

Janey L Wiggs

List of Publications by Year in descending order

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Version: 2024-02-01

202
papers

11,617
citations

28274

55
h-index

38395

95
g-index

223
all docs

223
docs citations

223
times ranked

10360
citing authors

#	ARTICLE	IF	CITATIONS
1	Detectable clonal mosaicism from birth to old age and its relationship to cancer. <i>Nature Genetics</i> , 2012, 44, 642-650.	21.4	511
2	Mutations in genes encoding melanosomal proteins cause pigmentary glaucoma in DBA/2J mice. <i>Nature Genetics</i> , 2002, 30, 81-85.	21.4	427
3	Primary open-angle glaucoma. <i>Nature Reviews Disease Primers</i> , 2016, 2, 16067.	30.5	319
4	A simple procedure for resolution of <i>Escherichia coli</i> RNA polymerase holoenzyme from core polymerase. <i>Archives of Biochemistry and Biophysics</i> , 1977, 182, 404-408.	3.0	315
5	Common Variants at 9p21 and 8q22 Are Associated with Increased Susceptibility to Optic Nerve Degeneration in Glaucoma. <i>PLoS Genetics</i> , 2012, 8, e1002654.	3.5	276
6	Genome-wide association analyses identify multiple loci associated with central corneal thickness and keratoconus. <i>Nature Genetics</i> , 2013, 45, 155-163.	21.4	269
7	Genetics of glaucoma. <i>Human Molecular Genetics</i> , 2017, 26, R21-R27.	2.9	266
8	Prediction of the Risk of Hereditary Retinoblastoma, Using DNA Polymorphisms within the Retinoblastoma Gene. <i>New England Journal of Medicine</i> , 1988, 318, 151-157.	27.0	249
9	Fried food consumption, genetic risk, and body mass index: gene-diet interaction analysis in three US cohort studies. <i>BMJ, The</i> , 2014, 348, g1610-g1610.	6.0	229
10	Panel-based genetic diagnostic testing for inherited eye diseases is highly accurate and reproducible, and more sensitive for variant detection, than exome sequencing. <i>Genetics in Medicine</i> , 2015, 17, 253-261.	2.4	216
11	Genome-wide analyses identify 68 new loci associated with intraocular pressure and improve risk prediction for primary open-angle glaucoma. <i>Nature Genetics</i> , 2018, 50, 778-782.	21.4	214
12	Genome-wide analysis of multi-ancestry cohorts identifies new loci influencing intraocular pressure and susceptibility to glaucoma. <i>Nature Genetics</i> , 2014, 46, 1126-1130.	21.4	212
13	Genome-wide association analysis identifies TXNRD2, ATXN2 and FOXC1 as susceptibility loci for primary open-angle glaucoma. <i>Nature Genetics</i> , 2016, 48, 189-194.	21.4	211
14	Prevalence of Mutations in TIGR/Myocilin in Patients with Adult and Juvenile Primary Open-Angle Glaucoma. <i>American Journal of Human Genetics</i> , 1998, 63, 1549-1552.	6.2	197
15	Genetic Etiologies of Glaucoma. <i>JAMA Ophthalmology</i> , 2007, 125, 30.	2.4	196
16	Genome-wide meta-analysis identifies 127 open-angle glaucoma loci with consistent effect across ancestries. <i>Nature Communications</i> , 2021, 12, 1258.	12.8	196
17	Multitrait analysis of glaucoma identifies new risk loci and enables polygenic prediction of disease susceptibility and progression. <i>Nature Genetics</i> , 2020, 52, 160-166.	21.4	192
18	Common variants near ABCA1, AFAP1 and GMDS confer risk of primary open-angle glaucoma. <i>Nature Genetics</i> , 2014, 46, 1120-1125.	21.4	186

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19	Angiotensin receptor TEK mutations underlie primary congenital glaucoma with variable expressivity. <i>Journal of Clinical Investigation</i> , 2016, 126, 2575-2587.	8.2	175
20	Common variants near CAV1 and CAV2 are associated with primary open-angle glaucoma in Caucasians from the USA. <i>Human Molecular Genetics</i> , 2011, 20, 4707-4713.	2.9	156
21	The gene, environment association studies consortium (GENEVA): maximizing the knowledge obtained from GWAS by collaboration across studies of multiple conditions. <i>Genetic Epidemiology</i> , 2010, 34, 364-372.	1.3	139
22	Glaucoma: genes, phenotypes, and new directions for therapy. <i>Journal of Clinical Investigation</i> , 2010, 120, 3064-3072.	8.2	121
23	New insights into the genetics of primary open-angle glaucoma based on meta-analyses of intraocular pressure and optic disc characteristics.. <i>Human Molecular Genetics</i> , 2017, 26, ddw399.	2.9	120
24	Distribution of WDR36 DNA Sequence Variants in Patients with Primary Open-Angle Glaucoma. , 2006, 47, 2542.		114
25	Genetic association study of exfoliation syndrome identifies a protective rare variant at LOXL1 and five new susceptibility loci. <i>Nature Genetics</i> , 2017, 49, 993-1004.	21.4	114
26	Genome-wide association study identifies seven novel susceptibility loci for primary open-angle glaucoma. <i>Human Molecular Genetics</i> , 2018, 27, 1486-1496.	2.9	111
27	Genetic Variants Associated with Optic Nerve Vertical Cup-to-Disc Ratio Are Risk Factors for Primary Open Angle Glaucoma in a US Caucasian Population. , 2011, 52, 1788.		109
28	Endothelial Nitric Oxide Synthase Gene Variants and Primary Open-Angle Glaucoma: Interactions with Sex and Postmenopausal Hormone Use. , 2010, 51, 971.		107
29	DNA sequence variants in the LOXL1 gene are associated with pseudoexfoliation glaucoma in a U.S. clinic-based population with broad ethnic diversity. <i>BMC Medical Genetics</i> , 2008, 9, 5.	2.1	105
30	Geographic and Climatic Factors Associated With Exfoliation Syndrome. <i>JAMA Ophthalmology</i> , 2011, 129, 1053.	2.4	105
31	A common variant near TGFBR3 is associated with primary open angle glaucoma. <i>Human Molecular Genetics</i> , 2015, 24, 3880-3892.	2.9	105
32	Characterization of Large Structural Genetic Mosaicism in Human Autosomes. <i>American Journal of Human Genetics</i> , 2015, 96, 487-497.	6.2	101
33	A Genomewide Scan Identifies Novel Early-Onset Primary Open-Angle Glaucoma Loci on 9q22 and 20p12. <i>American Journal of Human Genetics</i> , 2004, 74, 1314-1320.	6.2	100
34	A common variant mapping to CACNA1A is associated with susceptibility to exfoliation syndrome. <i>Nature Genetics</i> , 2015, 47, 387-392.	21.4	97
35	Nucleotide sequences of two <i>Bacillus subtilis</i> promoters used by <i>Bacillus subtilis</i> sigma-28 RNA polymerase. <i>Nucleic Acids Research</i> , 1981, 9, 5991-6000.	14.5	96
36	Genetic Linkage of Autosomal Dominant Juvenile Glaucoma to 1q21-q31 in Three Affected Pedigrees. <i>Genomics</i> , 1994, 21, 299-303.	2.9	95

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37	Angiotensin-1 is required for Schlemm's canal development in mice and humans. <i>Journal of Clinical Investigation</i> , 2017, 127, 4421-4436.	8.2	94
38	Genome-wide association study and meta-analysis of intraocular pressure. <i>Human Genetics</i> , 2014, 133, 41-57.	3.8	93
39	Molecular and Clinical Evaluation of a Patient Hemizygous for TIGR/MYOC. <i>JAMA Ophthalmology</i> , 2001, 119, 1674.	2.4	92
40	Association of CAV1/CAV2 Genomic Variants with Primary Open-Angle Glaucoma Overall and by Gender and Pattern of Visual Field Loss. <i>Ophthalmology</i> , 2014, 121, 508-516.	5.2	91
41	Meta-analysis of genome-wide association studies identifies novel loci that influence cupping and the glaucomatous process. <i>Nature Communications</i> , 2014, 5, 4883.	12.8	89
42	Biomechanical aspects of axonal damage in glaucoma: A brief review. <i>Experimental Eye Research</i> , 2017, 157, 13-19.	2.6	88
43	Patterns of functional vision loss in glaucoma determined with archetypal analysis. <i>Journal of the Royal Society Interface</i> , 2015, 12, 20141118.	3.4	87
44	Lack of Association of Mutations in Optineurin With Disease in Patients With Adult-onset Primary Open-angle Glaucoma. <i>JAMA Ophthalmology</i> , 2003, 121, 1181.	2.4	86
45	Early Adult-Onset POAG Linked to 15q11-13 Using Ordered Subset Analysis. , 2005, 46, 2002.		86
46	Distribution of Optineurin Sequence Variations in an Ethnically Diverse Population of Low-tension Glaucoma Patients From the United States. <i>Journal of Glaucoma</i> , 2006, 15, 358-363.	1.6	82
47	Association of Dietary Nitrate Intake With Primary Open-Angle Glaucoma. <i>JAMA Ophthalmology</i> , 2016, 134, 294.	2.5	81
48	Clinical Features of Five Pedigrees Genetically Linked to the Juvenile Glaucoma Locus on Chromosome 1q21-q31. <i>Ophthalmology</i> , 1995, 102, 1782-1789.	5.2	79
49	Discovery and Functional Annotation of SIX6 Variants in Primary Open-Angle Glaucoma. <i>PLoS Genetics</i> , 2014, 10, e1004372.	3.5	78
50	Demographic and Geographic Features of Exfoliation Glaucoma in 2 United States-Based Prospective Cohorts. <i>Ophthalmology</i> , 2012, 119, 27-35.	5.2	77
51	CDKN2B-AS1 Genotype's Glaucoma Feature Correlations in Primary Open-Angle Glaucoma Patients From the United States. <i>American Journal of Ophthalmology</i> , 2013, 155, 342-353.e5.	3.3	76
52	Four Susceptibility Loci for Gallstone Disease Identified in a Meta-analysis of Genome-Wide Association Studies. <i>Gastroenterology</i> , 2016, 151, 351-363.e28.	1.3	74
53	Investigation of Known Genetic Risk Factors for Primary Open Angle Glaucoma in Two Populations of African Ancestry. , 2013, 54, 6248.		73
54	Meta-analysis of Genome-Wide Association Studies Identifies Novel Loci Associated With Optic Disc Morphology. <i>Genetic Epidemiology</i> , 2015, 39, 207-216.	1.3	72

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55	Demographic, Systemic, and Ocular Factors Associated with Nonarteritic Anterior Ischemic Optic Neuropathy. <i>Ophthalmology</i> , 2016, 123, 2446-2455.	5.2	70
56	Solar Exposure and Residential Geographic History in Relation to Exfoliation Syndrome in the United States and Israel. <i>JAMA Ophthalmology</i> , 2014, 132, 1439.	2.5	66
57	Glaucoma Genes and Mechanisms. <i>Progress in Molecular Biology and Translational Science</i> , 2015, 134, 315-342.	1.7	65
58	Is Estrogen a Therapeutic Target for Glaucoma?. <i>Seminars in Ophthalmology</i> , 2016, 31, 140-146.	1.6	65
59	Endothelial Nitric Oxide Synthase Gene Variants and Primary Open-Angle Glaucoma. <i>JAMA Ophthalmology</i> , 2011, 129, 773.	2.4	63
60	Cross-ancestry genome-wide association analysis of corneal thickness strengthens link between complex and Mendelian eye diseases. <i>Nature Communications</i> , 2018, 9, 1864.	12.8	63
61	Mammalian Homolog of <i>Drosophila</i> retinal degeneration B Rescues the Mutant Fly Phenotype. <i>Journal of Neuroscience</i> , 1997, 17, 5881-5890.	3.6	62
62	<i>LOXL1</i> Promoter Haplotypes Are Associated with Exfoliation Syndrome in a U.S. Caucasian Population. , 2011, 52, 2372.		61
63	Biological aspects of axonal damage in glaucoma: A brief review. <i>Experimental Eye Research</i> , 2017, 157, 5-12.	2.6	61
64	Deep Learning of the Retina Enables Phenome- and Genome-Wide Analyses of the Microvasculature. <i>Circulation</i> , 2022, 145, 134-150.	1.6	57
65	The NEIGHBOR Consortium Primary Open-Angle Glaucoma Genome-wide Association Study. <i>Journal of Glaucoma</i> , 2013, 22, 517-525.	1.6	55
66	Phy-Mer: a novel alignment-free and reference-independent mitochondrial haplogroup classifier. <i>Bioinformatics</i> , 2015, 31, 1310-1312.	4.1	55
67	Soluble Guanylate Cyclase β -Deficient Mice: A Novel Murine Model for Primary Open Angle Glaucoma. <i>PLoS ONE</i> , 2013, 8, e60156.	2.5	55
68	Genome-Wide Analysis of Central Corneal Thickness in Primary Open-Angle Glaucoma Cases in the NEIGHBOR and GLAUGEN Consortia. , 2012, 53, 4468.		52
69	A comprehensive survey of genetic variation in 20,691 subjects from four large cohorts. <i>PLoS ONE</i> , 2017, 12, e0173997.	2.5	52
70	Common and Rare Genetic Risk Factors for Glaucoma. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2014, 4, a017244-a017244.	6.2	50
71	Systems genetics identifies a role for <i>Cacna2d1</i> regulation in elevated intraocular pressure and glaucoma susceptibility. <i>Nature Communications</i> , 2017, 8, 1755.	12.8	50
72	Association of Genetic Variants With Primary Open-Angle Glaucoma Among Individuals With African Ancestry. <i>JAMA - Journal of the American Medical Association</i> , 2019, 322, 1682.	7.4	50

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73	Genetic variation affects morphological retinal phenotypes extracted from UK Biobank optical coherence tomography images. <i>PLoS Genetics</i> , 2021, 17, e1009497.	3.5	50
74	Lack of Association between LOXL1 Variants and Primary Open-Angle Glaucoma in Three Different Populations. , 2008, 49, 3465.		48
75	Clinical implications of recent advances in primary open-angle glaucoma genetics. <i>Eye</i> , 2020, 34, 29-39.	2.1	48
76	The Cell and Molecular Biology of Complex Forms of Glaucoma: Updates on Genetic, Environmental, and Epigenetic Risk Factors. , 2012, 53, 2467.		47
77	Advances in the genomics of common eye diseases. <i>Human Molecular Genetics</i> , 2013, 22, R59-R65.	2.9	46
78	Comparison of Risk Factor Profiles for Primary Open-Angle Glaucoma Subtypes Defined by Pattern of Visual Field Loss: A Prospective Study. , 2015, 56, 2439.		45
79	Assessing the Association of Mitochondrial Genetic Variation With Primary Open-Angle Glaucoma Using Gene-Set Analyses. , 2016, 57, 5046.		44
80	DNA Methylation Variants at <i>HIF3A</i> Locus, B-Vitamin Intake, and Long-term Weight Change: Gene-Diet Interactions in Two U.S. Cohorts. <i>Diabetes</i> , 2015, 64, 3146-3154.	0.6	43
81	Diet quality and genetic association with body mass index: results from 3 observational studies. <i>American Journal of Clinical Nutrition</i> , 2018, 108, 1291-1300.	4.7	43
82	A Common Variant in <i>MIR182</i> Is Associated With Primary Open-Angle Glaucoma in the NEIGHBORHOOD Consortium. , 2016, 57, 4528.		42
83	Drug-induced Bilateral Secondary Angle-Closure Glaucoma. <i>Journal of Glaucoma</i> , 2016, 25, e99-e105.	1.6	41
84	Habitual coffee consumption and genetic predisposition to obesity: gene-diet interaction analyses in three US prospective studies. <i>BMC Medicine</i> , 2017, 15, 97.	5.5	41
85	Genome-wide association study of primary open-angle glaucoma in continental and admixed African populations. <i>Human Genetics</i> , 2018, 137, 847-862.	3.8	40
86	Estrogen pathway polymorphisms in relation to primary open angle glaucoma: an analysis accounting for gender from the United States. <i>Molecular Vision</i> , 2013, 19, 1471-81.	1.1	40
87	Epistatic Gene-Based Interaction Analyses for Glaucoma in eMERGE and NEIGHBOR Consortium. <i>PLoS Genetics</i> , 2016, 12, e1006186.	3.5	38
88	Disruption of the Blood-Aqueous Barrier and Lens Abnormalities in Mice Lacking Lysyl Oxidase-Like 1 (LOXL1). , 2014, 55, 856.		37
89	Non-Synonymous variants in premelanosome protein (PMEL) cause ocular pigment dispersion and pigmentary glaucoma. <i>Human Molecular Genetics</i> , 2019, 28, 1298-1311.	2.9	36
90	Association of Long-term Ambient Black Carbon Exposure and Oxidative Stress Allelic Variants With Intraocular Pressure in Older Men. <i>JAMA Ophthalmology</i> , 2019, 137, 129.	2.5	36

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91	Relation Between Time Spent Outdoors and Exfoliation Glaucoma or Exfoliation Glaucoma Suspect. <i>American Journal of Ophthalmology</i> , 2014, 158, 605-614.e1.	3.3	35
92	Intraocular Pressure, Glaucoma, and Dietary Caffeine Consumption. <i>Ophthalmology</i> , 2021, 128, 866-876.	5.2	35
93	Genome-wide association study identifies WNT7B as a novel locus for central corneal thickness in Latinos. <i>Human Molecular Genetics</i> , 2016, 25, ddw319.	2.9	34
94	Prospective Study of Oral Health and Risk of Primary Open-Angle Glaucoma in Men. <i>Ophthalmology</i> , 2016, 123, 2318-2327.	5.2	33
95	Analysis combining correlated glaucoma traits identifies five new risk loci for open-angle glaucoma. <i>Scientific Reports</i> , 2018, 8, 3124.	3.3	33
96	Hypothesis-independent pathway analysis implicates GABA and Acetyl-CoA metabolism in primary open-angle glaucoma and normal-pressure glaucoma. <i>Human Genetics</i> , 2014, 133, 1319-1330.	3.8	32
97	Genetics of Primary Inherited Disorders of the Optic Nerve: Clinical Applications. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2015, 5, a017277.	6.2	32
98	Association of a Primary Open-Angle Glaucoma Genetic Risk Score With Earlier Age at Diagnosis. <i>JAMA Ophthalmology</i> , 2019, 137, 1190.	2.5	32
99	Clinical Correlates of Computationally Derived Visual Field Defect Archetypes in Patients from a Glaucoma Clinic. <i>Current Eye Research</i> , 2017, 42, 568-574.	1.5	31
100	Genomic locus modulating corneal thickness in the mouse identifies POU6F2 as a potential risk of developing glaucoma. <i>PLoS Genetics</i> , 2018, 14, e1007145.	3.5	31
101	Nailfold Capillary Abnormalities in Primary Open-Angle Glaucoma: A Multisite Study. , 2015, 56, 7021.		30
102	DNA Copy Number Variants of Known Glaucoma Genes in Relation to Primary Open-Angle Glaucoma. <i>Investigative Ophthalmology and Visual Science</i> , 2014, 55, 8251-8258.	3.3	27
103	Characteristics of p.Gln368Ter Myocilin Variant and Influence of Polygenic Risk on Glaucoma Penetrance in the UK Biobank. <i>Ophthalmology</i> , 2021, 128, 1300-1311.	5.2	27
104	Genetic Testing for Inherited Eye Disease. <i>JAMA Ophthalmology</i> , 2013, 131, 1265.	2.5	26
105	Unusual Presentation of Presumed Posterior Polymorphous Dystrophy Associated With Iris Heterochromia, Band Keratopathy, and Keratoconus. <i>Cornea</i> , 2010, 29, 1180-1185.	1.7	25
106	The p53 Codon 72 PRO/PRO Genotype May Be Associated with Initial Central Visual Field Defects in Caucasians with Primary Open Angle Glaucoma. <i>PLoS ONE</i> , 2012, 7, e45613.	2.5	25
107	Association of Statin Use and High Serum Cholesterol Levels With Risk of Primary Open-Angle Glaucoma. <i>JAMA Ophthalmology</i> , 2019, 137, 756.	2.5	25
108	Childhood glaucoma genes and phenotypes: Focus on FOXC1 mutations causing anterior segment dysgenesis and hearing loss. <i>Experimental Eye Research</i> , 2020, 190, 107893.	2.6	23

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109	Association of <i>APOE</i> With Primary Open-Angle Glaucoma Suggests a Protective Effect for <i>APOE</i> ϵ 4. <i>Molecular Vision</i> , 2020, 61, 3.		23
110	DNA sequence variants in the tyrosinase-related protein 1 (TYRP1) gene are not associated with human pigmentary glaucoma. <i>Molecular Vision</i> , 2002, 8, 127-9.	1.1	23
111	The protective variant rs7173049 at LOXL1 locus impacts on retinoic acid signaling pathway in pseudoexfoliation syndrome. <i>Human Molecular Genetics</i> , 2019, 28, 2531-2548.	2.9	22
112	Multi-trait genome-wide association study identifies new loci associated with optic disc parameters. <i>Communications Biology</i> , 2019, 2, 435.	4.4	22
113	Reproductive factors and NOS3 variant interactions in primary open-angle glaucoma. <i>Molecular Vision</i> , 2011, 17, 2544-51.	1.1	22
114	Low prevalence of myocilin mutations in an African American population with primary open-angle glaucoma. <i>Molecular Vision</i> , 2012, 18, 2241-6.	1.1	22
115	No association between OPA1 polymorphisms and primary open-angle glaucoma in three different populations. <i>Molecular Vision</i> , 2007, 13, 2137-41.	1.1	22
116	Variations in <i>COL15A1</i> and <i>COL18A1</i> influence age of onset of primary open angle glaucoma. <i>Clinical Genetics</i> , 2013, 84, 167-174.	2.0	21
117	Lack of association of polymorphisms in homocysteine metabolism genes with pseudoexfoliation syndrome and glaucoma. <i>Molecular Vision</i> , 2008, 14, 2484-91.	1.1	21
118	Photoreceptor Layer Thinning Is an Early Biomarker for Age-Related Macular Degeneration. <i>Ophthalmology</i> , 2022, 129, 694-707.	5.2	21
119	The Genetics of Intraocular Pressure. <i>Seminars in Ophthalmology</i> , 2013, 28, 301-305.	1.6	20
120	Expression and Regulation of LOXL1 and Elastin-related Genes in Eyes With Exfoliation Syndrome. <i>Journal of Glaucoma</i> , 2014, 23, S62-S63.	1.6	20
121	Contribution of the Nurses' Health Study to the Epidemiology of Cataract, Age-Related Macular Degeneration, and Glaucoma. <i>American Journal of Public Health</i> , 2016, 106, 1684-1689.	2.7	19
122	Resting nailfold capillary blood flow in primary open-angle glaucoma. <i>British Journal of Ophthalmology</i> , 2019, 103, 203-207.	3.9	19
123	The genetics of glaucoma: Disease associations, personalised risk assessment and therapeutic opportunities—A review. <i>Clinical and Experimental Ophthalmology</i> , 2022, 50, 143-162.	2.6	19
124	Alcohol, Intraocular Pressure, and Open-Angle Glaucoma. <i>Ophthalmology</i> , 2022, 129, 637-652.	5.2	19
125	<i>EFEMP1</i> rare variants cause familial juvenile-onset open-angle glaucoma. <i>Human Mutation</i> , 2022, 43, 240-252.	2.5	19
126	Infectious Theories of Posner-Schlossman Syndrome. <i>International Ophthalmology Clinics</i> , 2011, 51, 105-115.	0.7	18

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127	Bupropion Use and Risk of Open-Angle Glaucoma among Enrollees in a Large U.S. Managed Care Network. PLoS ONE, 2015, 10, e0123682.	2.5	18
128	Genetic correlations between intraocular pressure, blood pressure and primary open-angle glaucoma: a multi-cohort analysis. European Journal of Human Genetics, 2017, 25, 1261-1267.	2.8	18
129	Genome-Wide Linkage Scan for Primary Open Angle Glaucoma: Influences of Ancestry and Age at Diagnosis. PLoS ONE, 2011, 6, e21967.	2.5	17
130	Myocilin Mutations in Patients With Normal-Tension Glaucoma. JAMA Ophthalmology, 2019, 137, 559.	2.5	17
131	Genes Associated with Human Glaucoma. Ophthalmology Clinics of North America, 2005, 18, 335-343.	1.8	17
132	Association of Rare <i>CYP39A1</i> Variants With Exfoliation Syndrome Involving the Anterior Chamber of the Eye. JAMA - Journal of the American Medical Association, 2021, 325, 753.	7.4	16
133	Association of clusterin (CLU) variants and exfoliation syndrome: An analysis in two Caucasian studies and a meta-analysis. Experimental Eye Research, 2015, 139, 115-122.	2.6	15
134	Juvenile-onset open-angle glaucoma – A clinical and genetic update. Survey of Ophthalmology, 2022, 67, 1099-1117.	4.0	15
135	The genetic basis for adult onset glaucoma: Recent advances and future directions. Progress in Retinal and Eye Research, 2022, 90, 101066.	15.5	15
136	Molecular and clinical characterization of a patient with a chromosome 4p deletion, Wolf-Hirschhorn syndrome, and congenital glaucoma. Ophthalmic Genetics, 2001, 22, 35-41.	1.2	14
137	Translating the Low Translaminal Cribrosa Pressure Gradient Hypothesis into the Clinical Care of Glaucoma. Seminars in Ophthalmology, 2016, 31, 131-139.	1.6	14
138	Genetic Susceptibility, Change in Physical Activity, and Long-term Weight Gain. Diabetes, 2017, 66, 2704-2712.	0.6	14
139	Sex hormone levels and risk of primary open-angle glaucoma in postmenopausal women. Menopause, 2018, 25, 1116-1123.	2.0	14
140	Testosterone Pathway Genetic Polymorphisms in Relation to Primary Open-Angle Glaucoma: An Analysis in Two Large Datasets. , 2018, 59, 629.		14
141	Association Between LOXL1 and Pseudoexfoliation. JAMA Ophthalmology, 2008, 126, 420.	2.4	13
142	LOXL1 Polymorphisms: Genetic Biomarkers that Presage Environmental Determinants of Exfoliation Syndrome. Journal of Glaucoma, 2018, 27, S20-S23.	1.6	13
143	DNAJC30 biallelic mutations extend mitochondrial complex I-deficient phenotypes to include recessive Leber's hereditary optic neuropathy. Journal of Clinical Investigation, 2021, 131, .	8.2	13
144	Association of Matrix Metalloproteinase-9 (MMP9) Variants with Primary Angle Closure and Primary Angle Closure Glaucoma. PLoS ONE, 2016, 11, e0157093.	2.5	13

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145	Fundus Densitometry Findings Suggest Optic Disc Hemorrhages in Primary Open-Angle Glaucoma Have an Arterial Origin. <i>American Journal of Ophthalmology</i> , 2018, 187, 108-116.	3.3	12
146	A Role for Clusterin in Exfoliation Syndrome and Exfoliation Glaucoma?. <i>Journal of Glaucoma</i> , 2018, 27, S61-S66.	1.6	12
147	Genetic Correlations Between Diabetes and Glaucoma: An Analysis of Continuous and Dichotomous Phenotypes. <i>American Journal of Ophthalmology</i> , 2019, 206, 245-255.	3.3	12
148	The Genetic Influence on Corticosteroid-Induced Ocular Hypertension: A Field Positioned for Discovery. <i>American Journal of Ophthalmology</i> , 2019, 202, 1-5.	3.3	12
149	Association of the SIX6 locus with primary open angle glaucoma in southern Chinese and Japanese. <i>Experimental Eye Research</i> , 2019, 180, 129-136.	2.6	12
150	Family-Based Genome-Wide Association Study of South Indian Pedigrees Supports <i>WNT7B</i> as a Central Corneal Thickness Locus. , 2018, 59, 2495.		11
151	Diagnostic genetic testing for patients with bilateral optic neuropathy and comparison of clinical features according to mutation status. <i>Molecular Vision</i> , 2017, 23, 548-560.	1.1	11
152	The Association between Serum Lipids and Intraocular Pressure in 2 Large United Kingdom Cohorts. <i>Ophthalmology</i> , 2022, 129, 986-996.	5.2	11
153	CPAMD8, a New Gene for Anterior Segment Dysgenesis and Childhood Glaucoma. <i>Ophthalmology</i> , 2020, 127, 767-768.	5.2	10
154	Association of the <i>CAV1</i> and <i>CAV2</i> locus with normal-tension glaucoma in Chinese and Japanese. <i>Clinical and Experimental Ophthalmology</i> , 2020, 48, 658-665.	2.6	10
155	Investigation of founder effects for the Thr377Met Myocilin mutation in glaucoma families from differing ethnic backgrounds. <i>Molecular Vision</i> , 2007, 13, 487-92.	1.1	10
156	Genome-Wide Association Study Identifies Two Common Loci Associated with Pigment Dispersion Syndrome/Pigmentary Glaucoma and Implicates Myopia in its Development. <i>Ophthalmology</i> , 2022, 129, 626-636.	5.2	10
157	Consideration for Gene-Environment Interactions as Novel Determinants of Exfoliation Syndrome. <i>International Ophthalmology Clinics</i> , 2014, 54, 29-41.	0.7	9
158	Quality Control for the Illumina HumanExome BeadChip. <i>Current Protocols in Human Genetics</i> , 2016, 90, 2.14.1-2.14.16.	3.5	9
159	Genomic loci modulating retinal ganglion cell death following elevated IOP in the mouse. <i>Experimental Eye Research</i> , 2018, 169, 61-67.	2.6	9
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