

CITATION REPORT

List of articles citing

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

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#	Paper	IF	Citations
61	Transforming growth factor- β : a biomarker in Marfan syndrome?. <i>Circulation</i> , 2009 , 120, 464-6	16.7	7
60	Biomarkers of abdominal aortic aneurysm progression. Part 2: inflammation. <i>Nature Reviews Cardiology</i> , 2009 , 6, 543-52	14.8	109
59	Drug-based therapies for vascular disease in Marfan syndrome: from mouse models to human patients. <i>Mount Sinai Journal of Medicine</i> , 2010 , 77, 366-73		9
58	Spontaneous arterial dissection: phenotype and molecular pathogenesis. <i>Cellular and Molecular Life Sciences</i> , 2010 , 67, 1799-815	10.3	32
57	Impact of Mendelian inheritance in cardiovascular disease. <i>Annals of the New York Academy of Sciences</i> , 2010 , 1214, 122-37	6.5	10
56	Adults with genetic syndromes. <i>International Journal on Disability and Human Development</i> , 2010 , 9,		
55	Does medical therapy for thoracic aortic aneurysms really work? Are beta-blockers truly indicated? <i>CON. Cardiology Clinics</i> , 2010 , 28, 261-9	2.5	13
54	Rationale and design of a randomized clinical trial (Marfan Sartan) of angiotensin II receptor blocker therapy versus placebo in individuals with Marfan syndrome. <i>Archives of Cardiovascular Diseases</i> , 2010 , 103, 317-25	2.7	59
53	Marfan Syndrome—An Echocardiographer’s Perspective. <i>Journal of Medical Ultrasound</i> , 2011 , 19, 1-6	0.8	2
52	Preventing the aortic complications of Marfan syndrome: a case-example of translational genomic medicine. <i>British Journal of Clinical Pharmacology</i> , 2011 , 72, 6-17	3.8	6
51	The role of the renin-angiotensin system in thoracic aortic aneurysms: clinical implications. <i>Pharmacology & Therapeutics</i> , 2011 , 131, 50-60	13.9	40
50	Genetic dissection of marfan syndrome and related connective tissue disorders: an update 2012. <i>Molecular Syndromology</i> , 2012 , 3, 47-58	1.5	29
49	Marfan syndrome: from gene to therapy. <i>Current Opinion in Pediatrics</i> , 2012 , 24, 498-504	3.2	44
48	Aortic event rate in the Marfan population: a cohort study. <i>Circulation</i> , 2012 , 125, 226-32	16.7	117
47	The Ghent Marfan Trial—a randomized, double-blind placebo controlled trial with losartan in Marfan patients treated with β blockers. <i>International Journal of Cardiology</i> , 2012 , 157, 354-8	3.2	51
46	An evidence-based hypothesis for beneficial effects of telmisartan on Marfan syndrome. <i>International Journal of Cardiology</i> , 2012 , 158, 101-2	3.2	4
45	Surgical management of patients with Marfan syndrome: evolution throughout the years. <i>Archives of Cardiovascular Diseases</i> , 2012 , 105, 84-90	2.7	4

44	Extra-aortic identifiers to guide genetic testing in familial thoracic aortic aneurysms and dissections syndromes: it is all about the company one keeps. <i>Journal of the American College of Cardiology</i> , 2012 , 60, 404-7	15.1	3
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42	Effects of atenolol, perindopril and verapamil on haemodynamic and vascular function in Marfan syndrome - a randomised, double-blind, crossover trial. <i>European Journal of Clinical Investigation</i> , 2012 , 42, 891-9	4.6	31
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40	Aortic dilatation and aortopathy in congenital heart diseases. <i>Journal of Cardiology</i> , 2013 , 61, 16-21	3	28
39	Genetic biomarkers in aortopathy. <i>Biomarkers in Medicine</i> , 2013 , 7, 547-63	2.3	6
38	Medical treatment for thoracic aortic aneurysm - much more work to be done. <i>Progress in Cardiovascular Diseases</i> , 2013 , 56, 103-8	8.5	8
37	High prevalence of eosinophilic esophagitis in patients with inherited connective tissue disorders. <i>Journal of Allergy and Clinical Immunology</i> , 2013 , 132, 378-86	11.5	114
36	Educational paper. Connective tissue disorders with vascular involvement: from gene to therapy. <i>European Journal of Pediatrics</i> , 2013 , 172, 997-1005	4.1	22
35	Proteomics in aortic aneurysm--what have we learnt so far?. <i>Proteomics - Clinical Applications</i> , 2013 , 7, 504-15	3.1	5
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33	Medical treatment of aortic aneurysms in Marfan syndrome and other heritable conditions. <i>Current Cardiology Reviews</i> , 2014 , 10, 161-71	2.4	27
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20	Histopathology of aortic complications in bicuspid aortic valve versus Marfan syndrome: relevance for therapy?. <i>Heart and Vessels</i> , 2016 , 31, 795-806	2.1	27
19	Editor's Choice - Management of Descending Thoracic Aorta Diseases: Clinical Practice Guidelines of the European Society for Vascular Surgery (ESVS). <i>European Journal of Vascular and Endovascular Surgery</i> , 2017 , 53, 4-52	2.3	487
18	'Precision and personalized medicine,' a dream that comes true?. <i>Journal of Cardiovascular Medicine</i> , 2017 , 18 Suppl 1, e1-e6	1.9	5
17	Beta-blockers for preventing aortic dissection in Marfan syndrome. <i>The Cochrane Library</i> , 2017 , 11, CD011103	1.03	15
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15	Current Status of Medical Therapy of Thoracic Aortic Aneurysm and Dissection. 2018 , 235-249		
14	Precise Therapy for Thoracic Aortic Aneurysm in Marfan Syndrome: A Puzzle Nearing Its Solution. <i>Progress in Cardiovascular Diseases</i> , 2018 , 61, 328-335	8.5	13
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9	The Future of Dupuytren's Research and Treatment. 2012 , 455-470		1

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6	Association between Oro-Facial Defects and Systemic Alterations in Children Affected by Marfan Syndrome. <i>Journal of Clinical and Diagnostic Research JCDR</i> , 2013 , 7, 700-3	0	7
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4	Loeys-Dietz Syndrome. <i>Advances in Experimental Medicine and Biology</i> , 2021 , 1348, 251-264	3.6	1
3	Aortic Wall Inflammation in the Pathogenesis, Diagnosis and Treatment of Aortic Aneurysms.. <i>Inflammation</i> , 2022 , 1	5.1	0
2	NADPH Oxidases in Aortic Aneurysms. 2022 , 11, 1830		0
1	Angiotensin receptor blockers and β blockers in Marfan syndrome: an individual patient data meta-analysis of randomised trials. 2022 , 400, 822-831		1