Muhammad Nasir

List of Publications by Year
in descending order

9	46	1937685 4	1720034 7
papers	citations	h-index	g-index

9	9	9	64
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Molecular analysis of lipoid proteinosis: identification of a novel nonsense mutation in the ECM1 gene in a Pakistani family. Diagnostic Pathology, 2011, 6, 69.	2.0	15
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. BMC Medical Genetics, 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. Diagnostic Pathology, 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. International Journal of Ophthalmology, 2016, 9, 662-8.	1.1	6
5	MC1R gene mutation and its association with oculocutaneous albinism type (OCA) phenotype in a consanguineous Pakistani family. Journal of Dermatological Science, 2013, 70, 68-70.	1.9	4
6	Identification of recurrent c.742G>T nonsense mutation in ECM1 in Pakistani families suffering from lipoid proteinosis. Molecular Biology Reports, 2014, 41, 2085-2092.	2.3	4
7	Characterization of Y-Chromosomal Short Tandem Repeat Markers in Pakistani Populations. Genetic Testing and Molecular Biomarkers, 2011, 15, 165-172.	0.7	2
8	In silico characterization of a novel pathogenic deletion mutation identified in XPA gene in a Pakistani family with severe xeroderma pigmentosum. Journal of Biomedical Science, 2013, 20, 70.	7.0	1
9	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