# Shazia Micheal

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3881554/publications.pdf

Version: 2024-02-01

| 14 | 505 | 933447 10 | 1058476 14 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 14 | 14 | 14 | 1187 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Genome-wide association study identifies five new susceptibility loci for primary angle closure glaucoma. Nature Genetics, 2016, 48, 556-562. | 21.4 | 147 |
| 2 | Genetic association study of exfoliation syndrome identifies a protective rare variant at LOXL1 and five new susceptibility loci. Nature Genetics, 2017, 49, 993-1004. | 21.4 | 114 |
| 3 | Mutations in CEP78 Cause Cone-Rod Dystrophy and Hearing Loss Associated with Primary-Cilia Defects. American Journal of Human Genetics, 2016, 99, 770-776. | 6.2 | 44 |
| 4 | Whole exome sequencing identifies a heterozygous missense variant in the PRDM5 gene in a family with Axenfeldâ€“Rieger syndrome. Neurogenetics, 2016, 17, 17-23. | 1.4 | 28 |
| 5 | Homozygosity Mapping and Targeted Sanger Sequencing Reveal Genetic Defects Underlying Inherited Retinal Disease in Families from Pakistan. PLoS ONE, 2015, 10, e0119806. | 2.5 | 27 |
| 6 | Association of known common genetic variants with primary open angle, primary angle closure, and pseudoexfoliation glaucoma in Pakistani cohorts. Molecular Vision, 2014, 20, 1471-9. | 1.1 | 27 |
| 7 | Identification of Novel Variants in LTBP2 and PXDN Using Whole-Exome Sequencing in Developmental and Congenital Glaucoma. PLoS ONE, 2016, 11, e0159259. | 2.5 | 24 |
| 8 | Identification of novel <scp>CYP1B</scp>1 gene mutations in patients with primary congenital and primary openâ€angle glaucoma. Clinical and Experimental Ophthalmology, 2015, 43, 31-39. | 2.6 | 22 |
| 9 | A Novel Homozygous Mutation in FOXC1 Causes Axenfeld Rieger Syndrome with Congenital Glaucoma. PLoS ONE, 2016, 11, e0160016. | 2.5 | 21 |
| 10 | Identification of Mutations in the PRDM5 Gene in Brittle Cornea Syndrome. Cornea, 2016, 35, 853-859. | 1.7 | 18 |
| 11 | Association of a Polymorphism in the BIRC6 Gene with Pseudoexfoliative Glaucoma. PLoS ONE, 2014, 9, e105023. | 2.5 | 10 |
| 12 | Variants in the ASB10 Gene Are Associated with Primary Open Angle Glaucoma. PLoS ONE, 2015, 10, e0145005. | 2.5 | 10 |
| 13 | Variants in the PRPF8 Gene are Associated with Glaucoma. Molecular Neurobiology, 2018, 55, 4504-4510. | 4.0 | 8 |
| 14 | Identification of TP53BP2 as a Novel Candidate Gene for Primary Open Angle Glaucoma by Whole Exome Sequencing in a Large Multiplex Family. Molecular Neurobiology, 2018, 55, 1387-1395. | 4.0 | 5 |