Lorenzo Paolo Giuliani

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6916797/publications.pdf

Version: 2024-02-01

24 papers

238 citations

1478280 6 h-index 996849 15 g-index

24 all docs

24 docs citations

times ranked

24

467 citing authors

#	Article	IF	CITATIONS
1	Differences between familial and sporadic dilated cardiomyopathy: ESC EORP Cardiomyopathy & ESC Myocarditis registry. ESC Heart Failure, 2021, 8, 95-105.	1.4	23
2	Oxalic Cardiomyopathy. Journal of the American College of Cardiology, 2021, 78, 998-999.	1.2	5
3	Spectrum of phenotype of ventricular noncompaction in adults. Progress in Pediatric Cardiology, 2021, 62, 101416.	0.2	1
4	Long COVID: long-term effects?. European Heart Journal Supplements, 2021, 23, E1-E5.	0.0	37
5	Diagnostic Criteria of Left Ventricular Dysfunction in Patients With Myotonic Dystrophy Type 1. Journal of Cardiac Failure, 2020, 26, 857-859.	0.7	1
6	Myths to debunk: the non-compacted myocardium. European Heart Journal Supplements, 2020, 22, L6-L10.	0.0	9
7	Hereditary muscle diseases and the heart: the cardiologist's perspective. European Heart Journal Supplements, 2020, 22, E13-E19.	0.0	2
8	Pathologic substrate of gastropathy in Anderson-Fabry disease. Orphanet Journal of Rare Diseases, 2020, 15, 156.	1.2	2
9	Renal and brain complications in GLA p.Phe113Leu Fabry disease. Comments on "Fabry disease caused by the GLA p.Phe113Leu (p.F113L) variant: Natural history in males―by Oliveira et al. (Eur. J. Med. Genet.) Tj ETQq1	100778431	l 4 rgBT /Ove
10	Genetic Basis of Myocarditis: Myth or Reality?. , 2020, , 45-89.		15
10	Genetic Basis of Myocarditis: Myth or Reality?., 2020, , 45-89. P5539Genetic heterogeneity of spontaneous coronary artery dissection (SCAD). European Heart Journal, 2019, 40, .	1.0	15
	P5539Genetic heterogeneity of spontaneous coronary artery dissection (SCAD). European Heart	1.0	
11	P5539Genetic heterogeneity of spontaneous coronary artery dissection (SCAD). European Heart Journal, 2019, 40, . 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		1
11 12	P5539Genetic heterogeneity of spontaneous coronary artery dissection (SCAD). European Heart Journal, 2019, 40, . 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, . Genetics and clinics: current applications, limitations, and future developments. European Heart	1.0	3
11 12 13	P5539Genetic heterogeneity of spontaneous coronary artery dissection (SCAD). European Heart Journal, 2019, 40, . 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, . Genetics and clinics: current applications, limitations, and future developments. European Heart Journal Supplements, 2019, 21, 87-814. P5723IEVA: Integration and Extraction of Variant Attributes in NGS analysis. European Heart Journal,	1.0	3
11 12 13	P5539Genetic heterogeneity of spontaneous coronary artery dissection (SCAD). European Heart Journal, 2019, 40,. 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,. Genetics and clinics: current applications, limitations, and future developments. European Heart Journal Supplements, 2019, 21, B7-B14. P5723IEVA: Integration and Extraction of Variant Attributes in NGS analysis. European Heart Journal, 2019, 40,. Common presentation of rare diseases: Aortic aneurysms & Common presentation of Variant Attributes in NGS analysis.	1.0	1 3 0
11 12 13 14	P5539Genetic heterogeneity of spontaneous coronary artery dissection (SCAD). European Heart Journal, 2019, 40, . 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, . Genetics and clinics: current applications, limitations, and future developments. European Heart Journal Supplements, 2019, 21, 87-814. P5723IEVA: Integration and Extraction of Variant Attributes in NGS analysis. European Heart Journal, 2019, 40, . Common presentation of rare diseases: Aortic aneurysms & Samp; valves. International Journal of Cardiology, 2018, 257, 358-365. Cardiac Phenotypes in HereditaryÂMuscleÂDisorders. Journal of the American College of Cardiology,	1.0 0.0 1.0 0.8	1 3 0 1

#	Article	IF	CITATIONS
19	Genetics of cardiomyopathies: left ventricular non-compaction. , 2018, , 697-699.		o
20	â€~Precision and personalized medicine,' a dream that comes true?. Journal of Cardiovascular Medicine, 2017, 18, e1-e6.	0.6	6
21	Fatal ventricular arrhythmias in a young male with unrecognized LQT3 and cardiolaminopathy. Journal of Cardiovascular Medicine, 2017, 18, e192-e194.	0.6	O
22	2835Complications and management of pregnancy in Danon disease. European Heart Journal, 2017, 38, .	1.0	0
23	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
24	Cardio-Oncology. Journal of the American College of Cardiology, 2016, 68, 1921-1923.	1.2	2