathology in Familial Alzheimer Disease. Archives of Neurology, 2006, 63, 370.	4.5	122
44	Support for the N -Methyl-D-Aspartate Receptor Hypofunction Hypothesis of Schizophrenia From Exome Sequencing in Multiplex Families. JAMA Psychiatry, 2013, 70, 582.	11.0	119
45	Propranolol for Disruptive Behaviors in Nursing Home Residents With Probable or Possible Alzheimer Disease. Alzheimer Disease and Associated Disorders, 2005, 19, 23-28.	1.3	118
46	Abnormal Auditory N100 Amplitude: A Heritable Endophenotype in First-Degree Relatives of Schizophrenia Probands. Biological Psychiatry, 2008, 64, 1051-1059.	1.3	115
47	Genome-Wide Linkage Analyses of 12 Endophenotypes for Schizophrenia From the Consortium on the Genetics of Schizophrenia. American Journal of Psychiatry, 2013, 170, 521-532.	7.2	114
48	Salivary cortisol and memory function in human aging. Neurobiology of Aging, 2006, 27, 1705-1714.	3.1	113
49	Criterion validity of the Short Mood and Feelings Questionnaire and one- and two-item depression screens in young adolescents. Child and Adolescent Psychiatry and Mental Health, 2010, 4, 8.	2.5	113
50	Missense variant in TREML2 protects against Alzheimer's disease. Neurobiology of Aging, 2014, 35, 1510.e19-1510.e26.	3.1	110
51	Effect of Vascular Lesions on Cognition in Alzheimer's Disease: A Communityâ€Based Study. Journal of the American Geriatrics Society, 2004, 52, 1442-1448.	2.6	107
52	RESEARCH ARTICLE: Empiric Refinement of the Pathologic Assessment of Lewyâ€Related Pathology in the Dementia Patient. Brain Pathology, 2008, 18, 220-224.	4.1	106
53	MicroRNA in Alzheimerâ€™s disease: an exploratory study in brain, cerebrospinal fluid and plasma. Biomarkers, 2013, 18, 455-466.	1.9	102
54	Alzheimerâ€™s Disease Risk Polymorphisms Regulate Gene Expression in the ZCWPW1 and the CELF1 Loci. PLoS ONE, 2016, 11, e0148717.	2.5	99

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55	Comprehensive Search for Alzheimer Disease Susceptibility Loci in the APOE Region. Archives of Neurology, 2012, 69, 1270.	4.5	97
56	Verbal working memory impairments in individuals with schizophrenia and their first-degree relatives: Findings from the Consortium on the Genetics of Schizophrenia. Schizophrenia Research, 2008, 103, 218-228.	2.0	96
57	An 8-year follow-up of patients with DSM-III-R psychotic depression, schizoaffective disorder, and schizophrenia. American Journal of Psychiatry, 1993, 150, 1182-1188.	7.2	95
58	Two rare AKAP9 variants are associated with Alzheimer's disease in African Americans. Alzheimer's and Dementia, 2014, 10, 609.	0.8	94
59	Deficient prepulse inhibition in schizophrenia detected by the multi-site COGS. Schizophrenia Research, 2014, 152, 503-512.	2.0	91
60	Inhibition of the P50 cerebral evoked response to repeated auditory stimuli: Results from the Consortium on Genetics of Schizophrenia. Schizophrenia Research, 2010, 119, 175-182.	2.0	89
61	The utility of P300 as a schizophrenia endophenotype and predictive biomarker: Clinical and socio-demographic modulators in COGS-2. Schizophrenia Research, 2015, 163, 53-62.	2.0	87
62	Anticholinergic Medication Burdenâ€"Associated Cognitive Impairment in Schizophrenia. American Journal of Psychiatry, 2021, 178, 838-847.	7.2	80
63	Genome-wide Association of Endophenotypes for Schizophrenia From the Consortium on the Genetics of Schizophrenia (COGS) Study. JAMA Psychiatry, 2019, 76, 1274.	11.0	78
64	Examination of genetic linkage of chromosome 15 to schizophrenia in a large Veterans Affairs Cooperative Study sample. American Journal of Medical Genetics Part A, 2001, 105, 662-668.	2.4	75
65	Multiple SNPs Within and Surrounding the Apolipoprotein E Gene Influence Cerebrospinal Fluid Apolipoprotein E Protein Levels. Journal of Alzheimer's Disease, 2008, 13, 255-266.	2.6	75
66	ABCA7 frameshift deletion associated with Alzheimer disease in African Americans. Neurology: Genetics, 2016, 2, e79.	1.9	74
67	Successful multi-site measurement of antisaccade performance deficits in schizophrenia. Schizophrenia Research, 2007, 89, 320-329.	2.0	72
68	ABCC9 gene polymorphism is associated with hippocampal sclerosis of aging pathology. Acta Neuropathologica, 2014, 127, 825-843.	7.7	70
69	Impact of Sample Selection on APOE â^4 Allele Frequency: A Comparison of Two Alzheimer's Disease Samples. Journal of the American Geriatrics Society, 1996, 44, 704-707.	2.6	67
70	Genetic assessment of additional endophenotypes from the Consortium on the Genetics of Schizophrenia Family Study. Schizophrenia Research, 2016, 170, 30-40.	2.0	65
71	Familial Dementia With Lewy Bodies. Archives of Neurology, 2002, 59, 1622.	4.5	64
72	Attention/vigilance in schizophrenia: Performance results from a large multi-site study of the Consortium on the Genetics of Schizophrenia (COGS). Schizophrenia Research, 2015, 163, 38-46.	2.0	62

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73	Familial Aggregation of Psychotic Symptoms in Huntingtonâ€™s Disease. American Journal of Psychiatry, 2000, 157, 1955-1959.	7.2	61
74	Multi-site studies of acoustic startle and prepulse inhibition in humans: Initial experience and methodological considerations based on studies by the Consortium on the Genetics of Schizophrenia. Schizophrenia Research, 2007, 92, 237-251.	2.0	61
75	LRRK2 mutations and risk variants in Japanese patients with Parkinson's disease. Movement Disorders, 2009, 24, 1034-1041.	3.9	60
76	Polygenic risk scores in familial Alzheimer disease. Neurology, 2017, 88, 1180-1186.	1.1	59
77	Shared genetic contribution to ischemic stroke and Alzheimer's disease. Annals of Neurology, 2016, 79, 739-747.	5.3	56
78	Sodium lactate and hypertonic sodium chloride induce equivalent panic incidence, panic symptoms, and hypernatremia in panic disorder. Biological Psychiatry, 1998, 44, 1007-1016.	1.3	55
79	Arguing against the proposed definition changes of PD. Movement Disorders, 2016, 31, 1619-1622.	3.9	55
80	Factor structure and heritability of endophenotypes in schizophrenia: Findings from the Consortium on the Genetics of Schizophrenia (COGS-1). Schizophrenia Research, 2015, 163, 73-79.	2.0	52
81	Deficient prepulse inhibition in schizophrenia in a multi-site cohort: Internal replication and extension. Schizophrenia Research, 2018, 198, 6-15.	2.0	52
82	Genetic association between the APOE * 4 allele and Lewy bodies in Alzheimer disease. Neurology, 2005, 64, 509-513.	1.1	48
83	Genetically predicted body mass index and Alzheimer's diseaseâ€“related phenotypes in three large samples: Mendelian randomization analyses. Alzheimer's and Dementia, 2015, 11, 1439-1451.	0.8	46
84	INFERNO: inferring the molecular mechanisms of noncoding genetic variants. Nucleic Acids Research, 2018, 46, 8740-8753.	14.5	46
85	The Utility of Apolipoprotein E Genotyping in the Diagnosis of Alzheimer Disease in a Community-Based Case Series. Archives of Neurology, 1999, 56, 1489.	4.5	42
86	Amyloid precursor protein (APP) processing genes and cerebrospinal fluid APP cleavage product levels in Alzheimer's disease. Neurobiology of Aging, 2011, 32, 556.e13-556.e23.	3.1	42
87	Gating Deficit Heritability and Correlation With Increased Clinical Severity in Schizophrenia Patients With Positive Family History. American Journal of Psychiatry, 2016, 173, 385-391.	7.2	42
88	Detection of probable dementia cases in undiagnosed patients using structured and unstructured electronic health records. BMC Medical Informatics and Decision Making, 2019, 19, 128.	3.0	42
89	Recent Advances in the Genetics of Alzheimer's Disease. Journal of Geriatric Psychiatry and Neurology, 1998, 11, 42-54.	2.3	41
90	Initial Assessment of the Pathogenic Mechanisms of the Recently Identified Alzheimer Risk Loci. Annals of Human Genetics, 2013, 77, 85-105.	0.8	41

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91	Rarity of the Alzheimer Diseaseâ€“ProtectiveAPPA673T Variant in the United States. JAMA Neurology, 2015, 72, 209.	9.0	41
92	Neurocognitive performance in family-based and case-control studies of schizophrenia. Schizophrenia Research, 2015, 163, 17-23.	2.0	37
93	Group and site differences on the California Verbal Learning Test in persons with schizophrenia and their first-degree relatives: Findings from the Consortium on the Genetics of Schizophrenia (COGS). Schizophrenia Research, 2011, 128, 102-110.	2.0	35
94	Biological markers and diagnostic accuracy in the genetics of posttraumatic stress disorder. Psychiatry Research, 2001, 102, 203-215.	3.3	34
95	Comparison of the Heritability of Schizophrenia and Endophenotypes in the COGS-1 Family Study. Schizophrenia Bulletin, 2014, 40, 1404-1411.	4.3	34
96	Evaluation of Selection Bias in an Incident-Based Dementia Autopsy Case Series. Alzheimer Disease and Associated Disorders, 2005, 19, 67-73.	1.3	33
97	Selective dendritic degeneration of medium spiny neurons in dementia with Lewy bodies. Neurology, 2006, 66, 1591-1593.	1.1	32
98	The Effect of Algorithms on Copy Number Variant Detection. PLoS ONE, 2010, 5, e14456.	2.5	32
99	Linkage of chromosome 13q32 to schizophrenia in a large veterans affairs cooperative study sample. American Journal of Medical Genetics Part A, 2002, 114, 598-604.	2.4	30
100	Familial Occurrence of Dementia With Lewy Bodies. American Journal of Geriatric Psychiatry, 2004, 12, 179-188.	1.2	30
101	Antisaccade performance in schizophrenia patients, their first-degree biological relatives, and community comparison subjects: Data from the COGS study. Psychophysiology, 2010, 47, 846-56.	2.4	30
102	Visual Hallucinations in Dementia: A Prospective Community-Based Study With Autopsy. American Journal of Geriatric Psychiatry, 2009, 17, 317-323.	1.2	29
103	Association Between Lifetime Cigarette Smoking and Lewy Body Accumulation. Brain Pathology, 2010, 20, 412-418.	4.1	29
104	Modest evidence for linkage and possible confirmation of association between NOTCH4 and schizophrenia in a large veterans affairs cooperative study sample. American Journal of Medical Genetics Part A, 2003, 118B, 8-15.	2.4	28
105	Evidence for involvement of GNB1L in autism. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 61-71.	1.7	28
106	Nextâ€generation sequencing in schizophrenia and other neuropsychiatric disorders. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 671-678.	1.7	28
107	Familial aggregation of schizophrenia-like symptoms in huntington's disease. American Journal of Medical Genetics Part A, 1998, 81, 323-327.	2.4	27
108	Familial Dementia with Lewy Bodies with an Atypical Clinical Presentation. Journal of Geriatric Psychiatry and Neurology, 2003, 16, 59-64.	2.3	27

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109	Verbal working memory in schizophrenia from the Consortium on the Genetics of Schizophrenia (COGS) Study: The moderating role of smoking status and antipsychotic medications. Schizophrenia Research, 2015, 163, 24-31.	2.0	26
110	Lewy body pathology in late-onset familial Alzheimer's disease: A clinicopathological case series. Journal of Alzheimer's Disease, 2006, 9, 235-242.	2.6	25
111	Enhancing the Power of Genetic Association Studies through the Use of Silver Standard Cases Derived from Electronic Medical Records. PLoS ONE, 2013, 8, e63481.	2.5	23
112	Genetics of dementia. Medical Clinics of North America, 2002, 86, 591-614.	2.5	22
113	ADAM10 expression and promoter haplotype in Alzheimer's disease. Neurobiology of Aging, 2012, 33, 2229.e1-2229.e9.	3.1	22
114	Genetic Variation in Genes Underlying Diverse Dementias May Explain a Small Proportion of Cases in the Alzheimerâ€™s Disease Sequencing Project. Dementia and Geriatric Cognitive Disorders, 2018, 45, 1-17.	1.5	22
115	Analysis of the LRRK2 G2019S Mutation in Alzheimer Disease. Archives of Neurology, 2006, 63, 156.	4.5	21
116	Nonlinear dynamics underlying sensory processing dysfunction in schizophrenia. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 3847-3852.	7.1	21
117	The effects of age and sex on cognitive impairment in schizophrenia: Findings from the Consortium on the Genetics of Schizophrenia (COGS) study. PLoS ONE, 2020, 15, e0232855.	2.5	21
118	Genetic counseling for psychiatric disorders. Current Psychiatry Reports, 2001, 3, 138-143.	4.5	20
119	Genome scan of schizophrenia families in a large Veterans Affairs Cooperative Study sample: Evidence for linkage to 18p11.32 and for racial heterogeneity on chromosomes 6 and 14. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2005, 139B, 91-100.	1.7	20
120	Predicting Lewy Body Pathology in a Community-Based Sample With Clinical Diagnosis of Alzheimerâ€™s Disease. Journal of Geriatric Psychiatry and Neurology, 2006, 19, 195-201.	2.3	20
121	Quantitation and Mapping of Cerebral Detergentâ€insoluble Proteins in the Elderly. Brain Pathology, 2009, 19, 365-374.	4.1	17
122	Sex Differences in Familiality Effects on Neurocognitive Performance in Schizophrenia. Biological Psychiatry, 2013, 73, 976-984.	1.3	17
123	APOE DNA methylation is altered in Lewy body dementia. Alzheimer's and Dementia, 2018, 14, 889-894.	0.8	17
124	Tau phosphorylation pathway genes and cerebrospinal fluid tau levels in Alzheimer's disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 874-883.	1.7	16
125	Robust differences in antisaccade performance exist between COGS schizophrenia cases and controls regardless of recruitment strategies. Schizophrenia Research, 2015, 163, 47-52.	2.0	16
126	Genetic association between APOE*4 and neuropsychiatric symptoms in patients with probable Alzheimer's disease is dependent on the psychosis phenotype. Behavioral and Brain Functions, 2012, 8, 62.	3.3	15

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127	Genetic variants associated with susceptibility to psychosis inÂlate-onset Alzheimerâ€™s disease families. Neurobiology of Aging, 2015, 36, 3116.e9-3116.e16.	3.1	14
128	Cerebrospinal fluid AÎ²₄₂ levels and APP processing pathway genes in Parkinson's disease. Movement Disorders, 2015, 30, 936-944.	3.9	14
129	Endophenotypes in Schizophrenia: Digging Deeper to Identify Genetic Mechanisms. Journal of Psychiatry and Brain Science, 2019, 4, .	0.5	14
130	Prioritizing schizophrenia endophenotypes for future genetic studies: An example using data from the COGS-1 family study. Schizophrenia Research, 2016, 174, 1-9.	2.0	13
131	Alzheimerâ€™s Disease and Alzheimerâ€™s Disease-Related Dementias in Older African American and White Veterans. Journal of Alzheimer's Disease, 2020, 75, 311-320.	2.6	13
132	California Verbal Learning Test-II performance in schizophrenia as a function of ascertainment strategy: Comparing the first and second phases of the Consortium on the Genetics of Schizophrenia (COGS). Schizophrenia Research, 2015, 163, 32-37.	2.0	12
133	Manifestations of Alzheimerâ€™s disease genetic risk in the blood are evident in a multiomic analysis in healthy adults aged 18 to 90. Scientific Reports, 2022, 12, 6117.	3.3	12
134	Is There an Association between Advanced Paternal Age and Endophenotype Deficit Levels in Schizophrenia?. PLoS ONE, 2014, 9, e88379.	2.5	11
135	Familial occurrence of dementia with Lewy bodies. American Journal of Geriatric Psychiatry, 2004, 12, 179-88.	1.2	10
136	A Genetic Study of Psychosis in Huntingtonâ€™s Disease: Evidence for the Involvement of Glutamate Signaling Pathways. Journal of Huntington's Disease, 2018, 7, 51-59.	1.9	9
137	Suicide and Lewy body dementia: Report of a Lewy body dementia association working group. International Journal of Geriatric Psychiatry, 2021, 36, 373-382.	2.7	9
138	The National Institute on Aging Lateâ€Onset Alzheimer's Disease Family Based Study: A resource for genetic discovery. Alzheimer's and Dementia, 2022, 18, 1889-1897.	0.8	9
139	Cognitive Impairment in Older Adults Without Dementia: Clinical and Pathologic Outcomes in a Community-Based Sample. Journal of Geriatric Psychiatry and Neurology, 2009, 22, 256-265.	2.3	8
140	Alzheimerâ€™s Disease Genetics. Current Behavioral Neuroscience Reports, 2014, 1, 191-196.	1.3	8
141	The Genetics of Alzheimerâ€™s Disease and Parkinsonâ€™s Disease. Advances in Neurobiology, 2011, , 695-755.	1.8	7
142	Testing the Validity of the Neurotic Depression Concept. Journal of Nervous and Mental Disease, 1992, 180, 446-450.	1.0	6
143	Association of cerebrospinal fluid AÎ²42 with A2M gene in cognitively normal subjects. Neurobiology of Aging, 2014, 35, 357-364.	3.1	6
144	The effect of apolipoprotein E genotype on expression of an autosomal dominant schizophreniform disorder with progressive dementia and neurofibrillary tangles. Biological Psychiatry, 1997, 41, 191-195.	1.3	5

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145	An Algorithm to Construct Genetically Similar Subsets of Families with the Use of Self-Reported Ethnicity Information. American Journal of Human Genetics, 2005, 77, 346-354.	6.2	5
146	The UCHL1 S18Y polymorphism and Parkinsonâ€™s disease in a Japanese population. Parkinsonism and Related Disorders, 2011, 17, 473-475.	2.2	5
147	Genetic factors in neurodegenerative diseases. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2017, 174, 3-4.	1.7	3
148	Heritability of acoustic startle magnitude and latency from the consortium on the genetics of schizophrenia. Schizophrenia Research, 2020, 224, 33-39.	2.0	3
149	Frontal-Complex Partial Status Epilepticus Misdiagnosed as Bipolar Affective Disorder in a 75-Year-Old Man. Journal of Geriatric Psychiatry and Neurology, 1997, 10, 158-160.	2.3	2
150	Inheritance Model Introduces Differential Bias in <scp>CNV</scp> Calls Between Parents and Offspring. Genetic Epidemiology, 2012, 36, 488-498.	1.3	2
151	Paternal age of schizophrenia probands and endophenotypic differences from unaffected siblings. Psychiatry Research, 2014, 219, 67-71.	3.3	2
152	Challenges of the Faculty Career for Womenby Maike Ingrid Philipsen. Foreword by Mary Deane Sorcinelli. Hoboken, N.J., Jossey-Bass, 2008, 368 pp., $40.00.. American Journal of Psychiatry, 2009, 166, 1192-1193.	7.2	1
153	P4â€559: DETECTION OF PROBABLE DEMENTIA CASES IN UNDIAGNOSED PATIENTS USING STRUCTURED AND UNSTRUCTURED ELECTRONIC HEALTH RECORDS. Alzheimer's and Dementia, 2019, 15, P1533.	0.8	1
154	Mild COVID-19 Disease Course With Protracted Delirium in a Cognitively Impaired Patient Over the Age of 85 Years. primary care companion for CNS disorders, The, 2020, 22, .	0.6	1
155	Response to the Validity of the Neurotic Depression Concept. Journal of Nervous and Mental Disease, 1993, 181, 459.	1.0	0
156	Genetics of Dementia. Journal of Geriatric Psychiatry and Neurology, 1998, 11, 41-41.	2.3	0
157	Dementia with Lewy Bodies. , 0, , 472-489.		0
158	Updates on the Genetics of Neurodegenerative Disorders. Journal of Geriatric Psychiatry and Neurology, 2010, 23, 211-212.	2.3	0
159	How Genes Influence Behaviorby FlintJonathan, GreenspanRalph J., and KendlerKenneth S., New York, Oxford University Press, 2010, 304 pp., $52.95.. American Journal of Psychiatry, 2011, 168, 656-657.	7.2	0
160	Visualization of Haplotype Sharing Patterns in Pedigree Samples. Human Heredity, 2014, 78, 1-8.	0.8	0
161	P1-059: MAPT haplotypes modify the association between head injury and risk of Alzheimer's disease. , 2015, 11, P361-P361.		0
162	P4â€326: Automated Machine Learning Methods to Dectect Undiagnosed Cognitive Impairment Using Electronic Medical Records. Alzheimer's and Dementia, 2016, 12, P1159.	0.8	0

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163	Cognitive trajectory changes in African American veterans with combat PTSD. Alzheimer's and Dementia, 2020, 16, e047359.	0.8	0
164	Mild disease course in cognitively impaired oldest old individuals with COVIDâ€19: A description of two cases. Alzheimer's and Dementia, 2020, 16, e047558.	0.8	0
165	The Role of Genetic Counseling. , 2004, , 325-336.		0
166	Objective home sleep profiles differentiate Alzheimer disease from Î±â€synucleinopathies. Alzheimer's and Dementia, 2021, 17, .	0.8	0
167	Title is missing!. , 2020, 15, e0232855.		0
168	Title is missing!. , 2020, 15, e0232855.		0
169	Title is missing!. , 2020, 15, e0232855.		0
170	Title is missing!. , 2020, 15, e0232855.		0