Ripoll-Vera, Tomas

List of Publications by Year
in descending order

43 papers	729 citations	777949 13 h-index	651938 25 g-index
48 all docs	48 docs citations	48 times ranked	1364 citing authors

#	Article	IF	Citations
1	Genotype-Phenotype Correlation in Hypertrophic Cardiomyopathy: New Variant p.Arg652Lys in MYH7. Genes, 2022, 13, 320.	1.0	2
2	Accessibility to Occupational Therapy Services for Hereditary Transthyretin Amyloidosis. International Journal of Environmental Research and Public Health, 2022, 19, 4464.	1.2	4
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.	1.0	41
4	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.	2.9	13
5	New Variant in Placophilin-2 Gene Causing Arrhythmogenic Myocardiopathy. Genes, 2022, 13, 782.	1.0	1
6	Inflammatory and Oxidative Stress Markers Related to Adherence to the Mediterranean Diet in Patients with Metabolic Syndrome. Antioxidants, 2022, 11, 901.	2.2	18
7	Impact of SARSâ€Covâ€2 infection in patients with hypertrophic cardiomyopathy: results of an international multicentre registry. ESC Heart Failure, 2022, 9, 2189-2198.	1.4	6
8	Muerte sÃºbita de jÃ³venes: rendimiento diagnÃ³stico de un programa autonÃ³mico de autopsia molecular con secuenciaciÃ³n masiva. Revista Espanola De Cardiologia, 2021, 74, 402-413.	0.6	4
9	SelecciÃ³n de lo mejor del aÃ±o 2020 en cardiopatÃas familiares y genÃ©tica cardiovascular. REC: CardioClinics, 2021, 56, 9-14.	0.1	0
10	AsociaciÃ³n entre estenosis aÃ³rtica y amiloidosis hereditaria por transtiretina. Revista Espanola De Cardiologia, 2021, 74, 185-187.	0.6	1
11	Predictores de riesgo en una cohorte espaÃ±ola con cardiolaminopatÃas. Registro REDLAMINA. Revista Espanola De Cardiologia, 2021, 74, 216-224.	0.6	19
12	Diagnostic Yield of Genetic Testing in Sudden Cardiac Death with Autopsy Findings of Uncertain Significance. Journal of Clinical Medicine, 2021, 10, 1806.	1.0	2
13	Val50Met hereditary transthyretin amyloidosis: not just a medical problem, but a psychosocial burden. Orphanet Journal of Rare Diseases, 2021, 16, 266.	1.2	10
14	Association of Left Ventricular Systolic Dysfunction Among Carriers of Truncating Variants in Filamin C With Frequent Ventricular Arrhythmia and End-stage Heart Failure. JAMA Cardiology, 2021, 6, 891.	3.0	36
15	Association of Genetic Variants With Outcomes in Patients With Nonischemic Dilated Cardiomyopathy. Journal of the American College of Cardiology, 2021, 78, 1682-1699.	1.2	55
16	Familial Dilated Cardiomyopathy and Sudden Cardiac Arrest: New Association with a SCN5A Mutation. Genes, 2021, 12, 1889.	1.0	4
17	Relationship between olive oil consumption and ankle-brachial pressure index in a population at high cardiovascular risk. Atherosclerosis, 2020, 314, 48-57.	0.4	6
18	Anticipation on age at onset in kindreds with hereditary ATTRV30M amyloidosis from the Majorcan cluster. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2020, 27, 254-258.	1.4	6

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19	ESC EORP Cardiomyopathy Registry: realâ€life practice of genetic counselling and testing in adult cardiomyopathy patients. ESC Heart Failure, 2020, 7, 3013-3021.	1.4	19
20	Multidisciplinary approach in the management of hATTR. European Journal of Clinical Investigation, 2020, 50, e13296.	1.7	4
21	Prognosis of Patients With Severe Aortic Stenosis After the Decision to Perform an Intervention. Revista Espanola De Cardiologia (English Ed), 2019, 72, 392-397.	0.4	4
22	Negative screening of Fabry disease in patients with conduction disorders requiring a pacemaker. Orphanet Journal of Rare Diseases, 2019, 14, 170.	1.2	6
23	Cardiac involvement in a large cohort of patients with Val30Met transthyretin amyloidosis from Majorca focus. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 15-16.	1.4	1
24	Cardiac involvement after liver transplantation in patients with Val30Met transthyretin amyloidosis from Majorca focus. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 18-19.	1.4	2
25	Amyloidotic breast nodule in hereditary transthyretin amyloidosis (hATTR): a case report. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 59-60.	1.4	1
26	Diagnostic yield of the implantable loop recorder in octogenarians. European Geriatric Medicine, 2019, 10, 129-133.	1.2	4
27	Cardiac Involvement in a Patient Cohort With Val30Met Mutation Transthyretin Amyloidosis. Revista Espanola De Cardiologia (English Ed), 2019, 72, 92-94.	0.4	0
28	DaÃ±o cardiaco en una cohorte de pacientes con amiloidosis por transtiretina por la mutaciÃ³n Val30Met. Revista Espanola De Cardiologia, 2019, 72, 92-94.	0.6	9
29	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.	1.0	94
30	Formin Homology 2 Domain Containing 3 (FHOD3) Is a Genetic Basis for Hypertrophic Cardiomyopathy. Journal of the American College of Cardiology, 2018, 72, 2457-2467.	1.2	59
31	Dilated Cardiomyopathy DueÂtoÂBLC2-Associated AthanogeneÂ3Â(BAG3)ÂMutations. Journal of the American College of Cardiology, 2018, 72, 2471-2481.	1.2	93
32	SÃndrome de tako-tsubo con presentaciÃ³n electrocardiogrÃ¡fica atÃpica. Cardiocore, 2017, 52, 35-36.	0.0	0
33	DiagnÃ³stico precoz en pacientes con polineuropatÃa amiloidÃ³tica familiar asociada a transtirretina. Estudio comparativo. Medicina ClÃnica, 2017, 148, 63-66.	0.3	2
34	Usefulness of Genetic Testing in Hypertrophic Cardiomyopathy: an Analysis Using Real-World Data. Journal of Cardiovascular Translational Research, 2017, 10, 35-46.	1.1	10
35	El volumen extracelular no se asocia a arritmias malignas en miocardiopatÃa hipertrÃ³fica de alto riesgo. Revista Espanola De Cardiologia, 2017, 70, 933-940.	0.6	2
36	Direct oral anticoagulants in patients with hypertrophic cardiomyopathy and atrial fibrillation. International Journal of Cardiology, 2017, 248, 232-238.	0.8	41

#	Article	IF	Citations
37	Comparison of 1-Year Outcome in Patients With Severe Aorta Stenosis Treated Conservatively or by Aortic Valve Replacement or by Percutaneous Transcatheter Aortic Valve Implantation (Data from a) Tj ETQq1 1 0.784314 rgBT /Overlo	0.4	27
38	Transthyretin familial amyloid polyneuropathy (TTRâ€FAP) in Mallorca: a comparison between lateâ€and earlyâ€onset disease. Journal of the Peripheral Nervous System, 2016, 21, 352-356.	1.4	12
39	Plan of Action for Inherited Cardiovascular Diseases: Synthesis of Recommendations and Action Algorithms. Revista Espanola De Cardiologia (English Ed), 2016, 69, 300-309.	0.4	14
40	Clinical and Prognostic Profiles of Cardiomyopathies Caused by Mutations in the Troponin T Gene. Revista Espanola De Cardiologia (English Ed), 2016, 69, 149-158.	0.4	17
41	Phenotypic Patterns of Cardiomyopathy Caused by Mutations in the Desmin Gene. A Clinical and Genetic Study in Two Inherited Heart Disease Units. Revista Espanola De Cardiologia (English Ed), 2015, 68, 1027-1029.	0.4	6
42	PatrÃ³n fenotÃpico de las miocardiopatÃas por mutaciones en el gen de la desmina. Estudio clÃnico y genÃ©tico en dos unidades de cardiopatÃas familiares. Revista Espanola De Cardiologia, 2015, 68, 1027-1029.	0.6	8
43	Frequency and Outcomes of Concomitant Use of Proton Pump Inhibitors and Clopidogrel after Hospital Discharge. Therapie, 2013, 68, 113-115.	0.6	1