

Robert H Henderson

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/11664521/publications.pdf>

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12
papers

547
citations

1040056

9
h-index

1372567

10
g-index

12
all docs

12
docs citations

12
times ranked

982
citing authors

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