

Monavvar Andarva

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/11645753/publications.pdf>

Version: 2024-02-01

9
papers

176
citations

1307594

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1474206

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times ranked

355
citing authors

#	ARTICLE	IF	CITATIONS
1	A Clinical and Molecular Genetic Study of 50 Families with Autosomal Recessive Parkinsonism Revealed Known and Novel Gene Mutations. <i>Molecular Neurobiology</i> , 2018, 55, 3477-3489.	4.0	67
2	RIT2 Polymorphisms: Is There a Differential Association?. <i>Molecular Neurobiology</i> , 2017, 54, 2234-2240.	4.0	31
3	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
4	The human RIT2 core promoter short tandem repeat predominant allele is species-specific in length: a selective advantage for human evolution?. <i>Molecular Genetics and Genomics</i> , 2017, 292, 611-617.	2.1	18
5	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
6	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
7	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
8	A novel c.240_241insGG mutation in NDP gene in a family with Norrie disease. <i>Australasian journal of optometry</i> , The, 2018, 101, 255-259.	1.3	4
9	Novel Mutations in Gene in Families with Gelatinous Drop-like Corneal Dystrophy (GDL). <i>International Journal of Molecular and Cellular Medicine</i> , 2017, 6, 204-211.	1.1	1