Valentina Favalli

List of Publications by Year
in descending order

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40 papers	1,791 citations	361045 20 h-index	315357 38 g-index
41 all docs	41 docs citations	41 times ranked	2788 citing authors

#	Article	IF	Citations
1	Interpretation and actionability of genetic variants in cardiomyopathies: a position statement from the European Society of Cardiology Council on cardiovascular genomics. European Heart Journal, 2022, 43, 1901-1916.	1.0	32
2	Age-specific reference values for carotid arterial stiffness estimated by ultrasonic wall tracking. Journal of Human Hypertension, 2020, 34, 214-222.	1.0	34
3	Myths to debunk: the non-compacted myocardium. European Heart Journal Supplements, 2020, 22, L6-L10.	0.0	9
4	Hereditary muscle diseases and the heart: the cardiologistâ€™s perspective. European Heart Journal Supplements, 2020, 22, E13-E19.	0.0	2
5	Pathologic substrate of gastropathy in Anderson-Fabry disease. Orphanet Journal of Rare Diseases, 2020, 15, 156.	1.2	2
6	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 ETQq0 0 0 rgBT /Overlock 10		
7	Genetics and clinics: current applications, limitations, and future developments. European Heart Journal Supplements, 2019, 21, B7-B14.	0.0	0
8	Extracellular Volume in DilatedÂCardiomyopathy. JACC: Cardiovascular Imaging, 2018, 11, 60-63.	2.3	5
9	Andersonâ€“Fabry disease. Journal of Cardiovascular Medicine, 2018, 19, e1-e5.	0.6	8
10	International External Validation Study of the 2014 European Society of Cardiology Guidelines on Sudden Cardiac Death Prevention in Hypertrophic Cardiomyopathy (EVIDENCE-HCM). Circulation, 2018, 137, 1015-1023.	1.6	149
11	Cardiac Phenotypes in HereditaryÂMuscleÂDisorders. Journal of the American College of Cardiology, 2018, 72, 2485-2506.	1.2	71
12	Inherited Cardiac Muscle Disease: Dilated Cardiomyopathy. , 2018, , 319-366.		1
13	â€˜Precision and personalized medicine,â€™ a dream that comes true?. Journal of Cardiovascular Medicine, 2017, 18, e1-e6.	0.6	6
14	Reply. Journal of the American College of Cardiology, 2017, 69, 1210-1211.	1.2	2
15	LMNA Mutations Associated With Mild and Late-Onset Phenotype. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	6
16	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
17	Left Ventricular Noncompaction. Journal of the American College of Cardiology, 2016, 68, 949-966.	1.2	206
18	Genetic causes of dilated cardiomyopathy. Heart, 2016, 102, 2004-2014.	1.2	22

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19	Cardio-Oncology. Journal of the American College of Cardiology, 2016, 68, 1921-1923.	1.2	2
20	Radiation Therapy for Head and Neck Cancer and Angioneogenesis. JACC: Cardiovascular Imaging, 2016, 9, 676-679.	2.3	1
21	Endomyocardial Biopsy in acute cardiogenic shock: Diagnosis of pheochromocytoma. International Journal of Cardiology, 2016, 202, 897-899.	0.8	3
22	European Cardiomyopathy Pilot Registry: EURObservational Research Programme of the European Society of Cardiology. European Heart Journal, 2016, 37, 164-173.	1.0	56
23	MOGE(S) nosology in low-to-middle-income countries. Nature Reviews Cardiology, 2014, 11, 307-307.	6.1	2
24	The Pathologic Basis of Recovery. Heart Failure Clinics, 2014, 10, S63-S74.	1.0	3
25	The MOGE(S) Classification of Cardiomyopathy for Clinicians. Journal of the American College of Cardiology, 2014, 64, 304-318.	1.2	158
26	Reply. Journal of the American College of Cardiology, 2014, 63, 2584-2586.	1.2	3
27	The MOGE(S) Classification for a Phenotypeâ€“Genotype Nomenclature of Cardiomyopathy. Journal of the American College of Cardiology, 2013, 62, 2046-2072.	1.2	203
28	Aortic root 3D parametric morphological model from 2D-echo images. Computers in Biology and Medicine, 2013, 43, 2196-2204.	3.9	20
29	The need to modify patient selection to improve the benefits of implantable cardioverter-defibrillator for primary prevention of sudden death in non-ischaemic dilated cardiomyopathy. Europace, 2013, 15, 1693-1701.	0.7	41
30	Autosomal Recessive Atrial Dilated Cardiomyopathy With Standstill Evolution Associated With Mutation of Natriuretic Peptide Precursor A. Circulation: Cardiovascular Genetics, 2013, 6, 27-36.	5.1	51
31	The MOGE(S) Classification for a Phenotypeâ€“Genotype Nomenclature of Cardiomyopathy: Endorsed by the World Heart Federation. Global Heart, 2013, 8, 355.	0.9	28
32	Familial dilated cardiomyopathy. Herz, 2012, 37, 822-829.	0.4	9
33	Structures of the lamin A/C R335W and E347K mutants: Implications for dilated cardiolaminopathies. Biochemical and Biophysical Research Communications, 2012, 418, 217-221.	1.0	21
34	Quantitative Expression of the Mutated Lamin A/C Gene in Patients With Cardiolaminopathy. Journal of the American College of Cardiology, 2012, 60, 1916-1920.	1.2	34
35	Diagnostic Work-Up and Risk Stratification in X-Linked Dilated Cardiomyopathies Caused by Dystrophin Defects. Journal of the American College of Cardiology, 2011, 58, 925-934.	1.2	73
36	Risk of dissection in thoracic aneurysms associated with mutations of smooth muscle alpha-actin 2 (ACTA2). Heart, 2011, 97, 321-326.	1.2	65

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37	Translational Bioinformatics: Challenges and Opportunities for Case-Based Reasoning and Decision Support. Lecture Notes in Computer Science, 2010, , 1-11.	1.0	1
38	Rationale and design of a trial evaluating the effects of losartan vs. nebivolol vs. the association of both on the progression of aortic root dilation in Marfan syndrome with FBN1 gene mutations. Journal of Cardiovascular Medicine, 2009, 10, 354-362.	0.6	66
39	Transcriptomic and proteomic analysis in the cardiovascular setting: unravelling the disease?. Journal of Cardiovascular Medicine, 2009, 10, 433-442.	0.6	8
40	Long-Term Outcome and Risk Stratification in Dilated Cardiolaminopathies. Journal of the American College of Cardiology, 2008, 52, 1250-1260.	1.2	335