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	A dual-Ca2+-sensor model for neurotransmitter release in a central synapse. Nature, 2007, 450, 676-682.	27.8	321
2	Acute Panretinal Structural and Functional Abnormalities After Intravitreous Ocriplasmin Injection. JAMA Ophthalmology, 2014, 132, 484.	2.5	92
3	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
4	Retinitis pigmentosa: recent advances and future directions in diagnosis and management. Current Opinion in Pediatrics, 2018, 30, 725-733.	2.0	58
5	Advancing Clinical Trials for Inherited Retinal Diseases: Recommendations from the Second Monaciano Symposium. Translational Vision Science and Technology, 2020, 9, 2.	2.2	56
6	Acute Ocriplasmin Retinopathy. Retina, 2015, 35, 1055-1058.	1.7	40
7	Peripheral fundus findings in X-linked retinoschisis. British Journal of Ophthalmology, 2017, 101, 1555-1559.	3.9	40
8	Diagnostic Fundus Autofluorescence Patterns in Achromatopsia. American Journal of Ophthalmology, 2013, 156, 1211-1219.e2.	3.3	38
9	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
10	The Role of X-Chromosome Inactivation in Retinal Development and Disease. Advances in Experimental Medicine and Biology, 2016, 854, 325-331.	1.6	29
11	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
12	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
13	Contrast sensitivity deficits in patients with mutation-proven inherited retinal degenerations. BMC Ophthalmology, 2018, 18, 313.	1.4	24
14	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
15	Development of a Gene Therapy Vector for RDH12-Associated Retinal Dystrophy. Human Gene Therapy, 2019, 30, 1325-1335.	2.7	19
16	Bullous X linked retinoschisis: clinical features and prognosis. British Journal of Ophthalmology, 2018, 102, 622-624.	3.9	18
17	Transgenic overexpression of a stable Plasminogen Activator Inhibitor-1 variant. Thrombosis Research, 2009, 123, 785-792.	1.7	16
18	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

#	Article	IF	Citations
19	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
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	Coats-like Exudative Vitreoretinopathy in Retinitis Pigmentosa. Ophthalmology Retina, 2021, 5, 86-96.	2.4	12
22	Characterization of the Spectrum of Ophthalmic Changes in Patients With Alagille Syndrome. , 2021, 62, 27.		11
23	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
24	Comparison of ophthalmic training in 6 English-speaking countries. Canadian Journal of Ophthalmology, 2016, 51, 212-218.	0.7	9
25	Natural History and Genotype-Phenotype Correlations in RDH12-Associated Retinal Degeneration. Advances in Experimental Medicine and Biology, 2019, 1185, 209-213.	1.6	7
26	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
27	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
28	Understanding the Adverse Effects of Ocriplasminâ€"Reply. JAMA Ophthalmology, 2015, 133, 230.	2.5	2
29	Double hyperautofluorescent ring on fundus autofluorescence in ABCA4. Ophthalmic Genetics, 2018, 39, 87-91.	1.2	2
30	Cystoid macular edema precipitated by altitude in a patient with X-linked retinitis pigmentosa. Ophthalmic Genetics, 2020, 41, 275-278.	1.2	1
31	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