DÃ³ra Nagy

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5374606/publications.pdf

Version: 2024-02-01

9	101	1478505 6	1588992 8
papers	citations	h-index	g-index

10	10	10	180
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Comprehensive Genetic Analysis of a Hungarian Amyotrophic Lateral Sclerosis Cohort. Frontiers in Genetics, 2019, 10, 732.	2.3	31
2	Genetic analysis of the SOD1 and C9ORF72 genes in Hungarian patients with amyotrophic lateral sclerosis. Neurobiology of Aging, 2017, 53, 195.e1-195.e5.	3.1	17
3	Copy number variants detection by microarray and multiplex ligation-dependent probe amplification in congenital heart diseases. Journal of Biotechnology, 2019, 299, 86-95.	3.8	11
4	Skin wipe test: A simple, inexpensive, and fast approach in the diagnosis of cystic fibrosis. Pediatric Pulmonology, 2020, 55, 1653-1660.	2.0	11
5	Genotype-Phenotype Associations in Patients With Type-1, Type-2, and Atypical NF1 Microdeletions. Frontiers in Genetics, 2021, 12, 673025.	2.3	11
6	Angiogenin mutations in Hungarian patients with amyotrophic lateral sclerosis: Clinical, genetic, computational, and functional analyses. Brain and Behavior, 2019, 9, e01293.	2.2	10
7	Genotype-Phenotype Comparison in POGZ-Related Neurodevelopmental Disorders by Using Clinical Scoring. Genes, 2022, 13, 154.	2.4	6
8	Further delineation of the phenotype of PAK3-associated x-linked intellectual disability: Identification of a novel missense mutation and review of literature. European Journal of Medical Genetics, 2020, 63, 103800.	1.3	4
9	Systemic Screening for 22q11.2 Copy Number Variations in Hungarian Pediatric and Adult Patients With Congenital Heart Diseases Identified Rare Pathogenic Patterns in the Region. Frontiers in Genetics, 2021, 12, 635480.	2.3	0