Maria-Christina Kotta

List of Publications by Year
in descending order

21	603	758635 12	839053 18
papers	citations	h-index	g-index

22	22	22	1108
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Identification of Cadherin 2 (CDH2) Mutations in Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	123
2	Calmodulin mutations and life-threatening cardiac arrhythmias: insights from the International Calmodulinopathy Registry. European Heart Journal, 2019, 40, 2964-2975.	1.0	116
3	Desmoplakin missense and non-missense mutations in arrhythmogenic right ventricular cardiomyopathy: Genotype-phenotype correlation. International Journal of Cardiology, 2017, 249, 268-273.	0.8	70
4	Enhancing rare variant interpretation in inherited arrhythmias through quantitative analysis of consortium disease cohorts and population controls. Genetics in Medicine, 2021, 23, 47-58.	1.1	57
5	The genetics underlying idiopathic ventricular fibrillation: A special role for catecholaminergic polymorphic ventricular tachycardia?. International Journal of Cardiology, 2018, 250, 139-145.	0.8	42
6	Exercise Training-Induced Repolarization Abnormalities Masquerading as Congenital Long QT Syndrome. Circulation, 2020, 142, 2405-2415.	1.6	36
7	Mutation location and I ÂKs 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.	1.0	26
8	Calmodulinopathy: A Novel, Life-Threatening Clinical Entity Affecting the Young. Frontiers in Cardiovascular Medicine, 2018, 5, 175.	1.1	25
9	Cadherin 2-Related Arrhythmogenic Cardiomyopathy. Circulation Genomic and Precision Medicine, 2021, 14, e003097.	1.6	21
10	Calmodulinopathy: Functional Effects of CALM Mutations and Their Relationship With Clinical Phenotypes. Frontiers in Cardiovascular Medicine, 2018, 5, 176.	1.1	19
11	Genetics of Peripartum Cardiomyopathy: Current Knowledge, Future Directions and Clinical Implications. Genes, 2021, 12, 103.	1.0	17
12	Abnormal myocardial expression of SAP97 is associated with arrhythmogenic risk. American Journal of Physiology - Heart and Circulatory Physiology, 2020, 318, H1357-H1370.	1.5	13
13	Estimating the Posttest Probability of Long QT Syndrome Diagnosis for Rare KCNH2 Variants. Circulation Genomic and Precision Medicine, 2021, 14, e003289.	1.6	10
14	The role of genetics in primary ventricular fibrillation, inherited channelopathies and cardiomyopathies. International Journal of Cardiology, 2017, 237, 45-48.	0.8	8
15	Phenotype reveals genotype in a Greek long QT syndrome family. Europace, 2006, 8, 241-244.	0.7	5
16	Novel sodium channel SCN5A mutations in Brugada syndrome patients from Greece. International Journal of Cardiology, 2010, 145, 45-48.	0.8	5
17	Sudden Infant Death Syndrome andÂGenetics. Journal of the American College of Cardiology, 2018, 71, 1228-1230.	1.2	5
18	Cardiac ion channel gene mutations in Greek long QT syndrome patients. Journal of Applied Genetics, 2010, 51, 515-518.	1.0	4

#	Article	IF	Citations
19	Is Careful Assessment of Rare Variants in the RYR2 Gene Piercing the Guidelinesâ€™ Strong Armor?. Circulation Genomic and Precision Medicine, 2018, 11, e002072.	1.6	1
20	Long and Short QT Syndromes. Cardiac and Vascular Biology, 2018, , 147-185.	0.2	0
21	Biventricular arrhythmogenic cardiomyopathy: a paradigmatic case. ScienceOpen Research, 2015, .	0.6	0