

Fernando Dominguez

List of Publications by Year in descending order

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Version: 2024-02-01

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	Truncating FLNC Mutations Are Associated With High-Risk Dilated and Arrhythmogenic Cardiomyopathies. <i>Journal of the American College of Cardiology</i> , 2016, 68, 2440-2451.	2.8	340
2	Clinical characteristics of wild-type transthyretin cardiac amyloidosis: disproving myths. <i>European Heart Journal</i> , 2017, 38, 1895-1904.	2.2	258
3	Genetic Variants Associated With Cancer Therapy-Induced Cardiomyopathy. <i>Circulation</i> , 2019, 140, 31-41.	1.6	195
4	International External Validation Study of the 2014 European Society of Cardiology Guidelines on Sudden Cardiac Death Prevention in Hypertrophic Cardiomyopathy (EVIDENCE-HCM). <i>Circulation</i> , 2018, 137, 1015-1023.	1.6	149
5	Association of Sleep Duration and Quality With Subclinical Atherosclerosis. <i>Journal of the American College of Cardiology</i> , 2019, 73, 134-144.	2.8	145
6	A mutation in the POT1 gene is responsible for cardiac angiosarcoma in TP53-negative Li-Fraumeni-like families. <i>Nature Communications</i> , 2015, 6, 8383.	12.8	124
7	Genetically Confirmed Familial Hypercholesterolemia in Patients With Acute Coronary Syndrome. <i>Journal of the American College of Cardiology</i> , 2017, 70, 1732-1740.	2.8	111
8	Dilated Cardiomyopathy Due to BCL2-Associated Athanogene (BAG3) Mutations. <i>Journal of the American College of Cardiology</i> , 2018, 72, 2471-2481.	2.8	93
9	Natural History and Prognostic Factors in Alcoholic Cardiomyopathy. <i>JACC: Heart Failure</i> , 2015, 3, 78-86.	4.1	78
10	The wide spectrum of POT1 gene variants correlates with multiple cancer types. <i>European Journal of Human Genetics</i> , 2017, 25, 1278-1281.	2.8	66
11	Infective Endocarditis in Patients With Bicuspid Aortic Valve or Mitral Valve Prolapse. <i>Journal of the American College of Cardiology</i> , 2018, 71, 2731-2740.	2.8	65
12	Severe Cardiac Dysfunction and Death Caused by Arrhythmogenic Right Ventricular Cardiomyopathy Type 5 Are Improved by Inhibition of Glycogen Synthase Kinase-3 β . <i>Circulation</i> , 2019, 140, 1188-1204.	1.6	62
13	Phenotypic clustering of dilated cardiomyopathy patients highlights important pathophysiological differences. <i>European Heart Journal</i> , 2021, 42, 162-174.	2.2	62
14	Association of Genetic Variants With Outcomes in Patients With Nonischemic Dilated Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2021, 78, 1682-1699.	2.8	55
15	Direct oral anticoagulants in patients with hypertrophic cardiomyopathy and atrial fibrillation. <i>International Journal of Cardiology</i> , 2017, 248, 232-238.	1.7	41
16	Prevalence of Cardiac Amyloidosis in Patients with Carpal Tunnel Syndrome. <i>Journal of Cardiovascular Translational Research</i> , 2019, 12, 507-513.	2.4	33
17	Prevalence of cardiac amyloidosis among elderly patients with systolic heart failure or conduction disorders. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2019, 26, 156-163.	3.0	33
18	Perfil clínico y evolución de la amiloidosis cardiaca en un centro español de referencia. <i>Revista Española De Cardiología</i> , 2021, 74, 149-158.	1.2	33

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19	Clinical characteristics and determinants of the phenotype in TMEM43 arrhythmogenic right ventricular cardiomyopathy type 5. <i>Heart Rhythm</i> , 2020, 17, 945-954.	0.7	28
20	Effect of the type of surgical indication on mortality in patients with infective endocarditis who are rejected for surgical intervention. <i>International Journal of Cardiology</i> , 2019, 282, 24-30.	1.7	27
21	Malignant ventricular arrhythmias in alcoholic cardiomyopathy. <i>International Journal of Cardiology</i> , 2015, 199, 99-105.	1.7	25
22	Systemic embolism in amyloid transthyretin cardiomyopathy. <i>European Journal of Heart Failure</i> , 2022, 24, 1387-1396.	7.1	23
23	Amiloidosis card�aca por transtiretina. <i>Medicina Cl�nica</i> , 2021, 156, 126-134.	0.6	22
24	Gentamicin may have no effect on mortality of staphylococcal prosthetic valve endocarditis. <i>Journal of Infection and Chemotherapy</i> , 2018, 24, 555-562.	1.7	21
25	Animal models of arrhythmogenic right ventricular cardiomyopathy: what have we learned and where do we go? Insight for therapeutics. <i>Basic Research in Cardiology</i> , 2017, 112, 50.	5.9	20
26	Infective endocarditis in hypertrophic cardiomyopathy. <i>Medicine (United States)</i> , 2016, 95, e4008.	1.0	15
27	Peripheral microRNA panels to guide the diagnosis of familial cardiomyopathy. <i>Translational Research</i> , 2020, 218, 1-15.	5.0	14
28	Prevalence and clinical outcomes of dystrophin�associated dilated cardiomyopathy without severe skeletal myopathy. <i>European Journal of Heart Failure</i> , 2021, 23, 1276-1286.	7.1	14
29	Combination of late gadolinium enhancement and genotype improves prediction of prognosis in non�ischaemic dilated cardiomyopathy. <i>European Journal of Heart Failure</i> , 2022, 24, 1183-1196.	7.1	13
30	Usefulness of Genetic Testing in Hypertrophic Cardiomyopathy: an Analysis Using Real-World Data. <i>Journal of Cardiovascular Translational Research</i> , 2017, 10, 35-46.	2.4	10
31	Role of echocardiography in the diagnosis and management of hypertrophic cardiomyopathy. <i>Heart</i> , 2018, 104, 261-273.	2.9	10
32	Clinical profile and outcome of cardiac amyloidosis in a Spanish referral center. <i>Revista Espanola De Cardiologia (English Ed)</i> , 2021, 74, 149-158.	0.6	10
33	Prognostic Impact and Predictors of Ejection Fraction Recovery in Patients With Alcoholic Cardiomyopathy. <i>Revista Espanola De Cardiologia (English Ed)</i> , 2018, 71, 612-619.	0.6	9
34	POT1 and Damage Response Malfunction Trigger Acquisition of Somatic Activating Mutations in the VEGF Pathway in Cardiac Angiosarcomas. <i>Journal of the American Heart Association</i> , 2019, 8, e012875.	3.7	8
35	Transthyretin amyloid cardiomyopathy. <i>Medicina Cl�nica (English Edition)</i> , 2021, 156, 126-134.	0.2	6
36	Follow-up and prognosis of HCM. <i>Global Cardiology Science & Practice</i> , 2018, 2018, 33.	0.4	4

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37	Erysipelas and Acute Myocarditis: An Unusual Combination. Canadian Journal of Cardiology, 2013, 29, 1138.e3-1138.e5.	1.7	3
38	Cardiac manifestations of Sneddon's syndrome. International Journal of Cardiology, 2015, 190, 275-276.	1.7	3
39	Saw-Tooth Cardiomyopathy. JACC: Case Reports, 2020, 2, 1210-1211.	0.6	3
40	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
41	Reversible transition from a hypertrophic to a dilated cardiomyopathy. ESC Heart Failure, 2016, 3, 138-142.	3.1	2
42	Endomyocardial biopsy-confirmed myocarditis and inflammatory cardiomyopathy: clinical profile and prognosis. Revista Espanola De Cardiologia (English Ed), 2022, , .	0.6	1
43	The Coronary Circulation in Cardiomyopathies and Cardiac Allografts. , 2017, , 119-135.		0
44	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