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	37	2682572 2	2053705 5
papers	citations	h-index	g-index

7	7	7	72
all docs	docs citations	times ranked	citing authors

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