# Sher Alam Khan

List of Publications by Year
in descending order



| | | | |
|---|---|---|---|
| 10 papers | 69 citations | 1684188 5 h-index | 1588992 8 g-index |
| 10 all docs | 10 docs citations | 10 times ranked | 142 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 | Genetic Spectrum of Syndromic and Non-Syndromic Hearing Loss in Pakistani Families. Genes, 2020, 11, 1329. | 2.4 | 7 |
| 4 | 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 |
| 5 | Homozygous variants of EDAR underlying hypohidrotic ectodermal dysplasia in three consanguineous families. European Journal of Dermatology, 2020, 30, 408-416. | 0.6 | 4 |
| 6 | 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 |
| 7 | Association of sequence variants in frizzled-6 with autosomal recessive nail dysplasia (NDNC-10) in Pashtun families. JPMA the Journal of the Pakistan Medical Association, 2019, 70, 1. | 0.2 | 0 |
| 8 | A novel mutation in the HPGD gene causing primary hypertrophic osteoarthropathy with digital clubbing in a Pakistani family. Annals of Human Genetics, 2018, 82, 171-176. | 0.8 | 9 |
| 9 | Novel sequence variants in the MKKS gene cause Bardetâ€Biedl syndrome with intraâ€•and interâ€familial variable phenotypes. Congenital Anomalies (discontinued), 2018, 58, 173-175. | 0.6 | 11 |
| 10 | A novel mutation in homeobox DNA binding domain of HOXC13 gene underlies pure hair and nail ectodermal dysplasia (ECTD9) in a Pakistani family. BMC Medical Genetics, 2017, 18, 42. | 2.1 | 10 |