

Maurizia Grasso

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

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	Atlas of the clinical genetics of human dilated cardiomyopathy. <i>European Heart Journal</i> , 2015 , 36, 1123-35	15.1	334
65	Long-term outcome and risk stratification in dilated cardiomyopathies. <i>Journal of the American College of Cardiology</i> , 2008 , 52, 1250-60	15.1	278
64	Autosomal dominant dilated cardiomyopathy with atrioventricular block: a lamin A/C defect-related disease. <i>Journal of the American College of Cardiology</i> , 2002 , 39, 981-90	15.1	257
63	Mitochondrial DNA mutations and mitochondrial abnormalities in dilated cardiomyopathy. <i>American Journal of Pathology</i> , 1998 , 153, 1501-10	5.8	190
62	Left Ventricular Noncompaction: A Distinct Genetic Cardiomyopathy?. <i>Journal of the American College of Cardiology</i> , 2016 , 68, 949-66	15.1	133
61	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. <i>Pediatrics</i> , 2009 , 123, 391-8	7.4	120
60	Desmin accumulation restrictive cardiomyopathy and atrioventricular block associated with desmin gene defects. <i>European Journal of Heart Failure</i> , 2006 , 8, 477-83	12.3	120
59	Coronary atherosclerotic plaques with and without thrombus in ischemic heart syndromes: a morphologic, immunohistochemical, and biochemical study. <i>American Journal of Cardiology</i> , 1991 , 68, 36B-50B	3	116
58	The MOGE(S) classification of cardiomyopathy for clinicians. <i>Journal of the American College of Cardiology</i> , 2014 , 64, 304-18	15.1	107
57	Restrictive cardiomyopathy, atrioventricular block and mild to subclinical myopathy in patients with desmin-immunoreactive material deposits. <i>Journal of the American College of Cardiology</i> , 1998 , 31, 645-53	15.1	98
56	Gender-specific differences in major cardiac events and mortality in lamin A/C mutation carriers. <i>European Journal of Heart Failure</i> , 2013 , 15, 376-84	12.3	97
55	Search for Coxsackievirus B3 RNA in idiopathic dilated cardiomyopathy using gene amplification by polymerase chain reaction. <i>American Journal of Cardiology</i> , 1992 , 69, 658-64	3	86
54	In-frame mutations in exon 1 of SKI cause dominant Shprintzen-Goldberg syndrome. <i>American Journal of Human Genetics</i> , 2012 , 91, 950-7	11	80
53	Identification of sixty-two novel and twelve known FBN1 mutations in eighty-one unrelated probands with Marfan syndrome and other fibrillinopathies. <i>Human Mutation</i> , 2005 , 26, 494	4.7	74
52	Prevalence and characteristics of dystrophin defects in adult male patients with dilated cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2000 , 35, 1760-8	15.1	70
51	Comparison of coronary lesions obtained by directional coronary atherectomy in unstable angina, stable angina, and restenosis after either atherectomy or angioplasty. <i>American Journal of Cardiology</i> , 1995 , 75, 675-82	3	65
50	Diagnostic work-up and risk stratification in X-linked dilated cardiomyopathies caused by dystrophin defects. <i>Journal of the American College of Cardiology</i> , 2011 , 58, 925-34	15.1	63

49	Two novel and one known mutation of the TGFBR2 gene in Marfan syndrome not associated with FBN1 gene defects. <i>European Journal of Human Genetics</i> , 2006 , 14, 34-8	5.3	54
48	Risk of dissection in thoracic aneurysms associated with mutations of smooth muscle alpha-actin 2 (ACTA2). <i>Heart</i> , 2011 , 97, 321-6	5.1	51
47	alphaB-crystallin mutation in dilated cardiomyopathies: low prevalence in a consecutive series of 200 unrelated probands. <i>Biochemical and Biophysical Research Communications</i> , 2006 , 346, 1115-7	3.4	46
46	Myocardial iron grading by endomyocardial biopsy. A clinico-pathologic study on iron overloaded patients. <i>European Journal of Haematology</i> , 1989 , 42, 382-8	3.8	45
45	Coronary thrombosis in non-cardiac death. <i>Coronary Artery Disease</i> , 1993 , 4, 751-9	1.4	45
44	KaposiB sarcoma in transplant and HIV-infected patients: an epidemiologic study in Italy and France. <i>Transplantation</i> , 2005 , 80, 1699-704	1.8	42
43	Cardiac Phenotypes in HereditaryMuscleDisorders: JACC State-of-the-Art Review. <i>Journal of the American College of Cardiology</i> , 2018 , 72, 2485-2506	15.1	42
42	The shortness of Pygmies is associated with severe under-expression of the growth hormone receptor. <i>Molecular Genetics and Metabolism</i> , 2009 , 98, 310-3	3.7	41
41	The morphologic spectrum of dilated cardiomyopathy and its relation to immune-response genes. <i>American Journal of Cardiology</i> , 1989 , 64, 991-5	3	41
40	Barth syndrome associated with compound hemizygoty and heterozygoty of the TAZ and LDB3 genes. <i>American Journal of Medical Genetics, Part A</i> , 2007 , 143A, 907-15	2.5	40
39	Cardiac immunocyte-derived (AL) amyloidosis: an endomyocardial biopsy study in 11 patients. <i>American Heart Journal</i> , 1995 , 130, 528-36	4.9	40
38	When should cardiologists suspect Anderson-Fabry disease?. <i>American Journal of Cardiology</i> , 2010 , 106, 1492-9	3	39
37	Expression of natriuretic peptide in ventricular myocardium of failing human hearts and its correlation with the severity of clinical and hemodynamic impairment. <i>American Journal of Cardiology</i> , 1990 , 66, 973-80	3	38
36	Genetic Screening of Anderson-Fabry Disease in Probands Referred From Multispecialty Clinics. <i>Journal of the American College of Cardiology</i> , 2016 , 68, 1037-50	15.1	37
35	Autosomal recessive atrial dilated cardiomyopathy with standstill evolution associated with mutation of Natriuretic Peptide Precursor A. <i>Circulation: Cardiovascular Genetics</i> , 2013 , 6, 27-36		35
34	The mitochondrial DNA mutation T12297C affects a highly conserved nucleotide of tRNA(Leu(CUN)) and is associated with dilated cardiomyopathy. <i>European Journal of Human Genetics</i> , 2001 , 9, 311-5	5.3	35
33	Atrial amyloid deposits in the failing human heart display both atrial and brain natriuretic peptide-like immunoreactivity. <i>Journal of Pathology</i> , 1991 , 165, 235-41	9.4	34
32	Localization of brain and atrial natriuretic peptide in human and porcine heart. <i>International Journal of Cardiology</i> , 1992 , 34, 237-47	3.2	32

31	Clinical Pre-genetic Screening for Stroke Monogenic Diseases: Results From Lombardia GENS Registry. <i>Stroke</i> , 2016 , 47, 1702-9	6.7	27
30	Quantitative expression of the mutated lamin A/C gene in patients with cardiolaminopathy. <i>Journal of the American College of Cardiology</i> , 2012 , 60, 1916-20	15.1	23
29	Human cytomegalovirus early infection, acute rejection, and major histocompatibility class II expression in transplanted lung. Molecular, immunocytochemical, and histopathologic investigations. <i>Transplantation</i> , 1996 , 61, 418-27	1.8	19
28	Frequency and characteristics of coronary thrombosis in the epicardial coronary arteries after cardiac transplantation. <i>American Journal of Cardiology</i> , 1996 , 78, 795-800	3	17
27	Expression of proliferating cell markers in normal and diseased human hearts. <i>American Journal of Cardiology</i> , 1993 , 72, 608-14	3	17
26	Genetic causes of dilated cardiomyopathy. <i>Heart</i> , 2016 , 102, 2004-2014	5.1	16
25	Enteroviral RNA and virus-like particles in the skeletal muscle of patients with idiopathic dilated cardiomyopathy. <i>American Journal of Cardiology</i> , 1997 , 80, 1188-93	3	16
24	RE: BRCA2 Polymorphic Stop Codon K3326X and the Risk of Breast, Prostate, and Ovarian Cancers. <i>Journal of the National Cancer Institute</i> , 2016 , 108,	9.7	14
23	Genome-wide association analysis in dilated cardiomyopathy reveals two new players in systolic heart failure on chromosomes 3p25.1 and 22q11.23. <i>European Heart Journal</i> , 2021 , 42, 2000-2011	9.5	14
22	Management of the axilla in patients with breast cancer and positive sentinel lymph node biopsy: An evidence-based update in a European breast center. <i>European Journal of Surgical Oncology</i> , 2020 , 46, 15-23	3.6	11
21	Autosomal recessive paediatric sick sinus syndrome associated with novel compound mutations in SCN5A. <i>International Journal of Cardiology</i> , 2013 , 167, 3078-80	3.2	10
20	A novel mutation of the glomulin gene in an Italian family with autosomal dominant cutaneous glomuvenous malformations. <i>Experimental Dermatology</i> , 2011 , 20, 1032-4	4	9
19	Morphologic changes induced by acetylcholine infusion in normal and atherosclerotic coronary arteries. <i>American Journal of Cardiology</i> , 1993 , 71, 1382-90	3	9
18	Glomuvenous malformations with smooth muscle and eccrine glands: unusual histopathologic features in a familial setting. <i>Journal of Cutaneous Pathology</i> , 2014 , 41, 308-15	1.7	8
17	Transcriptomic and proteomic analysis in the cardiovascular setting: unravelling the disease?. <i>Journal of Cardiovascular Medicine</i> , 2009 , 10, 433-42	1.9	8
16	H and L ferritins in myocardium in iron overload. <i>American Journal of Cardiology</i> , 1991 , 68, 1233-6	3	7
15	Hereditary Hyperferritinemia-Cataract Syndrome: Relationship Between Phenotypes and Specific Mutations in the Iron-Responsive Element of Ferritin Light-Chain mRNA. <i>Blood</i> , 1997 , 90, 814-821	2.2	5
14	Endomyocardial biopsy finding in two patients with idiopathic dilated cardiomyopathy receiving long-term treatment with amiodarone. <i>American Journal of Cardiology</i> , 1991 , 67, 661-2	3	4

13	Myths to debunk: the non-compacted myocardium. <i>European Heart Journal Supplements</i> , 2020 , 22, L6-L10.5	10.5	4
12	Incomplete penetrance of GLMN gene c.395-1G>C mutation in a family with glomuvenous malformations. <i>International Journal of Dermatology</i> , 2014 , 53, 1362-4	1.7	3
11	Cardio-Oncology: The Carney Complex Type I. <i>Journal of the American College of Cardiology</i> , 2016 , 68, 1921-1923	15.1	2
10	Reply: A Distinct Cardiomyopathy: HCN4 Syndrome Comprising Myocardial Noncompaction, Bradycardia, Mitral Valve Defects, and Aortic Dilation. <i>Journal of the American College of Cardiology</i> , 2017 , 69, 1210-1211	15.1	2
9	Letter by Maurizia Grasso et al. regarding article, "Restrictive cardiomyopathy with atrioventricular conduction block resulting from a desmin mutation". <i>International Journal of Cardiology</i> , 2008 , 131, 144-5; author reply 146-7	3.2	2
8	Early diagnosis of Wilson Disease in a six-year-old child. <i>Journal of Pediatrics</i> , 2006 , 148, 141	3.6	2
7	Hereditary muscle diseases and the heart: the cardiologist's perspective. <i>European Heart Journal Supplements</i> , 2020 , 22, E13-E19	1.5	2
6	Genetic counselling and high-penetrance susceptibility gene analysis reveal the novel CDKN2A p.D84V (c.251A>T) mutation in melanoma-prone families from Italy. <i>Melanoma Research</i> , 2017 , 27, 97-103	3.3	1
5	Clinical heterogeneity and reduced penetrance in DICER1 syndrome: a report of three families. <i>Tumori</i> , 2021 , 107, NP144-NP148	1.7	1
4	Genome wide association analysis in dilated cardiomyopathy reveals two new key players in systolic heart failure on chromosome 3p25.1 and 22q11.23		1
3	Spectrum of phenotype of ventricular noncompaction in adults. <i>Progress in Pediatric Cardiology</i> , 2021 , 62, 101416	0.4	1
2	Rare exon 10 deletion in POLH gene in a family with xeroderma pigmentosum variant correlating with protein expression by immunohistochemistry. <i>Giornale Italiano Di Dermatologia E Venereologia</i> , 2020 , 155, 349-354	0.8	0
1	Genetics and clinics: current applications, limitations, and future developments. <i>European Heart Journal Supplements</i> , 2019 , 21, B7-B14	1.5	