

# Malena Daich Varela

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9555970/publications.pdf>

Version: 2024-02-01

14  
papers

204  
citations

1307594

7  
h-index

1199594

12  
g-index

14  
all docs

14  
docs citations

14  
times ranked

164  
citing authors

#	ARTICLE	IF	CITATIONS
1	Treatments for dry age-related macular degeneration: therapeutic avenues, clinical trials and future directions. <i>British Journal of Ophthalmology</i> , 2022, 106, 297-304.	3.9	86
2	Leber congenital amaurosis/early-onset severe retinal dystrophy: current management and clinical trials. <i>British Journal of Ophthalmology</i> , 2022, 106, 445-451.	3.9	35
3	A rare canonical splice-site variant in VPS13B causes attenuated Cohen syndrome. <i>Ophthalmic Genetics</i> , 2022, 43, 110-115.	1.2	2
4	Functional evaluation in inherited retinal disease. <i>British Journal of Ophthalmology</i> , 2022, 106, 1479-1487.	3.9	9
5	RDH12 retinopathy: clinical features, biology, genetics and future directions. <i>Ophthalmic Genetics</i> , 2022, 43, 301-306.	1.2	4
6	A sialidosis type I cohort and a quantitative approach to multimodal ophthalmic imaging of the macular cherry-red spot. <i>British Journal of Ophthalmology</i> , 2021, 105, 838-843.	3.9	4
7	Clinical diagnosis of presumed SOX2 gonadosomatic mosaicism. <i>Ophthalmic Genetics</i> , 2021, 42, 320-325.	1.2	2
8	Structural evaluation in inherited retinal diseases. <i>British Journal of Ophthalmology</i> , 2021, 105, 1623-1631.	3.9	15
9	Reply. <i>Ophthalmology</i> , 2021, 128, e214-e215.	5.2	0
10	PDE6C: Novel Mutations, Atypical Phenotype, and Differences Among Children and Adults. , 2020, 61, 1.		8
11	A genetic and clinical study of individuals with nonsyndromic retinopathy consequent upon sequence variants in <i>HGSNAT</i> , the gene associated with Sanfilippo C mucopolysaccharidosis. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2020, 184, 631-643.	1.6	12
12	The peroxisomal disorder spectrum and Heimler syndrome: Deep phenotyping and review of the literature. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2020, 184, 618-630.	1.6	14
13	Ophthalmic genetics in South America. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2020, 184, 753-761.	1.6	5
14	Ocular and Systemic Findings in Adults with Uveal Coloboma. <i>Ophthalmology</i> , 2020, 127, 1772-1774.	5.2	8