

# Sandra Elfride Staffieri

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8680330/publications.pdf>

Version: 2024-02-01

40  
papers

1,233  
citations

430442

18  
h-index

395343

33  
g-index

46  
all docs

46  
docs citations

46  
times ranked

1974  
citing authors

| #  | ARTICLE  | IF  | CITATIONS |
|----|--|-----|-----------|
| 1  | Sex, gender, and retinoblastoma: analysis of 4351 patients from 153 countries. <i>Eye</i> , 2022, 36, 1571-1577.   | 1.1 | 9         |
| 2  | Quality of Life in Adults with Childhood Glaucoma. <i>Ophthalmology Glaucoma</i> , 2022, 5, 325-336.   | 0.9 | 8         |
| 3  | Defining High-risk Retinoblastoma. <i>JAMA Ophthalmology</i> , 2022, 140, 30.  | 1.4 | 9         |
| 4  | The Caregiver Experience in Childhood Glaucoma. <i>Ophthalmology Glaucoma</i> , 2022, 5, 531-543.  | 0.9 | 5         |
| 5  | Retinoblastoma management during the COVID-19 pandemic: A report by the Global Retinoblastoma Study Group including 194 centers from 94 countries. <i>Pediatric Blood and Cancer</i> , 2021, 68, e28584.                 | 0.8 | 8         |
| 6  | Parent satisfaction and acceptability of telehealth consultations in pediatric ophthalmology: initial experience during the COVID-19 pandemic. <i>Journal of AAPOS</i> , 2021, 25, 104-107.                              | 0.2 | 10        |
| 7  | Adherence to home-based videogame treatment for amblyopia in children and adults. <i>Australasian journal of optometry, The</i> , 2021, 104, 773-779.  | 0.6 | 12        |
| 8  | Childhood and Early Onset Glaucoma Classification and Genetic Profile in a Large Australasian Disease Registry. <i>Ophthalmology</i> , 2021, 128, 1549-1560.   | 2.5 | 20        |
| 9  | Living with heritable retinoblastoma and the perceived role of regular follow-up at a retinoblastoma survivorship clinic: "That is exactly what I have been missing"™. <i>BMJ Open Ophthalmology</i> , 2021, 6, e000760. | 0.8 | 2         |
| 10 | Establishing risk of vision loss in Leber hereditary optic neuropathy. <i>American Journal of Human Genetics</i> , 2021, 108, 2159-2170.   | 2.6 | 26        |
| 11 | Improving parents' knowledge of early signs of paediatric eye disease: A double-blind randomized controlled trial. <i>Clinical and Experimental Ophthalmology</i> , 2020, 48, 1250-1260.                                 | 1.3 | 3         |
| 12 | Recurrent Rare Copy Number Variants Increase Risk for Esotropia. , 2020, 61, 22.   |     | 8         |
| 13 | Preimplantation genetic diagnosis for retinoblastoma survivors: a cost-effectiveness study. <i>Reproductive Biomedicine and Society Online</i> , 2020, 10, 37-45.  | 0.9 | 5         |
| 14 | Global Retinoblastoma Presentation and Analysis by National Income Level. <i>JAMA Oncology</i> , 2020, 6, 685.   | 3.4 | 192       |
| 15 | The genetic and clinical landscape of nanophthalmos and posterior microphthalmos in an Australian cohort. <i>Clinical Genetics</i> , 2020, 97, 764-769.  | 1.0 | 17        |
| 16 | Biallelic CPAMD8 Variants Are a Frequent Cause of Childhood and Juvenile Open-Angle Glaucoma. <i>Ophthalmology</i> , 2020, 127, 758-766.   | 2.5 | 33        |
| 17 | Prevalence of <i>FOXC1</i> Variants in Individuals With a Suspected Diagnosis of Primary Congenital Glaucoma. <i>JAMA Ophthalmology</i> , 2019, 137, 348.  | 1.4 | 33        |
| 18 | Optical treatment of amblyopia in older children and adults is essential prior to enrolment in a clinical trial. <i>Ophthalmic and Physiological Optics</i> , 2018, 38, 129-143.   | 1.0 | 37        |

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|----|---|-----|-----------|
| 19 | Effectiveness of a Binocular Video Game vs Placebo Video Game for Improving Visual Functions in Older Children, Teenagers, and Adults With Amblyopia. <i>JAMA Ophthalmology</i> , 2018, 136, 172.                     | 1.4 | 106       |
| 20 | Crowd-sourced Ontology for Photoleukocoria: Identifying Common Internet Search Terms for a Potentially Important Pediatric Ophthalmic Sign. <i>Translational Vision Science and Technology</i> , 2018, 7, 18.         | 1.1 | 3         |
| 21 | Cross-ancestry genome-wide association analysis of corneal thickness strengthens link between complex and Mendelian eye diseases. <i>Nature Communications</i> , 2018, 9, 1864.                                       | 5.8 | 63        |
| 22 | Genome-Wide Association Study Identifies a Susceptibility Locus for Comitant Esotropia and Suggests a Parent-of-Origin Effect. , 2018, 59, 4054.  |     | 21        |
| 23 | Glaucoma spectrum and age-related prevalence of individuals with FOXC1 and PITX2 variants. <i>European Journal of Human Genetics</i> , 2017, 25, 839-847.   | 1.4 | 43        |
| 24 | High-Throughput Genetic Screening of 51 Pediatric Cataract Genes Identifies Causative Mutations in Inherited Pediatric Cataract in South Eastern Australia. <i>G3: Genes, Genomes, Genetics</i> , 2017, 7, 3257-3268. | 0.8 | 20        |
| 25 | Binocular treatment of amblyopia using videogames (BRAVO): study protocol for a randomised controlled trial. <i>Trials</i> , 2016, 17, 504.   | 0.7 | 31        |
| 26 | Managing fetuses at high risk of retinoblastoma: lesion detection on screening MRI. <i>Prenatal Diagnosis</i> , 2015, 35, 174-178.  | 1.1 | 16        |
| 27 | Clinical and molecular characterization of females affected by X-linked retinoschisis. <i>Clinical and Experimental Ophthalmology</i> , 2015, 43, 643-647.  | 1.3 | 7         |
| 28 | Superselective intra-arterial chemotherapy for advanced retinoblastoma complicated by metastatic disease. <i>Journal of AAPOS</i> , 2015, 19, 72-74.  | 0.2 | 14        |
| 29 | CYP1B1 copy number variation is not a major contributor to primary congenital glaucoma. <i>Molecular Vision</i> , 2015, 21, 160-4.  | 1.1 | 4         |
| 30 | Sequencing Analysis of the ATOH7 Gene in Individuals with Optic Nerve Hypoplasia. <i>Ophthalmic Genetics</i> , 2014, 35, 1-6.   | 0.5 | 7         |
| 31 | Genome-wide analysis of multi-ancestry cohorts identifies new loci influencing intraocular pressure and susceptibility to glaucoma. <i>Nature Genetics</i> , 2014, 46, 1126-1130.                                     | 9.4 | 212       |
| 32 | Incidence and predictors of glaucoma following surgery for congenital cataract in the first year of life in Victoria, Australia. <i>Clinical and Experimental Ophthalmology</i> , 2013, 41, 653-661.                  | 1.3 | 24        |
| 33 | Mutations in the EPHA2 Gene Are a Major Contributor to Inherited Cataracts in South-Eastern Australia. <i>PLoS ONE</i> , 2013, 8, e72518.   | 1.1 | 35        |
| 34 | Heritability of Strabismus: Genetic Influence Is Specific to Eso-Deviation and Independent of Refractive Error. <i>Twin Research and Human Genetics</i> , 2012, 15, 624-630.  | 0.3 | 24        |
| 35 | Familial retinal detachment associated with COL2A1 exon 2 and FZD4 mutations. <i>Clinical and Experimental Ophthalmology</i> , 2012, 40, 476-483.   | 1.3 | 9         |
| 36 | Best's macular dystrophy in Australia: phenotypic profile and identification of novel BEST1 mutations. <i>Eye</i> , 2011, 25, 208-217.  | 1.1 | 15        |

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|----|--|-----|-----------|
| 37 | Telemedicine model to prevent blindness from familial glaucoma. Clinical and Experimental Ophthalmology, 2011, 39, 760-765.  | 1.3 | 22        |
| 38 | Mitochondrial Oxidative Phosphorylation Compensation May Preserve Vision in Patients with OPA1-Linked Autosomal Dominant Optic Atrophy. PLoS ONE, 2011, 6, e21347. | 1.1 | 47        |
| 39 | Rock, paper and scissors? Traumatic paediatric cataract in Victoria 1992â€“2006. Clinical and Experimental Ophthalmology, 2010, 38, 237-241.                       | 1.3 | 31        |
| 40 | Retinoblastoma in Victoria, 1976-2000: changing management trends and outcomes. Clinical and Experimental Ophthalmology, 2004, 32, 354-359.                        | 1.3 | 23        |