Daniz Kooshavar

List of Publications by Year
in descending order

4 papers	27 citations	2258059 3 h-index	2550090 3 g-index
4 all docs	4 docs citations	4 times ranked	80 citing authors

#	Article	IF	Citations
1	Digenic inheritance in autosomal recessive non-syndromic hearing loss cases carrying GJB2 heterozygote mutations: Assessment of GJB4, GJA1, and GJC3. International Journal of Pediatric Otorhinolaryngology, 2013, 77, 189-193.	1.0	16
2	Targeted next generation sequencing identified a novel mutation in MYO7A causing Usher syndrome type 1 in an Iranian consanguineous pedigree. International Journal of Pediatric Otorhinolaryngology, 2018, 104, 10-13.	1.0	7
3	Identification of a Novel Mutation in the PAH Gene in an Iranian Phenylketonuria Family: A Case Report. Iranian Journal of Public Health, 2017, 46, 560-564.	0.5	4
4	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