Catarina Correia

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Functional impact of global rare copy number variation in autism spectrum disorders. Nature, 2010, 466, 368-372.	13.7	1,803
2	A genome-wide scan for common alleles affecting risk for autism. Human Molecular Genetics, 2010, 19, 4072-4082.	1.4	538
3	Individual common variants exert weak effects on the risk for autism spectrum disorders. Human Molecular Genetics, 2012, 21, 4781-4792.	1.4	334
4	A novel approach of homozygous haplotype sharing identifies candidate genes in autism spectrum disorder. Human Genetics, 2012, 131, 565-579.	1.8	180
5	Evidence for epistasis between SLC6A4 and ITGB3 in autism etiology and in the determination of platelet serotonin levels. Human Genetics, 2007, 121, 243-256.	1.8	135
6	Brief Report: High Frequency of Biochemical Markers for Mitochondrial Dysfunction in Autism: No Association with the Mitochondrial Aspartate/Glutamate Carrier SLC25A12 Gene. Journal of Autism and Developmental Disorders, 2006, 36, 1137-1140.	1.7	96
7	Oxytocin receptor (OXTR) does not play a major role in the aetiology of autism: Genetic and molecular studies. Neuroscience Letters, 2010, 474, 163-167.	1.0	90
8	Autoantibody repertoires to brain tissue in autism nuclear families. Journal of Neuroimmunology, 2004, 152, 176-182.	1.1	77
9	MECP2 coding sequence and 3′UTR variation in 172 unrelated autistic patients. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 475-483.	1.1	53
10	Association of the α4 integrin subunit gene (<i>ITGA4</i>) with autism. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2009, 150B, 1147-1151.	1.1	20
11	Protein Interaction Networks Reveal Novel Autism Risk Genes within GWAS Statistical Noise. PLoS ONE, 2014, 9, e112399.	1.1	14
12	Pharmacogenetics of risperidone response and induced side effects. Personalized Medicine, 2007, 4, 271-293.	0.8	13