

Manuel Hermida-Prieto

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

24
papers

847
citations

13
h-index

28
g-index

28
ext. papers

972
ext. citations

3.1
avg. IF

2.91
L-index

#	Paper	IF	Citations
24	AGT haplotype in ITGA4 gene is related to antibody-mediated rejection in heart transplant patients. <i>PLoS ONE</i> , 2019 , 14, e0219345	3.7	
23	Polymorphisms in genes related to the complement system and antibody-mediated cardiac allograft rejection. <i>Journal of Heart and Lung Transplantation</i> , 2018 , 37, 477-485	5.8	7
22	Allogeneic Adipose-Derived Mesenchymal Stem Cells (Horse Allo 20) for the Treatment of Osteoarthritis-Associated Lameness in Horses: Characterization, Safety, and Efficacy of Intra-Articular Treatment. <i>Stem Cells and Development</i> , 2018 , 27, 1147-1160	4.4	15
21	Allogeneic adipose-derived mesenchymal stem cell therapy in dogs with refractory atopic dermatitis: clinical efficacy and safety. <i>Veterinary Record</i> , 2018 , 183, 654	0.9	23
20	Donor Polymorphisms in Genes Related to B-Cell Biology Associated With Antibody-Mediated Rejection After Heart Transplantation. <i>Circulation Journal</i> , 2018 , 82, 1351-1359	2.9	3
19	Macrophagic enhancement in optical coherence tomography imaging by means of superparamagnetic iron oxide nanoparticles. <i>Cardiology Journal</i> , 2017 , 24, 459-466	1.4	2
18	Analysis of variants in the HCN4 gene and in three single nucleotide polymorphisms of the CYP3A4 gene for association with ivabradine reduction in heart rate: A preliminary report. <i>Cardiology Journal</i> , 2016 , 23, 573-582	1.4	3
17	Somatic MYH7, MYBPC3, TPM1, TNNT2 and TNNI3 mutations in sporadic hypertrophic cardiomyopathy. <i>Circulation Journal</i> , 2013 , 77, 2358-65	2.9	13
16	Insights into genotype-phenotype correlation in hypertrophic cardiomyopathy. Findings from 18 Spanish families with a single mutation in MYBPC3. <i>Heart</i> , 2010 , 96, 1980-4	5.1	39
15	Trastornos graves de la conducci3n cardiaca e implante de marcapasos en pacientes con miocardiopat3a hipertr3fica. <i>Revista Espanola De Cardiologia</i> , 2010 , 63, 985-988	1.5	16
14	Screening mutations in myosin binding protein C3 gene in a cohort of patients with Hypertrophic Cardiomyopathy. <i>BMC Medical Genetics</i> , 2010 , 11, 67	2.1	20
13	A homozygous MYBPC3 gene mutation associated with a severe phenotype and a high risk of sudden death in a family with hypertrophic cardiomyopathy. <i>Revista Espanola De Cardiologia (English Ed)</i> , 2009 , 62, 572-5	0.7	7
12	Mutaci3n en homocigosis en el gen MYBPC3 asociada a fenotipos severos y alto riesgo de muerte s3bita en una familia con miocardiopat3a hipertr3fica. <i>Revista Espanola De Cardiologia</i> , 2009 , 62, 572-575	1.5	12
11	Miocardiopat3a hipertr3fica. Estudio del gen de la troponina T en 127 familias espa3olas. <i>Revista Espanola De Cardiologia</i> , 2009 , 62, 1473-1477	1.5	13
10	Sudden death in a patient with lamin A/C gene mutation and near normal left ventricular systolic function. <i>International Journal of Cardiology</i> , 2008 , 126, 136-7	3.2	8
9	Identification of a Cardiac Allograft Rejection Marker Using Microarray Gene Expression Analysis in Lymphocytes From Heart Transplant Patients. <i>Revista Espanola De Cardiologia (English Ed)</i> , 2007 , 60, 217-218	0.7	
8	Prevalence of fabry disease in a cohort of 508 unrelated patients with hypertrophic cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2007 , 50, 2399-403	15.1	206

7	Avances en miocardiopatía dilatada idiopática: del genotipo al fenotipo clínico. <i>Revista Española De Cardiología Suplementos</i> , 2007 , 7, 2F-13F	0.2	
6	Mutational screening of phospholamban gene in hypertrophic and idiopathic dilated cardiomyopathy and functional study of the PLN -42 C>G mutation. <i>European Journal of Heart Failure</i> , 2007 , 9, 37-43	12.3	35
5	Mutation in the alpha-cardiac actin gene associated with apical hypertrophic cardiomyopathy, left ventricular non-compaction, and septal defects. <i>European Heart Journal</i> , 2007 , 28, 1953-61	9.5	202
4	Lack of cross-species transmission of porcine endogenous retrovirus (PERV) to transplant recipients and abattoir workers in contact with pigs. <i>Transplantation</i> , 2007 , 84, 548-50	1.8	18
3	Standard mutation nomenclature in hypertrophic cardiomyopathy: an urgent need. <i>Journal of the American College of Cardiology</i> , 2005 , 46, 380-1; author reply 381-2	15.1	
2	Lack of cross-species transmission of porcine endogenous retrovirus in pig-to-baboon xenotransplantation with sustained depletion of anti-alphagal antibodies. <i>Transplantation</i> , 2005 , 79, 777-82	1.8	32
1	Familial dilated cardiomyopathy and isolated left ventricular noncompaction associated with lamin A/C gene mutations. <i>American Journal of Cardiology</i> , 2004 , 94, 50-4	3	148