Haiba Kaul

List of Publications by Year
in descending order

24 papers	288 citations	1040056 9 h-index	888059 17 g-index
24 all docs	24 docs citations	24 times ranked	361 citing authors

#	Article	IF	Citations
1	Delineating Novel and Known Pathogenic Variants in TYR, OCA2 and HPS-1 Genes in Eight Oculocutaneous Albinism (OCA) Pakistani Families. Genes, 2022, 13, 503.	2.4	2
2	Clinical profile and screening of exon 6 and 14 of ABCB4 gene in obstetric cholestasis patients at a tertiary care hospital in Rawalpindi, Pakistan. JPMA the Journal of the Pakistan Medical Association, 2021, 71, 1-16.	0.2	0
3	SLC4A11 mutations causative of congenital hereditary endothelial dystrophy (CHED) progressing to Harboyan syndrome in consanguineous Pakistani families. Molecular Biology Reports, 2021, 48, 7467-7476.	2.3	4
4	Genetic mapping of autosomal recessive microspherophakia to chromosome 14q24.3 in a consanguineous Pakistani family and screening of exon 36 of LTBP2 gene. JPMA the Journal of the Pakistan Medical Association, 2020, 70, 1.	0.2	2
5	Association of Interleukin 10 (IL-10) Gene with Type 2 Diabetes Mellitus by Single Nucleotide Polymorphism of Its Promotor Region G/A 1082. Critical Reviews in Eukaryotic Gene Expression, 2020, 30, 285-289.	0.9	11
6	Cytochrome 2C19 and paraoxonase-1 polymorphisms and clopidogrel resistance in ischemic heart disease patients. Personalized Medicine, 2019, 16, 379-386.	1.5	5
7	Autosomal recessive congenital cataracts linked to HSF4 in a consanguineous Pakistani family. PLoS ONE, 2019, 14, e0225010.	2.5	5
8	Mutational analysis of the CYP1B1 gene in Pakistani primary congenital glaucoma patients: Identification of four known and a novel causative variant at the 3â€² splice acceptor site of intron 2. Congenital Anomalies (discontinued), 2019, 59, 152-161.	0.6	12
9	Mutations in TYR and OCA2 associated with oculocutaneous albinism in Pakistani families. Meta Gene, 2018, 17, 48-55.	0.6	9
10	In silico analysis of five missense mutations in CYP1B1 gene in Pakistani families affected with primary congenital glaucoma. International Ophthalmology, 2018, 38, 807-814.	1.4	8
11	Fine mapping of chromosome 9 locus associated with congenital cataract. International Ophthalmology, 2018, 38, 1187-1192.	1.4	2
12	Prevalence of stromal corneal dystrophies in Lahore. JPMA the Journal of the Pakistan Medical Association, 2018, 68, 663-665.	0.2	1
13	Determination of Serum Trace Elements (Zn, Cu, and Fe) in Pakistani Patients with Rheumatoid Arthritis. Biological Trace Element Research, 2017, 175, 10-16.	3.5	16
14	Interleukin 6 Receptor (IL6-R) Gene Polymorphisms Underlie Susceptibility to Rheumatoid Arthritis. Clinical Laboratory, 2017, 63, 1365-1369.	0.5	9
15	Mutation in LIM2 Is Responsible for Autosomal Recessive Congenital Cataracts. PLoS ONE, 2016, 11, e0162620.	2.5	17
16	Missense mutation in SLC4A11 in two Pakistani families affected with congenital hereditary endothelial dystrophy (CHED2). Australasian journal of optometry, The, 2016, 99, 73-77.	1.3	16
17	Homozygosity mapping of a consanguineous Pakistani family affected with oculocutaneous albinism to Tyrosinase gene. International Journal of Ophthalmology, 2016, 9, 794-6.	1.1	2
18	Ectopia Lentis in a Consanguineous Pakistani Family and a Novel Locus on Chromosome 8q. JAMA Ophthalmology, 2010, 128, 1046.	2.4	1

#	ARTICLE	IF	CITATIONS
19	A new locus for autosomal recessive congenital cataract identified in a Pakistani family. Molecular Vision, 2010, 16, 240-5.	1.1	25
20	Autosomal recessive congenital cataract linked to EPHA2 in a consanguineous Pakistani family. Molecular Vision, 2010, 16, 511-7.	1.1	81
21	Autosomal recessive congenital cataract in consanguineous Pakistani families is associated with mutations in GALK1. Molecular Vision, 2010, 16, 682-8.	1.1	16
22	Mapping of a novel locus associated with autosomal recessive congenital cataract to chromosome 8p. Molecular Vision, 2010, 16, 2911-5.	1.1	10
23	Novel SIL1 mutations in consanguineous Pakistani families mapping to chromosomes 5q31. Molecular Vision, 2009, 15, 1050-6.	1.1	13
24	Localization of autosomal recessive congenital cataracts in consanguineous Pakistani families to a new locus on chromosome 1p. Molecular Vision, 2007, 13, 1635-40.	1.1	21