

Haiba Kaul

List of Publications by Year in descending order

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Version: 2024-02-01

24
papers

288
citations

1040056

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888059

17
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24
all docs

24
docs citations

24
times ranked

361
citing authors

#	ARTICLE	IF	CITATIONS
1	Autosomal recessive congenital cataract linked to EPHA2 in a consanguineous Pakistani family. <i>Molecular Vision</i> , 2010, 16, 511-7.	1.1	81
2	A new locus for autosomal recessive congenital cataract identified in a Pakistani family. <i>Molecular Vision</i> , 2010, 16, 240-5.	1.1	25
3	Localization of autosomal recessive congenital cataracts in consanguineous Pakistani families to a new locus on chromosome 1p. <i>Molecular Vision</i> , 2007, 13, 1635-40.	1.1	21
4	Mutation in LIM2 Is Responsible for Autosomal Recessive Congenital Cataracts. <i>PLoS ONE</i> , 2016, 11, e0162620.	2.5	17
5	Missense mutation in SLC4A11 in two Pakistani families affected with congenital hereditary endothelial dystrophy (CHED2). <i>Australasian journal of optometry, The</i> , 2016, 99, 73-77.	1.3	16
6	Determination of Serum Trace Elements (Zn, Cu, and Fe) in Pakistani Patients with Rheumatoid Arthritis. <i>Biological Trace Element Research</i> , 2017, 175, 10-16.	3.5	16
7	Autosomal recessive congenital cataract in consanguineous Pakistani families is associated with mutations in GALK1. <i>Molecular Vision</i> , 2010, 16, 682-8.	1.1	16
8	Novel SIL1 mutations in consanguineous Pakistani families mapping to chromosomes 5q31. <i>Molecular Vision</i> , 2009, 15, 1050-6.	1.1	13
9	Mutational analysis of the <i>CYP1B1</i> gene in Pakistani primary congenital glaucoma patients: Identification of four known and a novel causative variant at the 3' splice acceptor site of intron 2. <i>Congenital Anomalies (discontinued)</i> , 2019, 59, 152-161.	0.6	12
10	Association of Interleukin 10 (IL-10) Gene with Type 2 Diabetes Mellitus by Single Nucleotide Polymorphism of Its Promotor Region G/A 1082. <i>Critical Reviews in Eukaryotic Gene Expression</i> , 2020, 30, 285-289.	0.9	11
11	Mapping of a novel locus associated with autosomal recessive congenital cataract to chromosome 8p. <i>Molecular Vision</i> , 2010, 16, 2911-5.	1.1	10
12	Mutations in TYR and OCA2 associated with oculocutaneous albinism in Pakistani families. <i>Meta Gene</i> , 2018, 17, 48-55.	0.6	9
13	Interleukin 6 Receptor (IL6-R) Gene Polymorphisms Underlie Susceptibility to Rheumatoid Arthritis. <i>Clinical Laboratory</i> , 2017, 63, 1365-1369.	0.5	9
14	In silico analysis of five missense mutations in CYP1B1 gene in Pakistani families affected with primary congenital glaucoma. <i>International Ophthalmology</i> , 2018, 38, 807-814.	1.4	8
15	<i>Cytochrome 2C19</i> and <i>paraoxonase-1</i> polymorphisms and clopidogrel resistance in ischemic heart disease patients. <i>Personalized Medicine</i> , 2019, 16, 379-386.	1.5	5
16	Autosomal recessive congenital cataracts linked to HSF4 in a consanguineous Pakistani family. <i>PLoS ONE</i> , 2019, 14, e0225010.	2.5	5
17	SLC4A11 mutations causative of congenital hereditary endothelial dystrophy (CHED) progressing to Harboyan syndrome in consanguineous Pakistani families. <i>Molecular Biology Reports</i> , 2021, 48, 7467-7476.	2.3	4
18	Fine mapping of chromosome 9 locus associated with congenital cataract. <i>International Ophthalmology</i> , 2018, 38, 1187-1192.	1.4	2

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19	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
20	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
21	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
22	Ectopia Lentis in a Consanguineous Pakistani Family and a Novel Locus on Chromosome 8q. JAMA Ophthalmology, 2010, 128, 1046.	2.4	1
23	Prevalence of stromal corneal dystrophies in Lahore. JPMA the Journal of the Pakistan Medical Association, 2018, 68, 663-665.	0.2	1
24	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