

# 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  
papers

466  
citations

933447

10  
h-index

940533

16  
g-index

19  
all docs

19  
docs citations

19  
times ranked

744  
citing authors

#	ARTICLE	IF	CITATIONS
1	Acute Panretinal Structural and Functional Abnormalities After Intravitreal Ocriplasmin Injection. <i>JAMA Ophthalmology</i> , 2014, 132, 484.	2.5	92
2	Inhibiting autophagy reduces retinal degeneration caused by protein misfolding. <i>Autophagy</i> , 2018, 14, 1226-1238.	9.1	81
3	ISCEV extended protocol for the photopic Onâ€“Off ERG. <i>Documenta Ophthalmologica</i> , 2018, 136, 199-206.	2.2	44
4	Autophagy-mediated catabolism of visual transduction proteins prevents retinal degeneration. <i>Autophagy</i> , 2016, 12, 2439-2450.	9.1	37
5	X-Chromosome Inactivation Is a Biomarker of Clinical Severity in Female Carriers of RPGR-Associated X-Linked Retinitis Pigmentosa. <i>Ophthalmology Retina</i> , 2020, 4, 510-520.	2.4	31
6	A specific phosphorylation regulates the protective role of Î±A-crystallin in diabetes. <i>JCI Insight</i> , 2018, 3, .	5.0	30
7	Real-world outcomes of voretigene neparvovec treatment in pediatric patients with RPE65-associated Leber congenital amaurosis. <i>Graefe's Archive for Clinical and Experimental Ophthalmology</i> , 2022, 260, 1543-1550.	1.9	27
8	Contrast sensitivity deficits in patients with mutation-proven inherited retinal degenerations. <i>BMC Ophthalmology</i> , 2018, 18, 313.	1.4	24
9	The Deubiquitinating Enzyme Ataxin-3 Regulates Ciliogenesis and Phagocytosis in the Retina. <i>Cell Reports</i> , 2020, 33, 108360.	6.4	23
10	Detailed clinical characterisation, unique features and natural history of autosomal recessive <i>RDH12</i> -associated retinal degeneration. <i>British Journal of Ophthalmology</i> , 2019, 103, bjophthalmol-2018-313580.	3.9	20
11	Loss of Raf-1 Kinase Inhibitory Protein Delays Early-Onset Severe Retinal Ciliopathy in <i>Cep290</i> <i>rd16</i> Mouse. , 2014, 55, 5788.		15
12	Deciphering the genetic architecture and ethnographic distribution of IRD in three ethnic populations by whole genome sequence analysis. <i>PLoS Genetics</i> , 2021, 17, e1009848.	3.5	13
13	IFT88 mutations identified in individuals with non-syndromic recessive retinal degeneration result in abnormal ciliogenesis. <i>Human Genetics</i> , 2018, 137, 447-458.	3.8	11
14	Inherited Retinal Degeneration: Genetics, Disease Characterization, and Outcome Measures. <i>Journal of Ophthalmology</i> , 2017, 2017, 1-2.	1.3	8
15	Comparison of Fundus-Guided Microperimetry and Multifocal Electroretinography for Evaluating Hydroxychloroquine Maculopathy. <i>Translational Vision Science and Technology</i> , 2019, 8, 19.	2.2	5
16	Macular hyperpigmentary changes in <i>ABCA4</i> -Stargardt disease. <i>International Journal of Retina and Vitreous</i> , 2019, 5, 9.	1.9	3
17	Calculation of test-retest variability in phase I/IIa clinical trials for Inherited Retinal Degenerations. <i>Ophthalmic Genetics</i> , 2021, 42, 283-290.	1.2	2
18	Rapid visual field constriction in a patient with retinitis pigmentosa and pituitary adenoma. <i>American Journal of Ophthalmology Case Reports</i> , 2020, 19, 100762.	0.7	0

#	ARTICLE	IF	CITATIONS
19	Adherence and satisfaction in Argus II prosthesis users: a self determination theory model. Ophthalmic Genetics, 2022, 43, 462-469.	1.2	0