

# Naveed Wasif

## List of Publications by Year in descending order

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27  
papers

453  
citations

687363

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713466

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docs citations

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times ranked

764  
citing authors

#	ARTICLE	IF	CITATIONS
1	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
2	Whole exome sequencing identified a novel zinc-finger gene<i>ZNF141</i> associated with autosomal recessive postaxial polydactyly type A. Journal of Medical Genetics, 2013, 50, 47-53.	3.2	51
3	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
4	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
5	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
6	A novel deletion mutation in <i>LIPH</i> gene causes autosomal recessive hypotrichosis (LAH2). Clinical Genetics, 2008, 74, 184-188.	2.0	28
7	Mutations in the<i>P2RY5</i> gene underlie autosomal recessive hypotrichosis in 13 Pakistani families. British Journal of Dermatology, 2009, 160, 1006-1010.	1.5	20
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	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
10	Variants in KIAA0825 underlie autosomal recessive postaxial polydactyly. Human Genetics, 2019, 138, 593-600.	3.8	16
11	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
12	Exome sequencing revealed a novel loss-of-function variant in the GLI3 transcriptional activator 2 domain underlies nonsyndromic postaxial polydactyly. Molecular Genetics & Genomic Medicine, 2019, 7, e00627.	1.2	15
13	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
14	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
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	A novel pathogenic missense variant in <i>CNNM4</i> underlying Jalili syndrome: Insights from molecular dynamics simulations. Molecular Genetics & Genomic Medicine, 2019, 7, e902.	1.2	11
17	A novel missense mutation in the <i>EVC</i> gene underlies Ellisâ€“van Creveld syndrome in a Pakistani family. Pediatrics International, 2010, 52, 240-246.	0.5	10
18	A Novel Nonsense Mutation in <i>RSP04</i> Gene Underlies Autosomal Recessive Congenital Anonychia in a Pakistani Family. Pediatric Dermatology, 2013, 30, 139-141.	0.9	10

#	ARTICLE	IF	CITATIONS
19	A Novel Homozygous Frameshift Variant in DYM Causing Dyggve-Melchior-Clausen Syndrome in Pakistani Patients. <i>Frontiers in Pediatrics</i> , 2020, 8, 383.	1.9	7
20	Two novel mutations in the gene <i>EDAR</i> causing autosomal recessive hypohidrotic ectodermal dysplasia. <i>Orthodontics and Craniofacial Research</i> , 2011, 14, 156-159.	2.8	6
21	A novel nonsense variant in SLC24A4 causing a rare form of amelogenesis imperfecta in a Pakistani family. <i>BMC Medical Genetics</i> , 2020, 21, 97.	2.1	6
22	Genetic analysis of Xp22.3 micro-deletions in seventeen families segregating isolated form of X-linked ichthyosis. <i>Journal of Dermatological Science</i> , 2015, 80, 214-217.	1.9	3
23	A Novel Splice Site Mutation in the EDAR Gene Underlies Autosomal Recessive Hypohidrotic Ectodermal Dysplasia in a Pakistani Family. <i>Pediatric Dermatology</i> , 2010, 27, 106-108.	0.9	2
24	A Novel Homozygous Frameshift Mutation in CCN6 Causing Progressive Pseudorheumatoid Dysplasia (PPRD) in a Consanguineous Yemeni Family. <i>Frontiers in Pediatrics</i> , 2019, 7, 245.	1.9	2
25	Novel homozygous nonsense variant in <i>MLPH</i> causing Griscelli syndrome type 3 in a consanguineous Pakistani family. <i>Journal of Dermatology</i> , 2020, 47, e382-e383.	1.2	2
26	Whole Exome Sequencing Confirms Molecular Diagnostics of Three Pakhtun Families With Autosomal Recessive Epidermolysis Bullosa. <i>Frontiers in Pediatrics</i> , 2021, 9, 727288.	1.9	2
27	The First Report of a Missense Variant in RFX2 Causing Non-Syndromic Tooth Agenesis in a Consanguineous Pakistani Family. <i>Frontiers in Genetics</i> , 2021, 12, 782653.	2.3	1