Abigail T Fahim

List of Publications by Year
in descending order

31	1,045	516710 16	477307 29
papers	citations	h-index	g-index

31	31	31	1360
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Real-world outcomes of voretigene neparvovec treatment in pediatric patients with RPE65-associated Leber congenital amaurosis. Graefe's Archive for Clinical and Experimental Ophthalmology, 2022, 260, 1543-1550.	1.9	27
2	The Michigan Vision-Related Anxiety Questionnaire: A Psychosocial Outcomes Measure for Inherited Retinal Degenerations. American Journal of Ophthalmology, 2021, 225, 137-146.	3.3	13
3	The Michigan Retinal Degeneration Questionnaire: A Patient-Reported Outcome Instrument for Inherited Retinal Degenerations. American Journal of Ophthalmology, 2021, 222, 60-68.	3.3	28
4	Coats-like Exudative Vitreoretinopathy in Retinitis Pigmentosa. Ophthalmology Retina, 2021, 5, 86-96.	2.4	12
5	Association of No-Cost Genetic Testing Program Implementation and Patient Characteristics With Access to Genetic Testing for Inherited Retinal Degenerations. JAMA Ophthalmology, 2021, 139, 449.	2.5	6
6	Characterization of the Spectrum of Ophthalmic Changes in Patients With Alagille Syndrome. , 2021, 62, 27.		11
7	X-Chromosome Inactivation Is a Biomarker of Clinical Severity in Female Carriers of RPGR-Associated X-Linked Retinitis Pigmentosa. Ophthalmology Retina, 2020, 4, 510-520.	2.4	31
8	Rapid visual field constriction in a patient with retinitis pigmentosa and pituitary adenoma. American Journal of Ophthalmology Case Reports, 2020, 19, 100762.	0.7	0
9	Advancing Clinical Trials for Inherited Retinal Diseases: Recommendations from the Second Monaciano Symposium. Translational Vision Science and Technology, 2020, 9, 2.	2.2	56
10	Genetic testing for inherited retinal degenerations: Triumphs and tribulations. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2020, 184, 571-577.	1.6	10
11	Cystoid macular edema precipitated by altitude in a patient with X-linked retinitis pigmentosa. Ophthalmic Genetics, 2020, 41, 275-278.	1.2	1
12	Development of a Gene Therapy Vector for RDH12-Associated Retinal Dystrophy. Human Gene Therapy, 2019, 30, 1325-1335.	2.7	19
13	Detailed clinical characterisation, unique features and natural history of autosomal recessive RDH12-associated retinal degeneration. British Journal of Ophthalmology, 2019, 103, bjophthalmol-2018-313580.	3.9	20
14	Natural History and Genotype-Phenotype Correlations in RDH12-Associated Retinal Degeneration. Advances in Experimental Medicine and Biology, 2019, 1185, 209-213.	1.6	7
15	Double hyperautofluorescent ring on fundus autofluorescence in ABCA4. Ophthalmic Genetics, 2018, 39, 87-91.	1.2	2
16	Bullous X linked retinoschisis: clinical features and prognosis. British Journal of Ophthalmology, 2018, 102, 622-624.	3.9	18
17	Retinitis pigmentosa: recent advances and future directions in diagnosis and management. Current Opinion in Pediatrics, 2018, 30, 725-733.	2.0	58
18	Contrast sensitivity deficits in patients with mutation-proven inherited retinal degenerations. BMC Ophthalmology, 2018, 18, 313.	1.4	24

#	Article	IF	Citations
19	Peripheral fundus findings in X-linked retinoschisis. British Journal of Ophthalmology, 2017, 101, 1555-1559.	3.9	40
20	Peripheral Visual Fields in ABCA4 Stargardt Disease and Correlation With Disease Extent on Ultra-widefield Fundus Autofluorescence. American Journal of Ophthalmology, 2017, 184, 181-188.	3.3	12
21	Comparison of ophthalmic training in 6 English-speaking countries. Canadian Journal of Ophthalmology, 2016, 51, 212-218.	0.7	9
22	The Role of X-Chromosome Inactivation in Retinal Development and Disease. Advances in Experimental Medicine and Biology, 2016, 854, 325-331.	1.6	29
23	Acute Ocriplasmin Retinopathy. Retina, 2015, 35, 1055-1058.	1.7	40
24	Understanding the Adverse Effects of Ocriplasminâ€"Reply. JAMA Ophthalmology, 2015, 133, 230.	2.5	2
25	Lateral Geniculate Lesions Causing Reversible Blindness in a Pre-eclamptic Patient With a Variant of Posterior Reversible Encephalopathy Syndrome. Journal of Neuro-Ophthalmology, 2014, 34, 372-376.	0.8	4
26	Acute Panretinal Structural and Functional Abnormalities After Intravitreous Ocriplasmin Injection. JAMA Ophthalmology, 2014, 132, 484.	2.5	92
27	Diagnostic Fundus Autofluorescence Patterns in Achromatopsia. American Journal of Ophthalmology, 2013, 156, 1211-1219.e2.	3.3	38
28	Polymorphic Variation of RPGRIP1L and IQCB1 as Modifiers of X-Linked Retinitis Pigmentosa Caused by Mutations in RPGR. Advances in Experimental Medicine and Biology, 2012, 723, 313-320.	1.6	16
29	Allelic Heterogeneity and Genetic Modifier Loci Contribute to Clinical Variation in Males with X-Linked Retinitis Pigmentosa Due to RPGR Mutations. PLoS ONE, 2011, 6, e23021.	2.5	83
30	Transgenic overexpression of a stable Plasminogen Activator Inhibitor-1 variant. Thrombosis Research, 2009, 123, 785-792.	1.7	16
31	A dual-Ca2+-sensor model for neurotransmitter release in a central synapse. Nature, 2007, 450, 676-682.	27.8	321