Kalliopi Pilichou

List of Publications by Year
in descending order

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73	4,907	168829	116156
papers	citations	31 h-index	66 g-index

74	74	74	4172
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Filamin-C variant-associated cardiomyopathy: AÂpooled analysis of individual patient data to evaluate the clinical profile and risk of sudden cardiac death. Heart Rhythm, 2022, 19, 235-243.	0.3	33
2	The 2020 â€œPadua Criteriaâ€•for Diagnosis and Phenotype Characterization of Arrhythmogenic Cardiomyopathy in Clinical Practice. Journal of Clinical Medicine, 2022, 11, 279.	1.0	9
3	Clinical management of a pregnant woman with Filamin C cardiomyopathy. Journal of Cardiovascular Medicine, 2022, 23, 198-202.	0.6	1
4	Autonomic dysfunction as first presentation of Glu54Gln transthyretin amyloidosis. Journal of the Neurological Sciences, 2022, 437, 120264.	0.3	0
5	Clinical profile and long-term follow-up of a cohort of patients with desmoplakin cardiomyopathy. Heart Rhythm, 2022, 19, 1315-1324.	0.3	22
6	Importance of genotype for risk stratification in arrhythmogenic right ventricular cardiomyopathy using the 2019 ARVC risk calculator. European Heart Journal, 2022, 43, 3053-3067.	1.0	41
7	Molecular genetic testing in athletes: Why and when a position statement from the Italian Society of Sports Cardiology. International Journal of Cardiology, 2022, 364, 169-177.	0.8	6
8	Papillary Muscles Abnormalities in Athletes With Otherwise Unexplained Tâ€Wave Inversion in the ECG Lateral Leads. Journal of the American Heart Association, 2021, 10, e019239.	1.6	5
9	Role of Exercise as a Modulating Factor in Arrhythmogenic Cardiomyopathy. Current Cardiology Reports, 2021, 23, 57.	1.3	17
10	Evidence-Based Assessment of Genes in Dilated Cardiomyopathy. Circulation, 2021, 144, 7-19.	1.6	213
11	International Evidence Based Reappraisal of Genes Associated With Arrhythmogenic Right Ventricular Cardiomyopathy Using the Clinical Genome Resource Framework. Circulation Genomic and Precision Medicine, 2021, 14, e003273.	1.6	112
12	The genetic architecture of Plakophilin 2 cardiomyopathy. Genetics in Medicine, 2021, 23, 1961-1968.	1.1	13
13	Arrhythmogenic cardiomyopathy: the ongoing search for mechanism-driven therapies meets extracellular vesicles. European Heart Journal, 2021, 42, 3572-3574.	1.0	6
14	Novel pathogenic role for galectin-3 in early disease stages of arrhythmogenic cardiomyopathy. Heart Rhythm, 2021, 18, 1394-1403.	0.3	8
15	Myocardial Tissue Characterization in Arrhythmogenic Cardiomyopathy. JACC: Cardiovascular Imaging, 2021, 14, 1675-1678.	2.3	13
16	Evolving Diagnostic Criteria for Arrhythmogenic Cardiomyopathy. Journal of the American Heart Association, 2021, 10, e021987.	1.6	60
17	Neuropeptide Y promotes adipogenesis of human cardiac mesenchymal stromal cells in arrhythmogenic cardiomyopathy. International Journal of Cardiology, 2021, 342, 94-102.	0.8	10
18	â€˜Hot phaseâ€™ clinical presentation in arrhythmogenic cardiomyopathy. Europace, 2021, 23, 907-917.	0.7	67

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19	Identification of Rare LRP5 Variants in a Cohort of Males with Impaired Bone Mass. International Journal of Molecular Sciences, 2021, 22, 10834.	1.8	5
20	A de novo ryanodine receptor 2 gene variant in a case of sudden cardiac death. International Journal of Legal Medicine, 2020, 134, 619-623.	1.2	4
21	Arrhythmogenic Cardiomyopathy. European Heart Journal, 2020, 41, 4457-4462.	1.0	12
22	The Role of MicroRNAs in Arrhythmogenic Cardiomyopathy: Biomarkers or Innocent Bystanders of Disease Progression?. International Journal of Molecular Sciences, 2020, 21, 6434.	1.8	5
23	Cardiac arrest at rest and during sport activity: causes and prevention. European Heart Journal Supplements, 2020, 22, E20-E24.	0.0	6
24	Diagnosis of arrhythmogenic cardiomyopathy: The Padua criteria. International Journal of Cardiology, 2020, 319, 106-114.	0.8	283
25	Development of a novel next-generation sequencing panel for diagnosis of quantitative spermatogenic impairment. Journal of Assisted Reproduction and Genetics, 2020, 37, 753-762.	1.2	13
26	Arrhythmogenic Right Ventricular Cardiomyopathy: Characterization of Left Ventricular Phenotype and Differential Diagnosis With Dilated Cardiomyopathy. Journal of the American Heart Association, 2020, 9, e014628.	1.6	92
27	A microRNA Expression Profile as Non-Invasive Biomarker in a Large Arrhythmogenic Cardiomyopathy Cohort. International Journal of Molecular Sciences, 2020, 21, 1536.	1.8	21
28	Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. European Heart Journal, 2020, 41, 1414-1429.	1.0	239
29	Update on cardiomyopathies and sudden cardiac death. Forensic Sciences Research, 2019, 4, 202-210.	0.9	17
30	Genetics in cardiovascular diseases. Italian Journal of Medicine, 2019, 13, 137-151.	0.2	3
31	Definition and treatment of arrhythmogenic cardiomyopathy: an updated expert panel report. European Journal of Heart Failure, 2019, 21, 955-964.	2.9	84
32	Heart failure in arrhythmogenic cardiomyopathy: is phenotypic variability justÂaÂmatter of genetics?. European Journal of Heart Failure, 2019, 21, 801-802.	2.9	1
33	Predictive value of exercise testing in athletes with ventricular ectopy evaluated by cardiac magnetic resonance. Heart Rhythm, 2019, 16, 239-248.	0.3	45
34	The complex molecular genetics of arrhythmogenic cardiomyopathy. International Journal of Cardiology, 2019, 284, 59-60.	0.8	3
35	Loss of cardiac Wnt/Î²-catenin signalling in desmoplakin-deficient AC8 zebrafish models is rescuable by genetic and pharmacological intervention. Cardiovascular Research, 2018, 114, 1082-1097.	1.8	39
36	Diagnostic Criteria, Genetics, and Molecular Basis of Arrhythmogenic Cardiomyopathy. Heart Failure Clinics, 2018, 14, 201-213.	1.0	27

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37	Arrhythmogenic Cardiomyopathy. , 2018, , 631-639.		0
38	Inherited Cardiac Muscle Disorders: Arrhythmogenic Right Ventricular Cardiomyopathy. , 2018, , 367-388.		0
39	Large Genomic Rearrangements of Desmosomal Genes in Italian Arrhythmogenic Cardiomyopathy Patients. Circulation: Arrhythmia and Electrophysiology, 2017, 10, .	2.1	35
40	Arrhythmogenic cardiomyopathy: pathology, genetics, and concepts in pathogenesis. Cardiovascular Research, 2017, 113, 1521-1531.	1.8	98
41	Sudden arrhythmic death and cardiomyopathies in the young: a molecular and pathology overview. Diagnostic Histopathology, 2017, 23, 486-498.	0.2	0
42	Arrhythmogenic Cardiomyopathy: History and Pathology. , 2016, , 5-33.		1
43	Response to Letters Regarding Article, â€œArrhythmic Mitral Valve Prolapse and Sudden Cardiac Deathâ€. Circulation, 2016, 133, e460.	1.6	3
44	Arrhythmogenic Right Ventricular Cardiomyopathy: Risk Stratification and Indications for Defibrillator Therapy. Current Cardiology Reports, 2016, 18, 57.	1.3	20
45	Morphofunctional Abnormalities of Mitral Annulus and Arrhythmic Mitral Valve Prolapse. Circulation: Cardiovascular Imaging, 2016, 9, e005030.	1.3	226
46	Nonischemic Left Ventricular Scar as a Substrate of Life-Threatening Ventricular Arrhythmias and Sudden Cardiac Death in Competitive Athletes. Circulation: Arrhythmia and Electrophysiology, 2016, 9, .	2.1	216
47	Arrhythmogenic cardiomyopathy. Orphanet Journal of Rare Diseases, 2016, 11, 33.	1.2	116
48	Phenotypic expression is a prerequisite for malignant arrhythmic events and sudden cardiac death in arrhythmogenic right ventricular cardiomyopathy. Europace, 2016, 18, 1086-1094.	0.7	50
49	The ARVD/C Genetic Variants Database: 2014 Update. Human Mutation, 2015, 36, 403-410.	1.1	77
50	Arrhythmic Mitral Valve Prolapse and Sudden Cardiac Death. Circulation, 2015, 132, 556-566.	1.6	422
51	Pharmacotherapy and Other Therapeutic Modalities for Managing Arrhythmogenic Right Ventricular Cardiomyopathy. Cardiovascular Drugs and Therapy, 2015, 29, 171-177.	1.3	12
52	Homozygous Desmocollin-2 Mutations and Arrhythmogenic Cardiomyopathy. American Journal of Cardiology, 2015, 116, 1245-1251.	0.7	38
53	TGF-beta1 pathway activation and adherens junction molecular pattern in nonsyndromic mitral valve prolapse. Cardiovascular Pathology, 2015, 24, 359-367.	0.7	25
54	Nonischemic Left Ventricular Scar. Circulation, 2014, 130, e180-2.	1.6	22

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55	Is it time for plakoglobin immune-histochemical diagnostic test for arrhythmogenic cardiomyopathy in the routine pathology practice?. Cardiovascular Pathology, 2013, 22, 312-313.	0.7	8
56	Pathologic Substrates of Sudden Cardiac Death During Sports. Cardiac Electrophysiology Clinics, 2013, 5, 1-11.	0.7	1
57	Compound and Digenic Heterozygosity Predicts Lifetime Arrhythmic Outcome and Sudden Cardiac Death in Desmosomal Geneâ€"Related Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Cardiovascular Genetics, 2013, 6, 533-542.	5.1	209
58	Assessing the Significance of Pathogenic Mutations and Autopsy Findings in the Light of 2010 Arrhythmogenic Right Ventricular Cardiomyopathy Diagnostic Criteria. Circulation: Cardiovascular Genetics, 2012, 5, 384-386.	5.1	1
59	Intercalated disc abnormalities, reduced Na+ current density, and conduction slowing in desmoglein-2 mutant mice prior to cardiomyopathic changes. Cardiovascular Research, 2012, 95, 409-418.	1.8	180
60	The changing spectrum of arrhythmogenic (right ventricular) cardiomyopathy. Cell and Tissue Research, 2012, 348, 319-323.	1.5	31
61	Molecular biology and clinical management of arrhythmogenic right ventricular cardiomyopathy/dysplasia. Heart, 2011, 97, 530-539.	1.2	120
62	Pathobiology of Arrhythmogenic Cardiomyopathy. Cardiac Electrophysiology Clinics, 2011, 3, 193-204.	0.7	5
63	Clinical phenotype and diagnosis of arrhythmogenic right ventricular cardiomyopathy in pediatric patients carrying desmosomal gene mutations. Heart Rhythm, 2011, 8, 1686-1695.	0.3	66
64	Arrhythmogenic Cardiomyopathy. Circulation: Cardiovascular Genetics, 2011, 4, 318-326.	5.1	35
65	Arrhythmogenic Right Ventricular Cardiomyopathy. , 2011, , 438-443.		0
66	Sudden arrhythmic death and the cardiomyopathies: Molecular genetics and pathology. Diagnostic Histopathology, 2010, 16, 31-42.	0.2	1
67	Compound and Digenic Heterozygosity Contributes to Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American College of Cardiology, 2010, 55, 587-597.	1.2	282
68	Sudden cardiac death with normal heart:. Cardiovascular Pathology, 2010, 19, 321-325.	0.7	119
69	Myocyte necrosis underlies progressive myocardial dystrophy in mouse dsg2-related arrhythmogenic right ventricular cardiomyopathy. Journal of Experimental Medicine, 2009, 206, 1787-1802.	4.2	184
70	Myocyte necrosis underlies progressive myocardial dystrophy in mouse dsg2-related arrhythmogenic right ventricular cardiomyopathy. Journal of Cell Biology, 2009, 186, i5-i5.	2.3	0
71	Homozygous SCN5A mutation in Brugada syndrome with monomorphic ventricular tachycardia and structural heart abnormalities. Europace, 2007, 9, 391-397.	0.7	41
72	Mutations in Desmoglein-2 Gene Are Associated With Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation, 2006, 113, 1171-1179.	1.6	509

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73	Ultrastructural evidence of intercalated disc remodelling in arrhythmogenic right ventricular cardiomyopathy: an electron microscopy investigation on endomyocardial biopsies. European Heart Journal, 2006, 27, 1847-1854.	1.0	219