Esther Zorio

List of Publications by Year
in descending order

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30 papers	863 citations	759233 12 h-index	501196 28 g-index
34 all docs	34 docs citations	34 times ranked	1541 citing authors

#	Article	IF	Citations
1	The EP300/TP53 pathway, a suppressor of the Hippo and canonical WNT pathways, is activated in human hearts with arrhythmogenic cardiomyopathy in the absence of overt heart failure. Cardiovascular Research, 2022, 118, 1466-1478.	3.8	20
2	An International Multicenter Cohort Study on Î²-Blockers for the Treatment of Symptomatic Children With Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation, 2022, 145, 333-344.	1.6	28
3	Importance of genotype for risk stratification in arrhythmogenic right ventricular cardiomyopathy using the 2019 ARVC risk calculator. European Heart Journal, 2022, 43, 3053-3067.	2.2	41
4	Impact of SARSâ€Covâ€2 infection in patients with hypertrophic cardiomyopathy: results of an international multicentre registry. ESC Heart Failure, 2022, 9, 2189-2198.	3.1	6
5	Cardiac phenotype in glycogen storage disease type XV: a rare cardiomyopathy to bear in mind. Revista Espanola De Cardiologia (English Ed), 2021, 74, 99-101.	0.6	1
6	Fenotipo cardiolÃ³gico de la glucogenosis tipo XV: una miocardiopatÃa muy infrecuente a tener en cuenta. Revista Espanola De Cardiologia, 2021, 74, 99-101.	1.2	0
7	Impaired Binding to Junctophilin-2 and Nanostructural Alteration in CPVT Mutation. Circulation Research, 2021, 129, e35-e52.	4.5	19
8	Classification model based on strain measurements to identify patients with arrhythmogenic cardiomyopathy with left ventricular involvement. Computer Methods and Programs in Biomedicine, 2020, 188, 105296.	4.7	1
9	RNA sequencing-based transcriptome profiling of cardiac tissue implicates novel putative disease mechanisms in FLNC-associated arrhythmogenic cardiomyopathy. International Journal of Cardiology, 2020, 302, 124-130.	1.7	23
10	Clinical Profile of Cardiac Involvement in Danon Disease. Circulation Genomic and Precision Medicine, 2020, 13, e003117.	3.6	29
11	Ranolazine as an Alternative Therapy to Flecainide for SCN5A V411M Long QT Syndrome Type 3 Patients. Frontiers in Pharmacology, 2020, 11, 580481.	3.5	4
12	Arrhythmogenic cardiomyopathy with left ventricular involvement versus ischemic heart disease: lessons learned from the family study and the reviewed autopsy of a young male. Forensic Sciences Research, 2019, 4, 274-279.	1.6	7
13	Left ventricular myocardial dysfunction in arrhythmogenic cardiomyopathy with left ventricular involvement: A door to improving diagnosis. International Journal of Cardiology, 2019, 274, 237-244.	1.7	7
14	Thickness and an Altered miRNA Expression in the Epicardial Adipose Tissue Is Associated With Coronary Heart Disease in Sudden Death Victims. Revista Espanola De Cardiologia (English Ed), 2019, 72, 30-39.	0.6	11
15	The Cardiomyopathy Registry of the EURObservational Research Programme of the European Society of Cardiology: baseline data and contemporary management of adult patients with cardiomyopathies. European Heart Journal, 2018, 39, 1784-1793.	2.2	94
16	RyR2R420Q catecholaminergic polymorphic ventricular tachycardia mutation induces bradycardia by disturbing the coupled clock pacemaker mechanism. JCI Insight, 2017, 2, .	5.0	24
17	Deregulated hepatic micro<scp>RNA</scp>s underlie the association between nonâ€alcoholic fatty liver disease and coronary artery disease. Liver International, 2016, 36, 1221-1229.	3.9	39
18	Truncating FLNC Mutations Are Associated With High-Risk Dilated and Arrhythmogenic Cardiomyopathies. Journal of the American College of Cardiology, 2016, 68, 2440-2451.	2.8	340

#	Article	IF	Citations
19	Sensitivity and Negative Predictive Value of Treadmill Exercise Stress Testing for the Diagnosis of Catecholaminergic Polymorphic Ventricular Tachycardia. Response. Revista Espanola De Cardiologia (English Ed), 2015, 68, 546-547.	0.6	0
20	A New Mutation in the Ryanodine Receptor 2 Gene (RYR2 C2277R) as a Cause Catecholaminergic Polymorphic Ventricular Tachycardia. Revista Espanola De Cardiologia (English Ed), 2015, 68, 71-73.	0.6	7
21	Una nueva mutaciÃ³n en el gen del receptor de la rianodina (RyR2 C2277R) como causa de taquicardia ventricular polimÃ³rfica catecolaminÃ©rgica. Revista Espanola De Cardiologia, 2015, 68, 71-73.	1.2	8
22	Non-ventricular, Clinical, and Functional Features of the RyR2R420Q Mutation Causing Catecholaminergic Polymorphic Ventricular Tachycardia. Revista Espanola De Cardiologia (English Ed) Tj ETQq0 0 0 rgBT /Overlock 10 Tf	0.6	10
23	El trasfondo genÃ©tico de la hipertrabeculaciÃ³n/miocardiopatÃa no compactada ventricular izquierda sigue sin estar claro. Respuesta. Revista Espanola De Cardiologia, 2015, 68, 167-168.	1.2	0
24	The Genetic Background of Left Ventricular Hypertrabeculation/Noncompaction Remains Vague. Response. Revista Espanola De Cardiologia (English Ed), 2015, 68, 167-168.	0.6	0
25	Sensibilidad y valor predictivo negativo de la ergometrÃa para el diagnÃ³stico de la taquicardia ventricular polimÃ³rfica catecolaminÃ©rgica. Respuesta. Revista Espanola De Cardiologia, 2015, 68, 546-547.	1.2	0
26	Familial Left Ventricular Noncompaction Associated With a Novel Mutation in the Alpha-cardiac Actin Gene. Revista Espanola De Cardiologia (English Ed), 2014, 67, 857-859.	0.6	4
27	MiocardiopatÃa no compactada familiar asociada con una mutaciÃ³n nueva en el gen de la alfa-actina cardiaca. Revista Espanola De Cardiologia, 2014, 67, 857-859.	1.2	7
28	Paradoxical Effect of Increased Diastolic Ca ²⁺ Release and Decreased Sinoatrial Node Activity in a Mouse Model of Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation, 2012, 126, 392-401.	1.6	77
29	Flecainide, a Therapeutic Option in a Patient With Long QT Syndrome Type 3 Caused by the Heterozygous V411M Mutation in the SCN5A Gene. Revista Espanola De Cardiologia (English Ed), 2012, 65, 1058-1059.	0.6	2
30	Left Dominant Arrhythmogenic Cardiomyopathy Caused by a Novel Nonsense Mutation in Desmoplakin. Revista Espanola De Cardiologia (English Ed), 2011, 64, 530-534.	0.6	8