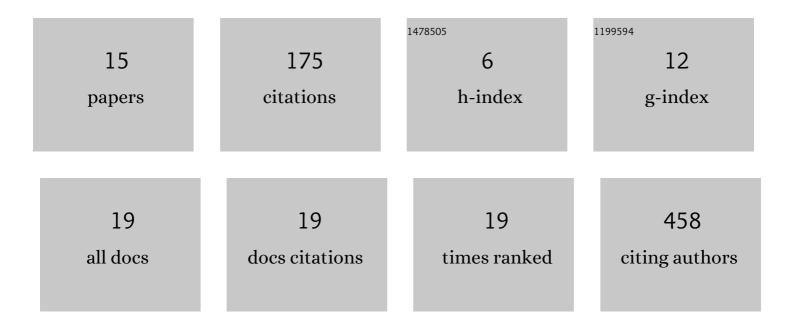
Asif Mir

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Hutchinson–Gilford Progeria Syndrome: A Premature Aging Disease. Molecular Neurobiology, 2017, 55, 4417-4427.	4.0	57
2	Identification of bioflavonoid as fusion inhibitor of dengue virus using molecular docking approach. Informatics in Medicine Unlocked, 2016, 3, 1-6.	3.4	25
3	Truncation of the E3 ubiquitin ligase component FBXO31 causes non-syndromic autosomal recessive intellectual disability in a Pakistani family. Human Genetics, 2014, 133, 975-984.	3.8	24
4	An update of pathogenic variants in <i>ASPM</i> , <i>WDR62, CDK5RAP2</i> , <i> STIL, CENPJ,</i> and <i> CEP135</i> underlying autosomal recessive primary microcephaly in 32 consanguineous families from Pakistan. Molecular Genetics & amp; Genomic Medicine, 2020, 8, e1408.	1.2	18
5	Adaptive evolution and elucidating the potential inhibitor against schizophrenia to target DAOA (G72) isoforms. Drug Design, Development and Therapy, 2015, 9, 3471.	4.3	12
6	Causal variants in Maturity Onset Diabetes of the Young (MODY) – A systematic review. BMC Endocrine Disorders, 2021, 21, 223.	2.2	10
7	ldentification and in silico characterization of a novel p.P208PfsX1 mutation in V-ATPase a3 subunit associated with autosomal recessive osteopetrosis in a Pakistani family. BMC Medical Genetics, 2017, 18, 148.	2.1	7
8	Identification and in silico characterization of p.G380R substitution in FGFR3, associated with achondroplasia in a non-consanguineous Pakistani family. Diagnostic Pathology, 2017, 12, 47.	2.0	7
9	ldentification of novel mutation in the <scp><i>HR</i></scp> gene responsible for atrichia with papular lesions in a <scp>P</scp> akistani family. Journal of Dermatology, 2013, 40, 927-928.	1.2	5
10	Whole-exome sequencing identifies a novel mutation in spermine synthase gene (SMS) associated with Snyder-Robinson Syndrome. BMC Medical Genetics, 2020, 21, 168.	2.1	3
11	Novel variants underlying autosomal recessive intellectual disability in Pakistani consanguineous families. BMC Medical Genetics, 2020, 21, 59.	2.1	3
12	Genetic characterization of suspected MODY patients in Pakistan by next generation sequencing—a pilot study. International Journal of Diabetes in Developing Countries, 2021, 41, 563-569.	0.8	2
13	Maturity Onset Diabetes of the Young – An Overview of Common Types. A Review. Romanian Journal of Diabetes Nutrition and Metabolic Diseases, 2018, 25, 209-213.	0.3	2
14	Exploring evolution of brain genes involved in microcephaly through phylogeny and synteny analysis. Theoretical Biology and Medical Modelling, 2013, 10, 61.	2.1	0
15	Identification and association of recurrent ALOXE3 mutation with nonâ€bullous congenital ichthyosiform erythroderma in two ethnically distinct Pakistani families. Congenital Anomalies (discontinued), 2019, 59, 93-98.	0.6	0