## Joseph H Lee

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3353471/publications.pdf

Version: 2024-02-01

201 papers 10,409 citations

44069 48 h-index 95 g-index

207 all docs

207 docs citations

times ranked

207

15445 citing authors

#	Article	IF	CITATIONS
1	Large-scale meta-analysis of genome-wide association data identifies six new risk loci for Parkinson's disease. Nature Genetics, 2014, 46, 989-993.	21.4	1,685
2	The neuronal sortilin-related receptor SORL1 is genetically associated with Alzheimer disease. Nature Genetics, 2007, 39, 168-177.	21.4	1,045
3	Localization of a gene for partial epilepsy to chromosome 10q. Nature Genetics, 1995, 10, 56-60.	21.4	342
4	Genome-Wide Association of Familial Late-Onset Alzheimer's Disease Replicates BIN1 and CLU and Nominates CUGBP2 in Interaction with APOE. PLoS Genetics, 2011, 7, e1001308.	3.5	223
5	Genome Scan for Human Obesity and Linkage to Markers in 20q13. American Journal of Human Genetics, 1999, 64, 196-209.	6.2	218
6	Prevalence of Mental Retardation and Developmental Disabilities: Estimates From the 1994/1995 National Health Interview Survey Disability Supplements. American Journal on Intellectual and Developmental Disabilites, 2001, 106, 231.	2.4	210
7	Shorter telomeres are associated with mortality in those withAPOEϵ4 and dementia. Annals of Neurology, 2006, 60, 181-187.	<b>5.</b> 3	176
8	Coding mutations in <scp><i>SORL</i> and <scp>A</scp> zheimer disease. Annals of Neurology, 2015, 77, 215-227.</scp>	<b>5.</b> 3	168
9	Health and function of participants in the Long Life Family Study: A comparison with other cohorts. Aging, 2011, 3, 63-76.	3.1	163
10	Identification of Novel Loci for Alzheimer Disease and Replication of CLU, PICALM, and BIN1 in Caribbean Hispanic Individuals. Archives of Neurology, 2011, 68, 320-8.	4.5	160
11	Prevalence of Chronic Medical Conditions in Adults With Mental Retardation: Comparison With the General Population. Mental Retardation, 1998, 36, 269-279.	1.0	158
12	Association of Shorter Leukocyte Telomere Repeat Length With Dementia and Mortality. Archives of Neurology, 2012, 69, 1332.	4.5	155
13	Meta-analysis of the Association Between Variants in SORL1 and Alzheimer Disease. Archives of Neurology, 2011, 68, 99.	<b>4.</b> 5	153
14	Genome-Wide association study identifies candidate genes for Parkinson's disease in an Ashkenazi Jewish population. BMC Medical Genetics, 2011, 12, 104.	2.1	149
15	Integrative genomics identifies APOE Îμ4 effectors in Alzheimer's disease. Nature, 2013, 500, 45-50.	27.8	148
16	The Association Between Genetic Variants in SORL1 and Alzheimer Disease in an Urban, Multiethnic, Community-Based Cohort. Archives of Neurology, 2007, 64, 501.	<b>4.</b> 5	141
17	Sequence variants in the $5\hat{a}\in^2$ flanking region of the leptin gene are associated with obesity in women. Annals of Human Genetics, 1999, 63, 227-234.	0.8	129
18	Rare coding mutations identified by sequencing of <scp>A</scp> Izheimer disease genomeâ€wide association studies loci. Annals of Neurology, 2015, 78, 487-498.	<b>5.</b> 3	126

#	Article	IF	CITATIONS
19	Analyses of the National Institute on Aging Late-Onset Alzheimer's Disease Family Study. Archives of Neurology, 2008, 65, 1518.	4.5	125
20	Relationship Between Plasma Lipids and Allâ€Cause Mortality in Nondemented Elderly. Journal of the American Geriatrics Society, 2005, 53, 219-226.	2.6	121
21	Increased risk of Alzheimer's disease in mothers of adults with Down's syndrome. Lancet, The, 1994, 344, 353-356.	13.7	112
22	Associations of Mitochondrial and Nuclear Mitochondrial Variants and Genes with Seven Metabolic Traits. American Journal of Human Genetics, 2019, 104, 112-138.	6.2	106
23	SORCS1 alters amyloid precursor protein processing and variants may increase Alzheimer's disease risk. Annals of Neurology, 2011, 69, 47-64.	<b>5.</b> 3	104
24	Outcome of pancreaticoduodenectomy and impact of adjuvant therapy for ampullary carcinomas. International Journal of Radiation Oncology Biology Physics, 2000, 47, 945-953.	0.8	99
25	Decline in cognitive and functional skills increases mortality risk in nondemented elderly. Neurology, 2005, 65, 1218-1226.	1.1	98
26	Early menopause in women with Down's syndrome. Journal of Intellectual Disability Research, 1997, 41, 264-267.	2.0	98
27	A Founder Mutation in Presenilin 1 Causing Early-Onset Alzheimer Disease in Unrelated Caribbean Hispanic Families. JAMA - Journal of the American Medical Association, 2001, 286, 2257.	7.4	95
28	Onset of dementia is associated with apolipoprotein E $\hat{l}\mu4$ in Down's syndrome. Annals of Neurology, 1996, 40, 799-801.	5.3	89
29	Multiancestry Genome-Wide Association Study of Lipid Levels Incorporating Gene-Alcohol Interactions. American Journal of Epidemiology, 2019, 188, 1033-1054.	3.4	85
30	Association between SORL1 and Alzheimer's disease in a genome-wide study. NeuroReport, 2007, 18, 1761-1764.	1.2	83
31	Familial risk ratios for extreme obesity: implications for mapping human obesity genes. International Journal of Obesity, 1997, 21, 935-940.	3.4	82
32	Tau Proteins Are Abnormally Expressed in Olfactory Epithelium of Alzheimer Patients and Developmentally Regulated in Human Fetal Spinal Cord. Experimental Neurology, 1993, 121, 93-105.	4.1	79
33	ASSOCIATION BETWEEN GENETIC VARIANTS IN <i>SORL1</i> Neurology, 2008, 70, 887-889.	1.1	73
34	Replication of the LINGO1 gene association with essential tremor in a North American population. European Journal of Human Genetics, 2010, 18, 838-843.	2.8	69
35	Heritability of Different Forms of Memory in the Late Onset Alzheimer's Disease Family Study. Journal of Alzheimer's Disease, 2011, 23, 249-255.	2.6	68
36	Chromosome-12 Mapping of Late-Onset Alzheimer Disease among Caribbean Hispanics. American Journal of Human Genetics, 2002, 70, 237-243.	6.2	66

#	Article	IF	CITATIONS
37	Clinical Indicators of Genetic Susceptibility to Epilepsy. Epilepsia, 1996, 37, 353-361.	5.1	65
38	Association studies of cholesterol metabolism genes (CH25H, ABCA1 and CH24H) in Alzheimer's disease. Neuroscience Letters, 2006, 391, 142-146.	2.1	64
39	Families Enriched for Exceptional Longevity also have Increased Health-Span: Findings from the Long Life Family Study. Frontiers in Public Health, 2013, 1, 38.	2.7	63
40	Heritability of telomere length in a study of long-lived families. Neurobiology of Aging, 2015, 36, 2785-2790.	3.1	61
41	Mediterranean diet and leukocyte telomere length in a multi-ethnic elderly population. Age, 2015, 37, 24.	3.0	61
42	Genome wide association and linkage analyses identified three loci—4q25, 17q23.2, and 10q11.21—associated with variation in leukocyte telomere length: the Long Life Family Study. Frontiers in Genetics, 2013, 4, 310.	2.3	60
43	Four New Families with Autosomal Dominant Partial Epilepsy with Auditory Features: Clinical Description and Linkage to Chromosome 10q24. Epilepsia, 2002, 43, 60-67.	5.1	59
44	Specificity of the fivefold increase in AD in mothers of adults with Down syndrome. Neurology, 2001, 57, 979-984.	1.1	58
45	Leptin resistance is associated with extreme obesity and aggregates in families. International Journal of Obesity, 2001, 25, 1471-1473.	3.4	55
46	Familial Alzheimer Disease Among Caribbean Hispanics. Archives of Neurology, 2002, 59, 87.	4.5	55
47	Fâ€box/ <scp>LRR</scp> â€repeat protein 7 is genetically associated with Alzheimer's disease. Annals of Clinical and Translational Neurology, 2015, 2, 810-820.	3.7	54
48	Reliability of seizure classification using a semistructured interview. Neurology, 1993, 43, 2526-2526.	1.1	53
49	Familial Alzheimer disease in Latinos: Interaction between APOE, stroke, and estrogen replacement. Neurology, 2006, 66, 35-40.	1.1	52
50	Gene-Wise Association of Variants in Four Lysosomal Storage Disorder Genes in Neuropathologically Confirmed Lewy Body Disease. PLoS ONE, 2015, 10, e0125204.	2.5	52
51	Glaucomatous Optic Neuropathy Associated with Nocturnal Dip in Blood Pressure. Ophthalmology, 2018, 125, 807-814.	5.2	52
52	Genetic Modifiers of Age at Onset in Carriers of the G206A Mutation in <i>PSEN1</i> With Familial Alzheimer Disease Among Caribbean Hispanics. JAMA Neurology, 2015, 72, 1043.	9.0	50
53	Are Generalized and Localization-Related Epilepsies Genetically Distinct?. Archives of Neurology, 1998, 55, 339.	4.5	49
54	Genome-Wide Survey of Large Rare Copy Number Variants in Alzheimer's Disease Among Caribbean Hispanics. G3: Genes, Genomes, Genetics, 2012, 2, 71-78.	1.8	49

#	Article	IF	CITATIONS
55	The Alzheimer's Biomarker Consortiumâ€Down Syndrome: Rationale and methodology. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2020, 12, e12065.	2.4	49
56	Genetic Evidence for Cognitive Reserve: Variations in Memory and Related Cognitive Functions. Journal of Clinical and Experimental Neuropsychology, 2003, 25, 594-613.	1.3	47
57	Fine mapping of 10q and 18q for familial Alzheimer's disease in Caribbean Hispanics. Molecular Psychiatry, 2004, 9, 1042-1051.	7.9	47
58	Early menopause in women with Down's syndrome. Journal of Intellectual Disability Research, 1997, 41, 264-267.	2.0	45
59	Mortality and Apolipoprotein E in Hispanic, African-American, and Caucasian elders. American Journal of Medical Genetics Part A, 2001, 103, 121-127.	2.4	45
60	APOE and APOC1 Promoter Polymorphisms and the Risk of Alzheimer Disease in African American and Caribbean Hispanic Individuals. Archives of Neurology, 2004, 61, 1434.	4.5	44
61	Association between genetic variants in sortilin-related receptor 1 (SORL1) and Alzheimer's disease in adults with Down syndrome. Neuroscience Letters, 2007, 425, 105-109.	2.1	44
62	A Recessive Gene for Primary Vesicoureteral Reflux Maps to Chromosome 12p11-q13. Journal of the American Society of Nephrology: JASN, 2009, 20, 1633-1640.	6.1	42
63	Variants in CYP17 and CYP19 Cytochrome P450 Genes are Associated with Onset of Alzheimer's Disease in Women with Down Syndrome. Journal of Alzheimer's Disease, 2012, 28, 601-612.	2.6	42
64	Excess Synaptojanin 1 Contributes to Place Cell Dysfunction and Memory Deficits in the Aging Hippocampus in Three Types of Alzheimer's Disease. Cell Reports, 2018, 23, 2967-2975.	6.4	41
65	The neuronal sortilin-related receptor gene SORL1 and late-onset Alzheimer's disease. Current Neurology and Neuroscience Reports, 2008, 8, 384-391.	4.2	39
66	Association of Long Runs of Homozygosity With Alzheimer Disease Among African American Individuals. JAMA Neurology, 2015, 72, 1313.	9.0	39
67	Evidence of Recessive Alzheimer Disease Loci in a Caribbean Hispanic Data Set. JAMA Neurology, 2013, 70, 1261-7.	9.0	37
68	Birth Cohort and Familial Risk of Epilepsy: The Effect of Diminished Recall in Studies of Lifetime Prevalence. American Journal of Epidemiology, 1995, 141, 235-241.	3.4	36
69	Polymorphisms in the Promoter of the Human APP Gene. Archives of Neurology, 2002, 59, 1793.	4.5	36
70	Apolipoprotein E $\hat{l}\mu 4$ and Age at Onset of Sporadic and Familial Alzheimer Disease in Caribbean Hispanics. Archives of Neurology, 2006, 63, 1586.	4.5	36
71	Further examination of the candidate genes in chromosome 12p13 locus for late-onset Alzheimer disease. Neurogenetics, 2008, 9, 127-138.	1.4	36
72	Diseaseâ€related mutations among Caribbean Hispanics with familial dementia. Molecular Genetics & Cenomic Medicine, 2014, 2, 430-437.	1.2	36

#	Article	IF	CITATIONS
73	Familial Aggregation of Dementia With Lewy Bodies. Archives of Neurology, 2011, 68, 90-3.	4.5	35
74	Expanded Genomewide Scan Implicates a Novel Locus at 3q28 Among Caribbean Hispanics With Familial Alzheimer Disease. Archives of Neurology, 2006, 63, 1591.	4.5	34
75	Relation of plasma lipids to all-cause mortality in Caucasian, African-American and Hispanic elders. Age and Ageing, 2008, 37, 207-213.	1.6	34
76	Substantial fat mass loss reduces low-grade inflammation and induces positive alteration in cardiometabolic factors in normal-weight individuals. Scientific Reports, 2019, 9, 3450.	3.3	34
77	Alzheimerâ€Related Cerebrovascular Disease in Down Syndrome. Annals of Neurology, 2020, 88, 1165-1177.	5.3	34
78	Genetic influences on memory performance in familial Alzheimer disease. Neurology, 2004, 62, 414-421.	1.1	33
79	Molecular Pathways Mediating Immunosuppression in Response to Prolonged Intensive Physical Training, Low-Energy Availability, and Intensive Weight Loss. Frontiers in Immunology, 2019, 10, 907.	4.8	33
80	A multi-ancestry genome-wide study incorporating gene–smoking interactions identifies multiple new loci for pulse pressure and mean arterial pressure. Human Molecular Genetics, 2019, 28, 2615-2633.	2.9	31
81	Mutations in the Parkinson's disease genes, Leucine Rich Repeat Kinase 2 (LRRK2) and Glucocerebrosidase (GBA), are not associated with essential tremor. Parkinsonism and Related Disorders, 2010, 16, 132-135.	2.2	30
82	Obesity related phenotypes in families selected for extreme obesity and leanness. International Journal of Obesity, 1998, 22, 406-413.	3.4	29
83	Comparison of Clinical Manifestations in Alzheimer Disease and Dementia With Lewy Bodies. Archives of Neurology, 2008, 65, 1634-9.	4.5	27
84	Incidence of dementia in elderly Latin Americans: Results of the Maracaibo Aging Study. , $2018, 14, 140-147$ .		27
85	Possible association between schizophrenia and a CAG repeat polymorphism in the spinocerebellar ataxia type 1 (SCA1) gene on human chromosome 6p23. Psychiatric Genetics, 1999, 9, 7-12.	1.1	26
86	Age-at-onset linkage analysis in Caribbean Hispanics with familial late-onset Alzheimer's disease. Neurogenetics, 2008, 9, 51-60.	1.4	26
87	Estrogen Receptor-α Variants Increase Risk of Alzheimer's Disease in Women with Down Syndrome. Dementia and Geriatric Cognitive Disorders, 2008, 25, 476-482.	1.5	26
88	Increased rate of sporadic and recurrent rare genic copy number variants in P arkinson's disease among A shkenazi J ews. Molecular Genetics & Enomic Medicine, 2013, 1, 142-154.	1.2	26
89	Comparison of CSF biomarkers in Down syndrome and autosomal dominant Alzheimer's disease: a cross-sectional study. Lancet Neurology, The, 2021, 20, 615-626.	10.2	26
90	Candidate genes for Alzheimer's disease are associated with individual differences in plasma levels of beta amyloid peptides in adults with Down syndrome. Neurobiology of Aging, 2015, 36, 2907.e1-2907.e10.	3.1	25

#	Article	IF	Citations
91	Telomere length is longer in women with late maternal age. Menopause, 2017, 24, 497-501.	2.0	25
92	White matter hyperintensities mediate the association of nocturnal blood pressure with cognition. Neurology, 2020, 94, e1803-e1810.	1.1	25
93	Aromatase Variants Modify Risk for Alzheimer's Disease in a Multiethnic Female Cohort. Dementia and Geriatric Cognitive Disorders, 2013, 35, 340-350.	1.5	24
94	Linkage analyses in Caribbean Hispanic families identify novel loci associated with familial lateâ€onset Alzheimer's disease. Alzheimer's and Dementia, 2015, 11, 1397-1406.	0.8	24
95	Prevalence and Associated Risk Factors of Chronic Kidney Disease in an Elderly Population from Eastern China. International Journal of Environmental Research and Public Health, 2019, 16, 4383.	2.6	24
96	Promising outcome measures of early Alzheimer's dementia in adults with Down syndrome. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2020, 12, e12044.	2.4	24
97	Genetic influences on life span and survival among elderly African-Americans, Caribbean Hispanics, and Caucasians. American Journal of Medical Genetics Part A, 2004, 128A, 159-164.	2.4	23
98	Plasma Total-Tau and Neurofilament Light Chain as Diagnostic Biomarkers of Alzheimer's Disease Dementia and Mild Cognitive Impairment in Adults with Down Syndrome. Journal of Alzheimer's Disease, 2021, 79, 671-681.	2.6	23
99	Estrogen Receptor-Beta Variants Are Associated with Increased Risk of Alzheimer's Disease in Women with Down Syndrome. Dementia and Geriatric Cognitive Disorders, 2011, 32, 241-249.	1.5	22
100	Estrogen Receptor α Variants Affect Age at Onset of Alzheimer's Disease in a Multiethnic Female Cohort. Dementia and Geriatric Cognitive Disorders, 2014, 38, 200-213.	1.5	22
101	Genome-wide association study identifies common loci influencing circulating glycated hemoglobin (HbA1c) levels in non-diabetic subjects: The Long Life Family Study (LLFS). Metabolism: Clinical and Experimental, 2014, 63, 461-468.	3.4	22
102	Candidate gene analysis for Alzheimer's disease in adults with Down syndrome. Neurobiology of Aging, 2017, 56, 150-158.	3.1	22
103	Preservation of cognitive and functional ability as markers of longevity. Neurobiology of Aging, 2004, 25, 1231-1240.	3.1	21
104	Association between variants in IDE-KIF11-HHEX and plasma amyloid $\hat{l}^2$ levels. Neurobiology of Aging, 2012, 33, 199.e13-199.e17.	3.1	21
105	Increased low-level chromosome 21 mosaicism in older individuals with Down syndrome. , 1997, 68, 147-151.		20
106	Inbreeding among Caribbean Hispanics from the Dominican Republic and its effects on risk of Alzheimer disease. Genetics in Medicine, 2015, 17, 639-643.	2.4	20
107	Risk Factors for Orthostatic Hypotension: Differences Between Elderly Men and Women. American Journal of Hypertension, 2018, 31, 797-803.	2.0	20
108	Rare Variants Imputation in Admixed Populations: Comparison Across Reference Panels and Bioinformatics Tools. Frontiers in Genetics, 2019, 10, 239.	2.3	20

#	Article	IF	CITATIONS
109	Segregation analysis of cryptogenic epilepsy and an empirical test of the validity of the results. American Journal of Human Genetics, 1997, 60, 667-75.	6.2	20
110	The Peroxisome Proliferator-Activated Receptor $\hat{I}^3$ 2 Pro12Ala Mutation Is Associated with Early Onset Extreme Obesity and Reduced Fasting Glucose. Molecular Genetics and Metabolism, 2000, 70, 159-161.	1.1	19
111	Polymorphisms in <i>HSD17B1</i> : Early Onset and Increased Risk of Alzheimer's Disease in Women with Down Syndrome. Current Gerontology and Geriatrics Research, 2012, 2012, 1-8.	1.6	18
112	Epidemiology of estrogen and dementia in women with Down syndrome. Free Radical Biology and Medicine, 2018, 114, 62-68.	2.9	18
113	Association between Atmospheric Particulate Pollutants and Mortality for Cardio-Cerebrovascular Diseases in Chinese Korean Population: A Case-Crossover Study. International Journal of Environmental Research and Public Health, 2018, 15, 2835.	2.6	18
114	Impact of Genetic Variation in SORCS1 on Memory Retention. PLoS ONE, 2011, 6, e24588.	2.5	17
115	Association of common variants, not rare mutations, in <i>IRF6</i> With nonsyndromic clefts in a honduran population. Laryngoscope, 2011, 121, 1756-1759.	2.0	17
116	Estrogen Receptor $\hat{l}^2$ Variants Modify Risk for Alzheimer's Disease in a Multiethnic Female Cohort. Journal of Alzheimer's Disease, 2014, 40, 83-93.	2.6	17
117	Candidate gene resequencing to identify rare, pedigree-specific variants influencing healthy aging phenotypes in the long life family study. BMC Geriatrics, 2016, 16, 80.	2.7	17
118	Possible Association between SORL1 and Alzheimer Disease?. Dementia and Geriatric Cognitive Disorders, 2008, 26, 482-482.	1.5	16
119	"Predicting―parental longevity from offspring endophenotypes: Data from the Long Life Family Study (LLFS). Mechanisms of Ageing and Development, 2010, 131, 215-222.	4.6	16
120	Short telomere length is associated with renal impairment in Japanese subjects with cardiovascular risk. PLoS ONE, 2017, 12, e0176138.	2.5	16
121	A balanced translocation truncates Neurotrimin in a family with intracranial and thoracic aortic aneurysm. Journal of Medical Genetics, 2012, 49, 621-629.	3.2	15
122	Genetic analysis of long-lived families reveals novel variants influencing high density-lipoprotein cholesterol. Frontiers in Genetics, 2014, 5, 159.	2.3	15
123	Cerebrospinal fluid biomarkers of Alzheimer's disease in a cohort of adults with Down syndrome. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2020, 12, e12057.	2.4	15
124	Proteomic profiles for Alzheimer's disease and mild cognitive impairment among adults with Down syndrome spanning serum and plasma: An Alzheimer's Biomarker Consortium–Down Syndrome (ABC–DS) study. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2020, 12, e12039.	2.4	13
125	Association studies between the plasmin genes and late-onset Alzheimer's disease. Neurobiology of Aging, 2007, 28, 1041-1043.	3.1	12
126	Overweight, high blood pressure and impaired fasting glucose in Uyghur, Han, and Kazakh Chinese children and adolescents. Ethnicity and Health, 2015, 20, 365-375.	2.5	12

#	Article	IF	CITATIONS
127	Real-Time Mobile Teleophthalmology for the Detection of Eye Disease in Minorities and Low Socioeconomics At-Risk Populations. Asia-Pacific Journal of Ophthalmology, 2021, 10, 461-472.	2.5	12
128	The Heritability of Abstract Reasoning in Caribbean Latinos with Familial Alzheimer Disease. Dementia and Geriatric Cognitive Disorders, 2007, 24, 411-417.	1.5	11
129	Novel IRF6 mutations in Honduran Van Der Woude syndrome patients. Molecular Medicine Reports, 2011, 4, 237-41.	2.4	11
130	The association between interferon regulatory factor 6 ( <i>IRF6</i> ) and nonsyndromic cleft lip with or without cleft palate in a Honduran population. Laryngoscope, 2009, 119, 1759-1764.	2.0	10
131	Risk Ratios for Obesity in Families of Obese African-American and Caucasian Women. Human Heredity, 2001, 51, 35-40.	0.8	9
132	Mortality and apolipoprotein E in African-American, and White Elders: An attempted replication. American Journal of Medical Genetics Part A, 2003, 119A, 141-146.	2.4	9
133	Proteomic profiles of prevalent mild cognitive impairment and Alzheimer's disease among adults with Down syndrome. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2020, 12, e12023.	2.4	9
134	Optic neuropathy and congenital glaucoma associated with probable Zika virus infection in Venezuelan patients. JMM Case Reports, 2018, 5, e005145.	1.3	9
135	Non-congenital severe ocular complications of Zika virus infection. JMM Case Reports, 2018, 5, e005152.	1.3	9
136	Ultra-rare mutations in <i>SRCAP</i> segregate in Caribbean Hispanic families with Alzheimer disease. Neurology: Genetics, 2017, 3, e178.	1.9	8
137	Linkage analysis of multiplex Caribbean Hispanic families loaded for unexplained earlyâ€onset cases identifies novel Alzheimer's disease loci. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2018, 10, 554-562.	2.4	8
138	The Association between Sex and Risk of Alzheimer's Disease in Adults with Down Syndrome. Journal of Clinical Medicine, 2021, 10, 2966.	2.4	8
139	Genetics of physiological dysregulation: findings from the long life family study using joint models. Aging, 2020, 12, 5920-5947.	3.1	8
140	A novel healthy metabolic phenotype developed among a cohort of families enriched for longevity. Metabolism: Clinical and Experimental, 2019, 94, 28-38.	3.4	7
141	Normal-tension glaucomatous optic neuropathy is related to blood pressure variability in the Maracaibo Aging Study. Hypertension Research, 2021, 44, 1105-1112.	2.7	7
142	Leukocyte Telomere Length Is Unrelated to Cognitive Performance Among Non-Demented and Demented Persons: An Examination of Long Life Family Study Participants. Journal of the International Neuropsychological Society, 2020, 26, 906-917.	1.8	6
143	Genome-wide linkage analysis of carotid artery traits in exceptionally long-lived families. Atherosclerosis, 2019, 291, 19-26.	0.8	5
144	Associations of increased physical performance and change in body composition with molecular pathways of heart disease and diabetes risk. American Journal of Physiology - Endocrinology and Metabolism, 2019, 316, E221-E229.	3.5	5

#	Article	IF	CITATIONS
145	Open-Angle Glaucomatous Optic Neuropathy Is Related to Dips Rather Than Increases in the Mean Arterial Pressure Over 24-H. American Journal of Hypertension, 2022, 35, 703-714.	2.0	5
146	Functional genomic characterization of the <i>FTO</i> locus in African Americans. Physiological Genomics, 2019, 51, 517-528.	2.3	4
147	Proteomic profiles of incident mild cognitive impairment and Alzheimer's disease among adults with Down syndrome. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2020, 12, e12033.	2.4	4
148	Prevalence of clinically actionable disease variants in exceptionally long-lived families. BMC Medical Genomics, 2020, 13, 61.	1.5	4
149	Total Plasma Homocysteine and Depressive Symptoms in Older Hispanics. Journal of Alzheimer's Disease, 2021, 82, S263-S269.	2.6	4
150	Linkage of Alzheimer disease families with Puerto Rican ancestry identifies a chromosome 9 locus. Neurobiology of Aging, 2021, 104, 115.e1-115.e7.	3.1	4
151	Probing the proteome to explore potential correlates of increased Alzheimer'sâ€related cerebrovascular disease in adults with Down syndrome. Alzheimer's and Dementia, 2022, 18, 1744-1753.	0.8	4
152	Estimating a Multivariate Familial Correlation Using Joint Models for Canonical Correlations: Application to Memory Score Analysis from Familial Hispanic Alzheimer's Disease Study. Biometrics, 2009, 65, 463-469.	1.4	3
153	F5â€01â€01: PROTEOMIC BIOMARKERS FOR DETECTING AND PREDICTING AD RISKÂAMONG ADULTS WITH DOW SYNDROME. Alzheimer's and Dementia, 2018, 14, P1624.	/N.8	3
154	Heterogeneity of the Predictive Polygenic Risk Scores for Coronary Heart Disease Age-at-Onset in Three Different Coronary Heart Disease Family-Based Ascertainments. Circulation Genomic and Precision Medicine, 2021, 14, e003201.	3.6	3
155	Sequence variants in the $5\hat{a} \in 2$ flanking region of the leptin gene are associated with obesity in women. , 0, .		3
156	Association of Genetic Polymorphisms with Complications of Implanted LVAD Devices in Patients with Congestive Heart Failure: A Kazakhstani Study. Journal of Personalized Medicine, 2022, 12, 744.	2.5	3
157	Genotypeâ€adjusted familial correlation analysis using three generalized estimating equations. Statistics in Medicine, 2008, 27, 5471-5483.	1.6	2
158	P3-016: SORBS2, SH3RF3, AND NPHP1 MODIFY AGE AT ONSET IN CARRIERS OF THE G206A MUTATION IN PSEN1 WITH FAMILIAL ALZHEIMER'S DISEASE. , 2014, 10, P632-P632.		2
159	P1-229: Neuropsychiatric Symptoms and Their Relationship with Progression to Severe Dementia and Death: Findings of the Maracaibo Aging Study (MAS). , 2016, 12, P495-P496.		2
160	An improved statistical model for taxonomic assignment of metagenomics. BMC Genetics, 2018, 19, 98.	2.7	2
161	Alzheimer's disease in aging Down syndrome. Developmental Neurobiology, 2019, 79, 611-612.	3.0	2
162	Nighttime Blood Pressure Interacts with APOE Genotype to Increase the Risk of Incident Dementia of the Alzheimer's Type in Hispanics. Journal of Alzheimer's Disease, 2020, 77, 569-579.	2.6	2

#	Article	IF	CITATIONS
163	Whole-genome sequencing data of Kazakh individuals. BMC Research Notes, 2021, 14, 45.	1.4	2
164	Patients with coronary heart disease, dilated cardiomyopathy and idiopathic ventricular tachycardia share overlapping patterns of pathogenic variation in cardiac risk genes. PeerJ, 2021, 9, e10711.	2.0	2
165	Genetic and dietary influences on life span. , 2020, , 671-685.		2
166	P1-268: Variants in candidate genes for Alzheimer's disease are associated with declining plasma abeta peptides in adults with down syndrome., 2015, 11, P458-P458.		1
167	Elevated plasma neurofilament light (NfL) is associated with incident Alzheimer's disease and accelerated cognitive decline in adults with Down syndrome. Alzheimer's and Dementia, 2020, 16, e045982.	0.8	1
168	NORMAL-TENSION GLAUCOMATOUS OPTIC NEUROPATHY IS RELATED TO BLOOD PRESSURE VARIABILITY IN THE MARACAIBO AGING STUDY. Journal of Hypertension, 2021, 39, e142.	0.5	1
169	Natural Experiments in Human Gene Mapping: The Intersection of Anthropological Genetics and Genetic Epidemiology., 0,, 38-76.		1
170	The Sortilin-Related Receptor SORL1 is Functionally and Genetically Associated with Alzheimer's Disease. Research and Perspectives in Alzheimer's Disease, 2009, , 157-165.	0.1	1
171	Sex differences in levels of plasma neurofilament light and total tau in adults with Down syndrome. Alzheimer's and Dementia, 2021, 17, .	0.8	1
172	Obstructive sleep apnea, cerebrovascular disease, and amyloid in older adults with Down syndrome across the Alzheimer's continuum. SLEEP Advances, 2022, 3, .	0.2	1
173	O3-02-08 Fine mapping of chromosome 10 using memory and related neuropsychological phenotypes in the Caribbean hispanics. Neurobiology of Aging, 2004, 25, S56.	3.1	0
174	P3-331: TOTAL PLASMA HOMOCYSTEINE CONCENTRATION AS A PREDICTOR OF WHITE MATTER HYPERINTENSITIES AMONG ELDERLY RESIDING IN MARACAIBO, VENEZUELA. , 2014, 10, P750-P751.		0
175	P2-002: GENETIC CHARACTERIZATION OF BETA AMYLOID PEPTIDES IN ADULTS WITH DOWN SYNDROME WITH A FOCUS ON CHROMOSOMES OTHER THAN 21., 2014, 10, P471-P471.		O
176	F1-01-02: Ambulatory blood pressure and cognitive decline in latinos: Findings from the maracaibo aging study., 2015, 11, P115-P115.		0
177	O3-13-05: Rare coding mutations identified by targeted sequencing of Alzheimer's disease loci detected in genome-wide association studies. , 2015, 11, P252-P253.		0
178	O3-05-05: Snps in cugbp2 influence the risk of Alzheimer disease in white and caribbean hispanic elderly and in adults with down syndrome. , 2015, 11, P229-P230.		0
179	P3-131: Total plasma homocysteine and depressive symptoms in older hispanics. , 2015, 11, P672-P674.		O
180	P3â€295: Blood Pressure Indices are Associated with Increased Reduced Hippocampal Volume and White Matter Hyperintensities in Hispanic Adults: the Maracaibo Aging Study. Alzheimer's and Dementia, 2016, 12, P954.	0.8	O

#	Article	IF	CITATIONS
181	[P3–434]: INDIVIDUAL DIFFERENCES IN LIFELONG ABILITIES MUST BE CONSIDERED FOR STAGING OF EARLY AD Alzheimer's and Dementia, 2017, 13, P1134.	0.8	О
182	P1â€435: WHITE MATTER HYPERINTENSITIES IN LATIN AMERICANS: FINDINGS FROM THE MARACAIBO AGING STUDY. Alzheimer's and Dementia, 2018, 14, P477.	0.8	0
183	P1â€340: FRONTOTEMPORAL DEMENTIAâ€LIKE PHENOTYPE IN SEVEN HISPANIC PATIENTS WITH PRESENILINâ€2. Alzheimer's and Dementia, 2018, 14, P423.	0.8	O
184	GENETICS OF CUMULATIVE MEASURE OF PHYSIOLOGICAL DYSREGULATION: EVIDENCE FROM LONG LIFE FAMILY STUDY. Innovation in Aging, 2018, 2, 405-405.	0.1	O
185	P3â€145: TELOMERE LENGTH IN HUMAN LEUKOCYTE SUBPOPULATIONS. Alzheimer's and Dementia, 2018, 14, P1123.	0.8	О
186	P4â€073: IN ABSENCE OF DEMENTIA, COGNITIVE PERFORMANCE DOES NOT RELATE TO THE BIOMARKER OF LEUKOCYTE TELOMERE LENGTH: AN EXAMINATION OF LONG LIFE FAMILY STUDY PARTICIPANTS. Alzheimer's and Dementia, 2018, 14, P1462.	0.8	0
187	A17443 Ambulatory Blood Pressure Variability and Glaucomatous Optic Neuropathy. Journal of Hypertension, 2018, 36, e249.	0.5	О
188	Reply. Ophthalmology, 2019, 126, e12-e13.	5.2	0
189	Plasma biomarkers AÎ <sup>2</sup> 42, AÎ <sup>2</sup> 40, and tau in Down syndrome dementia. Alzheimer's and Dementia, 2020, 16, e045698.	0.8	O
190	Cerebral small vessel diseases are better associated with ambulatory than office blood pressure measurements. Alzheimer's and Dementia, 2020, 16, e045987.	0.8	0
191	Cognitive Domains in Low Literacy Populations: The Experience of the Maracaibo Aging Study. , 2021, , 13-33.		O
192	Research on aging during the Venezuelan humanitarian crisis: the experience of the Maracaibo aging study. BMC Public Health, 2021, 21, 473.	2.9	0
193	Author Response: White Matter Hyperintensities Mediate the Association of Nocturnal Blood Pressure With Cognition. Neurology, 2021, 97, 46-46.	1.1	О
194	Genetics of Alzheimer's disease in adults with Down syndrome. , 2022, , 193-208.		0
195	APOE and APOC1 Promoter Polymorphisms and the Risk of Alzheimer Disease in African American and Caribbean Hispanic Individuals—Correction. Archives of Neurology, 2005, 62, 332.	4.5	O
196	P1â€013: INCIDENCE OF ALZHEIMER'S DISEASE IN HISPANICS: ROLE OF APOLIPOPROTEIN E GENOTYPES AND AMBULATORY BLOOD PRESSURE MONITORING. Alzheimer's and Dementia, 2018, 14, P269.	0.8	0
197	Genetics of Circulating HbA1c Long-Term Change over Time in Nondiabetic Subjects—The Long Life Family Study. Diabetes, 2018, 67, 1709-P.	0.6	O
198	1447-P: Epistatic Interactions of Known Gene Variants for Glycated Hemoglobin: Evidence from the Long Life Family Study. Diabetes, 2019, 68, 1447-P.	0.6	0

#	Article	IF	CITATIONS
199	CSF Biomarkers in Down Syndrome and Autosomal Dominant Alzheimer Disease. SSRN Electronic Journal, 0, , .	0.4	0
200	An interactive computer system for formulary management using cost-effectiveness analysis. M D Computing, 1995, 12, 59-65.	0.1	0
201	Genetics and its use in Alzheimer's disease. Psychopharmacology Bulletin, 2007, 40, 132-44.	0.0	0