

Chelsea Lowther

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/824485/publications.pdf>

Version: 2024-02-01

8
papers

461
citations

1307594

7
h-index

1720034

7
g-index

8
all docs

8
docs citations

8
times ranked

1317
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic contributors to risk of schizophrenia in the presence of a 22q11.2 deletion. <i>Molecular Psychiatry</i> , 2021, 26, 4496-4510.	7.9	87
2	Haploinsufficiency of vascular endothelial growth factor related signaling genes is associated with tetralogy of Fallot. <i>Genetics in Medicine</i> , 2019, 21, 1001-1007.	2.4	58
3	Molecular characterization of NRXN1 deletions from 19,263 clinical microarray cases identifies exons important for neurodevelopmental disease expression. <i>Genetics in Medicine</i> , 2017, 19, 53-61.	2.4	70
4	Genomic Disorders in Psychiatry—What Does the Clinician Need to Know?. <i>Current Psychiatry Reports</i> , 2017, 19, 82.	4.5	36
5	Rare Genome-Wide Copy Number Variation and Expression of Schizophrenia in 22q11.2 Deletion Syndrome. <i>American Journal of Psychiatry</i> , 2017, 174, 1054-1063.	7.2	77
6	Impact of IQ on the diagnostic yield of chromosomal microarray in a community sample of adults with schizophrenia. <i>Genome Medicine</i> , 2017, 9, 105.	8.2	30
7	MG-123—Exonic and intronic NRXN1 deletions: Novel genotype-phenotype correlations. <i>Journal of Medical Genetics</i> , 2015, 52, A9.1-A9.	3.2	0
8	Delineating the 15q13.3 microdeletion phenotype: a case series and comprehensive review of the literature. <i>Genetics in Medicine</i> , 2015, 17, 149-157.	2.4	103