

Sudha K Iyengar

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1191308/publications.pdf>

Version: 2024-02-01

69
papers

4,192
citations

117453

34
h-index

118652

62
g-index

79
all docs

79
docs citations

79
times ranked

6793
citing authors

#	ARTICLE	IF	CITATIONS
1	Seven new loci associated with age-related macular degeneration. <i>Nature Genetics</i> , 2013, 45, 433-439.	9.4	687
2	Genome-wide association meta-analysis highlights light-induced signaling as a driver for refractive error. <i>Nature Genetics</i> , 2018, 50, 834-848.	9.4	239
3	Genome-wide association analyses identify three new susceptibility loci for primary angle closure glaucoma. <i>Nature Genetics</i> , 2012, 44, 1142-1146.	9.4	196
4	Meta-Analysis of Genome-Wide Association Studies in African Americans Provides Insights into the Genetic Architecture of Type 2 Diabetes. <i>PLoS Genetics</i> , 2014, 10, e1004517.	1.5	191
5	School-Age Follow-Up of Children With Childhood Apraxia of Speech. <i>Language, Speech, and Hearing Services in Schools</i> , 2004, 35, 122-140.	0.7	167
6	Dissection of Genomewide-Scan Data in Extended Families Reveals a Major Locus and Oligogenic Susceptibility for Age-Related Macular Degeneration. <i>American Journal of Human Genetics</i> , 2004, 74, 20-39.	2.6	162
7	Genome-Wide Scans for Diabetic Nephropathy and Albuminuria in Multiethnic Populations: The Family Investigation of Nephropathy and Diabetes (FIND). <i>Diabetes</i> , 2007, 56, 1577-1585.	0.3	140
8	Markers of Inflammation, Oxidative Stress, and Endothelial Dysfunction and the 20-Year Cumulative Incidence of Early Age-Related Macular Degeneration. <i>JAMA Ophthalmology</i> , 2014, 132, 446.	1.4	122
9	Type 2 Diabetes Variants Disrupt Function of SLC16A11 through Two Distinct Mechanisms. <i>Cell</i> , 2017, 170, 199-212.e20.	13.5	121
10	Genome-Wide Association and Trans-ethnic Meta-Analysis for Advanced Diabetic Kidney Disease: Family Investigation of Nephropathy and Diabetes (FIND). <i>PLoS Genetics</i> , 2015, 11, e1005352.	1.5	118
11	Genetics of age-related macular degeneration (AMD). <i>Human Molecular Genetics</i> , 2017, 26, R45-R50.	1.4	109
12	The Genetic Basis of Complex Traits. <i>Methods in Molecular Biology</i> , 2007, 376, 71-84.	0.4	103
13	Genome-wide association study identifies three novel loci in Fuchs endothelial corneal dystrophy. <i>Nature Communications</i> , 2017, 8, 14898.	5.8	101
14	Prediction of Age-related Macular Degeneration in the General Population. <i>Ophthalmology</i> , 2013, 120, 2644-2655.	2.5	84
15	Harmonizing the Classification of Age-related Macular Degeneration in the Three-Continent AMD Consortium. <i>Ophthalmic Epidemiology</i> , 2014, 21, 14-23.	0.8	83
16	Lipids, Lipid Genes, and Incident Age-Related Macular Degeneration: The Three Continent Age-Related Macular Degeneration Consortium. <i>American Journal of Ophthalmology</i> , 2014, 158, 513-524.e3.	1.7	81
17	Electrocardiographic Prediction of Abnormal Genotype in Congenital Long QT Syndrome: Experience in 101 Related Family Members. <i>Journal of Cardiovascular Electrophysiology</i> , 2001, 12, 455-461.	0.8	79
18	A Multicenter Study to Map Genes for Fuchs Endothelial Corneal Dystrophy: Baseline Characteristics and Heritability. <i>Cornea</i> , 2012, 31, 26-35.	0.9	78

#	ARTICLE	IF	CITATIONS
19	Identification of a major locus for age-related cortical cataract on chromosome 6p12-q12 in the Beaver Dam Eye Study. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 14485-14490.	3.3	69
20	Combined Effects of Complement Factor H Genotypes, Fish Consumption, and Inflammatory Markers on Long-Term Risk for Age-related Macular Degeneration in a Cohort. American Journal of Epidemiology, 2008, 169, 633-641.	1.6	69
21	Linkage Analysis of Candidate Loci for End-Stage Renal Disease due to Diabetic Nephropathy. Journal of the American Society of Nephrology: JASN, 2003, 14, S195-S201.	3.0	63
22	Cigarette Smoking and the Natural History of Age-Related Macular Degeneration. Ophthalmology, 2014, 121, 1949-1955.	2.5	63
23	Genetic Susceptibility, Dietary Antioxidants, and Long-Term Incidence of Age-Related Macular Degeneration in Two Populations. Ophthalmology, 2014, 121, 667-675.	2.5	59
24	Multiethnic Genome-Wide Association Study of Diabetic Retinopathy Using Liability Threshold Modeling of Duration of Diabetes and Glycemic Control. Diabetes, 2019, 68, 441-456.	0.3	54
25	Rare and common variants in extracellular matrix gene Fibrillin 2 (FBN2) are associated with macular degeneration. Human Molecular Genetics, 2014, 23, 5827-5837.	1.4	52
26	Risk Alleles in CFH and ARMS2 and the Long-term Natural History of Age-Related Macular Degeneration. JAMA Ophthalmology, 2013, 131, 383.	1.4	48
27	A 32 kb Critical Region Excluding Y402H in CFH Mediates Risk for Age-Related Macular Degeneration. PLoS ONE, 2011, 6, e25598.	1.1	46
28	Joint Associations of Diet, Lifestyle, and Genes with Age-Related Macular Degeneration. Ophthalmology, 2015, 122, 2286-2294.	2.5	44
29	Exome Array Analysis Identifies CAV1/CAV2 as a Susceptibility Locus for Intraocular Pressure. Investigative Ophthalmology and Visual Science, 2015, 56, 544-551.	3.3	43
30	Association Between Vitamin D Status and Age-Related Macular Degeneration by Genetic Risk. JAMA Ophthalmology, 2015, 133, 1171.	1.4	43
31	Five-year progression of unilateral age-related macular degeneration to bilateral involvement: the Three Continent AMD Consortium report. British Journal of Ophthalmology, 2017, 101, 1185-1192.	2.1	38
32	A multi-ethnic genome-wide association study implicates collagen matrix integrity and cell differentiation pathways in keratoconus. Communications Biology, 2021, 4, 266.	2.0	36
33	Effect of the Y402H Variant in the Complement Factor H Gene on the Incidence and Progression of Age-Related Macular Degeneration. JAMA Ophthalmology, 2012, 130, 1169.	2.6	35
34	Meta-analysis of genome-wide association studies in multiethnic Asians identifies two loci for age-related nuclear cataract. Human Molecular Genetics, 2014, 23, 6119-6128.	1.4	35
35	Analyses of Cross Species Polymerase Chain Reaction Products to Infer the Ancestral State of Human Polymorphisms. DNA Sequence, 1998, 8, 317-327.	0.7	34
36	Sunlight Exposure, Pigmentation, and Incident Age-Related Macular Degeneration. , 2014, 55, 5855.		32

#	ARTICLE	IF	CITATIONS
37	Severity of Age-Related Macular Degeneration in 1 Eye and the Incidence and Progression of Age-Related Macular Degeneration in the Fellow Eye. <i>JAMA Ophthalmology</i> , 2015, 133, 125.	1.4	31
38	<i>APOL1</i> Risk Variants, Acute Kidney Injury, and Death in Participants With African Ancestry Hospitalized With COVID-19 From the Million Veteran Program. <i>JAMA Internal Medicine</i> , 2022, 182, 386.	2.6	31
39	Reading Outcomes for Individuals With Histories of Suspected Childhood Apraxia of Speech. <i>American Journal of Speech-Language Pathology</i> , 2019, 28, 1432-1447.	0.9	29
40	A Genome-Wide Search for Linkage of Estimated Glomerular Filtration Rate (eGFR) in the Family Investigation of Nephropathy and Diabetes (FIND). <i>PLoS ONE</i> , 2013, 8, e81888.	1.1	24
41	Mining the Genome for Susceptibility to Diabetic Nephropathy: The Role of Large-Scale Studies and Consortia. <i>Seminars in Nephrology</i> , 2007, 27, 208-222.	0.6	23
42	Psychosocial co-morbidities in adolescents and adults with histories of communication disorders. <i>Journal of Communication Disorders</i> , 2016, 61, 60-70.	0.8	20
43	Diversity of Ocular Surface Bacterial Microbiome Adherent to Worn Contact Lenses and Bacterial Communities Associated With Care Solution Use. <i>Eye and Contact Lens</i> , 2019, 45, 331-339.	0.8	20
44	Speech-Sound Disorders and Attention-Deficit/Hyperactivity Disorder Symptoms. <i>Topics in Language Disorders</i> , 2012, 32, 247-263.	0.9	18
45	Heritability and longitudinal outcomes of spelling skills in individuals with histories of early speech and language disorders. <i>Learning and Individual Differences</i> , 2018, 65, 1-11.	1.5	18
46	Small Drusen and Age-Related Macular Degeneration: The Beaver Dam Eye Study. <i>Journal of Clinical Medicine</i> , 2015, 4, 425-440.	1.0	17
47	A Phenome-Wide Association Study of genes associated with COVID-19 severity reveals shared genetics with complex diseases in the Million Veteran Program. <i>PLoS Genetics</i> , 2022, 18, e1010113.	1.5	16
48	Feature-driven classification reveals potential comorbid subtypes within childhood apraxia of speech. <i>BMC Pediatrics</i> , 2020, 20, 519.	0.7	15
49	Association of Kidney Comorbidities and Acute Kidney Failure With Unfavorable Outcomes After COVID-19 in Individuals With the Sickle Cell Trait. <i>JAMA Internal Medicine</i> , 0, , .	2.6	15
50	Approaches to understanding susceptibility to nephropathy: From genetics to genomics. <i>Kidney International</i> , 2002, 61, S61-S67.	2.6	14
51	The Quest for Genes Causing Complex Traits in Ocular Medicine. <i>JAMA Ophthalmology</i> , 2007, 125, 11.	2.6	14
52	Joint Contribution of Genetic Susceptibility and Modifiable Factors to the Progression of Age-Related Macular Degeneration over 10 Years. <i>Ophthalmology Retina</i> , 2018, 2, 684-693.	1.2	14
53	Differential Long-Term Outcomes for Individuals With Histories of Preschool Speech Sound Disorders. <i>American Journal of Speech-Language Pathology</i> , 2019, 28, 1582-1596.	0.9	13
54	Mitochondrial Polymorphism A10398G and Haplogroup I Are Associated With Fuchs' Endothelial Corneal Dystrophy. , 2014, 55, 4577.		12

#	ARTICLE	IF	CITATIONS
55	Associations Between Methylenetetrahydrofolate Reductase Polymorphisms, Serum Homocysteine Levels, and Incident Cortical Cataract. <i>JAMA Ophthalmology</i> , 2016, 134, 522.	1.4	11
56	Pooling Data and Linkage Analysis in the Chromosome 5q Candidate Region for Asthma. <i>Genetic Epidemiology</i> , 2001, 21, S103-8.	0.6	10
57	Common variants in SOX-2 and congenital cataract genes contribute to age-related nuclear cataract. <i>Communications Biology</i> , 2020, 3, 755.	2.0	10
58	Psychosocial Comorbidities in Adolescents With Histories of Childhood Apraxia of Speech. <i>American Journal of Speech-Language Pathology</i> , 2021, 30, 1-17.	0.9	9
59	Variation in PTCHD2, CRISP3, NAP1L4, FSCB, and AP3B2 associated with spherical equivalent. <i>Molecular Vision</i> , 2016, 22, 783-96.	1.1	8
60	Gene Set Enrichment Analyses Identify Pathways Involved in Genetic Risk for Diabetic Retinopathy. <i>American Journal of Ophthalmology</i> , 2022, 233, 111-123.	1.7	7
61	Association between genes regulating neural pathways for quantitative traits of speech and language disorders. <i>Npj Genomic Medicine</i> , 2021, 6, 64.	1.7	7
62	Improved Evidence for Linkage on 6p and 5p with Retrospective Pooling of Data from Three Asthma Genome Screens. <i>Genetic Epidemiology</i> , 2001, 21, S130-5.	0.6	5
63	Aldose Reductase Polymorphisms, Fasting Blood Glucose, and Age-Related Cortical Cataract. , 2018, 59, 4755.		5
64	The Application of the HapMap to Diabetic Nephropathy and Other Causes of Chronic Renal Failure. <i>Seminars in Nephrology</i> , 2007, 27, 223-236.	0.6	4
65	The fa leptin receptor mutation and the heritability of respiratory frequency in a Brown Norway and Zucker intercross. <i>Journal of Applied Physiology</i> , 2004, 97, 811-820.	1.2	3
66	Exome Array Analysis of Nuclear Lens Opacity. <i>Ophthalmic Epidemiology</i> , 2018, 25, 215-219.	0.8	3
67	Ancestral diversity improves discovery and fine-mapping of genetic loci for anthropometric traitsâ€™The Hispanic/Latino Anthropometry Consortium. <i>Human Genetics and Genomics Advances</i> , 2022, 3, 100099.	1.0	3
68	The Genetic Epidemiology of Age-Related Maculopathy. <i>International Journal of Human Genetics</i> , 2001, 1, 11-24.	0.1	2
69	Fuchsâ€™ endothelial corneal dystrophy: fostering change in clinical care using observational data. <i>Expert Review of Ophthalmology</i> , 2012, 7, 389-391.	0.3	0