MarÃ-a del Mar Abad-Grau

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1663022/publications.pdf

Version: 2024-02-01

26 papers 568 citations

933447 10 h-index 23 g-index

28 all docs

28 docs citations

times ranked

28

1114 citing authors

#	Article	IF	CITATIONS
1	A network model to predict the risk of death in sickle cell disease. Blood, 2007, 110, 2727-2735.	1.4	159
2	Multiple Sclerosis Risk Variant HLA-DRB1*1501 Associates with High Expression of DRB1 Gene in Different Human Populations. PLoS ONE, 2012, 7, e29819.	2.5	100
3	IL2RA/CD25 Gene Polymorphisms: Uneven Association with Multiple Sclerosis (MS) and Type 1 Diabetes (T1D). PLoS ONE, 2009, 4, e4137.	2.5	65
4	Identification of a functional variant in the <i>KIF5A-CYP27B1-METTL1-FAM119B</i> locus associated with multiple sclerosis. Journal of Medical Genetics, 2013, 50, 25-33.	3.2	59
5	A functional variant that affects exon-skipping and protein expression of (i>SP140 (i>as genetic mechanism predisposing to multiple sclerosis. Human Molecular Genetics, 2015, 24, 5619-5627.	2.9	43
6	A hierarchical and modular approach to the discovery of robust associations in genome-wide association studies from pooled DNA samples. BMC Genetics, 2008, 9, 6.	2.7	26
7	Operations strategy and flexibility: modeling with Bayesian classifiers. Industrial Management and Data Systems, 2006, 106, 460-484.	3.7	22
8	Robust Transmission/Disequilibrium Test for Incomplete Family Genotypes. Genetics, 2004, 168, 2329-2337.	2.9	18
9	Evolution and challenges in the design of computational systems for triage assistance. Journal of Biomedical Informatics, 2008, 41, 432-441.	4.3	15
10	Members 6B and 14 of the TNF receptor superfamily in multiple sclerosis predisposition. Genes and Immunity, 2011, 12, 145-148.	4.1	14
11	Building chromosome-wide LD maps. Bioinformatics, 2006, 22, 1933-1934.	4.1	8
12	Variant alleles of the mannose binding lectin 2 gene (<i>MBL2</i>) confer heterozygote advantage within Crohn's families. Scandinavian Journal of Gastroenterology, 2010, 45, 1129-1130.	1.5	6
13	Bayesian estimates of linkage disequilibrium. BMC Genetics, 2007, 8, 36.	2.7	5
14	Genome-wide association filtering using a highly locus-specific transmission/disequilibrium test. Human Genetics, 2010, 128, 325-344.	3.8	5
15	INCREASING POWER BY USING HAPLOTYPE SIMILARITY IN A MULTIMARKER TRANSMISSION/DISEQUILIBRIUM TEST. Journal of Bioinformatics and Computational Biology, 2013, 11, 1250014.	0.8	5
16	Sample Reproducibility of Genetic Association Using Different Multimarker TDTs in Genome-Wide Association Studies: Characterization and a New Approach. PLoS ONE, 2012, 7, e29613.	2.5	5
17	A comparison of genomic profiles of complex diseases under different models. BMC Medical Genomics, 2015, 9, 3.	1.5	4
18	GeneOnEarth: Fitting Genetic PC Plots on the Globe. IEEE/ACM Transactions on Computational Biology and Bioinformatics, 2013, 10, 1009-1016.	3.0	2

#	Article	IF	CITATIONS
19	Riskoweb: Web-Based Genetic Profiling to Complex Disease Using Genome-Wide SNP Markers. Advances in Intelligent and Soft Computing, $2011, 18$.	0.2	2
20	Bayesian Correction for SNP Ascertainment Bias. Lecture Notes in Computer Science, 2006, , 262-273.	1.3	1
21	HAPLOTYPE-BASED CLASSIFIERS TO PREDICT INDIVIDUAL SUSCEPTIBILITY TO COMPLEX DISEASES - An Example for Multiple Sclerosis. , 2012, , .		1
22	Discretization of Expression Quantitative Trait Loci in Association Analysis Between Genotypes and Expression Data [§] . Current Bioinformatics, 2015, 10, 144-164.	1.5	1
23	Fetal Hemoglobin (HbF) in Sickle Cell Anemia: Genome-Wide Association Studies Using Pooled DNA Samples Can Reveal Genetic Associations with HbF Concentration Blood, 2006, 108, 1221-1221.	1.4	O
24	3D VISUALIZATION OF HAPLOTYPE RISK MAPS., 2012,,.		0
25	Homogenizing Access to Highly Time-Consuming Biomedical Applications through a Web-Based Interface. Lecture Notes in Computer Science, 2012, , 33-42.	1.3	O
26	Multivariate Imputation of Genotype Data Using Short and Long Range Disequilibrium. , 2007, , 187-194.		0