Daniel Quiat

List of Publications by Citations

Source: https://exaly.com/author-pdf/4418577/daniel-quiat-publications-by-citations.pdf
Version: 2024-04-25

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.
The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

12 papers	1,075 citations	5 h-index	12 g-index
12 ext. papers	1,250 ext. citations	9.8 avg, IF	4.1 L-index

#	Paper	IF	Citations
12	A family of microRNAs encoded by myosin genes governs myosin expression and muscle performance. *Developmental Cell*, **2009**, 17, 662-73	10.2	737
11	MicroRNAs in cardiovascular disease: from pathogenesis to prevention and treatment. *Journal of Clinical Investigation*, **2013**, 123, 11-8	15.9	196
10	Concerted regulation of myofiber-specific gene expression and muscle performance by the transcriptional repressor Sox6. *Proceedings of the National Academy of Sciences of the United States of America*, **2011**, 108, 10196-201	11.5	106
9	Genomic frontiers in congenital heart disease. *Nature Reviews Cardiology*, **2021**,	14.8	15
8	Association of Damaging Variants in Genes With Increased Cancer Risk Among Patients With Congenital Heart Disease. *JAMA Cardiology*, **2021**, 6, 457-462	16.2	12
7	Discordant clinical features of identical hypertrophic cardiomyopathy twins. *Proceedings of the National Academy of Sciences of the United States of America*, **2021**, 118,	11.5	5
6	High-Throughput Screening of Kawasaki Disease Sera for Antiviral Antibodies. *Journal of Infectious Diseases*, **2020**, 222, 1853-1857	7	3
5	Neither cardiac mitochondrial DNA variation or copy number contribute to congenital heart disease risk.. *American Journal of Human Genetics*, **2022**,	11	1
4	Genome-Wide De Novo Variants in Congenital Heart Disease Are Not Associated With Maternal Diabetes or Obesity.. *Circulation Genomic and Precision Medicine*, **2022**, CIRCGEN121003500	5.2	0
3	Population Prevalence of Premature Truncating Variants in Plakophilin-2 and Association With Arrhythmogenic Right Ventricular Cardiomyopathy: a UK Biobank Analysis.. *Circulation Genomic and Precision Medicine*, **2022**, 101161CIRCGEN121003507	5.2	0
2	An ancient founder mutation located between and is responsible for increased microtia risk in Amerindigenous populations.. *Proceedings of the National Academy of Sciences of the United States of America*, **2022**, 119, e2203928119	11.5	0
1	Left Ventricular Outflow Tract Gradient Is Associated With Coronary Artery Obstruction in Children With Williams-Beuren Syndrome. *Journal of Cardiothoracic and Vascular Anesthesia*, **2021**, 35, 3677-3680	2.1	