

# Daniz Kooshavar

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5368247/publications.pdf>

Version: 2024-02-01

4  
papers

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citations

2258059

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2550090

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docs citations

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times ranked

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#	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	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	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