Guillaume Jondeau

List of Publications by Year
in descending order

59 papers	7,912 citations	136950 32 h-index	133252 59 g-index
60 all docs	60 docs citations	60 times ranked	7566 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.	2.2	32
2	The VASCERN European Reference Network: An overview. European Journal of Medical Genetics, 2022, 65, 104420.	1.3	3
3	Is physical activity a future therapy for patients with Marfan syndrome?. Orphanet Journal of Rare Diseases, 2022, 17, 46.	2.7	2
4	Non-Dissecting Distal Aortic and Peripheral Arterial Aneurysms in Patients With Marfan Syndrome. Frontiers in Cardiovascular Medicine, 2022, 9, 827357.	2.4	2
5	Arrhythmia and impaired myocardial function in heritable thoracic aortic disease: An international retrospective cohort study. European Journal of Medical Genetics, 2022, 65, 104503.	1.3	4
6	Staged hybrid repair of type II thoracoabdominal aneurysms. Journal of Vascular Surgery, 2021, 74, 20-27.	1.1	6
7	2020 ESC Guidelines for the management of adult congenital heart disease. European Heart Journal, 2021, 42, 563-645.	2.2	971
8	Clinical relevance of genotypeâ€“phenotype correlations beyond vascular events in a cohort study of 1500 Marfan syndrome patients with FBN1 pathogenic variants. Genetics in Medicine, 2021, 23, 1296-1304.	2.4	63
9	eHealth for patients with rare diseases: the eHealth Working Group of the European Reference Network on Rare Multisystemic Vascular Diseases (VASCERN). Orphanet Journal of Rare Diseases, 2021, 16, 164.	2.7	3
10	Coronavirus disease vaccination in heart failure: No time to waste. Archives of Cardiovascular Diseases, 2021, 114, 434-438.	1.6	6
11	Marfan syndrome. Nature Reviews Disease Primers, 2021, 7, 64.	30.5	99
12	Inhibition of HIPK2 Alleviates Thoracic Aortic Disease in Mice With Progressively Severe Marfan Syndrome. Arteriosclerosis, Thrombosis, and Vascular Biology, 2021, 41, 2483-2493.	2.4	4
13	Pregnancy outcome in thoracic aortic disease data from the Registry Of Pregnancy And Cardiac disease. Heart, 2021, 107, 1704-1709.	2.9	29
14	Preliminary Experience With Custom Made Hourglass Shaped Thoracic Stent Grafts for Endovascular Thoracic Aortic Coarctation Repair in Adults. European Journal of Vascular and Endovascular Surgery, 2021, 62, 1000-1001.	1.5	2
15	Optimising Aortic Endovascular Repair in Patients with Marfan Syndrome. European Journal of Vascular and Endovascular Surgery, 2020, 59, 577-585.	1.5	35
16	Incidence of cardiovascular events and risk markers in a prospective study of children diagnosed with Marfan syndrome. Archives of Cardiovascular Diseases, 2020, 113, 40-49.	1.6	12
17	Marfan sartan saga, episode X. European Heart Journal, 2020, 41, 4188-4190.	2.2	4
18	Pathogenic FBN1 Genetic Variation and Aortic Dissection in Patients With MarfanÂSyndrome. Journal of the American College of Cardiology, 2020, 75, 843-853.	2.8	38

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19	Quantifying the Genetic Basis of Marfan Syndrome Clinical Variability. Genes, 2020, 11, 574.	2.4	11
20	Pregnancy outcomes in women with cardiovascular disease: evolving trends over 10 years in the ESC Registry Of Pregnancy And Cardiac disease (ROPAC). European Heart Journal, 2019, 40, 3848-3855.	2.2	209
21	False lumen embolization in chronic aortic dissection promotes thoracic aortic remodeling at midterm follow-up. Journal of Vascular Surgery, 2019, 70, 710-717.	1.1	45
22	Reference Expression Profile of Three FBN1 Transcript Isoforms and Their Association with Clinical Variability in Marfan Syndrome. Genes, 2019, 10, 128.	2.4	6
23	European reference network for rare vascular diseases (VASCERN) consensus statement for the screening and management of patients with pathogenic ACTA2 variants. Orphanet Journal of Rare Diseases, 2019, 14, 264.	2.7	23
24	Risk of Ascending Aortic Aneurysm in Patients With Autosomal Dominant Polycystic Kidney Disease. American Journal of Cardiology, 2019, 123, 482-488.	1.6	16
25	Impact of an interatrial shunt device on survival and heart failure hospitalization in patients with preserved ejection fraction. ESC Heart Failure, 2019, 6, 62-69.	3.1	45
26	Systems pharmacologyâ€“based integration of human and mouse data for drug repurposing to treat thoracic aneurysms. JCI Insight, 2019, 4, .	5.0	21
27	From genetics to response to injury: vascular smooth muscle cells in aneurysms and dissections of the ascending aorta. Cardiovascular Research, 2018, 114, 578-589.	3.8	114
28	Skeletal evolution in Marfan syndrome: growth curves from a French national cohort. Pediatric Research, 2018, 83, 71-77.	2.3	6
29	Association of modifiers and other genetic factors explain Marfan syndrome clinical variability. European Journal of Human Genetics, 2018, 26, 1759-1772.	2.8	73
30	Marfan Syndrome Variability: Investigation of the Roles of Sarcolipin and Calcium as Potential Transregulator of FBN1 Expression. Genes, 2018, 9, 421.	2.4	4
31	Marfan Syndrome. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	2
32	Characterization of ¹⁸ F-Fluorodeoxyglucose Uptake Pattern in Noninfected Prosthetic Heart Valves. Circulation: Cardiovascular Imaging, 2017, 10, e005585.	2.6	75
33	Response by Mathieu et al to Letter Regarding Article, â€œCharacterization of 18 F-Fluorodeoxyglucose Uptake Pattern in Noninfected Prosthetic Heart Valvesâ€• Circulation: Cardiovascular Imaging, 2017, 10, .	2.6	3
34	Is Transesophageal Echocardiography Needed before Hospital Discharge in Patients after Bentall Surgery?. Journal of the American Society of Echocardiography, 2017, 30, 52-58.	2.8	1
35	LOX Mutations Predispose to Thoracic Aortic Aneurysms and Dissections. Circulation Research, 2016, 118, 928-934.	4.5	180
36	International Registry of Patients Carrying TGFBR1 or TGFBR2 Mutations. Circulation: Cardiovascular Genetics, 2016, 9, 548-558.	5.1	145

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37	The clinical presentation of Marfan syndrome is modulated by expression of wild-type FBN1 allele. Human Molecular Genetics, 2015, 24, 2764-2770.	2.9	57
38	Marfan Sartan: a randomized, double-blind, placebo-controlled trial. European Heart Journal, 2015, 36, 2160-2166.	2.2	179
39	Multimodality Imaging of Diseases of the Thoracic Aorta in Adults: From the American Society of Echocardiography and the European Association of Cardiovascular Imaging. Journal of the American Society of Echocardiography, 2015, 28, 119-182.	2.8	500
40	Comparative assessment of ascending aortic aneurysms in Marfan patients using ECG-gated computerized tomographic angiography versus trans-thoracic echocardiography. International Journal of Cardiology, 2015, 184, 22-27.	1.7	21
41	Beta-Blockers in Acute Heart Failure. JACC: Heart Failure, 2015, 3, 654-656.	4.1	8
42	Aortic Disease Presentation and Outcome Associated With ACTA2 Mutations. Circulation: Cardiovascular Genetics, 2015, 8, 457-464.	5.1	117
43	Loeysâ€“Dietz syndrome is a specific phenotype and not a concomitant of any mutation in a gene involved in TGF-Î² signaling. Genetics in Medicine, 2014, 16, 641-642.	2.4	18
44	MFAP5 Loss-of-Function Mutations Underscore the Involvement of Matrix Alteration in the Pathogenesis of Familial Thoracic Aortic Aneurysms and Dissections. American Journal of Human Genetics, 2014, 95, 736-743.	6.2	110
45	Aortic dilatation patterns and rates in adults with bicuspid aortic valves: a comparative study with Marfan syndrome and degenerative aortopathy. Heart, 2014, 100, 126-134.	2.9	190
46	Study of phenotype evolution during childhood in Marfan syndrome to improve clinical recognition. Genetics in Medicine, 2014, 16, 246-250.	2.4	45
47	Current aspects of the spectrum of acute heart failure syndromes in a realâ€life setting: the OFICA study. European Journal of Heart Failure, 2013, 15, 465-476.	7.1	135
48	Aortic Event Rate in the Marfan Population. Circulation, 2012, 125, 226-232.	1.6	165
49	TGFB2 mutations cause familial thoracic aortic aneurysms and dissections associated with mild systemic features of Marfan syndrome. Nature Genetics, 2012, 44, 916-921.	21.4	319
50	Genetics of Thoracic Aortic Aneurysms. Current Atherosclerosis Reports, 2012, 14, 219-226.	4.8	40
51	Prognosis Factors in Probands With an FBN1 Mutation Diagnosed Before the Age of 1 Year. Pediatric Research, 2011, 69, 265-270.	2.3	59
52	Nomograms for Aortic Root Diameters in Children Using Two-Dimensional Echocardiography. American Journal of Cardiology, 2010, 105, 888-894.	1.6	140
53	Cardiovascular manifestations in men and women carrying a FBN1 mutation. European Heart Journal, 2010, 31, 2223-2229.	2.2	133
54	The revised Ghent nosology for the Marfan syndrome. Journal of Medical Genetics, 2010, 47, 476-485.	3.2	1,677

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55	Comparison of Clinical Presentations and Outcomes Between Patients With TGFBR2 and FBN1 Mutations in Marfan Syndrome and Related Disorders. Circulation, 2009, 120, 2541-2549.	1.6	203
56	Identification of the minimal combination of clinical features in probands for efficient mutation detection in the FBN1 gene. European Journal of Human Genetics, 2009, 17, 1121-1128.	2.8	82
57	B-CONVINCED: Beta-blocker CONtinuation Vs. INterruption in patients with Congestive heart failure hospitalizED for a decompensation episode. European Heart Journal, 2009, 30, 2186-2192.	2.2	128
58	Executive summary of the guidelines on the diagnosis and treatment of acute heart failure: The Task Force on Acute Heart Failure of the European Society of Cardiology. European Heart Journal, 2005, 26, 384-416.	2.2	1,114
59	Central Pulse Pressure Is a Major Determinant of Ascending Aorta Dilation in Marfan Syndrome. Circulation, 1999, 99, 2677-2681.	1.6	178