Arezou Karamzade

List of Publications by Year
in descending order

9	58	2258059 3	1720034 7
papers	citations	h-index	g-index

9	9	9	137
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Next-generation sequencing identified novel truncating mutations in BBS9 causing Bardet Biedl syndrome in two Iranian consanguineous families.. Iranian Journal of Child Neurology, 2022, 16, 123-133.	0.3	0
2	Identification of a novel truncating variant in AHI1 gene and a brief review on mutations spectrum. Molecular Biology Reports, 2021, 48, 5339-5345.	2.3	3
3	Familial Hypomagnesemia with Hypercalciuria, Nephrocalcinosis, and Bilateral Chorioretinal Atrophy in a Patient with Homozygous p.G75S Variant in CLDN19. Journal of Pediatric Genetics, 2021, 10, 230-235.	0.7	0
4	Whole-Exome Sequencing Identified a Novel Variant (C.405_422+39del) in DSP Gene in an Iranian Pedigree with Familial Dilated Cardiomyopathy. Reports of Biochemistry and Molecular Biology, 2021, 10, 280-287.	1.4	2
5	X-linked SCID with a rare mutation. Allergy, Asthma and Clinical Immunology, 2021, 17, 107.	2.0	3
6	Dysfunction of the ciliary ARMC9/TOGARAM1 protein module causes Joubert syndrome. Journal of Clinical Investigation, 2020, 130, 4423-4439.	8.2	43
7	Succinate Dehydrogenase Deficiency: A Treatable Neurometabolic Disorder. Iranian Journal of Child Neurology, 2020, 14, 111-116.	0.3	1
8	CRB1-Related Leber Congenital Amaurosis: Reporting Novel Pathogenic Variants and a Brief Review on Mutations Spectrum. Iranian Biomedical Journal, 2019, 23, 362-8.	0.7	1
9	First report of Klein-Waardenburg Syndrome in Iran and a novel pathogenic splice site variant in PAX3 gene. International Journal of Pediatric Otorhinolaryngology, 2018, 113, 229-233.	1.0	5