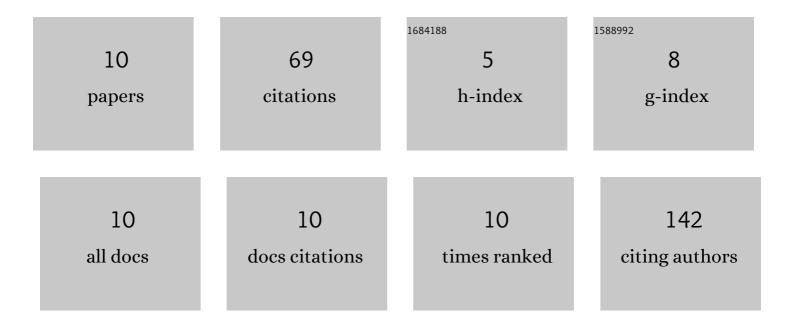
## Sher Alam Khan

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/208768/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	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
2	Novel sequence variants in the <i>MKKS</i> gene cause Bardetâ€Biedl syndrome with intra―and interâ€familial variable phenotypes. Congenital Anomalies (discontinued), 2018, 58, 173-175.	0.6	11
3	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
4	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
5	Genetic Spectrum of Syndromic and Non-Syndromic Hearing Loss in Pakistani Families. Genes, 2020, 11, 1329.	2.4	7
6	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
7	Homozygous variants of EDAR underlying hypohidrotic ectodermal dysplasia in three consanguineous families. European Journal of Dermatology, 2020, 30, 408-416.	0.6	4
8	Whole Exome Sequencing Confirms Molecular Diagnostics of Three Pakhtun Families With Autosomal Recessive Epidermolysis Bullosa. Frontiers in Pediatrics, 2021, 9, 727288.	1.9	2
9	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
10	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