Naveed Wasif

List of Publications by Year
in descending order

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Version: 2024-02-01

		687363	713466
27	453	13	21
papers	citations	h-index	g-index

27	27	27	764
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Whole Exome Sequencing Confirms Molecular Diagnostics of Three Pakhtun Families With Autosomal Recessive Epidermolysis Bullosa. Frontiers in Pediatrics, 2021, 9, 727288.	1.9	2
2	The First Report of a Missense Variant in RFX2 Causing Non-Syndromic Tooth Agenesis in a Consanguineous Pakistani Family. Frontiers in Genetics, 2021, 12, 782653.	2.3	1
3	A Novel Homozygous Frameshift Variant in DYM Causing Dyggve-Melchior-Clausen Syndrome in Pakistani Patients. Frontiers in Pediatrics, 2020, 8, 383.	1.9	7
4	Novel homozygous nonsense variant in MLPH causing Griscelli syndrome type 3 in a consanguineous Pakistani family. Journal of Dermatology, 2020, 47, e382-e383.	1.2	2
5	A novel nonsense variant in SLC24A4 causing a rare form of amelogenesis imperfecta in a Pakistani family. BMC Medical Genetics, 2020, 21, 97.	2.1	6
6	A Novel Homozygous Frameshift Mutation in CCN6 Causing Progressive Pseudorheumatoid Dysplasia (PPRD) in a Consanguineous Yemeni Family. Frontiers in Pediatrics, 2019, 7, 245.	1.9	2
7	A novel pathogenic missense variant in CNNM4 underlying Jalili syndrome: Insights from molecular dynamics simulations. Molecular Genetics & Genomic Medicine, 2019, 7, e902.	1.2	11
8	Deleterious Variants in WNT10A, EDAR, and EDA Causing Isolated and Syndromic Tooth Agenesis: A Structural Perspective from Molecular Dynamics Simulations. International Journal of Molecular Sciences, 2019, 20, 5282.	4.1	19
9	Exome sequencing revealed a novel lossâ€ofâ€function variant in the GLI3 transcriptional activator 2 domain underlies nonsyndromic postaxial polydactyly. Molecular Genetics & Genomic Medicine, 2019, 7, e00627.	1.2	15
10	Variants in KIAA0825 underlie autosomal recessive postaxial polydactyly. Human Genetics, 2019, 138, 593-600.	3.8	16
11	Genetic analysis of Xp22.3 micro-deletions in seventeen families segregating isolated form of X-linked ichthyosis. Journal of Dermatological Science, 2015, 80, 214-217.	1.9	3
12	Mutations in TBC1D24, a Gene Associated With Epilepsy, Also Cause Nonsyndromic Deafness DFNB86. American Journal of Human Genetics, 2014, 94, 144-152.	6.2	72
13	A Novel Nonsense Mutation in RSPO4 Gene Underlies Autosomal Recessive Congenital Anonychia in a Pakistani Family. Pediatric Dermatology, 2013, 30, 139-141.	0.9	10
14	Whole exome sequencing identified a novel zinc-finger geneZNF141associated with autosomal recessive postaxial polydactyly type A. Journal of Medical Genetics, 2013, 50, 47-53.	3.2	51
15	Congenital atrichia with papular lesions resulting from novel mutations in human hairless gene in four consanguineous families. Journal of Dermatology, 2011, 38, 755-760.	1.2	11
16	Mutations in the LPAR6 and LIPH genes underlie autosomal recessive hypotrichosis/woolly hair in 17 consanguineous families from Pakistan. Clinical and Experimental Dermatology, 2011, 36, 652-654.	1.3	18
17	Two novel mutations in the gene EDAR causing autosomal recessive hypohidrotic ectodermal dysplasia. Orthodontics and Craniofacial Research, 2011, 14, 156-159.	2.8	6
18	Novel mutations in the keratin-74 (KRT74) gene underlie autosomal dominant woolly hair/hypotrichosis in Pakistani families. Human Genetics, 2011, 129, 419-424.	3.8	29

#	Article	IF	Citations
19	Genetic mapping of a novel hypotrichosis locus to chromosome 7p21.3â€“p22.3 in a Pakistani family and screening of the candidate genes. Human Genetics, 2010, 128, 213-220.	3.8	13
20	A Novel Splice Site Mutation in theEDARGene Underlies Autosomal Recessive Hypohidrotic Ectodermal Dysplasia in a Pakistani Family. Pediatric Dermatology, 2010, 27, 106-108.	0.9	2
21	A novel missense mutation in the EVC gene underlies Ellisâ€van Creveld syndrome in a Pakistani family. Pediatrics International, 2010, 52, 240-246.	0.5	10
22	Mutations in theP2RY5gene underlie autosomal recessive hypotrichosis in 13 Pakistani families. British Journal of Dermatology, 2009, 160, 1006-1010.	1.5	20
23	Novel mutations in G protein-coupled receptor gene (P2RY5) in families with autosomal recessive hypotrichosis (LAH3). Human Genetics, 2008, 123, 515-519.	3.8	37
24	A novel insertion mutation in the cartilage-derived morphogenetic protein-1 (CDMP1) gene underlies Grebe-type chondrodysplasia in a consanguineous Pakistani family. BMC Medical Genetics, 2008, 9, 102.	2.1	32
25	A novel deletion mutation in LIPH gene causes autosomal recessive hypotrichosis (LAH2). Clinical Genetics, 2008, 74, 184-188.	2.0	28
26	A novel deletion mutation in the EDAR gene in a Pakistani family with autosomal recessive hypohidrotic ectodermal dysplasia. British Journal of Dermatology, 2007, 157, 207-209.	1.5	15
27	A novel 4-bp insertion mutation in EDA1 gene in a Pakistani family with X-linked hypohidrotic ectodermal dysplasia. European Journal of Dermatology, 2007, 17, 209-12.	0.6	15