Robert H Henderson

List of Publications by Year
in descending order

12	547	1040056 9	1372567 10
papers	citations	h-index	g-index

12	12	12	982
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Phenotypic variability in patients with retinal dystrophies due to mutations in CRB1. British Journal of Ophthalmology, 2011, 95, 811-817.	3.9	95
2	Novel mutations in MERTK associated with childhood onset rod-cone dystrophy. Molecular Vision, 2010, 16, 369-77.	1.1	73
3	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
4	An Assessment of the Apex Microarray Technology in Genotyping Patients with Leber Congenital Amaurosis and Early-Onset Severe Retinal Dystrophy. , 2007, 48, 5684.		56
5	Biallelic mutation of protocadherin-21 (PCDH21) causes retinal degeneration in humans. Molecular Vision, 2010, 16, 46-52.	1.1	50
6	Woundâ€related complications and clinical outcomes following open globe injury repair. Clinical and Experimental Ophthalmology, 2015, 43, 508-513.	2.6	48
7	RDH12 retinopathy: novel mutations and phenotypic description. Molecular Vision, 2011, 17, 2706-16.	1.1	47
8	Screening ofSPATA7in Patients with Leber Congenital Amaurosis and Severe Childhood-Onset Retinal Dystrophy Reveals Disease-Causing Mutations. , 2011, 52, 3032.		34
9	Immediate Sequential Bilateral Pediatric Vitreoretinal Surgery. Ophthalmology, 2016, 123, 1802-1808.	5.2	32
10	Leber Congenital Amaurosis Associated with AIPL1: Challenges in Ascribing Disease Causation, Clinical Findings, and Implications for Gene Therapy. PLoS ONE, 2012, 7, e32330.	2.5	28
11	Traumatic Retinal Detachment in Patients with Self-Injurious Behavior. Ophthalmology Retina, 2021, 5, 805-814.	2.4	9
12	New variants and in silico analyses in GRK1 associated Oguchi disease. Human Mutation, 2021, 42, 164-176.	2.5	7