Marjolijn Renard

List of Publications by Year
in descending order

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Version: 2024-02-01

18	932	1039880 9	794469 19
papers	citations	h-index	g-index

19	19	19	1570
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Molecular characterization and investigation of the role of genetic variation in phenotypic variability and response to treatment in a large pediatric Marfan syndrome cohort. Genetics in Medicine, 2022, 24, 1045-1053.	1.1	13
2	Pathogenic variants in THSD4, encoding the ADAMTS-like 6 protein, predispose to inherited thoracic aortic aneurysm. Genetics in Medicine, 2021, 23, 111-122.	1.1	25
3	MEK1/2 Inhibition in Murine Heart and Aorta After Oral Administration of Refametinib Supplemented Drinking Water. Frontiers in Pharmacology, 2020, 11, 1336.	1.6	4
4	Spontaneous Right Ventricular Pseudoaneurysms and Increased Arrhythmogenicity in a Mouse Model of Marfan Syndrome. International Journal of Molecular Sciences, 2020, 21, 7024.	1.8	3
5	Ambulatory Electrocardiographic Monitoring and Ectopic Beat Detection in Conscious Mice. Sensors, 2020, 20, 3867.	2.1	6
6	Variants in ADRB1 and CYP2C9: Association with Response to Atenolol and Losartan in Marfan Syndrome. Journal of Pediatrics, 2020, 222, 213-220.e5.	0.9	8
7	Slc2a10 knock-out mice deficient in ascorbic acid synthesis recapitulate aspects of arterial tortuosity syndrome and display mitochondrial respiration defects. Human Molecular Genetics, 2020, 29, 1476-1488.	1.4	5
8	SmgGDS, a new piece in the thoracic aortic aneurysm and dissection puzzle. Journal of Thoracic Disease, 2018, 10, S4133-S4136.	0.6	2
9	Expressed repetitive elements are broadly applicable reference targets for normalization of reverse transcription-qPCR data in mice. Scientific Reports, 2018, 8, 7642.	1.6	10
10	A heart for fibrillin: spatial arrangement in adult wild-type murine myocardial tissue. Histochemistry and Cell Biology, 2018, 150, 271-280.	0.8	11
11	Clinical Validity of Genes for Heritable Thoracic Aortic Aneurysm and Dissection. Journal of the American College of Cardiology, 2018, 72, 605-615.	1.2	190
12	Nitric oxide mediates aortic disease in mice deficient in the metalloprotease Adamts1 and in a mouse model of Marfan syndrome. Nature Medicine, 2017, 23, 200-212.	15.2	134
13	Sex, pregnancy and aortic disease in Marfan syndrome. PLoS ONE, 2017, 12, e0181166.	1.1	40
14	FBN1: The disease-causing gene for Marfan syndrome and other genetic disorders. Gene, 2016, 591, 279-291.	1.0	230
15	Gene panel sequencing in heritable thoracic aortic disorders and related entities â€“ results of comprehensive testing in a cohort of 264 patients. Orphanet Journal of Rare Diseases, 2015, 10, 9.	1.2	62
16	Intrinsic cardiomyopathy in Marfan syndrome: results from in-vivo and ex-vivo studies of the Fbn1C1039G/+ model and longitudinal findings in humans. Pediatric Research, 2015, 78, 256-263.	1.1	45
17	Absence of Cardiovascular Manifestations in a Haploinsufficient Tgfbr1 Mouse Model. PLoS ONE, 2014, 9, e89749.	1.1	9
18	Novel MYH11 and ACTA2 mutations reveal a role for enhanced TGFÎ² signaling in FTAAD. International Journal of Cardiology, 2013, 165, 314-321.	0.8	134