

Heather Mack

List of Publications by Year in descending order

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Version: 2024-02-01

53
papers

620
citations

932766

10
h-index

642321

23
g-index

53
all docs

53
docs citations

53
times ranked

839
citing authors

#	ARTICLE	IF	CITATIONS
1	Ocular Features in Alport Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2015, 10, 703-709.	2.2	147
2	Safety, tolerability, and efficacy of PBT2 in Huntington's disease: a phase 2, randomised, double-blind, placebo-controlled trial. Lancet Neurology, The, 2015, 14, 39-47.	4.9	112
3	Behavioral measures of cortical hyperexcitability assessed in people who experience visual snow. Neurology, 2017, 88, 1243-1249.	1.5	47
4	Isolated Metastases to the Retina or Optic Nerve. International Ophthalmology Clinics, 1997, 37, 251-260.	0.3	45
5	Retinal disease in the C3 glomerulopathies and the risk of impaired vision. Ophthalmic Genetics, 2016, 37, 369-376.	0.5	27
6	Orbital compartment syndrome following aneurysm surgery. Journal of Clinical Neuroscience, 2012, 19, 1032-1036.	0.8	16
7	Microlearning to improve <sc>CPD</sc> learning objectives. Clinical Teacher, 2020, 17, 695-699.	0.4	16
8	Becoming an expert: a review of adult learning theory and implications for vocational training in ophthalmology. Clinical and Experimental Ophthalmology, 2012, 40, 519-526.	1.3	13
9	Bull's eye and pigment maculopathy are further retinal manifestations of an abnormal Bruch's membrane in Alport syndrome. Ophthalmic Genetics, 2017, 38, 238-244.	0.5	12
10	Victorian evolution of inherited retinal diseases natural history registry (<sc>VENTURE</sc>) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 38 Ophthalmology, 2022, 50, 768-780.	1.3	12
11	Bull's eye maculopathy and subfoveal deposition in two mucopolysaccharidosis type I patients on long-term enzyme replacement therapy. American Journal of Ophthalmology Case Reports, 2018, 9, 1-6.	0.4	11
12	CPD? What happened to CME? CME and beyond. Medical Teacher, 2018, 40, 914-916.	1.0	10
13	A search for a mammalian homologue of the Drosophila photoreceptor development gene glass yields Zfp64, a zinc finger encoding gene which maps to the distal end of mouse chromosome 2. Gene, 1997, 185, 11-17.	1.0	9
14	Corneal endothelial cell abnormalities in X-linked Alport syndrome. Ophthalmic Genetics, 2020, 41, 13-19.	0.5	8
15	Genetic susceptibility to hydroxychloroquine retinal toxicity. Ophthalmic Genetics, 2020, 41, 159-170.	0.5	8
16	Perspectives of people with inherited retinal diseases on ocular gene therapy in Australia: protocol for a national survey. BMJ Open, 2021, 11, e048361.	0.8	8
17	Ocular complications of tumour necrosis factor alpha inhibitors. Australasian journal of optometry, The, 2020, 103, 148-154.	0.6	7
18	Alport Syndrome With Kidney Cysts Is Still Alport Syndrome. Kidney International Reports, 2022, 7, 339-342.	0.4	7

#	ARTICLE	IF	CITATIONS
19	Genetic Testing of Inherited Retinal Disease in Australian Private Tertiary Ophthalmology Practice. <i>Clinical Ophthalmology</i> , 2022, Volume 16, 1127-1138.	0.9	7
20	The Bartter-Gitelman Spectrum: 50-Year Follow-up With Revision of Diagnosis After Whole-Genome Sequencing. <i>Journal of the Endocrine Society</i> , 2022, 6, .	0.1	7
21	Bilateral Retrobulbar Optic Neuropathy in the Setting of Interferon Alpha-2a Therapy. <i>Case Reports in Ophthalmology</i> , 2014, 5, 270-276.	0.3	6
22	Transient bilateral optic disc oedema in mitochondrial encephalomyopathy, lactic acidosis, and stroke-like episodes (MELAS). <i>Canadian Journal of Ophthalmology</i> , 2018, 53, e208-e211.	0.4	6
23	Gitelman syndrome and ectopic calcification in the retina and joints. <i>CKJ: Clinical Kidney Journal</i> , 2021, 14, 2023-2028.	1.4	6
24	COVID-19 in ophthalmology: Implications for ophthalmologists, eye care, ophthalmic education and research. <i>Clinical and Experimental Ophthalmology</i> , 2021, 49, 9-11.	1.3	6
25	Preservation of residual field after surgical lowering of intraocular pressure. <i>Australian and New Zealand Journal of Ophthalmology</i> , 1992, 20, 343-345.	0.4	5
26	Hydroxychloroquine retinal toxicity in two patients with dermatological conditions. <i>Australasian Journal of Dermatology</i> , 2018, 59, e266-e268.	0.4	5
27	Retinal Neuronal Loss in Visually Asymptomatic Patients With Myoclonic Epilepsy With Ragged-Red Fibers. <i>Journal of Neuro-Ophthalmology</i> , 2019, 39, 18-22.	0.4	5
28	Acute Anterior Uveitis in a Patient Taking Fingolimod (FTY720) for Multiple Sclerosis. <i>Case Reports in Ophthalmology</i> , 2017, 7, 562-566.	0.3	5
29	Measurement Properties of the Attitudes to Gene Therapy for the Eye (AGT-Eye) Instrument for People With Inherited Retinal Diseases. <i>Translational Vision Science and Technology</i> , 2022, 11, 14.	1.1	5
30	Retinal findings in glomerulonephritis. <i>Australasian journal of optometry</i> , The, 2021, , 1-13.	0.6	5
31	Cystoid macular edema in a patient with Danon disease. <i>Indian Journal of Ophthalmology</i> , 2014, 62, 1161.	0.5	3
32	Ocular involvement in a patient with <i>Legionella longbeachae</i> 1 infection. <i>Clinical and Experimental Ophthalmology</i> , 2014, 42, 497-499.	1.3	3
33	Developing educators for continuing professional development. <i>Canadian Journal of Ophthalmology</i> , 2016, 51, 196-200.	0.4	3
34	Chronic Kidney Disease and Cataract: Seeing the Light. <i>American Journal of Nephrology</i> , 2017, 45, 522-523.	1.4	3
35	The case for adopting sustainability goals in ophthalmology. <i>Clinical and Experimental Ophthalmology</i> , 2019, 47, 837-839.	1.3	3
36	Bilateral Syphilitic Optic Neuropathy with Secondary Autoimmune Optic Neuropathy and Poor Visual Outcome. <i>Case Reports in Ophthalmology</i> , 2019, 10, 81-88.	0.3	3

#	ARTICLE	IF	CITATIONS
37	Cystoid Macular Edema. , 2022, , 693-731.		3
38	Retinal Drusen Are More Common and Larger in Systemic Lupus Erythematosus With Renal Impairment. <i>Kidney International Reports</i> , 2022, 7, 848-856.	0.4	3
39	Free Continuing Professional Development Resources for Low-Resource Settings. <i>Annals of Global Health</i> , 2018, 81, 731.	0.8	2
40	How to Add Metacognition to Your Continuing Professional Development: Scoping Review and Recommendations. <i>Asia-Pacific Journal of Ophthalmology</i> , 2019, 8, 256-263.	1.3	2
41	HaNDL with bilateral central venous occlusions. <i>BMJ Neurology Open</i> , 2020, 2, e000043.	0.7	2
42	Partnering to develop a continuing professional development program in a low-resource setting: Cambodia. <i>Canadian Journal of Ophthalmology</i> , 2017, 52, 379-384.	0.4	1
43	Clozapine-induced maculopathy. <i>Medical Journal of Australia</i> , 2017, 207, 316-316.	0.8	1
44	Paracentral acute middle maculopathy associated with Eisenmenger syndrome. <i>Clinical and Experimental Ophthalmology</i> , 2020, 48, 1106-1108.	1.3	1
45	Clinical teaching of CPD during the COVID pandemic. <i>Clinical Teacher</i> , 2021, 18, 84-86.	0.4	1
46	Expanding the phenotype of mucopolysaccharidosis type II retinopathy. <i>Ophthalmic Genetics</i> , 2021, 42, 631-636.	0.5	1
47	Late progression of visual loss from ocular quinine toxicity. <i>Canadian Journal of Ophthalmology</i> , 2021, 56, e116-e119.	0.4	1
48	Management of conjunctivitis and other causes of red eye during the COVID-19 pandemic. <i>Australian Journal of General Practice</i> , 2020, 49, 656-661.	0.3	1
49	Governing the reform of the medical internship. <i>Medical Journal of Australia</i> , 2016, 205, 380-380.	0.8	0
50	Functional screening devices for diabetic retinopathy. <i>Clinical and Experimental Ophthalmology</i> , 2018, 46, 573-575.	1.3	0
51	Technology driving advances in ophthalmology. <i>Australian Journal of General Practice</i> , 2019, 48, 501.	0.3	0
52	Taking a broader view of the health care needs of people with chronic kidney disease. <i>Medical Journal of Australia</i> , 2022, , .	0.8	0
53	Novel chorioretinal findings in two siblings with mucopolysaccharidosis type VI. <i>Ophthalmic Genetics</i> , 0, , 1-6.	0.5	0