Massimiliano Lorenzini

List of Publications by Year
in descending order

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Version: 2024-02-01

50 papers	2,892 citations	236612 25 h-index	214527 47 g-index
50 all docs	50 docs citations	50 times ranked	3400 citing authors

#	Article	IF	Citations
1	Cardiac magnetic resonance assessment of progressive myo-pericarditis due to cobalt cardiotoxicity. European Heart Journal Cardiovascular Imaging, 2021, 22, e71-e71.	0.5	0
2	Alpha-protein kinase 3 (ALPK3) truncating variants are a cause of autosomal dominant hypertrophic cardiomyopathy. European Heart Journal, 2021, 42, 3063-3073.	1.0	51
3	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
4	The complex interplay among atherosclerosis, inflammation, and degeneration in ascending thoracic aortic aneurysms. Journal of Thoracic and Cardiovascular Surgery, 2020, 160, 1434-1443.e6.	0.4	20
5	Mortality Among Referral Patients With Hypertrophic Cardiomyopathy vs the General European Population. JAMA Cardiology, 2020, 5, 73.	3.0	69
6	Clinical Phenotypes and Prognosis of Dilated Cardiomyopathy Caused by Truncating Variants in the TTN Gene. Circulation: Heart Failure, 2020, 13, e006832.	1.6	75
7	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
8	Penetrance of Hypertrophic Cardiomyopathy in Sarcomere Protein Mutation Carriers. Journal of the American College of Cardiology, 2020, 76, 550-559.	1.2	89
9	Clinical Features and Natural History of PRKAG2 Variant Cardiac Glycogenosis. Journal of the American College of Cardiology, 2020, 76, 186-197.	1.2	45
10	Carpal tunnel syndrome in cardiac amyloidosis: implications for early diagnosis and prognostic role across the spectrum of aetiologies. European Journal of Heart Failure, 2020, 22, 507-515.	2.9	106
11	Diagnostic performance of imaging investigations in detecting and differentiating cardiac amyloidosis: a systematic review and metaâ€analysis. ESC Heart Failure, 2019, 6, 1041-1051.	1.4	73
12	Prevalence of TTR variants detected by whole-exome sequencing in hypertrophic cardiomyopathy. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 243-247.	1.4	10
13	The complex interplay between fitness, genetics, lifestyle, and inflammation in the pathogenesis of coronary atherosclerosis: lessons from the Amazon rainforest. European Heart Journal Supplements, 2019, 21, B76-B79.	0.0	3
14	Tafamidis for the treatment of transthyretin amyloidosis. Future Cardiology, 2019, 15, 53-61.	0.5	3
15	Differences in cardiac phenotype and natural history of laminopathies with and without neuromuscular onset. Orphanet Journal of Rare Diseases, 2019, 14, 263.	1.2	12
16	Extracardiac imaging in amyloidosis: A long and winding (but possible) road. International Journal of Cardiology, 2018, 254, 351-352.	0.8	0
17	Redefining the histopathologic profile of acute aortic syndromes: Clinical and prognostic implications. Journal of Thoracic and Cardiovascular Surgery, 2018, 156, 1776-1785.e6.	0.4	22
18	Isolated aortic root dilation in homocystinuria. Journal of Inherited Metabolic Disease, 2018, 41, 109-115.	1.7	11

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19	Long-term Follow up of Patients with Acute Aortic Syndromes: Relevance of both Aortic and Non-aortic Events. European Journal of Vascular and Endovascular Surgery, 2018, 56, 200-208.	0.8	10
20	Phenotypic profile of Ile68Leu transthyretin amyloidosis: an underdiagnosed cause of heart failure. European Journal of Heart Failure, 2018, 20, 1417-1425.	2.9	36
21	Does the etiology of cardiac amyloidosis determine the myocardial uptake of [18F]-NaF PET/CT?. Journal of Nuclear Cardiology, 2017, 24, 746-749.	1.4	31
22	Clinical characteristics of wild-type transthyretin cardiac amyloidosis: disproving myths. European Heart Journal, 2017, 38, 1895-1904.	1.0	258
23	The complex interplay between systolic and diastolic function at rest and during exercise in heart failure: the case of cardiac amyloidosis. European Journal of Heart Failure, 2017, 19, 1466-1467.	2.9	4
24	Broadening the Phenotypic Spectrum andÂthe Diagnostic Needs of TTR-Related Cardiac Amyloidosis â^—. Journal of the American College of Cardiology, 2017, 70, 478-480.	1.2	2
25	Usefulness of Electrocardiographic Patterns at Presentation to Predict Long-term Risk of Cardiac Death in Patients With Hypertrophic Cardiomyopathy. American Journal of Cardiology, 2016, 118, 432-439.	0.7	45
26	Histological and Histometric Characterization of Myocardial Fibrosis in End-Stage Hypertrophic Cardiomyopathy. Circulation: Heart Failure, 2016, 9, .	1.6	103
27	The Authors Reply:. JACC: Cardiovascular Imaging, 2016, 9, 906.	2.3	0
28	Long-term prognostic role of cerebrovascular disease and peripheral arterial disease across the spectrum of acute coronary syndromes. Atherosclerosis, 2016, 245, 43-49.	0.4	13
29	Troponin T elevation in acute aortic syndromes: Frequency and impact on diagnostic delay and misdiagnosis. European Heart Journal: Acute Cardiovascular Care, 2016, 5, 61-71.	0.4	26
30	Coexistence of Degenerative Aortic Stenosis and Wild-Type Transthyretin-Related CardiacÂAmyloidosis. JACC: Cardiovascular Imaging, 2016, 9, 325-327.	2.3	89
31	Acute heart failure in patients with acute aortic syndrome: pathophysiology and clinicalâ€“prognostic implications. European Journal of Heart Failure, 2015, 17, 917-924.	2.9	9
32	Etiology of Amyloidosis Determines Myocardial 99mTc-DPD Uptake in Amyloidotic Cardiomyopathy. Clinical Nuclear Medicine, 2015, 40, 446-447.	0.7	6
33	Atrial fibrillation in amyloidotic cardiomyopathy: prevalence, incidence, risk factors and prognostic role. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 147-155.	1.4	115
34	Cardiac amyloidosis: the great pretender. Heart Failure Reviews, 2015, 20, 117-124.	1.7	147
35	Nuclear imaging for cardiac amyloidosis. Heart Failure Reviews, 2015, 20, 145-154.	1.7	15
36	Long-Term Outcomes and Causes of Death After Acute Coronary Syndrome in Patients in the Bologna, Italy, Area. American Journal of Cardiology, 2015, 115, 171-177.	0.7	11

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37	An unusual case of a congenital aorto-left atrial tunnel. Cardiovascular Pathology, 2014, 23, 241-243.	0.7	1
38	Role of 18F-FDG PET/CT in the diagnosis of infective endocarditis in patients with an implanted cardiac device: a prospective study. European Journal of Nuclear Medicine and Molecular Imaging, 2014, 41, 1617-1623.	3.3	79
39	Diagnostic performance of standard electrocardiogram for prediction of infarct related artery and site of coronary occlusion in unselected STEMI patients undergoing primary percutaneous coronary intervention. European Heart Journal: Acute Cardiovascular Care, 2014, 3, 326-339.	0.4	22
40	Identification ofÂTTR-Related Subclinical Amyloidosis WithÂ99mTc-DPD Scintigraphy. JACC: Cardiovascular Imaging, 2014, 7, 531-532.	2.3	91
41	Amyloid fibrils containing fragmented ATTR may be the standard fibril composition in ATTR amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2013, 20, 142-150.	1.4	106
42	Disease profile and differential diagnosis of hereditary transthyretin-related amyloidosis with exclusively cardiac phenotype: an Italian perspective. European Heart Journal, 2013, 34, 520-528.	1.0	252
43	Effects of myocardial fibrosis assessed by MRI on dynamic left ventricular outflow tract obstruction in patients with hypertrophic cardiomyopathy: a retrospective database analysis. BMJ Open, 2012, 2, e001267.	0.8	13
44	What is the acceptable rate of false positives for STEMI within a primary PCI network? Insights from a metropolitan system with direct ambulance-based access. International Journal of Cardiology, 2012, 154, 356-358.	0.8	5
45	Defining the Diagnosis in Echocardiographically Suspected Senile Systemic Amyloidosis. JACC: Cardiovascular Imaging, 2012, 5, 755-758.	2.3	33
46	Cardiac involvement in hereditary-transthyretin related amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2012, 19, 16-21.	1.4	24
47	Role of 99mTc-DPD Scintigraphy in Diagnosis and Prognosis of Hereditary Transthyretin-Related Cardiac Amyloidosis. JACC: Cardiovascular Imaging, 2011, 4, 659-670.	2.3	264
48	Significance of Magnetic Resonance Imaging in Apical Hypertrophic Cardiomyopathy. American Journal of Cardiology, 2010, 105, 1592-1596.	0.7	59
49	Transthyretin-related amyloidoses and the heart: a clinical overview. Nature Reviews Cardiology, 2010, 7, 398-408.	6.1	286
50	Prognostic Implications of the Doppler Restrictive Filling Pattern in Hypertrophic Cardiomyopathy. American Journal of Cardiology, 2009, 104, 1727-1731.	0.7	93