Monica Forzan

List of Publications by Year
in descending order

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

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