

# Parisa Mohamadynejad

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3038763/publications.pdf>

Version: 2024-02-01

8  
papers

20  
citations

2682572

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2272923

4  
g-index

8  
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8  
docs citations

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citing authors

#	ARTICLE	IF	CITATIONS
1	Significant Association of rs77493513 Polymorphism in 3'-UTR of the NRG1 Gene with the Risk of Multiple Sclerosis Disease. <i>Metabolic Brain Disease</i> , 2022, , 1.	2.9	2
2	Association study of polymorphism in Thrombomodulin gene [rs1042579] with cardiovascular disease.. <i>Acta Biomedica</i> , 2022, 92, e2021282.	0.3	2
3	PHLPP2 gene L1016S (rs61733127) and PIK3R1 gene Met326Ile (rs3730089) polymorphisms are associated with the risk of colon and breast cancers. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2021, 40, 767-778.	1.1	1
4	ErbB4 3'UTR Variant (c.*3622A>G) is Associated with ER/PR Negativity and Advanced Breast Cancer. <i>Indian Journal of Clinical Biochemistry</i> , 2020, 35, 115-120.	1.9	8
5	A genetic variant in the flanking region of miR-182 could decrease the susceptibility to the breast cancer risk in the Iranian population. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2020, 39, 806-817.	1.1	0
6	Evaluating the Immune Response of Recombinant H1N1 Hemagglutinin with MF59 Adjuvant in Animal Model as a Novel Alternative to the Influenza Vaccine. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2020, 19, 497-508.	0.4	1
7	Sexual dimorphism in the expression of GKN2 and FOXA2 genes in the human stomach. <i>Molecular Biology Reports</i> , 2019, 46, 2355-2362.	2.3	4
8	Identification of a novel missense mutation of PEX7 gene in an Iranian patient with rhizomelic chondrodysplasia punctata type 1. <i>Gene</i> , 2013, 518, 461-466.	2.2	2