Maurizia Grasso

List of Publications by Year in Descending Order

66 papers	3,280 citations	34 h-index	57 g-index
67 ext. papers	3,727 ext. citations	4.1 avg, IF	4.01 L-index

#	Paper	IF	Citations
66	Clinical heterogeneity and reduced penetrance in DICER1 syndrome: a report of three families. *Tumori*, **2021**, 107, NP144-NP148	1.7	1
65	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	9.5	14
64	Spectrum of phenotype of ventricular noncompaction in adults. *Progress in Pediatric Cardiology*, **2021**, 62, 101416	0.4	1
63	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	0
62	Myths to debunk: the non-compacted myocardium. *European Heart Journal Supplements*, **2020**, 22, L6-L10	1.5	4
61	Hereditary muscle diseases and the heart: the cardiologist's perspective. *European Heart Journal Supplements*, **2020**, 22, E13-E19	1.5	2
60	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	3.6	11
59	Genetics and clinics: current applications, limitations, and future developments. *European Heart Journal Supplements*, **2019**, 21, B7-B14	1.5	
58	Cardiac Phenotypes in Hereditary Muscle Disorders: JACC State-of-the-Art Review. *Journal of the American College of Cardiology*, **2018**, 72, 2485-2506	15.1	42
57	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	2.3	1
56	Reply: A Distinct Cardiomyopathy: HCN4 Syndrome Comprising Myocardial Noncompaction, Bradycardia, Mitral Valve Defects, and Aortic Dilation. *Journal of the American College of Cardiology*, **2017**, 69, 1210-1211	15.1	2
55	Genetic Screening of Anderson-Fabry Disease in Probands Referred From Multispecialty Clinics. *Journal of the American College of Cardiology*, **2016**, 68, 1037-50	15.1	37
54	RE: BRCA2 Polymorphic Stop Codon K3326X and the Risk of Breast, Prostate, and Ovarian Cancers. *Journal of the National Cancer Institute*, **2016**, 108,	9.7	14
53	Left Ventricular Noncompaction: A Distinct Genetic Cardiomyopathy?. *Journal of the American College of Cardiology*, **2016**, 68, 949-66	15.1	133
52	Genetic causes of dilated cardiomyopathy. *Heart*, **2016**, 102, 2004-2014	5.1	16
51	Cardio-Oncology: The Carney Complex Type I. *Journal of the American College of Cardiology*, **2016**, 68, 1921-1923	15.1	2
50	Clinical Pregenetic Screening for Stroke Monogenic Diseases: Results From Lombardia GENS Registry. *Stroke*, **2016**, 47, 1702-9	6.7	27

49	Atlas of the clinical genetics of human dilated cardiomyopathy. *European Heart Journal*, **2015**, 36, 1123-35a	9.9	334
48	The MOGE(S) classification of cardiomyopathy for clinicians. *Journal of the American College of Cardiology*, **2014**, 64, 304-18	15.1	107
47	Incomplete penetrance of GLMN gene c.395-1G>C mutation in a family with glomuvenous malformations. *International Journal of Dermatology*, **2014**, 53, 1362-4	1.7	3
46	Glomuvenous malformations with smooth muscle and eccrine glands: unusual histopathologic features in a familial setting. *Journal of Cutaneous Pathology*, **2014**, 41, 308-15	1.7	8
45	Autosomal recessive paediatric sick sinus syndrome associated with novel compound mutations in SCN5A. *International Journal of Cardiology*, **2013**, 167, 3078-80	3.2	10
44	Gender-specific differences in major cardiac events and mortality in lamin A/C mutation carriers. *European Journal of Heart Failure*, **2013**, 15, 376-84	12.3	97
43	Autosomal recessive atrial dilated cardiomyopathy with standstill evolution associated with mutation of Natriuretic Peptide Precursor A. *Circulation: Cardiovascular Genetics*, **2013**, 6, 27-36		35
42	In-frame mutations in exon 1 of SKI cause dominant Shprintzen-Goldberg syndrome. *American Journal of Human Genetics*, **2012**, 91, 950-7	11	80
41	Quantitative expression of the mutated lamin A/C gene in patients with cardiolaminopathy. *Journal of the American College of Cardiology*, **2012**, 60, 1916-20	15.1	23
40	Diagnostic work-up and risk stratification in X-linked dilated cardiomyopathies caused by dystrophin defects. *Journal of the American College of Cardiology*, **2011**, 58, 925-34	15.1	63
39	A novel mutation of the glomulin gene in an Italian family with autosomal dominant cutaneous glomuvenous malformations. *Experimental Dermatology*, **2011**, 20, 1032-4	4	9
38	Risk of dissection in thoracic aneurysms associated with mutations of smooth muscle alpha-actin 2 (ACTA2). *Heart*, **2011**, 97, 321-6	5.1	51
37	When should cardiologists suspect Anderson-Fabry disease?. *American Journal of Cardiology*, **2010**, 106, 1492-9	3	39
36	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, 391-8	7.4	120
35	The shortness of Pygmies is associated with severe under-expression of the growth hormone receptor. *Molecular Genetics and Metabolism*, **2009**, 98, 310-3	3.7	41
34	Transcriptomic and proteomic analysis in the cardiovascular setting: unravelling the disease?. *Journal of Cardiovascular Medicine*, **2009**, 10, 433-42	1.9	8
33	Long-term outcome and risk stratification in dilated cardiolaminopathies. *Journal of the American College of Cardiology*, **2008**, 52, 1250-60	15.1	278
32	Letter by Maurizia Grasso et al. regarding article, "Restrictive cardiomyopathy with atrioventricular conduction block resulting from a desmin mutation". *International Journal of Cardiology*, **2008**, 131, 144-5; author reply 146-7	3.2	2

31	Barth syndrome associated with compound hemizygosity and heterozygosity of the TAZ and LDB3 genes. *American Journal of Medical Genetics, Part A*, **2007**, 143A, 907-15	2.5	40
30	Desmin accumulation restrictive cardiomyopathy and atrioventricular block associated with desmin gene defects. *European Journal of Heart Failure*, **2006**, 8, 477-83	12.3	120
29	alphaB-crystallin mutation in dilated cardiomyopathies: low prevalence in a consecutive series of 200 unrelated probands. *Biochemical and Biophysical Research Communications*, **2006**, 346, 1115-7	3.4	46
28	Early diagnosis of Wilson Disease in a six-year-old child. *Journal of Pediatrics*, **2006**, 148, 141	3.6	2
27	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, 34-8	5.3	54
26	Kaposiß sarcoma in transplant and HIV-infected patients: an epidemiologic study in Italy and France. *Transplantation*, **2005**, 80, 1699-704	1.8	42
25	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	4.7	74
24	Autosomal dominant dilated cardiomyopathy with atrioventricular block: a lamin A/C defect-related disease. *Journal of the American College of Cardiology*, **2002**, 39, 981-90	15.1	257
23	The mitochondrial DNA mutation T12297C affects a highly conserved nucleotide of tRNA(Leu(CUN)) and is associated with dilated cardiomyopathy. *European Journal of Human Genetics*, **2001**, 9, 311-5	5.3	35
22	Prevalence and characteristics of dystrophin defects in adult male patients with dilated cardiomyopathy. *Journal of the American College of Cardiology*, **2000**, 35, 1760-8	15.1	70
21	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, 645-53	15.1	98
20	Mitochondrial DNA mutations and mitochondrial abnormalities in dilated cardiomyopathy. *American Journal of Pathology*, **1998**, 153, 1501-10	5.8	190
19	Enteroviral RNA and virus-like particles in the skeletal muscle of patients with idiopathic dilated cardiomyopathy. *American Journal of Cardiology*, **1997**, 80, 1188-93	3	16
18	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	2.2	5
17	Frequency and characteristics of coronary thrombosis in the epicardial coronary arteries after cardiac transplantation. *American Journal of Cardiology*, **1996**, 78, 795-800	3	17
16	Human cytomegalovirus early infection, acute rejection, and major histocompatibility class II expression in transplanted lung. Molecular, immunocytochemical, and histopathologic investigations. *Transplantation*, **1996**, 61, 418-27	1.8	19
15	Cardiac immunocyte-derived (AL) amyloidosis: an endomyocardial biopsy study in 11 patients. *American Heart Journal*, **1995**, 130, 528-36	4.9	40
14	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, 675-82	3	65

13	Coronary thrombosis in non-cardiac death. *Coronary Artery Disease*, **1993**, 4, 751-9	1.4	45
12	Expression of proliferating cell markers in normal and diseased human hearts. *American Journal of Cardiology*, **1993**, 72, 608-14	3	17
11	Morphologic changes induced by acetylcholine infusion in normal and atherosclerotic coronary arteries. *American Journal of Cardiology*, **1993**, 71, 1382-90	3	9
10	Localization of brain and atrial natriuretic peptide in human and porcine heart. *International Journal of Cardiology*, **1992**, 34, 237-47	3.2	32
9	Search for Coxsackievirus B3 RNA in idiopathic dilated cardiomyopathy using gene amplification by polymerase chain reaction. *American Journal of Cardiology*, **1992**, 69, 658-64	3	86
8	H and L ferritins in myocardium in iron overload. *American Journal of Cardiology*, **1991**, 68, 1233-6	3	7
7	Coronary atherosclerotic plaques with and without thrombus in ischemic heart syndromes: a morphologic, immunohistochemical, and biochemical study. *American Journal of Cardiology*, **1991**, 68, 36B-50B	3	116
6	Endomyocardial biopsy finding in two patients with idiopathic dilated cardiomyopathy receiving long-term treatment with amiodarone. *American Journal of Cardiology*, **1991**, 67, 661-2	3	4
5	Atrial amyloid deposits in the failing human heart display both atrial and brain natriuretic peptide-like immunoreactivity. *Journal of Pathology*, **1991**, 165, 235-41	9.4	34
4	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, 973-80	3	38
3	Myocardial iron grading by endomyocardial biopsy. A clinico-pathologic study on iron overloaded patients. *European Journal of Haematology*, **1989**, 42, 382-8	3.8	45
2	The morphologic spectrum of dilated cardiomyopathy and its relation to immune-response genes. *American Journal of Cardiology*, **1989**, 64, 991-5	3	41
1	Genome wide association analysis in dilated cardiomyopathy reveals two new key players in systolic heart failure on chromosome 3p25.1 and 22q11.23		1