Naheed W Khan

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5401637/publications.pdf

Version: 2024-02-01

19	466	933447	940533
papers	citations	10 h-index	16 g-index

19	19	19	744
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	Adherence and satisfaction in Argus II prosthesis users: a self determination theory model. Ophthalmic Genetics, 2022, 43, 462-469.	1.2	0
3	Calculation of test-retest variability in phase I/IIa clinical trials for Inherited Retinal Degenerations. Ophthalmic Genetics, 2021, 42, 283-290.	1.2	2
4	Deciphering the genetic architecture and ethnographic distribution of IRD in three ethnic populations by whole genome sequence analysis. PLoS Genetics, 2021, 17, e1009848.	3.5	13
5	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
6	The Deubiquitinating Enzyme Ataxin-3 Regulates Ciliogenesis and Phagocytosis in the Retina. Cell Reports, 2020, 33, 108360.	6.4	23
7	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
8	Comparison of Fundus-Guided Microperimetry and Multifocal Electroretinography for Evaluating Hydroxychloroquine Maculopathy. Translational Vision Science and Technology, 2019, 8, 19.	2.2	5
9	Macular hyperpigmentary changes in ABCA4-Stargardt disease. International Journal of Retina and Vitreous, 2019, 5, 9.	1.9	3
10	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
11	Contrast sensitivity deficits in patients with mutation-proven inherited retinal degenerations. BMC Ophthalmology, 2018, 18, 313.	1.4	24
12	Inhibiting autophagy reduces retinal degeneration caused by protein misfolding. Autophagy, 2018, 14, 1226-1238.	9.1	81
13	ISCEV extended protocol for the photopic Onâ€“Off ERG. Documenta Ophthalmologica, 2018, 136, 199-206.	2.2	44
14	IFT88 mutations identified in individuals with non-syndromic recessive retinal degeneration result in abnormal ciliogenesis. Human Genetics, 2018, 137, 447-458.	3.8	11
15	A specific phosphorylation regulates the protective role of Î±A-crystallin in diabetes. JCI Insight, 2018, 3, .	5.0	30
16	Inherited Retinal Degeneration: Genetics, Disease Characterization, and Outcome Measures. Journal of Ophthalmology, 2017, 2017, 1-2.	1.3	8
17	Autophagy-mediated catabolism of visual transduction proteins prevents retinal degeneration. Autophagy, 2016, 12, 2439-2450.	9.1	37
18	Loss of Raf-1 Kinase Inhibitory Protein Delays Early-Onset Severe Retinal Ciliopathy inCep290rd16Mouse. , 2014, 55, 5788.		15

#	Article	IF	Citations
19	Acute Panretinal Structural and Functional Abnormalities After Intravitreous Ocriplasmin Injection. JAMA Ophthalmology, 2014, 132, 484.	2.5	92