

Muhammad Nasir

List of Publications by Year in descending order

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#	ARTICLE	IF	CITATIONS
1	Identification and association of recurrent ALOXE3 mutation with non-bullous congenital ichthyosiform erythroderma in two ethnically distinct Pakistani families. <i>Congenital Anomalies (discontinued)</i> , 2019, 59, 93-98.	0.6	0
2	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. <i>BMC Medical Genetics</i> , 2017, 18, 148.	2.1	7
3	Identification and in silico characterization of p.G380R substitution in FGFR3, associated with achondroplasia in a non-consanguineous Pakistani family. <i>Diagnostic Pathology</i> , 2017, 12, 47.	2.0	7
4	In silico analysis of a disease-causing mutation in PCDH15 gene in a consanguineous Pakistani family with Usher phenotype. <i>International Journal of Ophthalmology</i> , 2016, 9, 662-8.	1.1	6
5	Identification of recurrent c.742G>T nonsense mutation in ECM1 in Pakistani families suffering from lipid proteinosis. <i>Molecular Biology Reports</i> , 2014, 41, 2085-2092.	2.3	4
6	MC1R gene mutation and its association with oculocutaneous albinism type (OCA) phenotype in a consanguineous Pakistani family. <i>Journal of Dermatological Science</i> , 2013, 70, 68-70.	1.9	4
7	In silico characterization of a novel pathogenic deletion mutation identified in XPA gene in a Pakistani family with severe xeroderma pigmentosum. <i>Journal of Biomedical Science</i> , 2013, 20, 70.	7.0	1
8	Molecular analysis of lipid proteinosis: identification of a novel nonsense mutation in the ECM1 gene in a Pakistani family. <i>Diagnostic Pathology</i> , 2011, 6, 69.	2.0	15
9	Characterization of Y-Chromosomal Short Tandem Repeat Markers in Pakistani Populations. <i>Genetic Testing and Molecular Biomarkers</i> , 2011, 15, 165-172.	0.7	2