

# Simin Rahimi-Aliabadi

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/11546081/publications.pdf>

Version: 2024-02-01

7  
papers

94  
citations

1478505

6  
h-index

1720034

7  
g-index

7  
all docs

7  
docs citations

7  
times ranked

207  
citing authors

#	ARTICLE	IF	CITATIONS
1	A genetic variant in miRNA binding site of glutamate receptor 4, metabotropic (GRM4) is associated with increased risk of major depressive disorder. <i>Journal of Affective Disorders</i> , 2017, 208, 218-222.	4.1	25
2	The analysis of association between SNCA, HUSEYO and CSMD1 gene variants and Parkinson's disease in Iranian population. <i>Neurological Sciences</i> , 2016, 37, 731-736.	1.9	20
3	Support for "Disease-Only" Genotypes and Excess of Homozygosity at the CYTH4 Primate-Specific GTTT-Repeat in Schizophrenia. <i>Genetic Testing and Molecular Biomarkers</i> , 2017, 21, 485-490.	0.7	15
4	A genetic variant in CAMKK2 gene is possibly associated with increased risk of bipolar disorder. <i>Journal of Neural Transmission</i> , 2016, 123, 323-328.	2.8	13
5	Incomplete penetrance of <i>CRX</i> gene for autosomal dominant form of cone-rod dystrophy. <i>Ophthalmic Genetics</i> , 2019, 40, 259-266.	1.2	11
6	c.376G>A mutation in WFS1 gene causes Wolfram syndrome without deafness. <i>European Journal of Medical Genetics</i> , 2016, 59, 65-69.	1.3	7
7	Association of Î²-Secretase Functional Polymorphism with Risk of Schizophrenia. <i>Genetic Testing and Molecular Biomarkers</i> , 2017, 21, 248-251.	0.7	3