# Asif Mir

List of Publications by Year
in descending order



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|---|---|---|---|
| 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 | Hutchinsonâ€“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 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 |
| 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) â€“ A systematic review. BMC Endocrine Disorders, 2021, 21, 223. | 2.2 | 10 |
| 7 | 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 |
| 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 | 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 |
| 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 sequencingâ€”a pilot study. International Journal of Diabetes in Developing Countries, 2021, 41, 563-569. | 0.8 | 2 |
| 13 | 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 |
| 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 |