Emanuele Monda

List of Publications by Year
in descending order

70	894	586496 16	591227 27
papers	citations	h-index	g-index

71	71	71	906
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Diagnostic issues faced by a rare disease healthcare network during Covid-19 outbreak: data from the Campania Rare Disease Registry. Journal of Public Health, 2022, 44, 586-594.	1.0	12
2	The Role of New Imaging Technologies in the Diagnosis of Cardiac Amyloidosis. Heart Failure Clinics, 2022, 18, 61-72.	1.0	8
3	Diagnosis and Management of Cardiovascular Involvement in Friedreich Ataxia. Heart Failure Clinics, 2022, 18, 31-37.	1.0	12
4	Diagnosis and Management of Cardiovascular Involvement in Fabry Disease. Heart Failure Clinics, 2022, 18, 39-49.	1.0	18
5	Cardiovascular Involvement in Transthyretin Cardiac Amyloidosis. Heart Failure Clinics, 2022, 18, 73-87.	1.0	12
6	Hypertrophic Cardiomyopathy in RASopathies. Heart Failure Clinics, 2022, 18, 19-29.	1.0	33
7	The Heart Muscle and Valve Involvement in Marfan Syndrome, Loeys-Dietz Syndromes, and Collagenopathies. Heart Failure Clinics, 2022, 18, 165-175.	1.0	7
8	The Risk of Sudden Unexpected Cardiac Death in Children. Heart Failure Clinics, 2022, 18, 115-123.	1.0	16
9	Rare Cardiovascular Diseases: From Genetics to Personalized Medicine. Heart Failure Clinics, 2022, 18, xix-xxi.	1.0	1
10	Rare Cardiovascular Diseases. Heart Failure Clinics, 2022, 18, i.	1.0	0
11	Natural history of left ventricular hypertrophy in infants of diabetic mothers. International Journal of Cardiology, 2022, 350, 77-82.	0.8	2
12	Multimodality Imaging in Cardiomyopathies with Hypertrophic Phenotypes. Journal of Clinical Medicine, 2022, 11, 868.	1.0	18
13	Diagnosis of Fabry Disease in a Patient with a Surgically Repaired Congenital Heart Defect: When Clinical History and Genetics Make the Difference. Neurology International, 2022, 12, 102-108.	0.2	1
14	Clinical and Molecular Characteristics of Patients with PLN R14del Cardiomyopathy: State-of-the-Art Review. Neurology International, 2022, 12, 112-121.	0.2	0
15	Pancarditis as the Clinical Presentation of Eosinophilic Granulomatosis with Polyangiitis: A Multimodality Approach to Diagnosis. Neurology International, 2022, 12, 133-141.	0.2	1
16	Diagnosis and Management of Rare Cardiomyopathies in Adult and Paediatric Patients. A Position Paper of the Italian Society of Cardiology (SIC) and Italian Society of Paediatric Cardiology (SICP). International Journal of Cardiology, 2022, 357, 55-71.	0.8	36
17	Bisoprolol for treatment of symptomatic patients with obstructive hypertrophic cardiomyopathy. The BASIC (bisoprolol AS therapy in hypertrophic cardiomyopathy) study. International Journal of Cardiology, 2022, 354, 22-28.	0.8	12
18	Implantable cardioverter defibrillator in hypertrophic cardiomyopathy: Time to avoid unnecessary procedure. International Journal of Cardiology, 2022, 355, 30-31.	0.8	0

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19	Modified Body Mass Index as a Novel Nutritional and Prognostic Marker in Patients with Cardiac Amyloidosis. Neurology International, 2022, 12, 185-197.	0.2	1
20	Thoracic Aortic Dilation: Implications for Physical Activity and Sport Participation. Diagnostics, 2022, 12, 1392.	1.3	5
21	Unexplained sudden cardiac arrest in children: clinical and genetic characteristics of survivors. European Journal of Preventive Cardiology, 2021, 28, 1134-1137.	0.8	20
22	Potential role of imaging markers in predicting future disease expression of arrhythmogenic cardiomyopathy. Future Cardiology, 2021, 17, 647-654.	0.5	8
23	Yield and clinical significance of genetic screening in elite and amateur athletes. European Journal of Preventive Cardiology, 2021, 28, 1081-1090.	0.8	35
24	Prognostic Implications of Declining Hemoglobin Content in Patients Hospitalized With Acute CoronaryÂSyndromes. Journal of the American College of Cardiology, 2021, 77, 375-388.	1.2	70
25	Rare case of Kawasaki disease with cardiac tamponade and giant coronary artery aneurysms. Cardiology in the Young, 2021, 31, 865-866.	0.4	3
26	Hypertrophic Cardiomyopathy in Children: Pathophysiology, Diagnosis, and Treatment of Non-sarcomeric Causes. Frontiers in Pediatrics, 2021, 9, 632293.	0.9	43
27	Access-Site Crossover in Patients With Acute Coronary Syndrome Undergoing Invasive Management. JACC: Cardiovascular Interventions, 2021, 14, 361-373.	1.1	25
28	Combined Effect of Mediterranean Diet and Aerobic Exercise on Weight Loss and Clinical Status in Obese Symptomatic Patients with Hypertrophic Cardiomyopathy. Heart Failure Clinics, 2021, 17, 303-313.	1.0	18
29	Impact of Regular Physical Activity on Aortic Diameter Progression in Paediatric Patients with Bicuspid Aortic Valve. Pediatric Cardiology, 2021, 42, 1133-1140.	0.6	9
30	Editorial: Paediatric Cardiomyopathies. Frontiers in Pediatrics, 2021, 9, 696443.	0.9	2
31	Troponin T Mutation as a Cause of Left Ventricular Systolic Dysfunction in a Young Patient with Previous Surgical Correction of Aortic Coarctation. Biomolecules, 2021, 11, 696.	1.8	4
32	Genetic evaluation in athletes and cascade family screening: reply. European Journal of Preventive Cardiology, 2021, , .	0.8	2
33	External validation of the increased wall thickness score for the diagnosis of cardiac amyloidosis. International Journal of Cardiology, 2021, 339, 99-101.	0.8	11
34	Advanced Heart Failure in Special Populationâ€"Pediatric Age. Heart Failure Clinics, 2021, 17, 673-683.	1.0	6
35	Lipoprotein(a): a genetic marker for cardiovascular disease and target for emerging therapies. Journal of Cardiovascular Medicine, 2021, 22, 151-161.	0.6	53
36	Cardiovascular Involvement in mtDNA Disease. Heart Failure Clinics, 2021, 18, 51-60.	1.0	11

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37	Clinical Manifestations of 22q11.2 Deletion Syndrome. Heart Failure Clinics, 2021, 18, 155-164.	1.0	15
38	Left ventricular rotational mechanics in cardiac amyloidosis - reply. International Journal of Cardiology, 2021, 345, 152.	0.8	1
39	Multidisciplinary In-Depth Investigation in a Young Athlete Suffering from Syncope Caused by Myocardial Bridge. Diagnostics, 2021, 11, 2144.	1.3	11
40	585â€ƒNatural history of left ventricular hypertrophy in infants of diabetic mothers. European Heart Journal Supplements, 2021, 23, .	0.0	0
41	Global Left Ventricular Myocardial Work Efficiency in Heart Failure Patients with Cardiac Amyloidosis: Pathophysiological Implications and Role in Differential Diagnosis.. Journal of Cardiovascular Echography, 2021, 31, 157-164.	0.1	2
42	Prediction of radial crossover in acute coronary syndromes: derivation and validation of the MATRIX score. EuroIntervention, 2021, 17, e971-e980.	1.4	13
43	596â€ƒMultidisciplinary in-depth investigation in a young athlete suffering from syncope caused by myocardial bridge. European Heart Journal Supplements, 2021, 23, .	0.0	0
44	Prevalence and direct costs of patients with neuromyelitis optica: data from Campania region in southern Italy. Future Rare Diseases, 2021, 1, .	0.1	3
45	589â€ƒExternal validation of the increased wall thickness score for the diagnosis of cardiac amyloidosis. European Heart Journal Supplements, 2021, 23, .	0.0	0
46	A complex unit for a complex disease: the HCM-Family Unit. Monaldi Archives for Chest Disease, 2021, , .	0.3	0
47	577â€ƒBisoprolol for the treatment of symptomatic patients with obstructive hypertrophic cardiomyopathy. European Heart Journal Supplements, 2021, 23, .	0.0	0
48	Prevalence and clinical significance of red flags in patients with hypertrophic cardiomyopathy. International Journal of Cardiology, 2020, 299, 186-191.	0.8	58
49	Prevalence and clinical implications of hyperhomocysteinaemia in patients with hypertrophic cardiomyopathy and MTHFR C6777T polymorphism. European Journal of Preventive Cardiology, 2020, 27, 1906-1908.	0.8	16
50	Beta-blockers in heart failure prognosis: Lessons learned by MECKI Score Group papers. European Journal of Preventive Cardiology, 2020, 27, 65-71.	0.8	4
51	The Hidden Fragility in the Heart of the Athletes: A Review of Genetic Biomarkers. International Journal of Molecular Sciences, 2020, 21, 6682.	1.8	14
52	Molecular Basis of Inflammation in the Pathogenesis of Cardiomyopathies. International Journal of Molecular Sciences, 2020, 21, 6462.	1.8	38
53	The hospitalizations in hypertrophic cardiomyopathy: â€œThe dark side of the moonâ€• International Journal of Cardiology, 2020, 318, 101-102.	0.8	10
54	Low-Dose Ticagrelor in Patients With High Ischemic Risk and Previous Myocardial Infarction: A Multicenter Prospective Real-World Observational Study. Journal of Cardiovascular Pharmacology, 2020, 76, 173-180.	0.8	31

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55	Combined PTPN11 and MYBPC3 Gene Mutations in an Adult Patient with Noonan Syndrome and Hypertrophic Cardiomyopathy. Genes, 2020, 11, 947.	1.0	18
56	Clinical Profile of Cardiac Involvement in Danon Disease. Circulation Genomic and Precision Medicine, 2020, 13, e003117.	1.6	29
57	Aortopathies in mouse models of Pompe, Fabry and Mucopolysaccharidosis IIIB lysosomal storage diseases. PLoS ONE, 2020, 15, e0233050.	1.1	13
58	Genetic analysis resolves differential diagnosis of a familial syndromic dilated cardiomyopathy: A new case of AlstrÃ¶m syndrome. Molecular Genetics & Genomic Medicine, 2020, 8, e1260.	0.6	22
59	Genotype-Phenotype Correlation: A Triple DNA Mutational Event in a Boy Entering Sport Conveys an Additional Pathogenicity Risk. Genes, 2020, 11, 524.	1.0	15
60	Beyond cholesterol metabolism: The pleiotropic effects of proprotein convertase subtilisin/kexin type 9 (PCSK9). Genetics, mutations, expression, and perspective for longâ€term inhibition. BioFactors, 2020, 46, 367-380.	2.6	46
61	Clinical significance of family history and bicuspid aortic valve in children and young adult patients with Marfan syndrome. Cardiology in the Young, 2020, 30, 663-667.	0.4	5
62	Myocarditis in Children. , 2020, , 243-260.		2
63	Title is missing!. , 2020, 15, e0233050.		0
64	Title is missing!. , 2020, 15, e0233050.		0
65	Title is missing!. , 2020, 15, e0233050.		0
66	Title is missing!. , 2020, 15, e0233050.		0
67	Title is missing!. , 2020, 15, e0233050.		0
68	Title is missing!. , 2020, 15, e0233050.		0
69	ECG analysis in patients with acute coronary syndrome undergoing invasive management: rationale and design of the electrocardiography sub-study of the MATRIX trial. Journal of Electrocardiology, 2019, 57, 44-54.	0.4	7
70	Ranolazine treatment for refractory angina in a patient with Hutchinson-Gilford progeria syndrome and end stage aortic stenosis. Neurology International, 2019, 9, .	0.2	1