# Malena Daich Varela

List of Publications by Year
in descending order



| 14 | 204 | 1306789 7 | 1199166 12 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 14 | 14 | 14 | 164 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Treatments for dry age-related macular degeneration: therapeutic avenues, clinical trials and future directions. British Journal of Ophthalmology, 2022, 106, 297-304. | 2.1 | 86 |
| 2 | Leber congenital amaurosis/early-onset severe retinal dystrophy: current management and clinical trials. British Journal of Ophthalmology, 2022, 106, 445-451. | 2.1 | 35 |
| 3 | Structural evaluation in inherited retinal diseases. British Journal of Ophthalmology, 2021, 105, 1623-1631. | 2.1 | 15 |
| 4 | The peroxisomal disorder spectrum and Heimler syndrome: Deep phenotyping and review of the literature. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2020, 184, 618-630. | 0.7 | 14 |
| 5 | A genetic and clinical study of individuals with nonsyndromic retinopathy consequent upon sequence variants in <scp>HGSNAT</scp>, the gene associated with Sanfilippo C mucopolysaccharidosis. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2020, 184, 631-643. | 0.7 | 12 |
| 6 | Functional evaluation in inherited retinal disease. British Journal of Ophthalmology, 2022, 106, 1479-1487. | 2.1 | 9 |
| 7 | PDE6C: Novel Mutations, Atypical Phenotype, and Differences Among Children and Adults. , 2020, 61, 1. | | 8 |
| 8 | Ocular and Systemic Findings in Adults with Uveal Coloboma. Ophthalmology, 2020, 127, 1772-1774. | 2.5 | 8 |
| 9 | Ophthalmic genetics in South America. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2020, 184, 753-761. | 0.7 | 5 |
| 10 | A sialidosis type I cohort and a quantitative approach to multimodal ophthalmic imaging of the macular cherry-red spot. British Journal of Ophthalmology, 2021, 105, 838-843. | 2.1 | 4 |
| 11 | RDH12 retinopathy: clinical features, biology, genetics and future directions. Ophthalmic Genetics, 2022, 43, 301-306. | 0.5 | 4 |
| 12 | Clinical diagnosis of presumed SOX2 gonadosomatic mosaicism. Ophthalmic Genetics, 2021, 42, 320-325. | 0.5 | 2 |
| 13 | A rare canonical splice-site variant in VPS13B causes attenuated Cohen syndrome. Ophthalmic Genetics, 2022, 43, 110-115. | 0.5 | 2 |
| 14 | Reply. Ophthalmology, 2021, 128, e214-e215. | 2.5 | 0 |