Robert Henderson

List of Publications by Year
in descending order

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	Recommended protocol for performing oral fundus fluorescein angiography (FFA) in children. Eye, 2022, 36, 234-236.	1.1	14
2	Misaligned foveal morphology and sector retinal dysfunction in AKT1-mosaic Proteus syndrome. Documenta Ophthalmologica, 2021, 142, 119-126.	1.0	1
3	New variants and in silico analyses in GRK1 associated Oguchi disease. Human Mutation, 2021, 42, 164-176.	1.1	7
4	Retinal vasculopathy in STING-associated vasculitis of infancy (SAVI). Rheumatology, 2021, 60, e351-e353.	0.9	5
5	Evolving outcomes of surgery for retinal detachment in retinopathy of prematurity: the need for a national service in the United Kingdom. Eye, 2021, , .	1.1	1
6	An ERG and OCT study of neuronal ceroid lipofuscinosis CLN2 Battens retinopathy. Eye, 2021, 35, 2438-2448.	1.1	6
7	SRD5A3-CDG: Emerging Phenotypic Features of an Ultrarare CDG Subtype. Frontiers in Genetics, 2021, 12, 737094.	1.1	9
8	Expanding the phenotypic spectrum consequent upon de novo <scp>WDR37</scp> missense variants. Clinical Genetics, 2020, 98, 191-197.	1.0	8
9	Diffuse bear-track retina: profound, bilateral, grouped congenital pigmentation of the retinal pigment epithelium in an infant. Journal of AAPOS, 2020, 24, 384-386.	0.2	1
10	Impact of sight and hearing loss in patients with Norrie disease: advantages of Dual Sensory clinics in patient care. BMJ Paediatrics Open, 2020, 4, e000781.	0.6	4
11	64â€...Walker Warburg syndrome (WWS) with ISPD genetic mutation- a case report. , 2020, , .		1
12	The Oculome Panel Test. Ophthalmology, 2019, 126, 888-907.	2.5	77
13	Accuracy of scleral transillumination techniques to identify infant ciliary body for sclerostomy and intravitreal injections. Clinical and Experimental Ophthalmology, 2019, 47, 478-483.	1.3	7
14	Missense variants in the X-linked genePRPS1cause retinal degeneration in females. Human Mutation, 2018, 39, 80-91.	1.1	23
15	Clinical Characterization of CNGB1-Related Autosomal Recessive Retinitis Pigmentosa. JAMA Ophthalmology, 2017, 135, 137.	1.4	23
16	Biallelic Mutation of ARHGEF18, Involved in the Determination of Epithelial Apicobasal Polarity, Causes Adult-Onset Retinal Degeneration. American Journal of Human Genetics, 2017, 100, 334-342.	2.6	26
17	Comprehensive Rare Variant Analysis via Whole-Genome Sequencing to Determine the Molecular Pathology of Inherited Retinal Disease. American Journal of Human Genetics, 2017, 100, 75-90.	2.6	343
18	Immediate Sequential Bilateral Pediatric Vitreoretinal Surgery. Ophthalmology, 2016, 123, 1802-1808.	2.5	32

#	Article	IF	Citations
19	Woundâ€related complications and clinical outcomes following open globe injury repair. Clinical and Experimental Ophthalmology, 2015, 43, 508-513.	1.3	48
20	Group A streptococcal endophthalmitis complicating a sore throat in a 2-year-old child. BMJ Case Reports, 2015, 2015, bcr2014208168-bcr2014208168.	0.2	4
21	A Homozygous Mutation in the<scp>TUB</scp>Gene Associated with Retinal Dystrophy and Obesity. Human Mutation, 2014, 35, 289-293.	1.1	63
22	A Novel Technique for High-Density Silicone Oil Removal. Retina, 2012, 32, 1672-1673.	1.0	4
23	Early Onset Retinal Dystrophy Due to Mutations inLRAT: Molecular Analysis and Detailed Phenotypic Study. , 2012, 53, 3927.		38
24	Leber Congenital Amaurosis Associated with AIPL1: Challenges in Ascribing Disease Causation, Clinical Findings, and Implications for Gene Therapy. PLoS ONE, 2012, 7, e32330.	1.1	28
25	Screening ofSPATA7in Patients with Leber Congenital Amaurosis and Severe Childhood-Onset Retinal Dystrophy Reveals Disease-Causing Mutations. , 2011, 52, 3032.		34
26	Phenotypic variability in patients with retinal dystrophies due to mutations in CRB1. British Journal of Ophthalmology, 2011, 95, 811-817.	2.1	95
27	RDH12 retinopathy: novel mutations and phenotypic description. Molecular Vision, 2011, 17, 2706-16.	1.1	47
28	HISTOPATHOLOGICALLY PROVEN MUCINOUS CYSTADENOCARCINOMA METASTATIC TO THE CHOROID. Retinal Cases and Brief Reports, 2010, 4, 181-183.	0.3	0
29	Biallelic mutation of protocadherin-21 (PCDH21) causes retinal degeneration in humans. Molecular Vision, 2010, 16, 46-52.	1.1	50
30	Novel mutations in MERTK associated with childhood onset rod-cone dystrophy. Molecular Vision, 2010, 16, 369-77.	1.1	73
31	A rare de novo nonsense mutation in OTX2 causes early onset retinal dystrophy and pituitary dysfunction. Molecular Vision, 2009, 15, 2442-7.	1.1	68
32	Effect of Gene Therapy on Visual Function in Leber's Congenital Amaurosis. New England Journal of Medicine, 2008, 358, 2231-2239.	13.9	1,793
33	An Assessment of the Apex Microarray Technology in Genotyping Patients with Leber Congenital Amaurosis and Early-Onset Severe Retinal Dystrophy. , 2007, 48, 5684.		56