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List of Publications by Year in descending order

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687363 996975 4,533 14 13 15 citations g-index h-index papers 15 15 15 8646 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Functional impact of global rare copy number variation in autism spectrum disorders. Nature, 2010, 466, 368-372.	27.8	1,803
2	Convergence of Genes and Cellular Pathways Dysregulated in Autism Spectrum Disorders. American Journal of Human Genetics, 2014, 94, 677-694.	6.2	819
3	A genome-wide scan for common alleles affecting risk for autism. Human Molecular Genetics, 2010, 19, 4072-4082.	2.9	538
4	wishful thinking Encodes a BMP Type II Receptor that Regulates Synaptic Growth in Drosophila. Neuron, 2002, 33, 545-558.	8.1	469
5	Individual common variants exert weak effects on the risk for autism spectrum disorders. Human Molecular Genetics, 2012, 21, 4781-4792.	2.9	334
6	Geospatial Resolution of Human and Bacterial Diversity with City-Scale Metagenomics. Cell Systems, 2015, 1, 72-87.	6.2	241
7	Unexplained early onset epileptic encephalopathy: Exome screening and phenotype expansion. Epilepsia, 2016, 57, e12-7.	5.1	164
8	Association of Genetic Variants with Self-Assessed Color Categories in Brazilians. PLoS ONE, 2014, 9, e83926.	2.5	38
9	Modern Methods for Delineating Metagenomic Complexity. Cell Systems, 2015, 1, 6-7.	6.2	20
10	Personalized Cardioâ€Metabolic Responses to an Antiâ€Inflammatory Nutrition Intervention in Obese Adolescents: A Randomized Controlled Crossover Trial. Molecular Nutrition and Food Research, 2018, 62, e1701008.	3.3	20
11	Investigation of $15q11-q13$, $16p11.2$ and $22q13$ CNVs in Autism Spectrum Disorder Brazilian Individuals with and without Epilepsy. PLoS ONE, 2014, 9, e107705.	2.5	17
12	HGDP and HapMap Analysis by Ancestry Mapper Reveals Local and Global Population Relationships. PLoS ONE, 2012, 7, e49438.	2.5	13
13	Transcriptional control in embryonic Drosophila midline guidance assessed through a whole genome approach. BMC Neuroscience, 2007, 8, 59.	1.9	9
14	Ancestry of the Timorese: age-related macular degeneration associated genotype and allele sharing among human populations from throughout the world. Frontiers in Genetics, 2015, 6, 238.	2.3	9