

# Mukesh Tanwar

## List of Publications by Year in descending order

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Version: 2024-02-01

17  
papers

306  
citations

1163117

8  
h-index

1058476

14  
g-index

17  
all docs

17  
docs citations

17  
times ranked

559  
citing authors

#	ARTICLE	IF	CITATIONS
1	VSX1 gene analysis in keratoconus. <i>Molecular Vision</i> , 2010, 16, 2395-401.	1.1	47
2	Mutation spectrum of CYP1B1 in North Indian congenital glaucoma patients. <i>Molecular Vision</i> , 2009, 15, 1200-9.	1.1	44
3	Axenfeld-Rieger Syndrome Associated with Congenital Glaucoma and Cytochrome P4501B1 Gene Mutations. <i>Case Reports in Medicine</i> , 2010, 2010, 1-6.	0.7	41
4	Effect of <i>Tectona grandis</i> Linn. on dexamethasone-induced insulin resistance in mice. <i>Journal of Ethnopharmacology</i> , 2009, 122, 304-307.	4.1	37
5	Mitochondrial DNA analysis in primary congenital glaucoma. <i>Molecular Vision</i> , 2010, 16, 518-33.	1.1	35
6	Homozygosity Mapping and Genetic Analysis of Autosomal Recessive Retinal Dystrophies in 144 Consanguineous Pakistani Families. , 2017, 58, 2218.		34
7	Identification of four novel cytochrome P4501B1 mutations (p.I94X, p.H279D, p.Q340H, and p.K433K) in primary congenital glaucoma patients. <i>Molecular Vision</i> , 2009, 15, 2926-37.	1.1	18
8	MYOC and FOXC1 gene analysis in primary congenital glaucoma. <i>Molecular Vision</i> , 2010, 16, 1996-2006.	1.1	15
9	Preventive effect of <i>Sphaeranthus indicus</i> during progression of glucocorticoid-induced insulin resistance in mice. <i>Pharmaceutical Biology</i> , 2010, 48, 1371-1375.	2.9	10
10	Sturge-Weber Syndrome With Congenital Glaucoma and Cytochrome P450 (CYP1B1) Gene Mutations. <i>Journal of Glaucoma</i> , 2010, 19, 398-404.	1.6	8
11	PAX6 gene analysis in irido-fundal coloboma. <i>Molecular Vision</i> , 2011, 17, 1414-9.	1.1	6
12	Novel occurrence of axenfeld: Rieger syndrome in a patient with blepharophimosis ptosis epicanthus inversus syndrome. <i>Indian Journal of Ophthalmology</i> , 2014, 62, 358.	1.1	4
13	Rieger syndrome with multiple chromosomal breaks and chromosome 4 deletion. <i>BMJ Case Reports</i> , 2009, 2009, bcr0620080297-bcr0620080297.	0.5	3
14	Genetic association of <i>ARID5B</i> with the risk of colorectal cancer within Jammu and Kashmir, India. <i>Genes and Genetic Systems</i> , 2021, 96, 187-191.	0.7	3
15	Cytogenetic and Clinical Assessment of a Family with Treacher Collins Syndrome. <i>Case Reports in Medicine</i> , 2011, 2011, 1-5.	0.7	1
16	Microbiome of the Human Eye. , 2021, , 517-517.		0
17	Polymorphism in the TP63 gene imparts a potential risk for leukemia in the North Indian population. <i>African Health Sciences</i> , 2021, 21, 1243-1249.	0.7	0