

Ayaz Khan

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4666632/publications.pdf>

Version: 2024-02-01

7
papers

135
citations

1684188

5
h-index

1720034

7
g-index

7
all docs

7
docs citations

7
times ranked

438
citing authors

#	ARTICLE	IF	CITATIONS
1	Modifier Genes in Microcephaly: A Report on WDR62, CEP63, RAD50 and PCNT Variants Exacerbating Disease Caused by Biallelic Mutations of ASPM and CENPJ. <i>Genes</i> , 2021, 12, 731.	2.4	8
2	Updates on Clinical and Genetic Heterogeneity of ASPM in 12 Autosomal Recessive Primary Microcephaly Families in Pakistani Population. <i>Frontiers in Pediatrics</i> , 2021, 9, 695133.	1.9	5
3	A Homozygous AKNA Frameshift Variant Is Associated with Microcephaly in a Pakistani Family. <i>Genes</i> , 2021, 12, 1494.	2.4	3
4	Identification of a novel variant in GPR56/ADGRC1 gene through whole exome sequencing in a consanguineous Pakistani family. <i>Journal of Clinical Neuroscience</i> , 2021, 94, 8-12.	1.5	1
5	Whole exome sequencing identifies novel variant underlying hereditary spastic paraplegia in consanguineous Pakistani families. <i>Journal of Clinical Neuroscience</i> , 2019, 67, 19-23.	1.5	8
6	Mutations in multiple components of the nuclear pore complex cause nephrotic syndrome. <i>Journal of Clinical Investigation</i> , 2018, 128, 4313-4328.	8.2	89
7	Homozygous GRID2 missense mutation predicts a shift in the D-serine binding domain of GluD2 in a case with generalized brain atrophy and unusual clinical features. <i>BMC Medical Genetics</i> , 2017, 18, 144.	2.1	21