Shi-Ying Li

List of Publications by Year
in descending order

40	953	840585 11	477173 29
papers	citations	h-index	g-index

50	50	50	1118
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Characteristics of punctate inner choroidopathy complicated by choroidal neovascularisation on Multispectral Imaging in comparison with other imaging modalities. Ocular Immunology and Inflammation, 2022, 30, 402-408.	1.0	2
2	Human Umbilical Cord Blood-Derived CD133+CD34+ Cells Protect Retinal Endothelial Cells and Ganglion Cells in X-Irradiated Rats through Angioprotective and Neurotrophic Factors. Frontiers in Cell and Developmental Biology, 2022, 10, 801302.	1.8	2
3	ISCEV Standard for full-field clinical electroretinography (2022 update). Documenta Ophthalmologica, 2022, 144, 165-177.	1.0	179
4	Ocular abnormalities in a large patient cohort with retinitis pigmentosa in Western China. BMC Ophthalmology, 2021, 21, 43.	0.6	11
5	Quantitative assessment of visual pathway function in blind retinitis pigmentosa patients. Clinical Neurophysiology, 2021, 132, 392-403.	0.7	10
6	Ocular Characteristics of Patients With Bardetâ€“Biedl Syndrome Caused by Pathogenic BBS Gene Variation in a Chinese Cohort. Frontiers in Cell and Developmental Biology, 2021, 9, 635216.	1.8	12
7	Electrophysiology as a prognostic indicator of visual recovery in diabetic patients undergoing cataract surgery. Graefe's Archive for Clinical and Experimental Ophthalmology, 2021, 259, 1879-1887.	1.0	4
8	Molecular genetics with clinical characteristics of Leber congenital amaurosis in the Han population of western China. Ophthalmic Genetics, 2021, 42, 392-401.	0.5	6
9	Intravitreally injected ranibizumab versus photodynamic therapy for CNV secondary to choroidal osteoma: a 7-year follow-up case report. International Journal of Ophthalmology, 2021, 14, 940-944.	0.5	2
10	A phase I clinical trial of human embryonic stem cellâ€derived retinal pigment epithelial cells for earlyâ€stage Stargardt macular degeneration: 5â€years' followâ€up. Cell Proliferation, 2021, 54, e13100.	2.4	33
11	Comparative Study of a Modified Sub-Tenonâ€™s Capsule Injection of Triamcinolone Acetonide and the Intravenous Infusion of Umbilical Cord Mesenchymal Stem Cells in Retinitis Pigmentosa Combined With Macular Edema. Frontiers in Pharmacology, 2021, 12, 694225.	1.6	1
12	ISCEV standard for clinical multifocal electroretinography (mfERG) (2021 update). Documenta Ophthalmologica, 2021, 142, 5-16.	1.0	81
13	Long-term follow-up of a Chinese patient with KCNV2-retinopathy. Ophthalmic Genetics, 2021, 42, 144-149.	0.5	0
14	Ocular Characteristics of Patients with Leber Congenital Amaurosis 6 Caused by Pathogenic RPGRIP1 Gene Variation in a Chinese Cohort. Journal of Ophthalmology, 2021, 2021, 1-8.	0.6	3
15	Intravenous Infusion of Umbilical Cord Mesenchymal Stem Cells Maintains and Partially Improves Visual Function in Patients with Advanced Retinitis Pigmentosa. Stem Cells and Development, 2020, 29, 1029-1037.	1.1	28
16	Predictors of health-related quality of life in Chinese patients receiving treatment for neovascular age-related macular degeneration: a prospective longitudinal study. BMC Ophthalmology, 2020, 20, 291.	0.6	4
17	Clinical and genetic characteristics of Stargardt disease in a large Western China cohort: Report 1. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2020, 184, 694-707.	0.7	7
18	Nonsyndromic retinitis pigmentosa caused by two novel variants in the HGSNAT gene in a Chinese family. Ophthalmic Genetics, 2020, 41, 390-393.	0.5	1

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19	Clinical and Genetic Characteristics of 15 Affected Patients From 12 Japanese Families with GUCY2D-Associated Retinal Disorder. Translational Vision Science and Technology, 2020, 9, 2.	1.1	15
20	Oguchi disease caused by a homozygous novel SAG splicing alteration associated with the multiple evanescent white dot syndrome: A 15-month follow-up. Documenta Ophthalmologica, 2020, 141, 217-226.	1.0	4
21	Electrophysiological and Structural Changes in Chinese Patients with LHON. Journal of Ophthalmology, 2020, 2020, 1-9.	0.6	5
22	Identification of 13 novel USH2A mutations in Chinese retinitis pigmentosa and Usher syndrome patients by targeted next-generation sequencing. Bioscience Reports, 2020, 40, .	1.1	7
23	Genetic analysis in a cohort of patients with hereditary optic neuropathies in Southwest of China. Mitochondrion, 2019, 46, 327-333.	1.6	6
24	Validation and Safety of Visual Restoration by Ectopic Expression of Human Melanopsin in Retinal Ganglion Cells. Human Gene Therapy, 2019, 30, 714-726.	1.4	4
25	Hookworm in the eye. Lancet Infectious Diseases, The, 2018, 18, 582.	4.6	1
26	ISCEV guide to visual electrodiagnostic procedures. Documenta Ophthalmologica, 2018, 136, 1-26.	1.0	248
27	Toxocariasis of the eye. IDCases, 2018, 12, e3.	0.4	0
28	Domains of health-related quality of life in age-related macular degeneration: a qualitative study in the Chinese cultural context. BMJ Open, 2018, 8, e018756.	0.8	10
29	Cases of visual impairment caused by cerebral venous sinus occlusion-induced intracranial hypertension in the absence of headache. BMC Neurology, 2018, 18, 159.	0.8	7
30	Treatment of Punctate Inner Choroidopathy with Choroidal Neovascularization Using Corticosteroid and Intravitreal Ranibizumab. BioMed Research International, 2018, 2018, 1-7.	0.9	11
31	Efficacy and Safety of Autologous Bone Marrow Mesenchymal Stem Cell Transplantation in Patients with Diabetic Retinopathy. Cellular Physiology and Biochemistry, 2018, 49, 40-52.	1.1	50
32	Improved retinal function in RCS rats after suppressing the over-activation of mGluR5. Scientific Reports, 2017, 7, 3546.	1.6	17
33	PRPF3-Associated Autosomal Dominant Retinitis Pigmentosa and CYP4V2-Associated Biettiâ€™s Crystalline Corneoretinal Dystrophy Coexist in a Multigenerational Chinese Family. Journal of Ophthalmology, 2017, 2017, 1-10.	0.6	1
34	Contribution of GABAa, GABAc and glycine receptors to rat dark-adapted oscillatory potentials in the time and frequency domain. Oncotarget, 2017, 8, 77696-77709.	0.8	12
35	Usher Syndrome in Chinese and Japanese Population. Essentials in Ophthalmology, 2017, , 437-447.	0.0	0
36	Detecting novel genetic mutations in Chinese Usher syndrome families using next-generation sequencing technology. Molecular Genetics and Genomics, 2015, 290, 353-363.	1.0	10

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37	Systemic Administration of Erythropoietin Inhibits Retinopathy in RCS Rats. PLoS ONE, 2014, 9, e104759.	1.1	21
38	A novel mutation in C5L2 gene was associated with hyperlipidemia and retinitis pigmentosa in a Chinese family. Lipids in Health and Disease, 2014, 13, 75.	1.2	3
39	Tyrosine phosphorylation of VE-cadherin and claudin-5 is associated with TGF-Î²1-induced permeability of centrally derived vascular endothelium. European Journal of Cell Biology, 2011, 90, 323-332.	1.6	82
40	Retinal vascular changes after glial disruption in rats. Journal of Neuroscience Research, 2010, 88, 1485-1499.	1.3	52