Asif Mir

List of Publications by Year
in descending order

15 papers	175 citations	1478505 6 h-index	1199594 12 g-index
19 all docs	19 docs citations	19 times ranked	458 citing authors

#	Article	IF	Citations
1	Genetic characterization of suspected MODY patients in Pakistan by next generation sequencingâ€"a pilot study. International Journal of Diabetes in Developing Countries, 2021, 41, 563-569.	0.8	2
2	Causal variants in Maturity Onset Diabetes of the Young (MODY) â€" A systematic review. BMC Endocrine Disorders, 2021, 21, 223.	2.2	10
3	An update of pathogenic variants in ASPM, WDR62, CDK5RAP2, STIL, CENPJ, and CEP135 underlying autosomal recessive primary microcephaly in 32 consanguineous families from Pakistan. Molecular Genetics & Genomic Medicine, 2020, 8, e1408.	1.2	18
4	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
5	Novel variants underlying autosomal recessive intellectual disability in Pakistani consanguineous families. BMC Medical Genetics, 2020, 21, 59.	2.1	3
6	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
7	Maturity Onset Diabetes of the Young â€" An Overview of Common Types. A Review. Romanian Journal of Diabetes Nutrition and Metabolic Diseases, 2018, 25, 209-213.	0.3	2
8	Hutchinsonâ€"Gilford Progeria Syndrome: A Premature Aging Disease. Molecular Neurobiology, 2017, 55, 4417-4427.	4.0	57
9	Identification 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
10	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
11	Identification of bioflavonoid as fusion inhibitor of dengue virus using molecular docking approach. Informatics in Medicine Unlocked, 2016, 3, 1-6.	3.4	25
12	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
13	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
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 of novel mutation in the <scp>HR</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