

Elham Davoudi-Dehaghani

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7865109/publications.pdf>

Version: 2024-02-01

7
papers

37
citations

2682572

2
h-index

2053705

5
g-index

7
all docs

7
docs citations

7
times ranked

72
citing authors

#	ARTICLE	IF	CITATIONS
1	Regulatory Mutation Study in Cases with Unsolved Hypochromic Microcytic Anemia and $\hat{\alpha}$ -Major Regulatory Element Haplotype Analysis in Iran. <i>Hemoglobin</i> , 2021, 45, 37-40.	0.8	0
2	Bi-allelic variants in OGDHL cause a neurodevelopmental spectrum disease featuring epilepsy, hearing loss, visual impairment, and ataxia. <i>American Journal of Human Genetics</i> , 2021, 108, 2368-2384.	6.2	12
3	The Spectrum of Pathogenic Variants in Iranian Families with Hemophilia A. <i>Archives of Iranian Medicine</i> , 2021, 24, 887-896.	0.6	2
4	Once in a Blue Moon, a Very Rare Coexistence of Glutaric Acidemia Type I and Mucopolysaccharidosis Type IIIB in a Patient. <i>Iranian Biomedical Journal</i> , 2020, 24, 201-205.	0.7	2
5	Characterization of Niemann-Pick diseases genes mutation spectrum in Iran and identification of a novel mutation in gene. <i>Medical Journal of the Islamic Republic of Iran</i> , 2019, 33, 126.	0.9	1
6	Homozygosity mapping and CDH23 mutation analysis in Iranian deaf families. <i>Hearing, Balance and Communication</i> , 2016, 14, 189-193.	0.4	0
7	A transversion mutation in non-coding exon 3 of the TMC1 gene in two ethnically related Iranian deaf families from different geographical regions; evidence for founder effect. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2013, 77, 821-826.	1.0	20