Maria-Christina Kotta

List of Publications by Year in Descending Order

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	Cadherin 2-Related Arrhythmogenic Cardiomyopathy: Prevalence and Clinical Features. *Circulation Genomic and Precision Medicine*, **2021**, 14, e003097	5.2	8
17	Enhancing rare variant interpretation in inherited arrhythmias through quantitative analysis of consortium disease cohorts and population controls. *Genetics in Medicine*, **2021**, 23, 47-58	8.1	13
16	Genetics of Peripartum Cardiomyopathy: Current Knowledge, Future Directions and Clinical Implications. *Genes*, **2021**, 12,	4.2	3
15	Estimating the Posttest Probability of Long QT Syndrome Diagnosis for Rare Variants. *Circulation Genomic and Precision Medicine*, **2021**, 14, e003289	5.2	0
14	Mutation location and IKs regulation in the arrhythmic risk of long QT syndrome type 1: the importance of the KCNQ1 S6 region. *European Heart Journal*, **2021**, 42, 4743-4755	9.5	4
13	Abnormal myocardial expression of SAP97 is associated with arrhythmogenic risk. *American Journal of Physiology - Heart and Circulatory Physiology*, **2020**, 318, H1357-H1370	5.2	5
12	Exercise Training-Induced Repolarization Abnormalities Masquerading as Congenital Long QT Syndrome. *Circulation*, **2020**, 142, 2405-2415	16.7	8
11	Calmodulin mutations and life-threatening cardiac arrhythmias: insights from the International Calmodulinopathy Registry. *European Heart Journal*, **2019**, 40, 2964-2975	9.5	61
10	The genetics underlying idiopathic ventricular fibrillation: A special role for catecholaminergic polymorphic ventricular tachycardia?. *International Journal of Cardiology*, **2018**, 250, 139-145	3.2	22
9	Calmodulinopathy: A Novel, Life-Threatening Clinical Entity Affecting the Young. *Frontiers in Cardiovascular Medicine*, **2018**, 5, 175	5.4	15
8	Calmodulinopathy: Functional Effects of CALM Mutations and Their Relationship With Clinical Phenotypes. *Frontiers in Cardiovascular Medicine*, **2018**, 5, 176	5.4	12
7	Long and Short QT Syndromes. *Cardiac and Vascular Biology*, **2018**, 147-185	0.2	
6	Identification of Cadherin 2 () Mutations in Arrhythmogenic Right Ventricular Cardiomyopathy. *Circulation: Cardiovascular Genetics*, **2017**, 10,		94
5	Desmoplakin missense and non-missense mutations in arrhythmogenic right ventricular cardiomyopathy: Genotype-phenotype correlation. *International Journal of Cardiology*, **2017**, 249, 268-273	3.2	46
4	The role of genetics in primary ventricular fibrillation, inherited channelopathies and cardiomyopathies. *International Journal of Cardiology*, **2017**, 237, 45-48	3.2	8
3	Novel sodium channel SCN5A mutations in Brugada syndrome patients from Greece. *International Journal of Cardiology*, **2010**, 145, 45-8	3.2	4
2	Cardiac ion channel gene mutations in Greek long QT syndrome patients. *Journal of Applied Genetics*, **2010**, 51, 515-8	2.5	4

1	Phenotype reveals genotype in a Greek long QT syndrome family. *Europace*, **2006**, 8, 241-4	3.9	5