Mukesh Tanwar

List of Publications by Year
in descending order

17	306	1163117 8	1058476 14
papers	citations	h-index	g-index

17	17	17	559
all docs	docs citations	times ranked	citing authors

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