

# Robert Henderson

## List of Publications by Year in descending order

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33  
papers

2,990  
citations

566801

15  
h-index

500791

28  
g-index

36  
all docs

36  
docs citations

36  
times ranked

3891  
citing authors

#	ARTICLE	IF	CITATIONS
1	Effect of Gene Therapy on Visual Function in Leber's Congenital Amaurosis. <i>New England Journal of Medicine</i> , 2008, 358, 2231-2239.	13.9	1,793
2	Comprehensive Rare Variant Analysis via Whole-Genome Sequencing to Determine the Molecular Pathology of Inherited Retinal Disease. <i>American Journal of Human Genetics</i> , 2017, 100, 75-90.	2.6	343
3	Phenotypic variability in patients with retinal dystrophies due to mutations in CRB1. <i>British Journal of Ophthalmology</i> , 2011, 95, 811-817.	2.1	95
4	The Oculome Panel Test. <i>Ophthalmology</i> , 2019, 126, 888-907.	2.5	77
5	Novel mutations in MERTK associated with childhood onset rod-cone dystrophy. <i>Molecular Vision</i> , 2010, 16, 369-77.	1.1	73
6	A rare de novo nonsense mutation in OTX2 causes early onset retinal dystrophy and pituitary dysfunction. <i>Molecular Vision</i> , 2009, 15, 2442-7.	1.1	68
7	A Homozygous Mutation in the <i>TUB</i> Gene Associated with Retinal Dystrophy and Obesity. <i>Human Mutation</i> , 2014, 35, 289-293.	1.1	63
8	An Assessment of the Apex Microarray Technology in Genotyping Patients with Leber Congenital Amaurosis and Early-Onset Severe Retinal Dystrophy. , 2007, 48, 5684.		56
9	Biallelic mutation of protocadherin-21 (PCDH21) causes retinal degeneration in humans. <i>Molecular Vision</i> , 2010, 16, 46-52.	1.1	50
10	Wound-related complications and clinical outcomes following open globe injury repair. <i>Clinical and Experimental Ophthalmology</i> , 2015, 43, 508-513.	1.3	48
11	RDH12 retinopathy: novel mutations and phenotypic description. <i>Molecular Vision</i> , 2011, 17, 2706-16.	1.1	47
12	Early Onset Retinal Dystrophy Due to Mutations in <i>LRAT</i> : Molecular Analysis and Detailed Phenotypic Study. , 2012, 53, 3927.		38
13	Screening of <i>SPATA7</i> in Patients with Leber Congenital Amaurosis and Severe Childhood-Onset Retinal Dystrophy Reveals Disease-Causing Mutations. , 2011, 52, 3032.		34
14	Immediate Sequential Bilateral Pediatric Vitreoretinal Surgery. <i>Ophthalmology</i> , 2016, 123, 1802-1808.	2.5	32
15	Leber Congenital Amaurosis Associated with AIPL1: Challenges in Ascribing Disease Causation, Clinical Findings, and Implications for Gene Therapy. <i>PLoS ONE</i> , 2012, 7, e32330.	1.1	28
16	Biallelic Mutation of ARHGEF18, Involved in the Determination of Epithelial Apicobasal Polarity, Causes Adult-Onset Retinal Degeneration. <i>American Journal of Human Genetics</i> , 2017, 100, 334-342.	2.6	26
17	Clinical Characterization of <i>CNGB1</i> -Related Autosomal Recessive Retinitis Pigmentosa. <i>JAMA Ophthalmology</i> , 2017, 135, 137.	1.4	23
18	Missense variants in the X-linked gene <i>PRPS1</i> cause retinal degeneration in females. <i>Human Mutation</i> , 2018, 39, 80-91.	1.1	23

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19	Recommended protocol for performing oral fundus fluorescein angiography (FFA) in children. <i>Eye</i> , 2022, 36, 234-236.	1.1	14
20	SRD5A3-CDG: Emerging Phenotypic Features of an Ultrarare CDG Subtype. <i>Frontiers in Genetics</i> , 2021, 12, 737094.	1.1	9
21	Expanding the phenotypic spectrum consequent upon de novo <scp><i>WDR37</i></scp> missense variants. <i>Clinical Genetics</i> , 2020, 98, 191-197.	1.0	8
22	Accuracy of scleral transillumination techniques to identify infant ciliary body for sclerostomy and intravitreal injections. <i>Clinical and Experimental Ophthalmology</i> , 2019, 47, 478-483.	1.3	7
23	New variants and in silico analyses in GRK1 associated Oguchi disease. <i>Human Mutation</i> , 2021, 42, 164-176.	1.1	7
24	An ERG and OCT study of neuronal ceroid lipofuscinosis CLN2 Battens retinopathy. <i>Eye</i> , 2021, 35, 2438-2448.	1.1	6
25	Retinal vasculopathy in STING-associated vasculitis of infancy (SAVI). <i>Rheumatology</i> , 2021, 60, e351-e353.	0.9	5
26	A Novel Technique for High-Density Silicone Oil Removal. <i>Retina</i> , 2012, 32, 1672-1673.	1.0	4
27	Group A streptococcal endophthalmitis complicating a sore throat in a 2-year-old child. <i>BMJ Case Reports</i> , 2015, 2015, bcr2014208168-bcr2014208168.	0.2	4
28	Impact of sight and hearing loss in patients with Norrie disease: advantages of Dual Sensory clinics in patient care. <i>BMJ Paediatrics Open</i> , 2020, 4, e000781.	0.6	4
29	Misaligned foveal morphology and sector retinal dysfunction in AKT1-mosaic Proteus syndrome. <i>Documenta Ophthalmologica</i> , 2021, 142, 119-126.	1.0	1
30	Evolving outcomes of surgery for retinal detachment in retinopathy of prematurity: the need for a national service in the United Kingdom. <i>Eye</i> , 2021, , .	1.1	1
31	Diffuse bear-track retina: profound, bilateral, grouped congenital pigmentation of the retinal pigment epithelium in an infant. <i>Journal of AAPOS</i> , 2020, 24, 384-386.	0.2	1
32	64â€¦Walker Warburg syndrome (WWS) with ISPD genetic mutation- a case report. , 2020, , .		1
33	HISTOPATHOLOGICALLY PROVEN MUCINOUS CYSTADENOCARCINOMA METASTATIC TO THE CHOROID. <i>Retinal Cases and Brief Reports</i> , 2010, 4, 181-183.	0.3	0