

# Shazia Micheal

## List of Publications by Year in descending order

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Version: 2024-02-01

14  
papers

505  
citations

933447

10  
h-index

1058476

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14  
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14  
docs citations

14  
times ranked

1187  
citing authors

#	ARTICLE	IF	CITATIONS
1	Variants in the PRPF8 Gene are Associated with Glaucoma. <i>Molecular Neurobiology</i> , 2018, 55, 4504-4510.	4.0	8
2	Identification of TP53BP2 as a Novel Candidate Gene for Primary Open Angle Glaucoma by Whole Exome Sequencing in a Large Multiplex Family. <i>Molecular Neurobiology</i> , 2018, 55, 1387-1395.	4.0	5
3	Genetic association study of exfoliation syndrome identifies a protective rare variant at LOXL1 and five new susceptibility loci. <i>Nature Genetics</i> , 2017, 49, 993-1004.	21.4	114
4	Identification of Novel Variants in LTBP2 and PXDN Using Whole-Exome Sequencing in Developmental and Congenital Glaucoma. <i>PLoS ONE</i> , 2016, 11, e0159259.	2.5	24
5	A Novel Homozygous Mutation in FOXC1 Causes Axenfeld Rieger Syndrome with Congenital Glaucoma. <i>PLoS ONE</i> , 2016, 11, e0160016.	2.5	21
6	Identification of Mutations in the PRDM5 Gene in Brittle Cornea Syndrome. <i>Cornea</i> , 2016, 35, 853-859.	1.7	18
7	Genome-wide association study identifies five new susceptibility loci for primary angle closure glaucoma. <i>Nature Genetics</i> , 2016, 48, 556-562.	21.4	147
8	Mutations in CEP78 Cause Cone-Rod Dystrophy and Hearing Loss Associated with Primary-Cilia Defects. <i>American Journal of Human Genetics</i> , 2016, 99, 770-776.	6.2	44
9	Whole exome sequencing identifies a heterozygous missense variant in the PRDM5 gene in a family with Axenfeld-Rieger syndrome. <i>Neurogenetics</i> , 2016, 17, 17-23.	1.4	28
10	Homozygosity Mapping and Targeted Sanger Sequencing Reveal Genetic Defects Underlying Inherited Retinal Disease in Families from Pakistan. <i>PLoS ONE</i> , 2015, 10, e0119806.	2.5	27
11	Variants in the ASB10 Gene Are Associated with Primary Open Angle Glaucoma. <i>PLoS ONE</i> , 2015, 10, e0145005.	2.5	10
12	Identification of novel <i>CYP11B</i> gene mutations in patients with primary congenital and primary open angle glaucoma. <i>Clinical and Experimental Ophthalmology</i> , 2015, 43, 31-39.	2.6	22
13	Association of a Polymorphism in the BIRC6 Gene with Pseudoexfoliative Glaucoma. <i>PLoS ONE</i> , 2014, 9, e105023.	2.5	10
14	Association of known common genetic variants with primary open angle, primary angle closure, and pseudoexfoliation glaucoma in Pakistani cohorts. <i>Molecular Vision</i> , 2014, 20, 1471-9.	1.1	27