Maurizia Grasso

List of Publications by Year
in descending order

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65	4,115	109264	114418
papers	citations	35 h-index	63 g-index

67	67	67	5343
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genome-wide association analysis in dilated cardiomyopathy reveals two new players in systolic heart failure on chromosomes 3p25.1 and 22q11.23. European Heart Journal, 2021, 42, 2000-2011.	1.0	49
2	Spectrum of phenotype of ventricular noncompaction in adults. Progress in Pediatric Cardiology, 2021, 62, 101416.	0.2	1
3	Clinical heterogeneity and reduced penetrance in DICER1 syndrome: a report of three families. Tumori, 2021, 107, NP144-NP148.	0.6	2
4	Management of the axilla in patients with breast cancer and positive sentinel lymph node biopsy: An evidence-based update in a European breast center. European Journal of Surgical Oncology, 2020, 46, 15-23.	0.5	24
5	Myths to debunk: the non-compacted myocardium. European Heart Journal Supplements, 2020, 22, L6-L10.	0.0	9
6	Hereditary muscle diseases and the heart: the cardiologistâ€™s perspective. European Heart Journal Supplements, 2020, 22, E13-E19.	0.0	2
7	Rare exon 10 deletion in POLH gene in a family with xeroderma pigmentosum variant correlating with protein expression by immunohistochemistry. Giornale Italiano Di Dermatologia E Venereologia, 2020, 155, 349-354.	0.8	1
8	Genetics and clinics: current applications, limitations, and future developments. European Heart Journal Supplements, 2019, 21, B7-B14.	0.0	0
9	Cardiac Phenotypes in HereditaryÂMuscleÂDisorders. Journal of the American College of Cardiology, 2018, 72, 2485-2506.	1.2	71
10	Genetic counselling and high-penetrance susceptibility gene analysis reveal the novel CDKN2A p.D84V (c.251A>T) mutation in melanoma-prone families from Italy. Melanoma Research, 2017, 27, 97-103.	0.6	1
11	Reply. Journal of the American College of Cardiology, 2017, 69, 1210-1211.	1.2	2
12	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
13	RE: BRCA2 Polymorphic Stop Codon K3326X and the Risk of Breast, Prostate, and Ovarian Cancers. Journal of the National Cancer Institute, 2016, 108, djw172.	3.0	15
14	Left Ventricular Noncompaction. Journal of the American College of Cardiology, 2016, 68, 949-966.	1.2	206
15	Genetic causes of dilated cardiomyopathy. Heart, 2016, 102, 2004-2014.	1.2	22
16	Cardio-Oncology. Journal of the American College of Cardiology, 2016, 68, 1921-1923.	1.2	2
17	Clinical Pregenetic Screening for Stroke Monogenic Diseases. Stroke, 2016, 47, 1702-1709.	1.0	34
18	Atlas of the clinical genetics of human dilated cardiomyopathy. European Heart Journal, 2015, 36, 1123-1135.	1.0	456

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19	Incomplete penetrance ofGLMNgene c.395-1G>C mutation in a family with glomuvenous malformations. International Journal of Dermatology, 2014, 53, 1362-1364.	0.5	3
20	Glomuvenous malformations with smooth muscle and eccrine glands: unusual histopathologic features in a familial setting. Journal of Cutaneous Pathology, 2014, 41, 308-315.	0.7	9
21	The MOGE(S) Classification of Cardiomyopathy for Clinicians. Journal of the American College of Cardiology, 2014, 64, 304-318.	1.2	158
22	Autosomal recessive paediatric sick sinus syndrome associated with novel compound mutations in SCN5A. International Journal of Cardiology, 2013, 167, 3078-3080.	0.8	11
23	Genderâ€specific differences in major cardiac events and mortality in lamin A/C mutation carriers. European Journal of Heart Failure, 2013, 15, 376-384.	2.9	120
24	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
25	In-Frame Mutations in Exon 1 of SKI Cause Dominant Shprintzen-Goldberg Syndrome. American Journal of Human Genetics, 2012, 91, 950-957.	2.6	95
26	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
27	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
28	A novel mutation of the glomulin gene in an Italian family with autosomal dominant cutaneous glomuvenous malformations. Experimental Dermatology, 2011, 20, 1032-1034.	1.4	11
29	Risk of dissection in thoracic aneurysms associated with mutations of smooth muscle alpha-actin 2 (ACTA2). Heart, 2011, 97, 321-326.	1.2	65
30	When Should Cardiologists Suspect Anderson-Fabry Disease?. American Journal of Cardiology, 2010, 106, 1492-1499.	0.7	46
31	Clinical and Molecular Study of 320 Children With Marfan Syndrome and Related Type I Fibrillinopathies in a Series of 1009 Probands With Pathogenic FBN1 Mutations. Pediatrics, 2009, 123, 391-398.	1.0	146
32	The shortness of Pygmies is associated with severe under-expression of the growth hormone receptor. Molecular Genetics and Metabolism, 2009, 98, 310-313.	0.5	44
33	Transcriptomic and proteomic analysis in the cardiovascular setting: unravelling the disease?. Journal of Cardiovascular Medicine, 2009, 10, 433-442.	0.6	8
34	Long-Term Outcome and Risk Stratification in Dilated Cardiolaminopathies. Journal of the American College of Cardiology, 2008, 52, 1250-1260.	1.2	335
35	Letter by Maurizia Grasso et al. regarding article, â€œRestrictive cardiomyopathy with atrioventricular conduction block resulting from a desmin mutationâ€. International Journal of Cardiology, 2008, 131, 144-145.	0.8	2
36	Barth syndrome associated with compound hemizygosity and heterozygosity of theTAZ andLDB3 genes. American Journal of Medical Genetics, Part A, 2007, 143A, 907-915.	0.7	41

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37	Î±B-Crystallin mutation in dilated cardiomyopathies: Low prevalence in a consecutive series of 200 unrelated probands. Biochemical and Biophysical Research Communications, 2006, 346, 1115-1117.	1.0	52
38	Early Diagnosis of Wilson Disease in a Six-year-old Child. Journal of Pediatrics, 2006, 148, 141.	0.9	4
39	Two novel and one known mutation of the TGFBR2 gene in Marfan syndrome not associated with FBN1 gene defects. European Journal of Human Genetics, 2006, 14, 34-38.	1.4	62
40	Desmin accumulation restrictive cardiomyopathy and atrioventricular block associated with desmin gene defects. European Journal of Heart Failure, 2006, 8, 477-483.	2.9	153
41	Kaposiâ€™s Sarcoma in Transplant and HIV-infected Patients: An Epidemiologic Study in Italy and France. Transplantation, 2005, 80, 1699-1704.	0.5	50
42	Identification of sixty-two novel and twelve known FBN1 mutations in eighty-one unrelated probands with Marfan syndrome and other fibrillinopathies. Human Mutation, 2005, 26, 494-494.	1.1	83
43	Autosomal dominant dilated cardiomyopathy with atrioventricular block: a lamin A/C defect-related disease. Journal of the American College of Cardiology, 2002, 39, 981-990.	1.2	306
44	The mitochondrial DNA mutation T12297C affects a highly conserved nucleotide of tRNALeu(CUN) and is associated with dilated cardiomyopathy. European Journal of Human Genetics, 2001, 9, 311-315.	1.4	40
45	Prevalence and characteristics of dystrophin defects in adult male patients with dilated cardiomyopathy. Journal of the American College of Cardiology, 2000, 35, 1760-1768.	1.2	83
46	Restrictive Cardiomyopathy, Atrioventricular Block and Mild to Subclinical Myopathy in Patients With Desmin-Immunoreactive Material Deposits. Journal of the American College of Cardiology, 1998, 31, 645-653.	1.2	117
47	Mitochondrial DNA Mutations and Mitochondrial Abnormalities in Dilated Cardiomyopathy. American Journal of Pathology, 1998, 153, 1501-1510.	1.9	225
48	Enteroviral RNA and Virus-Like Particles in the Skeletal Muscle of Patients With Idiopathic Dilated Cardiomyopathy. American Journal of Cardiology, 1997, 80, 1188-1193.	0.7	19
49	Hereditary Hyperferritinemia-Cataract Syndrome: Relationship Between Phenotypes and Specific Mutations in the Iron-Responsive Element of Ferritin Light-Chain mRNA. Blood, 1997, 90, 814-821.	0.6	5
50	Frequency and characteristics of coronary thrombosis in the epicardial coronary arteries after cardiac transplantation. American Journal of Cardiology, 1996, 78, 795-800.	0.7	19
51	HUMAN CYTOMEGALOVIRUS EARLY INFECTION, ACUTE REJECTION, AND MAJOR HISTOCOMPATIBILITY CLASS II EXPRESSION IN TRANSPLANTED LUNG. Transplantation, 1996, 61, 418-427.	0.5	22
52	Comparison of coronary lesions obtained by directional coronary atherectomy in unstable angina, stable angina, and restenosis after either atherectomy or angioplasty. American Journal of Cardiology, 1995, 75, 675-682.	0.7	72
53	Cardiac immunocyte-derived (AL) amyloidosis: An endomyocardial biopsy study in 11 patients. American Heart Journal, 1995, 130, 528-536.	1.2	50
54	Expression of proliferating cell markers in normal and diseased human hearts. American Journal of Cardiology, 1993, 72, 608-614.	0.7	21

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55	Morphologic changes induced by acetylcholine infusion in normal and atherosclerotic coronary arteries. American Journal of Cardiology, 1993, 71, 1382-1390.	0.7	10
56	Coronary thrombosis in non-cardiac death. Coronary Artery Disease, 1993, 4, 751-760.	0.3	58
57	Localization of brain and atrial natriuretic peptide in human and porcine heart. International Journal of Cardiology, 1992, 34, 237-247.	0.8	34
58	Search for Coxsackievirus B3 RNA in idiopathic dilated cardiomyopathy using gene amplification by polymerase chain reaction. American Journal of Cardiology, 1992, 69, 658-664.	0.7	97
59	H and L ferritins in myocardium in iron overload. American Journal of Cardiology, 1991, 68, 1233-1236.	0.7	10
60	Coronary atherosclerotic plaques with and without thrombus in ischemic heart syndromes: A morphologic, immunohistochemical, and biochemical study. American Journal of Cardiology, 1991, 68, B36-B50.	0.7	136
61	Endomyocardial biopsy finding in two patients with idiopathic dilated cardiomyopathy receiving long-term treatment with amiodarone. American Journal of Cardiology, 1991, 67, 661-662.	0.7	5
62	Atrial amyloid deposits in the failing human heart display both atrial and brain natriuretic peptide-like immunoreactivity. Journal of Pathology, 1991, 165, 235-241.	2.1	40
63	Expression of natriuretic peptide in ventricular myocardium of failing human hearts and its correlation with the severity of clinical and hemodynamic impairment. American Journal of Cardiology, 1990, 66, 973-980.	0.7	41
64	The morphologic spectrum of dilated cardiomyopathy and its relation to immune-response genes. American Journal of Cardiology, 1989, 64, 991-995.	0.7	44
65	Myocardial iron grading by endomyocardial biopsy. A clinicoâ€pathologic study on iron overloaded patients. European Journal of Haematology, 1989, 42, 382-388.	1.1	57