

Monica Forzan

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8451499/publications.pdf>

Version: 2024-02-01

18
papers

2,651
citations

623574

14
h-index

839398

18
g-index

18
all docs

18
docs citations

18
times ranked

6615
citing authors

#	ARTICLE	IF	CITATIONS
1	Shared genetic risk between eating disorder and substance use-related phenotypes: Evidence from genome-wide association studies. <i>Addiction Biology</i> , 2021, 26, e12880.	1.4	28
2	Hybrid Minigene Assay: An Efficient Tool to Characterize mRNA Splicing Profiles of NF1 Variants. <i>Cancers</i> , 2021, 13, 999.	1.7	7
3	Setup and Validation of a Targeted Next-Generation Sequencing Approach for the Diagnosis of Lysosomal Storage Disorders. <i>Journal of Molecular Diagnostics</i> , 2020, 22, 488-502.	1.2	15
4	Genome-wide association study identifies eight risk loci and implicates metabo-psychiatric origins for anorexia nervosa. <i>Nature Genetics</i> , 2019, 51, 1207-1214.	9.4	641
5	Associations Between Attention-Deficit/Hyperactivity Disorder and Various Eating Disorders: A Swedish Nationwide Population Study Using Multiple Genetically Informative Approaches. <i>Biological Psychiatry</i> , 2019, 86, 577-586.	0.7	43
6	The Arg1038Gly missense variant in the <i>NF1</i> gene causes a mild phenotype without neurofibromas. <i>Molecular Genetics & Genomic Medicine</i> , 2019, 7, e616.	0.6	26
7	Analysis of shared heritability in common disorders of the brain. <i>Science</i> , 2018, 360, .	6.0	1,085
8	ATXN2 trinucleotide repeat length correlates with risk of ALS. <i>Neurobiology of Aging</i> , 2017, 51, 178.e1-178.e9.	1.5	86
9	Significant Locus and Metabolic Genetic Correlations Revealed in Genome-Wide Association Study of Anorexia Nervosa. <i>American Journal of Psychiatry</i> , 2017, 174, 850-858.	4.0	410
10	Catechol-O-methyltransferase (COMT) Val158Met Polymorphism and Eating Disorders: Data From a New Biobank and Meta-Analysis of Previously Published Studies. <i>European Eating Disorders Review</i> , 2017, 25, 524-532.	2.3	6
11	Functional connectivity correlates of response inhibition impairment in anorexia nervosa. <i>Psychiatry Research - Neuroimaging</i> , 2016, 247, 9-16.	0.9	40
12	Clinical and genetic correlates of decision making in anorexia nervosa. <i>Journal of Clinical and Experimental Neuropsychology</i> , 2016, 38, 327-337.	0.8	22
13	p.Arg1809Cys substitution in neurofibromin is associated with a distinctive NF1 phenotype without neurofibromas. <i>European Journal of Human Genetics</i> , 2015, 23, 1068-1071.	1.4	113
14	Neural signatures of the interaction between the 5-HTTLPR genotype and stressful life events in healthy women. <i>Psychiatry Research - Neuroimaging</i> , 2014, 223, 157-163.	0.9	14
15	Catechol-O-methyltransferase genotype modifies executive functioning and prefrontal functional connectivity in women with anorexia nervosa. <i>Journal of Psychiatry and Neuroscience</i> , 2013, 38, 241-248.	1.4	42
16	No Difference in 5-HTTLPR and Stin2 Polymorphisms Frequency Between Premature Ejaculation Patients and Controls. <i>Journal of Sexual Medicine</i> , 2012, 9, 1659-1668.	0.3	28
17	Is CFTR 621+3 A>G a cystic fibrosis causing mutation?. <i>Journal of Human Genetics</i> , 2010, 55, 23-26.	1.1	14
18	Association study of AMH and AMHR1I polymorphisms with unexplained infertility. <i>Fertility and Sterility</i> , 2010, 94, 1244-1248.	0.5	31