Mireia Alcalde

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5851973/publications.pdf

Version: 2024-02-01

794141 840119 20 361 11 19 citations h-index g-index papers 21 21 21 704 all docs docs citations times ranked citing authors

#	Article	IF	Citations
1	Clinical impact of rare variants associated with inherited channelopathies: a 5-year update. Human Genetics, 2022, 141, 1579-1589.	1.8	11
2	Premature Termination Codon in 5′ Region of Desmoplakin and Plakoglobin Genes May Escape Nonsense-Mediated Decay through the Reinitiation of Translation. International Journal of Molecular Sciences, 2022, 23, 656.	1.8	2
3	Discerning the Ambiguous Role of Missense TTN Variants in Inherited Arrhythmogenic Syndromes. Journal of Personalized Medicine, 2022, 12, 241.	1.1	2
4	Prevalence of Pathogenic Variants in Cardiomyopathy-Associated Genes in Myocarditis. Circulation Genomic and Precision Medicine, 2022, 15, 101161CIRCGEN121003408.	1.6	13
5	Rare Variants Associated with Arrhythmogenic Cardiomyopathy: Reclassification Five Years Later. Journal of Personalized Medicine, 2021, 11, 162.	1.1	13
6	Inflammation in the Pathogenesis of Arrhythmogenic Cardiomyopathy: Secondary Event or Active Driver?. Frontiers in Cardiovascular Medicine, 2021, 8, 784715.	1.1	14
7	Lamotrigine induced Brugada-pattern in a patient with genetic epilepsy associated with a novel variant in SCN9A. Gene, 2020, 754, 144847.	1.0	14
8	Sudden Cardiac Death and Copy Number Variants: What Do We Know after 10 Years of Genetic Analysis?. Forensic Science International: Genetics, 2020, 47, 102281.	1.6	20
9	Personalized Interpretation and Clinical Translation of Genetic Variants Associated With Cardiomyopathies. Frontiers in Genetics, 2019, 10, 450.	1.1	6
10	A Genetically Vulnerable Myocardium May Predispose to Myocarditis. Journal of the American College of Cardiology, 2015, 66, 2913-2914.	1.2	41
11	Genetic and toxicologic investigation of Sudden Cardiac Death in a patient with Arrhythmogenic Right Ventricular Cardiomyopathy (ARVC) under cocaine and alcohol effects. International Journal of Legal Medicine, 2015, 129, 89-96.	1.2	10
12	Clinical interpretation of genetic variants in arrhythmogenic right ventricular cardiomyopathy. Clinical Research in Cardiology, 2015, 104, 288-303.	1.5	13
13	Sequenom MassARRAY approach in the arrhythmogenic right ventricular cardiomyopathy post-mortem setting: clinical and forensic implications. International Journal of Legal Medicine, 2015, 129, 1-10.	1.2	18
14	Stop-Gain Mutations in PKP2 Are Associated with a Later Age of Onset of Arrhythmogenic Right Ventricular Cardiomyopathy. PLoS ONE, 2014, 9, e100560.	1.1	22
15	P389Role of truncated plakophilin-2 in arrhythmogenic right ventricular cardiomyopathy. Cardiovascular Research, 2014, 103, S71.3-S71.	1.8	O
16	The role of clinical, genetic and segregation evaluation in sudden infant death. Forensic Science International, 2014, 242, 9-15.	1.3	19
17	Role of novel DSP_p.Q986X genetic variation in arrhythmogenic right ventricular cardiomyopathy. European Journal of Medical Genetics, 2013, 56, 541-545.	0.7	3
18	Genetics of arrhythmogenic right ventricular cardiomyopathy. Journal of Medical Genetics, 2013, 50, 280-289.	1.5	56

#	Article	IF	CITATIONS
19	Arrhythmogenic right ventricular cardiomyopathy: severe structural alterations are associated with inflammation. Journal of Clinical Pathology, 2012, 65, 1077-1083.	1.0	69
20	Genetic testing of candidate genes in arrhythmogenic right ventricular cardiomyopathy/dysplasia. European Journal of Medical Genetics, 2012, 55, 225-234.	0.7	15