Orsolya Nagy

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6615812/publications.pdf

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7	70	1937685 4	1872680 6
papers	citations	h-index	g-index

7	7	7	153
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	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
2	Case Report: Expressive Speech Disorder in a Family as a Hallmark of 7q31 Deletion Involving the FOXP2 Gene. Frontiers in Pediatrics, 2021, 9, 664548.	1.9	3
3	22q13 Microduplication Syndrome in Siblings with Mild Clinical Phenotype: Broadening the Clinical and Behavioral Spectrum. Molecular Syndromology, 2020, 11, 146-152.	0.8	4
4	The importance of the multiplex ligation-dependent probe amplification in the identification of a novel two-exon deletion of the NR5A1 gene in a patient with 46,XY differences of sex development. Molecular Biology Reports, 2019, 46, 5595-5601.	2.3	5
5	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
6	The role of microRNAs in congenital heart disease. Electronic Journal of the International Federation of Clinical Chemistry and Laboratory Medicine, 2019, 30, 165-178.	0.7	17
7	Elevated Factor VIII and von Willebrand Factor Levels Predict Unfavorable Outcome in Stroke Patients Treated with Intravenous Thrombolysis. Frontiers in Neurology, 2017, 8, 721.	2.4	30