

# MarÃ-a del Mar Abad-Grau

## List of Publications by Year in descending order

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26  
papers

568  
citations

933447

10  
h-index

642732

23  
g-index

28  
all docs

28  
docs citations

28  
times ranked

1114  
citing authors

#	ARTICLE	IF	CITATIONS
1	A comparison of genomic profiles of complex diseases under different models. BMC Medical Genomics, 2015, 9, 3.	1.5	4
2	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
3	Discretization of Expression Quantitative Trait Loci in Association Analysis Between Genotypes and Expression Data. Current Bioinformatics, 2015, 10, 144-164.	1.5	1
4	GeneOnEarth: Fitting Genetic PC Plots on the Globe. IEEE/ACM Transactions on Computational Biology and Bioinformatics, 2013, 10, 1009-1016.	3.0	2
5	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
6	INCREASING POWER BY USING HAPLOTYPE SIMILARITY IN A MULTIMARKER TRANSMISSION/DISEQUILIBRIUM TEST. Journal of Bioinformatics and Computational Biology, 2013, 11, 1250014.	0.8	5
7	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
8	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
9	HAPLOTYPE-BASED CLASSIFIERS TO PREDICT INDIVIDUAL SUSCEPTIBILITY TO COMPLEX DISEASES - An Example for Multiple Sclerosis. , 2012, , .		1
10	3D VISUALIZATION OF HAPLOTYPE RISK MAPS. , 2012, , .		0
11	Homogenizing Access to Highly Time-Consuming Biomedical Applications through a Web-Based Interface. Lecture Notes in Computer Science, 2012, , 33-42.	1.3	0
12	Members 6B and 14 of the TNF receptor superfamily in multiple sclerosis predisposition. Genes and Immunity, 2011, 12, 145-148.	4.1	14
13	Riskoweb: Web-Based Genetic Profiling to Complex Disease Using Genome-Wide SNP Markers. Advances in Intelligent and Soft Computing, 2011, , 1-8.	0.2	2
14	Genome-wide association filtering using a highly locus-specific transmission/disequilibrium test. Human Genetics, 2010, 128, 325-344.	3.8	5
15	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
16	IL2RA/CD25 Gene Polymorphisms: Uneven Association with Multiple Sclerosis (MS) and Type 1 Diabetes (T1D). PLoS ONE, 2009, 4, e4137.	2.5	65
17	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
18	Evolution and challenges in the design of computational systems for triage assistance. Journal of Biomedical Informatics, 2008, 41, 432-441.	4.3	15

#	ARTICLE	IF	CITATIONS
19	A network model to predict the risk of death in sickle cell disease. <i>Blood</i> , 2007, 110, 2727-2735.	1.4	159
20	Bayesian estimates of linkage disequilibrium. <i>BMC Genetics</i> , 2007, 8, 36.	2.7	5
21	Multivariate Imputation of Genotype Data Using Short and Long Range Disequilibrium. , 2007, , 187-194.		0
22	Operations strategy and flexibility: modeling with Bayesian classifiers. <i>Industrial Management and Data Systems</i> , 2006, 106, 460-484.	3.7	22
23	Building chromosome-wide LD maps. <i>Bioinformatics</i> , 2006, 22, 1933-1934.	4.1	8
24	Bayesian Correction for SNP Ascertainment Bias. <i>Lecture Notes in Computer Science</i> , 2006, , 262-273.	1.3	1
25	Fetal Hemoglobin (HbF) in Sickle Cell Anemia: Genome-Wide Association Studies Using Pooled DNA Samples Can Reveal Genetic Associations with HbF Concentration.. <i>Blood</i> , 2006, 108, 1221-1221.	1.4	0
26	Robust Transmission/Disequilibrium Test for Incomplete Family Genotypes. <i>Genetics</i> , 2004, 168, 2329-2337.	2.9	18