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 7 h-index	1474206 9 g-index
9 all docs	9 docs citations	9 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. Molecular Neurobiology, 2018, 55, 3477-3489.	4.0	67
2	A novel c.240_241insGG mutation in NDP gene in a family with Norrie disease. Australasian journal of optometry, The, 2018, 101, 255-259.	1.3	4
3	RIT2 Polymorphisms: Is There a Differential Association?. Molecular Neurobiology, 2017, 54, 2234-2240.	4.0	31
4	The human RIT2 core promoter short tandem repeat predominant allele is species-specific in length: a selective advantage for human evolution?. Molecular Genetics and Genomics, 2017, 292, 611-617.	2.1	18
5	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
6	Novel Mutations in Gene in Families with Gelatinous Drop-like Corneal Dystrophy (GDLD). International Journal of Molecular and Cellular Medicine, 2017, 6, 204-211.	1.1	1
7	c.376G>A mutation in WFS1 gene causes Wolfram syndrome without deafness. European Journal of Medical Genetics, 2016, 59, 65-69.	1.3	7
8	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
9	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