## List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Increased LCN2 (lipocalin 2) in the RPE decreases autophagy and activates inflammasome-ferroptosis processes in a mouse model of dry AMD. Autophagy, 2023, 19, 92-111.	4.3	41
2	Review of the Current Literature and Our Experience on the Value of OCT-angiography in White Dot Syndromes. Ocular Immunology and Inflammation, 2022, 30, 364-378.	1.0	9
3	Natural history of patients with Leber hereditary optic neuropathy—results from the REALITY study. Eye, 2022, 36, 818-826.	1.1	37
4	Early and late stage gene therapy interventions for inherited retinal degenerations. Progress in Retinal and Eye Research, 2022, 86, 100975.	7.3	85
5	Choriocapillaris: Fundamentals and advancements. Progress in Retinal and Eye Research, 2022, 87, 100997.	7.3	56
6	Near infrared autofluorescence imaging of retinal pigmented epithelial cells using 663 nm excitation. Eye, 2022, 36, 1878-1883.	1.1	2
7	Clinical Features and Multimodal Imaging in Atypical Posterior Uveitis Secondary to <i>Bartonella Henselae</i> Infection. Ocular Immunology and Inflammation, 2022, 30, 2047-2054.	1.0	1
8	Design of a radial multi-offset detection pattern for in vivo phase contrast imaging of the inner retina in humans. Biomedical Optics Express, 2022, 13, 117.	1.5	11
9	Simultaneous perception of prosthetic and natural vision in AMD patients. Nature Communications, 2022, 13, 513.	5.8	60
10	Planar polarity in primate cone photoreceptors: a potential role in Stiles Crawford effect phototropism. Communications Biology, 2022, 5, 89.	2.0	11
11	Indocyanine Green Angiography Features in Acute Syphilitic Posterior Placoid Chorioretinitis. American Journal of Ophthalmology, 2022, 241, 40-46.	1.7	4
12	Evaluation of neuroprotective and immunomodulatory properties of mesenchymal stem cells in an ex vivo retinal explant model. Journal of Neuroinflammation, 2022, 19, 63.	3.1	11
13	Three‥ear Safety Results of SAR422459 (EIAVâ€ABCA4) Gene Therapy in Patients With ABCA4â€Associated Stargardt Disease: An Open‣abel Doseâ€Escalation Phase I/IIa Clinical Trial, Cohorts 1â€5. American Journal of Ophthalmology, 2022, 240, 285-301.	1.7	24
14	Driving behaviour and visual compensation in glaucoma patients: Evaluation on a driving simulator. Clinical and Experimental Ophthalmology, 2022, , .	1.3	1
15	Pathology of the Retina and Vitreous. , 2022, , 6315-6379.		0
16	Macular Dystrophies. , 2022, , 3967-3995.		0
17	New Editing Tools for Gene Therapy in Inherited Retinal Dystrophies. CRISPR Journal, 2022, 5, 377-388.	1.4	9
18	Retinal Phenotype of Patients with <i>CLRN1</i> -Associated Usher 3A Syndrome in French Light4Deaf Cohort. , 2022, 63, 25.		0

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19	Large Benefit from Simple Things: High-Dose Vitamin A Improves RBP4-Related Retinal Dystrophy. International Journal of Molecular Sciences, 2022, 23, 6590.	1.8	4
20	Dynamic full-field optical coherence tomography allows live imaging of retinal pigment epithelium stress model. Communications Biology, 2022, 5, .	2.0	10
21	Absence of lenadogene nolparvovec DNA in a brain tumor biopsy from a patient in the REVERSE clinical study, a case report. BMC Neurology, 2022, 22, .	0.8	4
22	Quality of Evidence in Ophthalmology: An Overview of Cochrane Reviews. Ophthalmology, 2021, 128, 330-332.	2.5	5
23	<scp><i>WDR34</i></scp> , a candidate gene for nonâ€syndromic rodâ€cone dystrophy. Clinical Genetics, 2021, 99, 298-302.	1.0	7
24	Role of glia in optic nerve. Progress in Retinal and Eye Research, 2021, 81, 100886.	7.3	23
25	Gene Therapies for the Treatment of Leber Hereditary Optic Neuropathy. International Ophthalmology Clinics, 2021, 61, 195-208.	0.3	14
26	Macular Dystrophies. , 2021, , 1-29.		0
27	Delivery of Genetic Information: Viral Vector and Nonviral Vector Gene Therapies. International Ophthalmology Clinics, 2021, 61, 35-57.	0.3	3
28	Near-infrared fundus autofluorescence alterations correlate with swept-source optical coherence tomography angiography findings in patients with retinitis pigmentosa. Scientific Reports, 2021, 11, 3180.	1.6	7
29	Safety of Intravitreal Gene Therapy for Treatment of Subjects with Leber Hereditary Optic Neuropathy due to Mutations in the Mitochondrial ND4 Gene: The REVEAL Study. BioDrugs, 2021, 35, 201-214.	2.2	25
30	<i>CHM</i> mutation spectrum and disease: An update at the time of human therapeutic trials. Human Mutation, 2021, 42, 323-341.	1.1	8
31	Of fluid and tubes. Journal Francais D'Ophtalmologie, 2021, 44, 277-278.	0.2	0
32	βA1-crystallin regulates glucose metabolism and mitochondrial function in mouse retinal astrocytes by modulating PTP1B activity. Communications Biology, 2021, 4, 248.	2.0	10
33	Retinal Phenotype of Patients With Isolated Retinal Degeneration Due to <i>CLN3</i> Pathogenic Variants in a French Retinitis Pigmentosa Cohort. JAMA Ophthalmology, 2021, 139, 278.	1.4	21
34	Cone-Enriched Cultures from the Retina of Chicken Embryos to Study Rod to Cone Cellular Interactions. Journal of Visualized Experiments, 2021, , .	0.2	2
35	Control of Microbial Opsin Expression in Stem Cell Derived Cones for Improved Outcomes in Cell Therapy. Frontiers in Cellular Neuroscience, 2021, 15, 648210.	1.8	10
36	Restoration of mGluR6 Localization Following AAV-Mediated Delivery in a Mouse Model of Congenital Stationary Night Blindness. , 2021, 62, 24.		10

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37	Reproducing diabetic retinopathy features using newly developed human inducedâ€pluripotent stem cellâ€derived retinal Müller glial cells. Glia, 2021, 69, 1679-1693.	2.5	11
38	A New Mouse Model for Complete Congenital Stationary Night Blindness Due to Gpr179 Deficiency. International Journal of Molecular Sciences, 2021, 22, 4424.	1.8	3
39	DEEP PHENOTYPING AND FURTHER INSIGHTS INTO ITM2B-RELATED RETINAL DYSTROPHY. Retina, 2021, 41, 872-881.	1.0	2
40	Tackling the Challenges of Product Development Through a Collaborative Rare Disease Network: The Foundation Fighting Blindness Consortium. Translational Vision Science and Technology, 2021, 10, 23.	1.1	3
41	Efficacy and Safety of Intravitreal Gene Therapy for Leber Hereditary Optic Neuropathy Treated within 6 Months of Disease Onset. Ophthalmology, 2021, 128, 649-660.	2.5	87
42	Partial recovery of visual function in a blind patient after optogenetic therapy. Nature Medicine, 2021, 27, 1223-1229.	15.2	335
43	Intravitreal Gene Therapy vs. Natural History in Patients With Leber Hereditary Optic Neuropathy Carrying the m.11778G>A ND4 Mutation: Systematic Review and Indirect Comparison. Frontiers in Neurology, 2021, 12, 662838.	1.1	42
44	<i>CNGB1</i> â€related rodâ€cone dystrophy: A mutation review and update. Human Mutation, 2021, 42, 641-666.	1.1	16
45	Mobile brain/body imaging of landmarkâ€based navigation with highâ€density EEG. European Journal of Neuroscience, 2021, 54, 8256-8282.	1.2	28
46	Functional ultrasound imaging of the spreading activity following optogenetic stimulation of the rat visual cortex. Scientific Reports, 2021, 11, 12603.	1.6	11
47	Novel TTLL5 Variants Associated with Cone-Rod Dystrophy and Early-Onset Severe Retinal Dystrophy. International Journal of Molecular Sciences, 2021, 22, 6410.	1.8	9
48	Mutated CCDC51 Coding for a Mitochondrial Protein, MITOK Is a Candidate Gene Defect for Autosomal Recessive Rod-Cone Dystrophy. International Journal of Molecular Sciences, 2021, 22, 7875.	1.8	3
49	βA3/A1-crystallin regulates apical polarity and EGFR endocytosis in retinal pigmented epithelial cells. Communications Biology, 2021, 4, 850.	2.0	13
50	Cystoid maculopathy is a frequent feature of Cohen syndrome-associated retinopathy. Scientific Reports, 2021, 11, 16412.	1.6	7
51	Characteristics of Retinitis Pigmentosa Associated with ADGRV1 and Comparison with USH2A in Patients from a Multicentric Usher Syndrome Study Treatrush. International Journal of Molecular Sciences, 2021, 22, 10352.	1.8	3
52	A2E-induced inflammation and angiogenesis in RPE cells in vitro are modulated by PPAR-α, -β/Ĩ´, -γ, and RXR antagonists and by norbixin. Aging, 2021, 13, 22040-22058.	1.4	12
53	Substantial restoration of night vision in adult mice with congenital stationary night blindness. Molecular Therapy - Methods and Clinical Development, 2021, 22, 15-25.	1.8	10
54	Glare and Mobility Performance in Glaucoma. Journal of Glaucoma, 2021, Publish Ahead of Print, 963-970.	0.8	2

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55	Assessing Photoreceptor Status in Retinal Dystrophies: From High-Resolution Imaging to Functional Vision. American Journal of Ophthalmology, 2021, 230, 12-47.	1.7	19
56	BNIP3L-mediated mitophagy is required for mitochondrial remodeling during the differentiation of optic nerve oligodendrocytes. Autophagy, 2021, 17, 3140-3159.	4.3	37
57	A Splice Variant in SLC16A8 Gene Leads to Lactate Transport Deficit in Human iPS Cell-Derived Retinal Pigment Epithelial Cells. Cells, 2021, 10, 179.	1.8	12
58	Optogenetic therapy: high spatiotemporal resolution and pattern discrimination compatible with vision restoration in non-human primates. Communications Biology, 2021, 4, 125.	2.0	65
59	Long-Term Follow-Up After Unilateral Intravitreal Gene Therapy for Leber Hereditary Optic Neuropathy: The RESTORE Study. Journal of Neuro-Ophthalmology, 2021, 41, 309-315.	0.4	30
60	Improved performance and safety from Argus II retinal prosthesis postâ€approval study in France. Acta Ophthalmologica, 2021, 99, e1212-e1221.	0.6	8
61	Association of Smoking, Alcohol Consumption, Blood Pressure, Body Mass Index, and Glycemic Risk Factors With Age-Related Macular Degeneration. JAMA Ophthalmology, 2021, 139, 1299.	1.4	29
62	The metabolic signaling of the nucleoredoxin-like 2 gene supports brain function. Redox Biology, 2021, 48, 102198.	3.9	7
63	A New Method for Visualizing Drusen and Their Progression in Flood-Illumination Adaptive Optics Ophthalmoscopy. Translational Vision Science and Technology, 2021, 10, 19.	1.1	9
64	Age-related preference for geometric spatial cues during real-world navigation. Nature Human Behaviour, 2020, 4, 88-99.	6.2	44
65	High-Resolution Imaging of Retinal Vasculitis by Flood Illumination Adaptive Optics Ophthalmoscopy: A Follow-up Study. Ocular Immunology and Inflammation, 2020, 28, 1171-1180.	1.0	6
66	PHENOTYPIC CHARACTERISTICS OF ROD–CONE DYSTROPHY ASSOCIATED WITH MYO7A MUTATIONS IN A LARGE FRENCH COHORT. Retina, 2020, 40, 1603-1615.	1.0	16
67	Clinical-grade production and safe delivery of human ESC derived RPE sheets in primates and rodents. Biomaterials, 2020, 230, 119603.	5.7	21
68	IL-1β induces rod degeneration through the disruption of retinal glutamate homeostasis. Journal of Neuroinflammation, 2020, 17, 1.	3.1	172
69	Behavioural responses to a photovoltaic subretinal prosthesis implanted in non-human primates. Nature Biomedical Engineering, 2020, 4, 172-180.	11.6	55
70	Correlations Between Subjective Evaluation of Quality of Life, Visual Field Loss, and Performance in Simulated Activities of Daily Living in Glaucoma Patients. Journal of Glaucoma, 2020, 29, 970-974.	0.8	18
71	Topical treatment with a mu opioid receptor agonist alleviates corneal allodynia and corneal nerve sensitization in mice. Biomedicine and Pharmacotherapy, 2020, 132, 110794.	2.5	12
72	The 10q26 Risk Haplotype of Age-Related Macular Degeneration Aggravates Subretinal Inflammation by Impairing Monocyte Elimination. Immunity, 2020, 53, 429-441.e8.	6.6	47

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73	ATTENUATION OUTER RETINAL BANDS ON OPTICAL COHERENCE TOMOGRAPHY FOLLOWING MACULAR EDEMA. Retina, 2020, 40, 2232-2239.	1.0	3
74	Insulin inhibits inflammation-induced cone death in retinal detachment. Journal of Neuroinflammation, 2020, 17, 358.	3.1	9
75	Towards optogenetic vision restoration with high resolution. PLoS Computational Biology, 2020, 16, e1007857.	1.5	20
76	VEGF is an autocrine/paracrine neuroprotective factor for injured retinal ganglion neurons. Scientific Reports, 2020, 10, 12409.	1.6	48
77	Generation of a Transplantable Population of Human iPSC-Derived Retinal Ganglion Cells. Frontiers in Cell and Developmental Biology, 2020, 8, 585675.	1.8	30
78	Phototoxic damage to cone photoreceptors can be independent of the visual pigment: the porphyrin hypothesis. Cell Death and Disease, 2020, 11, 711.	2.7	16
79	Dynamic full-field optical coherence tomography: 3D live-imaging of retinal organoids. Light: Science and Applications, 2020, 9, 140.	7.7	71
80	Bilateral visual improvement with unilateral gene therapy injection for Leber hereditary optic neuropathy. Science Translational Medicine, 2020, 12, .	5.8	128
81	Ophthalmology Practice During the Coronavirus Disease 2019 Pandemic: The University of Pittsburgh Experience in Promoting Clinic Safety and Embracing Video Visits. Ophthalmology and Therapy, 2020, 9, 1-9.	1.0	53
82	Microstructure of the retinal pigment epithelium near-infrared autofluorescence in healthy young eyes and in patients with AMD. Scientific Reports, 2020, 10, 9561.	1.6	19
83	Incorporating Video Visits into Ophthalmology Practice: A Retrospective Analysis and Patient Survey to Assess Initial Experiences and Patient Acceptability at an Academic Eye Center. Ophthalmology and Therapy, 2020, 9, 549-562.	1.0	24
84	Functional ultrasound imaging of deep visual cortex in awake nonhuman primates. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 14453-14463.	3.3	44
85	Metabolic and Redox Signaling of the Nucleoredoxin-Like-1 Gene for the Treatment of Genetic Retinal Diseases. International Journal of Molecular Sciences, 2020, 21, 1625.	1.8	20
86	Faster Sensitivity Loss around Dense Scotomas than for Overall Macular Sensitivity in Stargardt Disease: ProgStar Report No. 14. American Journal of Ophthalmology, 2020, 216, 219-225.	1.7	20
87	AAV-Mediated Gene Delivery to 3D Retinal Organoids Derived from Human Induced Pluripotent Stem Cells. International Journal of Molecular Sciences, 2020, 21, 994.	1.8	51
88	Opsins for vision restoration. Biochemical and Biophysical Research Communications, 2020, 527, 325-330.	1.0	22
89	Identification and characterization of novel TRPM1 autoantibodies from serum of patients with melanoma-associated retinopathy. PLoS ONE, 2020, 15, e0231750.	1.1	12
90	Baseline Visual Field Findings in the RUSH2A Study: Associated Factors and Correlation With Other Measures of Disease Severity. American Journal of Ophthalmology, 2020, 219, 87-100.	1.7	22

# ARTICLE IF CITATIONS Systemic administration of the di-apocarotenoid norbixin (BIO201) is neuroprotective, preserves photoreceptor function and inhibits A2E and lipofuscin accumulation in animal models of age-related 1.4 macular degeneration and Stargardt disease. Aging, 2020, 12, 6151-6171. Pathology of the Retina and Vitreous., 2020, , 1-66. 92 0 93 AAV-Mediated Gene to Foveal Cones. Methods in Molecular Biology, 2020, 2173, 101-112. 0.4 Curved-field optical coherence tomography: large-field imaging of human corneal cells and nerves. 94 4.8 19 Optica, 2020, 7, 872. Towards optogenetic vision restoration with high resolution., 2020, 16, e1007857. 96 Towards optogenetic vision restoration with high resolution., 2020, 16, e1007857. 0 Towards optogenetic vision restoration with high resolution., 2020, 16, e1007857. Towards optogenetic vision restoration with high resolution., 2020, 16, e1007857. 98 0 Towards optogenetic vision restoration with high resolution., 2020, 16, e1007857. 100 Towards optogenetic vision restoration with high resolution., 2020, 16, e1007857. 0 Detailed genetic characteristics of an international large cohort of patients with Stargardt disease: 2.1 ProgStar study report 8. British Journal of Ophthalmology, 2019, 103, 390-397. Scotopic Microperimetric Assessment of Rod Function in Stargardt Disease (SMART) Study: Design and 102 1.0 26 Baseline Characteristics (Report No. 1). Ophthalmic Research, 2019, 61, 36-43. A Comparison of the Dexamethasone Implant (Ozurdex®) and Inferior Fornix-Based Sub-Tenon Triamcinolone Acetonide for Treatment of Inflammatory Ocular Diseases. Ocular Immunology and 1.0 Inflammation, 2019, 27, 319-329. Progression of Stargardt Disease as Determined by Fundus Autofluorescence Over a 12-Month Period. 104 1.4 57 JAMA Ophthalmology, 2019, 137, 1134. Full-field electroretinography, visual acuity and visual fields in Usher syndrome: a multicentre European study. Documenta Ophthalmologica, 2019, 139, 151-160. Odysight: A Mobile Medical Application Designed for Remote Monitoringâ€"A Prospective Study 106 1.0 24 Comparison with Standard Clinical Eye Tests. Ophthalmology and Therapy, 2019, 8, 461-476. Mo-derived perivascular macrophage recruitment protects against endothelial cell death in retinal 3.1 vein occlusion. Journal of Neuroinflammation, 2019, 16, 157 Prevalence of ABCA4 Deep-Intronic Variants and Related Phenotype in An Unsolved "One-Hit―Cohort 108 1.8 26 with Stargardt Disease. International Journal of Molecular Sciences, 2019, 20, 5053.

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109	Outer Retinal Alterations Associated With Visual Outcomes in Best Vitelliform Macular Dystrophy. American Journal of Ophthalmology, 2019, 208, 429-437.	1.7	14
110	Ocular injuries caused by less-lethal weapons in France. Lancet, The, 2019, 394, 1616-1617.	6.3	14
111	Phenotype Analysis of Retinal Dystrophies in Light of the Underlying Genetic Defects: Application to Cone and Cone-Rod Dystrophies. International Journal of Molecular Sciences, 2019, 20, 4854.	1.8	20
112	Restoration of visual function by transplantation of optogenetically engineered photoreceptors. Nature Communications, 2019, 10, 4524.	5.8	92
113	Depicting brighter possibilities for treating blindness. Science Translational Medicine, 2019, 11, .	5.8	24
114	Longitudinal Clinical Follow-up and Genetic Spectrum of Patients With Rod-Cone Dystrophy Associated With Mutations in <i>PDE6A</i> and <i>PDE6B</i> . JAMA Ophthalmology, 2019, 137, 669.	1.4	32
115	Where are the missing gene defects in inherited retinal disorders? Intronic and synonymous variants contribute at least to 4% of <i>CACNA1F</i> â€mediated inherited retinal disorders. Human Mutation, 2019, 40, 765-787.	1.1	24
116	Functional ultrasound imaging of the brain reveals propagation of task-related brain activity in behaving primates. Nature Communications, 2019, 10, 1400.	5.8	90
117	A Mathematical Analysis of Aerobic Glycolysis Triggered by Glucose Uptake in Cones. Scientific Reports, 2019, 9, 4162.	1.6	18
118	The development of white matter structural changes during the process of deterioration of the visual field. Scientific Reports, 2019, 9, 2085.	1.6	12
119	Probing dynamic processes in the eye at multiple spatial and temporal scales with multimodal full field OCT. Biomedical Optics Express, 2019, 10, 731.	1.5	34
120	Generation of human induced pluripotent stem cell lines from a patient with ITM2B-related retinal dystrophy and a non mutated brother. Stem Cell Research, 2019, 41, 101625.	0.3	4
121	The primate model for understanding and restoring vision. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 26280-26287.	3.3	73
122	Six-Month Safety and Efficacy of the Intelligent Retinal Implant System II Device in Retinitis Pigmentosa. Ophthalmology, 2019, 126, 637-639.	2.5	31
123	Effects of corneal injury on ciliary nerve fibre activity and corneal nociception in mice: A behavioural and electrophysiological study. European Journal of Pain, 2019, 23, 589-602.	1.4	22
124	Whole exome sequencing resolves complex phenotype and identifies <i>CC2D2A</i> mutations underlying nonâ€syndromic rodâ€cone dystrophy. Clinical Genetics, 2019, 95, 329-333.	1.0	19
125	AUTOSOMAL DOMINANT VITREORETINOCHOROIDOPATHY. Retina, 2019, 39, 867-878.	1.0	6
126	Choroidal vasculature imaging with laser Doppler holography. Biomedical Optics Express, 2019, 10, 995.	1.5	27

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127	Waveform analysis of human retinal and choroidal blood flow with laser Doppler holography. Biomedical Optics Express, 2019, 10, 4942.	1.5	18
128	CD36 Deficiency Inhibits Retinal Inflammation and Retinal Degeneration in Cx3cr1 Knockout Mice. Frontiers in Immunology, 2019, 10, 3032.	2.2	9
129	Comparative Biochemical Outcomes, Effectiveness and Tolerance of Densiron 68 and Oxane HD for the Management of Complicated Retinal Detachment. Türk Oftalmoloji Dergisi, 2019, 49, 334-341.	0.4	6
130	Visual restoration by an infrared photovoltaic implant and by optogenetic therapy: validation in non-human primates. Journal of Vision, 2019, 19, 48.	0.1	0
131	Implications of monocular vision for racing drivers. , 2019, 14, e0226308.		0
132	Implications of monocular vision for racing drivers. , 2019, 14, e0226308.		0
133	Implications of monocular vision for racing drivers. , 2019, 14, e0226308.		0
134	Implications of monocular vision for racing drivers. , 2019, 14, e0226308.		0
135	Implications of monocular vision for racing drivers. , 2019, 14, e0226308.		0
136	Implications of monocular vision for racing drivers. , 2019, 14, e0226308.		0
137	Light action spectrum on oxidative stress and mitochondrial damage in A2E-loaded retinal pigment epithelium cells. Cell Death and Disease, 2018, 9, 287.	2.7	92
138	Safety of rAAV2/2-ND4 Gene Therapy for Leber Hereditary Optic Neuropathy. Ophthalmology, 2018, 125, 945-947.	2.5	82
139	A novel nonsense variant in <i>REEP6</i> is involved in a sporadic rod one dystrophy case. Clinical Genetics, 2018, 93, 707-711.	1.0	7
140	<i>MERTK</i> mutation update in inherited retinal diseases. Human Mutation, 2018, 39, 887-913.	1.1	41
141	Multimodal imaging including semiquantitative short-wavelength and near-infrared autofluorescence in achromatopsia. Scientific Reports, 2018, 8, 5665.	1.6	10
142	Retinal Prostheses: Other Therapies and Future Directions. Essentials in Ophthalmology, 2018, , 105-125.	0.0	1
143	Adapted Surgical Procedure for Argus II Retinal Implantation: Feasibility, Safety, Efficiency, and Postoperative Anatomic Findings. Ophthalmology Retina, 2018, 2, 276-287.	1.2	20
144	Dietary, environmental, and genetic risk factors of Extensive Macular Atrophy with Pseudodrusen, a severe bilateral macular atrophy of middle-aged patients. Scientific Reports, 2018, 8, 6840.	1.6	12

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145	Chronic exposure to tumor necrosis factor alpha induces retinal pigment epithelium cell dedifferentiation. Journal of Neuroinflammation, 2018, 15, 85.	3.1	25
146	Otx2-Genetically Modified Retinal Pigment Epithelial Cells Rescue Photoreceptors after Transplantation. Molecular Therapy, 2018, 26, 219-237.	3.7	19
147	TREATMENT OF MACULAR FOLDS COMPLICATING RETINAL DETACHMENT SURGERY USING AIR FOR RETINAL UNFOLDING. Retinal Cases and Brief Reports, 2018, 12, 228-230.	0.3	15
148	Noninvasive gene delivery to foveal cones for vision restoration. JCI Insight, 2018, 3, .	2.3	102
149	Correlation Between Visual Function and Performance of Simulated Daily Living Activities in Glaucomatous Patients. Journal of Glaucoma, 2018, 27, 1017-1024.	0.8	24
150	Facing hatred. Science, 2018, 362, 621-621.	6.0	0
151	Optogenetic Light Sensors in Human Retinal Organoids. Frontiers in Neuroscience, 2018, 12, 789.	1.4	48
152	Defined Xeno-free and Feeder-free Culture Conditions for the Generation of Human iPSC-derived Retinal Cell Models. Journal of Visualized Experiments, 2018, , .	0.2	10
153	Translational Retinal Research and Therapies. Translational Vision Science and Technology, 2018, 7, 8.	1.1	11
154	In vivo laser Doppler holography of the human retina. Biomedical Optics Express, 2018, 9, 4113.	1.5	26
155	Visual brain plasticity induced by central and peripheral visual field loss. Brain Structure and Function, 2018, 223, 3473-3485.	1.2	13
156	Restoring vision. Nature, 2018, 557, 359-367.	13.7	108
157	In vivo high resolution human corneal imaging using full-field optical coherence tomography. Biomedical Optics Express, 2018, 9, 557.	1.5	79
158	In vivo high-resolution human retinal imaging with wavefront-correctionless full-field OCT. Optica, 2018, 5, 409.	4.8	37
159	Expanding the Mutation Spectrum in ABCA4: Sixty Novel Disease Causing Variants and Their Associated Phenotype in a Large French Stargardt Cohort. International Journal of Molecular Sciences, 2018, 19, 2196.	1.8	22
160	Usher Syndrome and Color Vision. Current Eye Research, 2018, 43, 1295-1301.	0.7	3
161	Increase in Ocular Syphilis Cases at Ophthalmologic Reference Center, France, 2012–2015. Emerging Infectious Diseases, 2018, 24, 193-200.	2.0	36
162	Maintaining Cone Function in Rod-Cone Dystrophies. Advances in Experimental Medicine and Biology, 2018, 1074, 499-509.	0.8	15

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163	Phenotypic Characteristics of a French Cohort of Patients with X-Linked Retinoschisis. Ophthalmology, 2018, 125, 1587-1596.	2.5	25
164	Characterization and Transplantation of CD73-Positive Photoreceptors Isolated from Human iPSC-Derived Retinal Organoids. Stem Cell Reports, 2018, 11, 665-680.	2.3	128
165	Longitudinal Changes of Fixation Location and Stability Within 12 Months in Stargardt Disease: ProgStar Report No. 12. American Journal of Ophthalmology, 2018, 193, 54-61.	1.7	24
166	In vivo imaging through the entire thickness of human cornea by full-field optical coherence tomography. , 2018, , .		1
167	Vers la restauration visuelle, un point d'étape. Bulletin De L'Academie Nationale De Medecine, 2018, 202, 2059-2068.	0.0	0
168	Generation of Storable Retinal Organoids and Retinal Pigmented Epithelium from Adherent Human iPS Cells in Xeno-Free and Feeder-Free Conditions. Stem Cells, 2017, 35, 1176-1188.	1.4	186
169	Reorganization of early visual cortex functional connectivity following selective peripheral and central visual loss. Scientific Reports, 2017, 7, 43223.	1.6	24
170	Complement Factor H Inhibits CD47-Mediated Resolution of Inflammation. Immunity, 2017, 46, 261-272.	6.6	132
171	Impact of Retinitis Pigmentosa on Quality of Life, Mental Health, and Employment Among Young Adults. American Journal of Ophthalmology, 2017, 177, 169-174.	1.7	50
172	3D functional ultrasound imaging of the cerebral visual system in rodents. NeuroImage, 2017, 149, 267-274.	2.1	82
173	Multimodal imaging findings in â€ <sup>-</sup> hyper-early' stage MEWDS. British Journal of Ophthalmology, 2017, 101, 1381-1385.	2.1	18
174	Retinitis Pigmentosa and Other Dystrophies. Developments in Ophthalmology, 2017, 58, 191-201.	0.1	19
175	Usher syndrome type 1–associated cadherins shape the photoreceptor outer segment. Journal of Cell Biology, 2017, 216, 1849-1864.	2.3	47
176	Multimodal imaging and functional correlations identify unusual cases of macular retinal pigment epithelium hypopigmentation occurring without functional loss. Documenta Ophthalmologica, 2017, 135, 77-83.	1.0	3
177	Visual Acuity Change over 12 Months in the Prospective Progression of Atrophy Secondary to Stargardt Disease (ProgStar) Study. Ophthalmology, 2017, 124, 1640-1651.	2.5	43
178	On phagocytes and macular degeneration. Progress in Retinal and Eye Research, 2017, 61, 98-128.	7.3	121
179	Col4a1 mutation generates vascular abnormalities correlated with neuronal damage in a mouse model of HANAC syndrome. Neurobiology of Disease, 2017, 100, 52-61.	2.1	9
180	Lebecetin, a Câ€ŧype lectin, inhibits choroidal and retinal neovascularization. FASEB Journal, 2017, 31, 1107-1119.	0.2	17

#	Article	IF	CITATIONS
181	<i>ARL2BP</i> mutations account for 0.1% of autosomal recessive rod one dystrophies with the report of a novel splice variant. Clinical Genetics, 2017, 92, 109-111.	1.0	7
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