Fernando Dominguez

List of Publications by Year
in descending order

44 papers	2,360 citations	304743 22 h-index	243625 44 g-index
48 all docs	48 docs citations	48 times ranked	4281 citing authors

#	Article	IF	Citations
1	Combination of late gadolinium enhancement and genotype improves prediction of prognosis in nonâ€ischaemic dilated cardiomyopathy. European Journal of Heart Failure, 2022, 24, 1183-1196.	7.1	13
2	Endomyocardial biopsy-confirmed myocarditis and inflammatory cardiomyopathy: clinical profile and prognosis. Revista Espanola De Cardiologia (English Ed), 2022, , .	0.6	1
3	Systemic embolism in amyloid transthyretin cardiomyopathy. European Journal of Heart Failure, 2022, 24, 1387-1396.	7.1	23
4	Predicting pacemaker implantation in cardiac amyloidosis: let's start with an <scp>ECG</scp>. European Journal of Heart Failure, 2022, 24, 1237-1238.	7.1	0
5	Clinical profile and outcome of cardiac amyloidosis in a Spanish referral center. Revista Espanola De Cardiologia (English Ed), 2021, 74, 149-158.	0.6	10
6	Phenotypic clustering of dilated cardiomyopathy patients highlights important pathophysiological differences. European Heart Journal, 2021, 42, 162-174.	2.2	62
7	Transthyretin amyloid cardiomyopathy. Medicina ClÃnica (English Edition), 2021, 156, 126-134.	0.2	6
8	Perfil clÃnico y evoluciÃ³n de la amiloidosis cardiaca en un centro espaÃ±ol de referencia. Revista Espanola De Cardiologia, 2021, 74, 149-158.	1.2	33
9	Amiloidosis cardÃaca por transtiretina. Medicina ClÃnica, 2021, 156, 126-134.	0.6	22
10	Prevalence and clinical outcomes of dystrophinâ€associated dilated cardiomyopathy without severe skeletal myopathy. European Journal of Heart Failure, 2021, 23, 1276-1286.	7.1	14
11	Early Preventive Treatment With Enalapril Improves Cardiac Function and Delays Mortality in Mice With Arrhythmogenic Right Ventricular Cardiomyopathy Type 5. Circulation: Heart Failure, 2021, 14, e007616.	3.9	3
12	Association of Genetic Variants With Outcomes in Patients With Nonischemic Dilated Cardiomyopathy. Journal of the American College of Cardiology, 2021, 78, 1682-1699.	2.8	55
13	Saw-Tooth Cardiomyopathy. JACC: Case Reports, 2020, 2, 1210-1211.	0.6	3
14	Peripheral microRNA panels to guide the diagnosis of familial cardiomyopathy. Translational Research, 2020, 218, 1-15.	5.0	14
15	Clinical characteristics and determinants of the phenotype in TMEM43 arrhythmogenic right ventricular cardiomyopathy type 5. Heart Rhythm, 2020, 17, 945-954.	0.7	28
16	Severe Cardiac Dysfunction and Death Caused by Arrhythmogenic Right Ventricular Cardiomyopathy Type 5 Are Improved by Inhibition of Glycogen Synthase Kinase-3Î². Circulation, 2019, 140, 1188-1204.	1.6	62
17	POT1 and Damage Response Malfunction Trigger Acquisition of Somatic Activating Mutations in the VEGF Pathway in Cardiac Angiosarcomas. Journal of the American Heart Association, 2019, 8, e012875.	3.7	8
18	Prevalence of Cardiac Amyloidosis in Patients with Carpal Tunnel Syndrome. Journal of Cardiovascular Translational Research, 2019, 12, 507-513.	2.4	33

#	Article	IF	Citations
19	Prevalence of cardiac amyloidosis among elderly patients with systolic heart failure or conduction disorders. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 156-163.	3.0	33
20	Genetic Variants Associated With Cancer Therapyâ€“Induced Cardiomyopathy. Circulation, 2019, 140, 31-41.	1.6	195
21	Effect of the type of surgical indication on mortality in patients with infective endocarditis who are rejected for surgical intervention. International Journal of Cardiology, 2019, 282, 24-30.	1.7	27
22	Association of Sleep Duration and Quality With Subclinical Atherosclerosis. Journal of the American College of Cardiology, 2019, 73, 134-144.	2.8	145
23	Gentamicin may have no effect on mortality of staphylococcal prosthetic valve endocarditis. Journal of Infection and Chemotherapy, 2018, 24, 555-562.	1.7	21
24	Prognostic Impact and Predictors of Ejection Fraction Recovery in Patients With Alcoholic Cardiomyopathy. Revista Espanola De Cardiologia (English Ed), 2018, 71, 612-619.	0.6	9
25	Role of echocardiography in the diagnosis and management of hypertrophic cardiomyopathy. Heart, 2018, 104, 261-273.	2.9	10
26	International External Validation Study of the 2014 European Society of Cardiology Guidelines on Sudden Cardiac Death Prevention in Hypertrophic Cardiomyopathy (EVIDENCE-HCM). Circulation, 2018, 137, 1015-1023.	1.6	149
27	Follow-up and prognosis of HCM. Global Cardiology Science & Practice, 2018, 2018, 33.	0.4	4
28	Dilated Cardiomyopathy DueÂtoÂBLC2-Associated AthanogeneÂ3Â(BAG3)ÂMutations. Journal of the American College of Cardiology, 2018, 72, 2471-2481.	2.8	93
29	Infective Endocarditis in Patients With Bicuspid Aortic Valve or MitralÂValveÂProlapse. Journal of the American College of Cardiology, 2018, 71, 2731-2740.	2.8	65
30	Usefulness of Genetic Testing in Hypertrophic Cardiomyopathy: an Analysis Using Real-World Data. Journal of Cardiovascular Translational Research, 2017, 10, 35-46.	2.4	10
31	Clinical characteristics of wild-type transthyretin cardiac amyloidosis: disproving myths. European Heart Journal, 2017, 38, 1895-1904.	2.2	258
32	Genetically Confirmed Familial Hypercholesterolemia in Patients With Acute Coronary Syndrome. Journal of the American College of Cardiology, 2017, 70, 1732-1740.	2.8	111
33	The wide spectrum of POT1 gene variants correlates with multiple cancer types. European Journal of Human Genetics, 2017, 25, 1278-1281.	2.8	66
34	Direct oral anticoagulants in patients with hypertrophic cardiomyopathy and atrial fibrillation. International Journal of Cardiology, 2017, 248, 232-238.	1.7	41
35	Animal models of arrhythmogenic right ventricular cardiomyopathy: what have we learned and where do we go? Insight for therapeutics. Basic Research in Cardiology, 2017, 112, 50.	5.9	20
36	The Coronary Circulation in Cardiomyopathies and Cardiac Allografts. , 2017, , 119-135.		0

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37	Reversible transition from a hypertrophic to a dilated cardiomyopathy. ESC Heart Failure, 2016, 3, 138-142.	3.1	2
38	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
39	Infective endocarditis in hypertrophic cardiomyopathy. Medicine (United States), 2016, 95, e4008.	1.0	15
40	Natural History and Prognostic Factors in Alcoholic Cardiomyopathy. JACC: Heart Failure, 2015, 3, 78-86.	4.1	78
41	Malignant ventricular arrhythmias in alcoholic cardiomyopathy. International Journal of Cardiology, 2015, 199, 99-105.	1.7	25
42	Cardiac manifestations of Sneddon's syndrome. International Journal of Cardiology, 2015, 190, 275-276.	1.7	3
43	A mutation in the POT1 gene is responsible for cardiac angiosarcoma in TP53-negative Liâ€“Fraumeni-like families. Nature Communications, 2015, 6, 8383.	12.8	124
44	Erysipelas and Acute Myocarditis: An Unusual Combination. Canadian Journal of Cardiology, 2013, 29, 1138.e3-1138.e5.	1.7	3