

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

22
papers

171
citations

8
h-index

12
g-index

24
ext. papers

249
ext. citations

4.1
avg, IF

2.91
L-index

#	Paper	IF	Citations
22	Comprehensive Molecular Screening in Chinese Usher Syndrome Patients 2018 , 59, 1229-1237		31
21	Screening of BEST1 Gene in a Chinese Cohort With Best Vitelliform Macular Dystrophy or Autosomal Recessive Bestrophinopathy 2017 , 58, 3366-3375		22
20	Mutation screening of mitochondrial DNA as well as OPA1 and OPA3 in a Chinese cohort with suspected hereditary optic atrophy. <i>Investigative Ophthalmology and Visual Science</i> , 2014 , 55, 6987-95		16
19	Genome-Wide Association and Functional Studies Identify and as Novel Susceptibility Genes for Coronary Artery Disease. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2018 , 38, 964-975	9.4	15
18	Genetic and clinical findings in a Chinese cohort with Leber congenital amaurosis and early onset severe retinal dystrophy. <i>British Journal of Ophthalmology</i> , 2020 , 104, 932-937	5.5	15
17	Genetic and Functional Evidence Supports LPAR1 as a Susceptibility Gene for Hypertension. <i>Hypertension</i> , 2015 , 66, 641-6	8.5	13
16	Detection of serum anti-retinal antibodies in the Chinese patients with presumed autoimmune retinopathy. <i>Graefes Archive for Clinical and Experimental Ophthalmology</i> , 2019 , 257, 1759-1764	3.8	9
15	Sector Retinitis Pigmentosa caused by mutations of the RHO gene. <i>Eye</i> , 2019 , 33, 592-599	4.4	9
14	Novel mutations of the RS1 gene in a cohort of Chinese families with X-linked retinoschisis. <i>Molecular Vision</i> , 2014 , 20, 132-9	2.3	7
13	Clinical findings and genotype in 90 Chinese families with X-linked retinoschisis. <i>Molecular Vision</i> , 2020 , 26, 291-298	2.3	7
12	GUCY2D mutations in a Chinese cohort with autosomal dominant cone or cone-rod dystrophies. <i>Documenta Ophthalmologica</i> , 2015 , 131, 105-14	2.2	6
11	Comprehensive screening of in a cohort of Chinese patients with Bietti crystalline dystrophy. <i>Molecular Vision</i> , 2018 , 24, 700-711	2.3	6
10	Comparisons of the seizure-free outcome and visual field deficits between anterior temporal lobectomy and selective amygdalohippocampectomy: A systematic review and meta-analysis. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2020 , 81, 228-235	3.2	6
9	Mutation spectrum of and clinical findings in 95 Chinese patients with aniridia. <i>Molecular Vision</i> , 2020 , 26, 226-234	2.3	4
8	Variant Profiling of a Large Cohort of 138 Chinese Families With Autosomal Dominant Retinitis Pigmentosa. <i>Frontiers in Cell and Developmental Biology</i> , 2020 , 8, 629994	5.7	3
7	Generation of an induced pluripotent stem cell line BIOi002-A from a patient with autosomal dominant optic atrophy. <i>Stem Cell Research</i> , 2021 , 53, 102278	1.6	1
6	Prognostic value of histopathologic pattern for long-term surgical outcomes of 198 patients with confirmed mesial temporal lobe epilepsy. <i>Human Pathology</i> , 2021 , 115, 47-55	3.7	1

- 5 Mitochondrial Mutations in Ethambutol-Induced Optic Neuropathy. *Frontiers in Cell and Developmental Biology*, **2021**, 9, 754676 5·7 ○
- 4 Phenotype-Based Genetic Analysis Reveals Missing Heritability of ABCA4-Related Retinopathy: Deep Intronic Variants and Copy Number Variations **2022**, 63, 5 ○
- 3 Genotype Profile of Global EYS-Associated Inherited Retinal Dystrophy and Clinical Findings in a Large Chinese Cohort. *Frontiers in Cell and Developmental Biology*, **2021**, 9, 634220 5·7
- 2 genotypic and ocular phenotypic characterisation in a Chinese cohort with ocular albinism. *Ophthalmic Genetics*, **2021**, 42, 717-724 1·2
- 1 Artificial Intelligence Uncovers Natural MMP Inhibitor Crocin as a Potential Treatment of Thoracic Aortic Aneurysm and Dissection.. *Frontiers in Cardiovascular Medicine*, **2022**, 9, 871486 5·4