F Yesim Demirci

List of Publications by Year
in descending order

61	5,153	236612	143772
papers	citations	25 h-index	57 g-index

62	62	62	9065
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Common variants at MS4A4/MS4A6E, CD2AP, CD33 and EPHA1 are associated with late-onset Alzheimer's disease. Nature Genetics, 2011, 43, 436-441.	9.4	1,676
2	Association of Systemic Lupus Erythematosus withC8orf13â€“BLKandITGAMâ€“ITGAX. New England Journal of Medicine, 2008, 358, 900-909.	13.9	848
3	Rare coding variants in the phospholipase D3 gene confer risk for Alzheimerâ€™s disease. Nature, 2014, 505, 550-554.	13.7	425
4	Differential Genetic Associations for Systemic Lupus Erythematosus Based on Antiâ€“dsDNA Autoantibody Production. PLoS Genetics, 2011, 7, e1001323.	1.5	206
5	Specificity of the STAT4 Genetic Association for Severe Disease Manifestations of Systemic Lupus Erythematosus. PLoS Genetics, 2008, 4, e1000084.	1.5	180
6	Assessment of the genetic variance of late-onset Alzheimer's disease. Neurobiology of Aging, 2016, 41, 200.e13-200.e20.	1.5	174
7	Effects of Multiple Genetic Loci on Age at Onset in Late-Onset Alzheimer Disease. JAMA Neurology, 2014, 71, 1394.	4.5	166
8	Transethnic genomeâ€wide scan identifies novel Alzheimer's disease loci. Alzheimer's and Dementia, 2017, 13, 727-738.	0.4	166
9	Novel late-onset Alzheimer disease loci variants associate with brain gene expression. Neurology, 2012, 79, 221-228.	1.5	144
10	X-Linked Recessive Atrophic Macular Degeneration from RPGR Mutation. Genomics, 2002, 80, 166-171.	1.3	124
11	Lupus Nephritis Susceptibility Loci in Women with Systemic Lupus Erythematosus. Journal of the American Society of Nephrology: JASN, 2014, 25, 2859-2870.	3.0	117
12	Association of CLU and PICALM variants with Alzheimer's disease. Neurobiology of Aging, 2012, 33, 518-521.	1.5	67
13	Genetic Determinants of Disease Progression in Alzheimer's Disease. Journal of Alzheimer's Disease, 2014, 43, 649-655.	1.2	53
14	Genome-wide association study of brain amyloid deposition as measured by Pittsburgh Compound-B (PiB)-PET imaging. Molecular Psychiatry, 2021, 26, 309-321.	4.1	47
15	Connecting the Dots: Potential of Data Integration to Identify Regulatory SNPs in Late-Onset Alzheimer's Disease GWAS Findings. PLoS ONE, 2014, 9, e95152.	1.1	43
16	Rarity of the Alzheimer Diseaseâ€“ProtectiveAPPA673T Variant in the United States. JAMA Neurology, 2015, 72, 209.	4.5	41
17	More evidence for association of a rare TREM2 mutation (R47H) with Alzheimer's disease risk. Neurobiology of Aging, 2015, 36, 2443.e21-2443.e26.	1.5	39
18	No association betweenCALHM1variation and risk of Alzheimer disease. Human Mutation, 2009, 30, E566-E569.	1.1	37

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19	Functional significance of lipoprotein lipase HindIII polymorphism associated with the risk of coronary artery disease. Atherosclerosis, 2008, 200, 102-108.	0.4	33
20	Genetic Variation in Imprinted Genes is Associated with Risk of Late-Onset Alzheimer's Disease. Journal of Alzheimer's Disease, 2015, 44, 989-994.	1.2	32
21	Identification of a New Susceptibility Locus for Systemic Lupus Erythematosus on Chromosome 12 in Individuals of European Ancestry. Arthritis and Rheumatology, 2016, 68, 174-183.	2.9	30
22	Integrative Exome and Transcriptome Analysis of Conjunctival Melanoma and Its Potential Application for Personalized Therapy. JAMA Ophthalmology, 2019, 137, 1444.	1.4	29
23	Association analysis of PON2 genetic variants with serum paraoxonase activity and systemic lupus erythematosus. BMC Medical Genetics, 2011, 12, 7.	2.1	28
24	Investigation of an amyloid precursor protein protective mutation (A673T) in a North American case-control sample of late-onset Alzheimer's disease. Neurobiology of Aging, 2014, 35, 1779.e15-1779.e16.	1.5	28
25	Association of 32 type 1 diabetes risk loci in Pakistani patients. Diabetes Research and Clinical Practice, 2015, 108, 137-142.	1.1	28
26	A Multiethnic Replication Study of Plasma Lipoprotein Levels-Associated SNPs Identified in Recent GWAS. PLoS ONE, 2013, 8, e63469.	1.1	25
27	Lipoprotein lipase gene sequencing and plasma lipid profile. Journal of Lipid Research, 2014, 55, 85-93.	2.0	24
28	Comprehensive Evaluation of the Association of APOE Genetic Variation with Plasma Lipoprotein Traits in U.S. Whites and African Blacks. PLoS ONE, 2014, 9, e114618.	1.1	23
29	Population-based genome-wide association study of cognitive decline in older adults free of dementia: identification of a novel locus for the attention domain. Neurobiology of Aging, 2019, 84, 239.e15-239.e24.	1.5	21
30	Genome-Wide Association Study of Antiphospholipid Antibodies. Autoimmune Diseases, 2013, 2013, 1-11.	2.7	20
31	Resequencing of the CETP gene in American whites and African blacks: Association of rare and common variants with HDL-cholesterol levels. Metabolism: Clinical and Experimental, 2016, 65, 36-47.	1.5	19
32	Whole-Exome Sequencing Analysis of Alzheimerâ€™s Disease in Non-APOE*4 Carriers. Journal of Alzheimer's Disease, 2020, 76, 1553-1565.	1.2	18
33	Beta-amyloid toxicity modifier genes and the risk of Alzheimer's disease. American Journal of Neurodegenerative Disease, 2012, 1, 191-8.	0.1	18
34	Genetic link of type 1 diabetes susceptibility loci with rheumatoid arthritis in Pakistani patients. Immunogenetics, 2015, 67, 277-282.	1.2	17
35	A Rare Duplication on Chromosome 16p11.2 Is Identified in Patients with Psychosis in Alzheimer's Disease. PLoS ONE, 2014, 9, e111462.	1.1	16
36	Impact of Genetic Variants in Human Scavenger Receptor Class B Type I (SCARB1) on Plasma Lipid Traits. Circulation: Cardiovascular Genetics, 2014, 7, 838-847.	5.1	16

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37	Genetic contribution of SCARB1 variants to lipid traits in African Blacks: a candidate gene association study. BMC Medical Genetics, 2015, 16, 106.	2.1	16
38	Apolipoprotein E-C1-C4-C2 gene cluster region and inter-individual variation in plasma lipoprotein levels: a comprehensive genetic association study in two ethnic groups. PLoS ONE, 2019, 14, e0214060.	1.1	16
39	Association studies of 22 candidate SNPs with lateâ€onset Alzheimer's disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2009, 150B, 520-526.	1.1	13
40	Multiple signals at the extended 8p23 locus are associated with susceptibility to systemic lupus erythematosus. Journal of Medical Genetics, 2017, 54, 381-389.	1.5	13
41	Genetic Determinants of Survival in Patients with Alzheimer's Disease. Journal of Alzheimer's Disease, 2015, 45, 651-658.	1.2	11
42	Investigating the GWAS-Implicated Loci for Rheumatoid Arthritis in the Pakistani Population. Disease Markers, 2020, 2020, 1-9.	0.6	11
43	Resequencing of LPL in African Blacks and associations with lipoproteinâ€“lipid levels. European Journal of Human Genetics, 2015, 23, 1244-1253.	1.4	10
44	Immunotherapy for Conjunctival Squamous Cell Carcinoma with Orbital Extension. Ophthalmology, 2021, 128, 801-804.	2.5	10
45	APOEGene Polymorphism and Risk of Coronary Stenosis in Pakistani Population. BioMed Research International, 2015, 2015, 1-5.	0.9	9
46	A sequencing study of CTLA4 in Pakistani rheumatoid arthritis cases. PLoS ONE, 2020, 15, e0239426.	1.1	9
47	Replication of European Rheumatoid Arthritis Loci in a Pakistani Population. Journal of Rheumatology, 2013, 40, 401-407.	1.0	8
48	Functional and genetic characterization of the promoter region of apolipoprotein H (Î²₂â€glycoprotein I). FEBS Journal, 2010, 277, 951-963.	2.2	7
49	Assessment of genetic risk of type 2 diabetes among Pakistanis based on GWAS-implicated loci. Gene, 2021, 783, 145563.	1.0	7
50	Association Study of Coronary Artery Disease-Associated Genome-Wide Significant SNPs with Coronary Stenosis in Pakistani Population. Disease Markers, 2020, 2020, 1-7.	0.6	6
51	Exploration of shared genetic susceptibility loci between type 1 diabetes and rheumatoid arthritis in the Pakistani population. BMC Research Notes, 2019, 12, 544.	0.6	5
52	Association of VPREB1 Gene Copy Number Variation and Rheumatoid Arthritis Susceptibility. Disease Markers, 2020, 2020, 1-5.	0.6	5
53	Association of Three Lipoprotein Lipase Polymorphisms with Coronary Artery Disease in Chinese and Asian Indians. International Journal of Cardiology, 2010, 144, 142-143.	0.8	3
54	Gene Expression and Cardiometabolic Phenotypes of Vitamin D-Deficient Overweight and Obese Black Children. Nutrients, 2019, 11, 2016.	1.7	3

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55	Hepatic lipase (LIPC) sequencing in individuals with extremely high and low high-density lipoprotein cholesterol levels. PLoS ONE, 2020, 15, e0243919.	1.1	3
56	Refinement of the Physical Location and the Genomic Characterization of the CRSP2 (EXLM1) Gene on Xp11.4. DNA Sequence, 2003, 14, 123-127.	0.7	1
57	Association analysis of 23 susceptibility loci with risk of dementia in a Pakistani population. Psychiatry Research, 2015, 225, 223-224.	1.7	0
58	Title is missing!. , 2020, 15, e0243919.		0
59	Title is missing!. , 2020, 15, e0243919.		0
60	Title is missing!. , 2020, 15, e0243919.		0
61	Title is missing!. , 2020, 15, e0243919.		0