

Maria-Christina Kotta

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.

18
papers

315
citations

8
h-index

17
g-index

22
ext. papers

487
ext. citations

5.2
avg, IF

2.79
L-index

#	Paper	IF	Citations
18	Identification of Cadherin 2 () Mutations in Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2017 , 10,		94
17	Calmodulin mutations and life-threatening cardiac arrhythmias: insights from the International Calmodulinopathy Registry. <i>European Heart Journal</i> , 2019 , 40, 2964-2975	9.5	61
16	Desmoplakin missense and non-missense mutations in arrhythmogenic right ventricular cardiomyopathy: Genotype-phenotype correlation. <i>International Journal of Cardiology</i> , 2017 , 249, 268-273 ²	3.2	46
15	The genetics underlying idiopathic ventricular fibrillation: A special role for catecholaminergic polymorphic ventricular tachycardia?. <i>International Journal of Cardiology</i> , 2018 , 250, 139-145	3.2	22
14	Calmodulinopathy: A Novel, Life-Threatening Clinical Entity Affecting the Young. <i>Frontiers in Cardiovascular Medicine</i> , 2018 , 5, 175	5.4	15
13	Enhancing rare variant interpretation in inherited arrhythmias through quantitative analysis of consortium disease cohorts and population controls. <i>Genetics in Medicine</i> , 2021 , 23, 47-58	8.1	13
12	Calmodulinopathy: Functional Effects of CALM Mutations and Their Relationship With Clinical Phenotypes. <i>Frontiers in Cardiovascular Medicine</i> , 2018 , 5, 176	5.4	12
11	The role of genetics in primary ventricular fibrillation, inherited channelopathies and cardiomyopathies. <i>International Journal of Cardiology</i> , 2017 , 237, 45-48	3.2	8
10	Exercise Training-Induced Repolarization Abnormalities Masquerading as Congenital Long QT Syndrome. <i>Circulation</i> , 2020 , 142, 2405-2415	16.7	8
9	Cadherin 2-Related Arrhythmogenic Cardiomyopathy: Prevalence and Clinical Features. <i>Circulation Genomic and Precision Medicine</i> , 2021 , 14, e003097	5.2	8
8	Abnormal myocardial expression of SAP97 is associated with arrhythmogenic risk. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2020 , 318, H1357-H1370	5.2	5
7	Phenotype reveals genotype in a Greek long QT syndrome family. <i>Europace</i> , 2006 , 8, 241-4	3.9	5
6	Novel sodium channel SCN5A mutations in Brugada syndrome patients from Greece. <i>International Journal of Cardiology</i> , 2010 , 145, 45-8	3.2	4
5	Cardiac ion channel gene mutations in Greek long QT syndrome patients. <i>Journal of Applied Genetics</i> , 2010 , 51, 515-8	2.5	4
4	Mutation location and IKs regulation in the arrhythmic risk of long QT syndrome type 1: the importance of the KCNQ1 S6 region. <i>European Heart Journal</i> , 2021 , 42, 4743-4755	9.5	4
3	Genetics of Peripartum Cardiomyopathy: Current Knowledge, Future Directions and Clinical Implications. <i>Genes</i> , 2021 , 12,	4.2	3
2	Estimating the Posttest Probability of Long QT Syndrome Diagnosis for Rare Variants. <i>Circulation Genomic and Precision Medicine</i> , 2021 , 14, e003289	5.2	0

1 Long and Short QT Syndromes. *Cardiac and Vascular Biology*, **2018**, 147-185

0.2