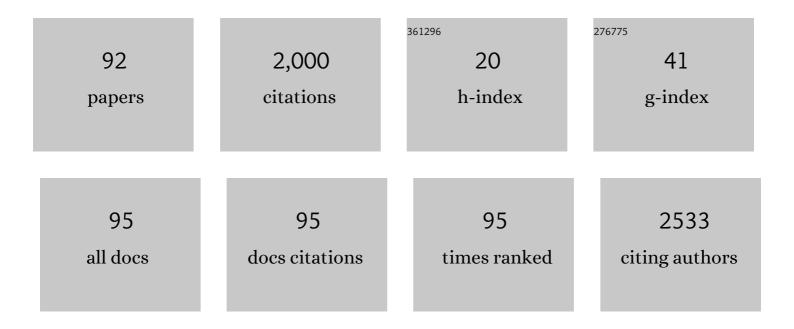
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Prevalence of refractive error in Europe: the European Eye Epidemiology (E3) Consortium. European Journal of Epidemiology, 2015, 30, 305-315.	2.5	306
2	Meta-analysis of 542,934 subjects of European ancestry identifies new genes and mechanisms predisposing to refractive error and myopia. Nature Genetics, 2020, 52, 401-407.	9.4	180
3	Differentiating drusen: Drusen and drusen-like appearances associated with ageing, age-related macular degeneration, inherited eye disease and other pathological processes. Progress in Retinal and Eye Research, 2016, 53, 70-106.	7.3	159
4	Genetic Basis of Inherited Retinal Disease in a Molecularly Characterized Cohort of More Than 3000 Families from the United Kingdom. Ophthalmology, 2020, 127, 1384-1394.	2.5	131
5	Complications of cataract surgery. Australasian journal of optometry, The, 2010, 93, 379-389.	0.6	121
6	Recovery of the human photopic electroretinogram after bleaching exposures: estimation of pigment regeneration kinetics. Journal of Physiology, 2004, 554, 417-437.	1.3	77
7	The X-linked retinopathies: Physiological insights, pathogenic mechanisms, phenotypic features and novel therapies. Progress in Retinal and Eye Research, 2021, 82, 100898.	7.3	65
8	Retinal Ganglion Cells—Diversity of Cell Types and Clinical Relevance. Frontiers in Neurology, 2021, 12, 661938.	1.1	53
9	A clinical and molecular characterisation of CRB1-associated maculopathy. European Journal of Human Genetics, 2018, 26, 687-694.	1.4	51
10	Transient Smartphone "Blindness― New England Journal of Medicine, 2016, 374, 2502-2504.	13.9	41
11	Dark adaptation of human rod bipolar cells measured from theb-wave of the scotopic electroretinogram. Journal of Physiology, 2006, 575, 507-526.	1.3	39
12	SPECTRAL DOMAIN OPTICAL COHERENCE TOMOGRAPHY FINDINGS IN LONG-TERM SILICONE OIL–RELATED VISUAL LOSS. Retina, 2015, 35, 555-563.	1.0	38
13	FUNCTIONAL AND ANATOMICAL OUTCOMES OF CHOROIDAL NEOVASCULARIZATION COMPLICATING BEST1-RELATED RETINOPATHY. Retina, 2017, 37, 1360-1370.	1.0	34
14	Macular spectral domain optical coherence tomography findings in Tanzanian endemic optic neuropathy. Brain, 2013, 136, 3418-3426.	3.7	32
15	Patient-reported benefit from oculoplastic surgery. Eye, 2012, 26, 1418-1423.	1.1	29
16	Extremely rapid recovery of human cone circulating current at the extinction of bleaching exposures. Journal of Physiology, 2005, 567, 95-112.	1.3	28
17	Effect of varying skin surface electrode position on electroretinogram responses recorded using a handheld stimulating and recording system. Documenta Ophthalmologica, 2018, 137, 79-86.	1.0	25
18	Characteristics of rhegmatogenous retinal detachment in pseudophakic and phakic eyes. Eye, 2012, 26, 1114-1121	1.1	24

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19	Common Mechanisms Underlying Refractive Error Identified in Functional Analysis of Gene Lists From Genome-Wide Association Study Results in 2 European British Cohorts. JAMA Ophthalmology, 2014, 132, 50.	1.4	23
20	Do twins share the same dress code? Quantifying relative genetic and environmental contributions to subjective perceptions of "the dress―in a classical twin study. Journal of Vision, 2017, 17, 29.	0.1	22
21	Negative electroretinograms: genetic and acquired causes, diagnostic approaches and physiological insights. Eye, 2021, 35, 2419-2437.	1.1	22
22	Phenotypic and genotypic correlation between myopia and intelligence. Scientific Reports, 2017, 7, 45977.	1.6	20
23	X-Linked Retinoschisis. Ophthalmology, 2022, 129, 542-551.	2.5	19
24	Photophobia in migraine: A symptom cluster?. Cephalalgia, 2021, 41, 1240-1248.	1.8	18
25	KCNV2-Associated Retinopathy: Genetics, Electrophysiology, and Clinical Course—KCNV2 Study Group Report 1. American Journal of Ophthalmology, 2021, 225, 95-107.	1.7	17
26	ISCEV extended protocol for derivation and analysis of the strong flash rod-isolated ERG a-wave. Documenta Ophthalmologica, 2020, 140, 5-12.	1.0	16
27	<i>CNGB1</i> â€related rodâ€cone dystrophy: A mutation review and update. Human Mutation, 2021, 42, 641-666.	1.1	16
28	Risk Factors for Visual Impairment in Patients with Sickle Cell Disease in London. European Journal of Ophthalmology, 2016, 26, 431-435.	0.7	15
29	Changes in quality of life shortly after routine cataract surgery. Canadian Journal of Ophthalmology, 2016, 51, 282-287.	0.4	15
30	Quantifying the Separation Between the Retinal Pigment Epithelium and Bruch's Membrane using Optical Coherence Tomography in Patients with Inherited Macular Degeneration. Translational Vision Science and Technology, 2020, 9, 26.	1.1	15
31	Re: Abegg et al.: Microcystic macular edema: retrograde maculopathy caused by optic neuropathy (Ophthalmology 2014;121:142-9). Ophthalmology, 2014, 121, e40.	2.5	14
32	Phenotype and genotype of 197 British patients with <scp>McArdle</scp> disease: An observational singleâ€centre study. Journal of Inherited Metabolic Disease, 2021, 44, 1409-1418.	1.7	14
33	High Heritability of Posterior Corneal Tomography, as Measured by Scheimpflug Imaging, in a Twin Study. Investigative Ophthalmology and Visual Science, 2014, 55, 8359-8364.	3.3	12
34	Outcomes of ptosis surgery assessed using a patient-reported outcome measure: an exploration of time effects. British Journal of Ophthalmology, 2014, 98, 387-390.	2.1	12
35	Relative Genetic and Environmental Contributions to Variations in Human Retinal Electrical Responses Quantified in a Twin Study. Ophthalmology, 2017, 124, 1175-1185.	2.5	12
36	Retinopathy Associated with Biallelic Mutations in PYGM (McArdle Disease). Ophthalmology, 2019, 126, 320-322.	2.5	12

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37	A genetic and clinical study of individuals with nonsyndromic retinopathy consequent upon sequence variants in <scp><i>HGSNAT</i></scp> , the gene associated with Sanfilippo C mucopolysaccharidosis. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2020, 184, 631-643.	0.7	12
38	Modelling the initial phase of the human rod photoreceptor response to the onset of steady illumination. Documenta Ophthalmologica, 2012, 124, 125-131.	1.0	11
39	The Heritability of the Ring-Like Distribution of Macular Pigment Assessed in a Twin Study. , 2014, 55, 2214.		11
40	A twin study of cilioretinal arteries, tilted discs and situs inversus. Graefe's Archive for Clinical and Experimental Ophthalmology, 2018, 256, 333-340.	1.0	11
41	KCNV2-Associated Retinopathy: Detailed Retinal Phenotype and Structural Endpoints—KCNV2 Study Group Report 2. American Journal of Ophthalmology, 2021, 230, 1-11.	1.7	11
42	Interocular Asymmetries in Axial Length and Refractive Error in 4 Cohorts. Ophthalmology, 2015, 122, 648-649.	2.5	10
43	Genome-Wide Association Study Identifies Two Common Loci Associated with Pigment Dispersion Syndrome/Pigmentary Glaucoma and Implicates Myopia in its Development. Ophthalmology, 2022, 129, 626-636.	2.5	10
44	Slowed recovery of human photopic ERG a-wave amplitude following intense bleaches: a slowing of cone pigment regeneration?. Documenta Ophthalmologica, 2012, 125, 137-147.	1.0	9
45	Comparison of ophthalmic training in 6 English-speaking countries. Canadian Journal of Ophthalmology, 2016, 51, 212-218.	0.4	9
46	Retinal thickness measurements in sickle cell patients with HbSS and HbSC genotype. Canadian Journal of Ophthalmology, 2018, 53, 420-424.	0.4	9
47	Clinical Features of a Retinopathy Associated With a Dominant Allele of the <i>RGR</i> Gene. , 2018, 59, 4812.		9
48	Choroidal macrovessels: multimodal imaging findings and review of the literature. British Journal of Ophthalmology, 2022, 106, 568-575.	2.1	9
49	Spectral Domain Optical Coherence Tomography Findings in a Case Series of Patients with Bilateral Diffuse Uveal Melanocytic Proliferation. Ocular Immunology and Inflammation, 2014, 22, 490-493.	1.0	8
50	Exploring Sex and Laterality Imbalances in Patients Undergoing Laser Retinopexy. JAMA Ophthalmology, 2015, 133, 1334.	1.4	8
51	Clinical and Genetic Findings in CTNNA1-Associated Macular Pattern Dystrophy. Ophthalmology, 2021, 128, 952-955.	2.5	8
52	Electrical responses from human retinal cone pathways associate with a common genetic polymorphism implicated in myopia. Proceedings of the National Academy of Sciences of the United States of America, 2022, 119, .	3.3	8
53	Jalili Syndrome: Cross-sectional and Longitudinal Features of Seven Patients With Cone-Rod Dystrophy and Amelogenesis Imperfecta. American Journal of Ophthalmology, 2018, 188, 123-130.	1.7	7
54	The Role of Chromosome X in Intraocular Pressure Variation and Sex-Specific Effects. , 2020, 61, 20.		7

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55	New variants and in silico analyses in GRK1 associated Oguchi disease. Human Mutation, 2021, 42, 164-176.	1.1	7
56	Swept Source OCT Image of a Choroidal Macrovessel. Ophthalmology Retina, 2017, 1, 281.	1.2	6
57	Pupil area and photopigment spectral sensitivity are relevant to study of migraine photophobia. Brain, 2017, 140, e2-e2.	3.7	6
58	Axial Length Distributions in Patients With Genetically Confirmed Inherited Retinal Diseases. , 2022, 63, 15.		6
59	Choice of Analytic Approach for Eye-Specific Outcomes: One Eye or Two?. American Journal of Ophthalmology, 2012, 153, 781-782.	1.7	5
60	Roth Spots in Infective Endocarditis. New England Journal of Medicine, 2014, 370, e38.	13.9	5
61	Electroretinography can provide objective assessment of inner retinal function prior to atrophic change on OCT. Eye, 2015, 29, 1513-1513.	1.1	5
62	Cilioretinal Artery Territory Infarction Associated With Papilledema in a Patient With Neurofibromatosis Type 2. Journal of Neuro-Ophthalmology, 2016, 36, 58-60.	0.4	5
63	Retinal findings in a patient with mutations in ABCC6 and ABCA4. Eye, 2018, 32, 1542-1543.	1.1	5
64	Reanalysis of Association of Pro50Leu Substitution in Guanylate Cyclase Activating Protein-1 With Dominant Retinal Dystrophy. JAMA Ophthalmology, 2020, 138, 200.	1.4	5
65	A clinical study of patients with novel CDHR1 genotypes associated with late-onset macular dystrophy. Eye, 2021, 35, 1482-1489.	1.1	5
66	Association Between Medication-Taking and Refractive Error in a Large General Population-Based Cohort. , 2021, 62, 15.		5
67	Exploratory Study of the Association between the Severity of Idiopathic Intracranial Hypertension and Electroretinogram Photopic Negative Response Amplitude Obtained Using a Handheld Device. Life, 2021, 11, 437.	1.1	5
68	Confirmation That Angioid Streaks Are Not Common in Ehlers-Danlos Syndrome. JAMA Ophthalmology, 2019, 137, 1463.	1.4	4
69	Inherited Retinal Disease Panels—Caveat Emptor—Truly Know Your Inherited Retinal Disease Panel. Retina, 2022, 42, 1-3.	1.0	4
70	A large animal model of <i>RDH5</i> -associated retinopathy recapitulates important features of the human phenotype. Human Molecular Genetics, 2022, 31, 1263-1277.	1.4	4
71	Human retinal dark adaptation tracked <i>in vivo</i> with the electroretinogram: insights into processes underlying recovery of cone―and rodâ€mediated vision. Journal of Physiology, 2022, 600, 4603-4621.	1.3	4
72	Using Cyanoacrylate Glue for Corneal Perforations. Cornea, 2013, 32, e193.	0.9	3

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73	Prevalence of electronegative electroretinograms in a healthy adult cohort. BMJ Open Ophthalmology, 2021, 6, e000751.	0.8	3
74	Exploring correlations between change in visual acuity following routine cataract surgery and improvement in quality of life assessed with the Glasgow Benefit Inventory. Eye, 2018, 32, 1549-1550.	1.1	2
75	Awareness of olfactory impairment in a cohort of patients with CNGB1-associated retinitis pigmentosa. Eye, 2020, 34, 783-784.	1.1	2
76	CASE REPORT: RETINOPATHY IN A PATIENT WITH COLD HEMAGGLUTININ DISEASE. Retinal Cases and Brief Reports, 2011, 5, 254-255.	0.3	1
77	Central visual disturbance associated with transient disruption of photoreceptor inner-outer segment junction. Canadian Journal of Ophthalmology, 2012, 47, e19-e20.	0.4	1
78	Outcomes of Trabeculectomy With Transconjunctival Application Versus Subconjunctival Application of Mitomycin C. Journal of Glaucoma, 2016, 25, 467-471.	0.8	1
79	Pediatric Ocular Tuberculosis – Choroidal Tubercles. Journal of Pediatrics, 2016, 169, 323.	0.9	1
80	Transient smartphone "blindness―arises from interocular differences in retinal adaptational states. Canadian Journal of Ophthalmology, 2017, 52, 425.	0.4	1
81	Repeatability of the macular pigment spatial profile: A comparison of objective versus subjective classification. Acta Ophthalmologica, 2018, 96, e797-e803.	0.6	1
82	Re: You etÂal.: Evidence of Müller glial dysfunction in patients with aquaporin-4 immunoglobulin G–positive neuromyelitis optica spectrum disorder (Ophthalmology. 2019;126:801–810). Ophthalmology, 2019, 126, e63-e64.	2.5	1
83	Re: Bhatti etÂal.: Microcystic macular edema in optic nerveÂglioma (Ophthalmology. 2020;127:930). Ophthalmology, 2021, 128, e9.	2.5	1
84	Segmented Macular Layer Volumes from Spectral Domain Optical Coherence Tomography in 184 Adult Twins: Associations With Age and Heritability. , 2020, 61, 44.		1
85	Electrophysiological Assessment in Birdshot Chorioretinopathy: Flicker Electroretinograms Recorded With a Handheld Device. Translational Vision Science and Technology, 2022, 11, 23.	1.1	1
86	Limbus vs. Fornix-Based Trabeculectomy Results. Ophthalmology, 2012, 119, 2196.	2.5	0
87	Potential effects of laterality and learning on misalignment in marking horizontal meridian. Journal of Cataract and Refractive Surgery, 2013, 39, 966.	0.7	0
88	â€~Dilatation' and â€~dilation': trends in use on both sides of the Atlantic. British Journal of Ophthalmology, 2014, 98, 845-846.	2.1	0
89	Ellipsoid Zone Change According to Glaucoma-Stage Advancement. American Journal of Ophthalmology, 2019, 197, 183.	1.7	0
90	Re: Lee etÂal.: Longitudinal changes in peripapillary retinal nerve fiber layer thickness in high myopia: a prospective, observational study (Ophthalmology. 2019;126:522-528). Ophthalmology, 2020, 127, e9-e10.	2.5	0

#	Article	IF	CITATIONS
91	Money, motivation, and medicine. BMJ: British Medical Journal, 2010, 341, c6830-c6830.	2.4	Ο
92	A genome-wide analysis of 340 318 participants identifies four novel loci associated with the age of first spectacle wear. Human Molecular Genetics, 2022, , .	1.4	0