Simin Rahimi-Aliabadi

List of Publications by Year
in descending order

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