Lorenzo Paolo Giuliani

List of Publications by Year
in descending order

24 papers	238 citations	1478280 6 h-index	996849 15 g-index
24 all docs	24 docs citations	24 times ranked	467 citing authors

#	Article	IF	Citations
1	Cardiac Phenotypes in HereditaryÂMuscleÂDisorders. Journal of the American College of Cardiology, 2018, 72, 2485-2506.	1.2	71
2	Genetic Screening of Anderson-Fabry Disease in Probands Referred From Multispecialty Clinics. Journal of the American College of Cardiology, 2016, 68, 1037-1050.	1.2	50
3	Long COVID: long-term effects?. European Heart Journal Supplements, 2021, 23, E1-E5.	0.0	37
4	Differences between familial and sporadic dilated cardiomyopathy: ESC EORP Cardiomyopathy & Myocarditis registry. ESC Heart Failure, 2021, 8, 95-105.	1.4	23
5	Genetic Basis of Myocarditis: Myth or Reality?. , 2020, , 45-89.		15
6	Myths to debunk: the non-compacted myocardium. European Heart Journal Supplements, 2020, 22, L6-L10.	0.0	9
7	â€˜Precision and personalized medicine,â€™ a dream that comes true?. Journal of Cardiovascular Medicine, 2017, 18, e1-e6.	0.6	6
8	Oxalic Cardiomyopathy. Journal of the American College of Cardiology, 2021, 78, 998-999.	1.2	5
9	Common presentation of rare diseases: Aortic aneurysms & valves. International Journal of Cardiology, 2018, 257, 358-365.	0.8	4
10	6128Losartan vs Nebivolol vs the association of both on the progression of aortic root dilation in genotyped Marfan Syndrome: 48 months open label randomized controlled phase III trial. European Heart Journal, 2019, 40, .	1.0	3
11	Renal and brain complications in GLA p.Phe113Leu Fabry disease. Comments on â€œFabry disease caused by the GLA p.Phe113Leu (p.F113L) variant: Natural history in malesâ€•by Oliveira et al. (Eur. J. Med. Genet.) Tj ETQq1 1 0.784314 rgBT /Ov		
12	Cardio-Oncology. Journal of the American College of Cardiology, 2016, 68, 1921-1923.	1.2	2
13	Hereditary muscle diseases and the heart: the cardiologistâ€™s perspective. European Heart Journal Supplements, 2020, 22, E13-E19.	0.0	2
14	Pathologic substrate of gastropathy in Anderson-Fabry disease. Orphanet Journal of Rare Diseases, 2020, 15, 156.	1.2	2
15	When Genes, More Than Phenotype, Identify Different Diseases. Journal of the American College of Cardiology, 2018, 72, 616-619.	1.2	1
16	P5539Genetic heterogeneity of spontaneous coronary artery dissection (SCAD). European Heart Journal, 2019, 40, .	1.0	1
17	P5723IEVA: Integration and Extraction of Variant Attributes in NGS analysis. European Heart Journal, 2019, 40, .	1.0	1
18	Diagnostic Criteria of Left Ventricular Dysfunction in Patients With Myotonic Dystrophy Type 1. Journal of Cardiac Failure, 2020, 26, 857-859.	0.7	1

#	Article	IF	Citations
19	Spectrum of phenotype of ventricular noncompaction in adults. Progress in Pediatric Cardiology, 2021, 62, 101416.	0.2	1
20	Inherited Cardiac Muscle Disease: Dilated Cardiomyopathy. , 2018, , 319-366.		1
21	Fatal ventricular arrhythmias in a young male with unrecognized LQT3 and cardiolaminopathy. Journal of Cardiovascular Medicine, 2017, 18, e192-e194.	0.6	0
22	2835Complications and management of pregnancy in Danon disease. European Heart Journal, 2017, 38, .	1.0	0
23	Genetics and clinics: current applications, limitations, and future developments. European Heart Journal Supplements, 2019, 21, B7-B14.	0.0	0
24	Genetics of cardiomyopathies: left ventricular non-compaction. , 2018, , 697-699.		0