

Sher Alam Khan

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/208768/publications.pdf>

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10
papers

69
citations

1684188

5
h-index

1588992

8
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10
times ranked

142
citing authors

#	ARTICLE	IF	CITATIONS
1	Deleterious Variants in WNT10A, EDAR, and EDA Causing Isolated and Syndromic Tooth Agenesis: A Structural Perspective from Molecular Dynamics Simulations. <i>International Journal of Molecular Sciences</i> , 2019, 20, 5282.	4.1	19
2	Novel sequence variants in the <i>MKKS</i> gene cause Bardet-Biedl syndrome with intra- and inter-familial variable phenotypes. <i>Congenital Anomalies (discontinued)</i> , 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. <i>BMC Medical Genetics</i> , 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. <i>Annals of Human Genetics</i> , 2018, 82, 171-176.	0.8	9
5	Genetic Spectrum of Syndromic and Non-Syndromic Hearing Loss in Pakistani Families. <i>Genes</i> , 2020, 11, 1329.	2.4	7
6	A novel nonsense variant in SLC24A4 causing a rare form of amelogenesis imperfecta in a Pakistani family. <i>BMC Medical Genetics</i> , 2020, 21, 97.	2.1	6
7	Homozygous variants of EDAR underlying hypohidrotic ectodermal dysplasia in three consanguineous families. <i>European Journal of Dermatology</i> , 2020, 30, 408-416.	0.6	4
8	Whole Exome Sequencing Confirms Molecular Diagnostics of Three Pakhtun Families With Autosomal Recessive Epidermolysis Bullosa. <i>Frontiers in Pediatrics</i> , 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. <i>Frontiers in Genetics</i> , 2021, 12, 782653.	2.3	1
10	Association of sequence variants in frizzled-6 with autosomal recessive nail dysplasia (NDNC-10) in Pashtun families. <i>JPMA the Journal of the Pakistan Medical Association</i> , 2019, 70, 1.	0.2	0