Cynthia A James

List of Publications by Year
in descending order

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163 papers	10,895 citations	28274 55 h-index	33894 99 g-index
166 all docs	166 docs citations	166 times ranked	6608 citing authors

#	Article	IF	Citations
1	Exercise Increases Age-Related Penetrance and Arrhythmic Risk in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathyâ€"Associated Desmosomal Mutation Carriers. Journal of the American College of Cardiology, 2013, 62, 1290-1297.	2.8	553
2	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy. Heart Rhythm, 2019, 16, e301-e372.	0.7	494
3	Arrhythmogenic Right Ventricular Dysplasia. Circulation, 2005, 112, 3823-3832.	1.6	434
4	Clinical Presentation, Long-Term Follow-Up, and Outcomes of 1001 Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy Patients and Family Members. Circulation: Cardiovascular Genetics, 2015, 8, 437-446.	5.1	370
5	Impact of genotype on clinical course in arrhythmogenic right ventricular dysplasia/cardiomyopathy-associated mutation carriers. European Heart Journal, 2015, 36, 847-855.	2.2	338
6	Ethical and Practical Guidelines for Reporting Genetic Research Results to Study Participants. Circulation: Cardiovascular Genetics, 2010, 3, 574-580.	5.1	328
7	Electrocardiographic Features of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy According to Disease Severity. Circulation, 2004, 110, 1527-1534.	1.6	261
8	Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. European Heart Journal, 2020, 41, 1414-1429.	2.2	239
9	Long-Term Efficacy of Catheter Ablation of Ventricular Tachycardia in Patients With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of the American College of Cardiology, 2007, 50, 432-440.	2.8	236
10	Incidence and Predictors of Implantable Cardioverter-Defibrillator Therapy in Patients With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy Undergoing Implantable Cardioverter-Defibrillator Implantation for Primary Prevention. Journal of the American College of Cardiology, 2011, 58, 1485-1496.	2.8	226
11	Clinical Features of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy Associated With Mutations in Plakophilin-2. Circulation, 2006, 113, 1641-1649.	1.6	225
12	Identification of a New Modulator of the Intercalated Disc in a Zebrafish Model of Arrhythmogenic Cardiomyopathy. Science Translational Medicine, 2014, 6, 240ra74.	12.4	222
13	Evidence-Based Assessment of Genes in Dilated Cardiomyopathy. Circulation, 2021, 144, 7-19.	1.6	213
14	DSG2 Mutations Contribute to Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Human Genetics, 2006, 79, 136-142.	6.2	206
15	Misdiagnosis of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of Cardiovascular Electrophysiology, 2004, 15, 300-306.	1.7	199
16	Implantable Cardioverter-Defibrillators in patients with arrhythmogenic right ventricular Dysplasia/Cardiomyopathy. Journal of the American College of Cardiology, 2004, 43, 1843-1852.	2.8	197
17	Comprehensive Desmosome Mutation Analysis in North Americans With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Circulation: Cardiovascular Genetics, 2009, 2, 428-435.	5.1	195
18	Outcomes of Catheter Ablation of Ventricular Tachycardia in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2012, 5, 499-505.	4.8	175

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19	Exercise has a Disproportionate Role in the Pathogenesis of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy in Patients Without Desmosomal Mutations. Journal of the American Heart Association, 2014, 3, e001471.	3.7	158
20	2020 APHRS/HRS expert consensus statement on the investigation of decedents with sudden unexplained death and patients with sudden cardiac arrest, and of their families. Heart Rhythm, 2021, 18, e1-e50.	0.7	151
21	Mutationâ€Positive Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy: The Triangle of Dysplasia Displaced. Journal of Cardiovascular Electrophysiology, 2013, 24, 1311-1320.	1.7	148
22	Multilevel analyses of SCN5A mutations in arrhythmogenic right ventricular dysplasia/cardiomyopathy suggest non-canonical mechanisms for disease pathogenesis. Cardiovascular Research, 2017, 113, 102-111.	3.8	148
23	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy: Executive summary. Heart Rhythm, 2019, 16, e373-e407.	0.7	135
24	Comparison of Novel Echocardiographic Parameters of Right Ventricular Function with Ejection Fraction by Cardiac Magnetic Resonance. Journal of the American Society of Echocardiography, 2007, 20, 1058-1064.	2.8	130
25	Penetrance of Mutations in Plakophilin-2 Among Families With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of the American College of Cardiology, 2006, 48, 1416-1424.	2.8	122
26	How does the mode of inheritance of a genetic condition influence families? A study of guilt, blame, stigma, and understanding of inheritance and reproductive risks in families with X-linked and autosomal recessive diseases. Genetics in Medicine, 2006, 8, 234-242.	2.4	116
27	Predictors of appropriate implantable defibrillator therapies in patients with arrhythmogenic right ventricular dysplasia. Heart Rhythm, 2005, 2, 1188-1194.	0.7	114
28	Incremental Value of Cardiac Magnetic Resonance Imaging in Arrhythmic Risk Stratification of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathyâ€“Associated Desmosomal Mutation Carriers. Journal of the American College of Cardiology, 2013, 62, 1761-1769.	2.8	112
29	Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy and Cardiac Sarcoidosis. Circulation: Arrhythmia and Electrophysiology, 2014, 7, 230-236.	4.8	112
30	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.	3.6	112
31	Outcomes and ventricular tachycardia recurrence characteristics after epicardial ablation of ventricular tachycardia in arrhythmogenic right ventricular dysplasia/cardiomyopathy. Heart Rhythm, 2015, 12, 716-725.	0.7	101
32	Arrhythmogenic cardiomyopathy: pathology, genetics, and concepts in pathogenesis. Cardiovascular Research, 2017, 113, 1521-1531.	3.8	98
33	Risk Stratification in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathyâ€“Associated Desmosomal Mutation Carriers. Circulation: Arrhythmia and Electrophysiology, 2013, 6, 569-578.	4.8	94
34	The Responsibility to Recontact Research Participants after Reinterpretation of Genetic and Genomic Research Results. American Journal of Human Genetics, 2019, 104, 578-595.	6.2	91
35	Electrocardiographic Features of Arrhythmogenic Right Ventricular Dysplasia. Circulation, 2009, 120, 477-487.	1.6	88
36	Yield of Serial Evaluation in At-Risk Family Members of Patients With ARVD/C. Journal of the American College of Cardiology, 2014, 64, 293-301.	2.8	88

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37	Morphologic Variants of Familial Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of the American College of Cardiology, 2009, 53, 1289-1299.	2.8	84
38	Sudden Cardiac Death Prediction in Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2021, 14, e008509.	4.8	82
39	Predicting arrhythmic risk in arrhythmogenic right ventricular cardiomyopathy: A systematic review and meta-analysis. Heart Rhythm, 2018, 15, 1097-1107.	0.7	79
40	Recessive arrhythmogenic right ventricular dysplasia due to novel cryptic splice mutation inPKP2. Human Mutation, 2006, 27, 1157-1157.	2.5	77
41	Cardiac Transplantation in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of the American College of Cardiology, 2012, 59, 289-290.	2.8	76
42	Safety of American Heart Association-recommended minimum exercise for desmosomal mutation carriers. Heart Rhythm, 2016, 13, 199-207.	0.7	76
43	Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy in the PediatricÂPopulation. JACC: Clinical Electrophysiology, 2015, 1, 551-560.	3.2	74
44	Utility of Tissue Doppler and Strain Echocardiography in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Cardiology, 2007, 100, 507-512.	1.6	73
45	Feasibility and Variability of Three Dimensional Echocardiography in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Cardiology, 2006, 97, 703-709.	1.6	71
46	Prevalence of atrial arrhythmias in arrhythmogenic right ventricular dysplasia/cardiomyopathy. Heart Rhythm, 2013, 10, 1661-1668.	0.7	71
47	Approach to family screening in arrhythmogenic right ventricular dysplasia/cardiomyopathy. European Heart Journal, 2016, 37, 755-763.	2.2	68
48	Implantable Cardioverterâ€Defibrillator Therapy in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy: Predictors of Appropriate Therapy, Outcomes, and Complications. Journal of the American Heart Association, 2017, 6, .	3.7	68
49	Exercise Testing in Asymptomatic Gene Carriers Exposes a Latent Electrical Substrate of Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American College of Cardiology, 2013, 62, 1772-1779.	2.8	64
50	High Prevalence of Catecholamine-facilitated Focal Ventricular Tachycardia in Patients With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2013, 6, 160-166.	4.8	64
51	Findings on magnetic resonance imaging of idiopathic right ventricular outflow tachycardia. American Journal of Cardiology, 2004, 94, 1441-1445.	1.6	61
52	General and Disease-Specific Psychosocial Adjustment in Patients With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy With Implantable Cardioverter Defibrillators: A Large Cohort Study. Circulation: Cardiovascular Genetics, 2012, 5, 18-24.	5.1	60
53	Evolving Role of Multidetector Computed Tomography in Evaluation of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Cardiology, 2007, 100, 99-105.	1.6	57
54	Plasma BIN1 correlates with heart failure and predicts arrhythmia in patients with arrhythmogenic right ventricular cardiomyopathy. Heart Rhythm, 2012, 9, 961-967.	0.7	56

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55	An electrocardiographic scoring system for distinguishing right ventricular outflow tract arrhythmias in patients with arrhythmogenic right ventricular cardiomyopathy from idiopathic ventricular tachycardia. Heart Rhythm, 2013, 10, 477-482.	0.7	56
56	Impact of Exercise Restriction on Arrhythmic Risk Among Patients With Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American Heart Association, 2018, 7, .	3.7	55
57	The role of genetics in cardiovascular disease: arrhythmogenic cardiomyopathy. European Heart Journal, 2020, 41, 1393-1400.	2.2	54
58	Evaluation of Structural Progression in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. JAMA Cardiology, 2017, 2, 293.	6.1	53
59	Heart Failure Is Common and Under-Recognized in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Circulation: Heart Failure, 2017, 10, .	3.9	53
60	Cardiac MR Findings and Potential Diagnostic Pitfalls in Patients Evaluated for Arrhythmogenic Right Ventricular Cardiomyopathy. Radiographics, 2014, 34, 1553-1570.	3.3	52
61	Pregnancy course and outcomes in women with arrhythmogenic right ventricular cardiomyopathy. Heart, 2016, 102, 303-312.	2.9	50
62	Feature tracking CMR reveals abnormal strain in preclinical arrhythmogenic right ventricular dysplasia/ cardiomyopathy: a multisoftware feasibility and clinical implementation study. Journal of Cardiovascular Magnetic Resonance, 2016, 19, 66.	3.3	50
63	Malignant Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy with a normal 12-lead electrocardiogram: A rare but underrecognized clinical entity. Heart Rhythm, 2013, 10, 1484-1491.	0.7	47
64	Cardiac phenotype and long-term prognosis of arrhythmogenic right ventricular cardiomyopathy/dysplasia patients with late presentation. Heart Rhythm, 2017, 14, 883-891.	0.7	47
65	Exercise triggers CAPN1-mediated AIF truncation, inducing myocyte cell death in arrhythmogenic cardiomyopathy. Science Translational Medicine, 2021, 13, .	12.4	46
66	Electronic health record phenotype in subjects with genetic variants associated with arrhythmogenic right ventricular cardiomyopathy: a study of 30,716 subjects with exome sequencing. Genetics in Medicine, 2017, 19, 1245-1252.	2.4	43
67	FLNC truncations cause arrhythmogenic right ventricular cardiomyopathy. Journal of Medical Genetics, 2020, 57, 254-257.	3.2	43
68	Phenotypic Expression, Natural History, and Risk Stratification of Cardiomyopathy Caused by Filamin C Truncating Variants. Circulation, 2021, 144, 1600-1611.	1.6	43
69	Cardiac sympathectomy for refractory ventricular tachycardia in arrhythmogenic right ventricular cardiomyopathy. Heart Rhythm, 2019, 16, 1003-1010.	0.7	42
70	Prevalence and Electronic Health Record-Based Phenotype of Loss-of-Function Genetic Variants in Arrhythmogenic Right Ventricular Cardiomyopathy-Associated Genes. Circulation Genomic and Precision Medicine, 2019, 12, e002579.	3.6	42
71	Ankyrin-B dysfunction predisposes to arrhythmogenic cardiomyopathy and is amenable to therapy. Journal of Clinical Investigation, 2019, 129, 3171-3184.	8.2	42
72	Clinical characteristics and risk stratification of desmoplakin cardiomyopathy. Europace, 2022, 24, 268-277.	1.7	41

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73	Right ventricular strain by MR quantitatively identifies regional dysfunction in patients with arrhythmogenic right ventricular cardiomyopathy. Journal of Magnetic Resonance Imaging, 2016, 43, 1132-1139.	3.4	40
74	Diagnosing arrhythmogenic right ventricular cardiomyopathy by 2010 Task Force Criteria: clinical performance and simplified practical implementation. Europace, 2020, 22, 787-796.	1.7	40
75	Adults With X-Linked Agammaglobulinemia. Medicine (United States), 2008, 87, 253-258.	1.0	38
76	Arrhythmogenic Right Ventricular Cardiomyopathy-Associated Desmosomal Variants Are Rarely De Novo. Circulation Genomic and Precision Medicine, 2019, 12, e002467.	3.6	38
77	Arrhythmogenic Right Ventricular Cardiomyopathy Presenting as Clinical Myocarditis in Women. American Journal of Cardiology, 2021, 145, 128-134.	1.6	38
78	Informed consent for exome sequencing research in families with genetic disease: The emerging issue of incidental findings. American Journal of Medical Genetics, Part A, 2014, 164, 2745-2752.	1.2	37
79	Characterizing the Molecular Pathology of Arrhythmogenic Cardiomyopathy in Patient Buccal Mucosa Cells. Circulation: Arrhythmia and Electrophysiology, 2016, 9, e003688.	4.8	35
80	Comparison of Features of Fatal Versus Nonfatal Cardiac Arrest in Patients With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Cardiology, 2017, 120, 111-117.	1.6	35
81	A new prediction model for ventricular arrhythmias in arrhythmogenic right ventricular cardiomyopathy. European Heart Journal, 2022, 43, e1-e9.	2.2	35
82	Prevalence and Pathophysiologic Attributes of Ventricular Dyssynchrony in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of the American College of Cardiology, 2009, 54, 445-451.	2.8	34
83	Diagnostic and therapeutic strategies for arrhythmogenic right ventricular dysplasia/cardiomyopathy patient. Europace, 2019, 21, 9-21.	1.7	33
84	Arrhythmic risk prediction in arrhythmogenic right ventricular cardiomyopathy: external validation of the arrhythmogenic right ventricular cardiomyopathy risk calculator. European Heart Journal, 2022, 43, 3041-3052.	2.2	32
85	Regional Differences in Systolic and Diastolic Function in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy Using Magnetic Resonance Imaging. American Journal of Cardiology, 2005, 95, 1507-1511.	1.6	31
86	Electroanatomic Correlates of Depolarization Abnormalities in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of Cardiovascular Electrophysiology, 2016, 27, 443-452.	1.7	31
87	Choices for return of primary and secondary genomic research results of 790 members of families with Mendelian disease. European Journal of Human Genetics, 2017, 25, 530-537.	2.8	31
88	Communication of genetic information to families with inherited rhythm disorders. Heart Rhythm, 2018, 15, 780-786.	0.7	30
89	Perceptions of reproductive risk and carrier testing among adolescent sisters of males with chronic granulomatous disease. American Journal of Medical Genetics Part A, 2003, 119C, 60-69.	2.4	28
90	Arrhythmogenic right ventricular cardiomyopathy and sports activity: from molecular pathways in diseased hearts to new insights into the athletic heart mimicry. European Heart Journal, 2021, 42, 1231-1243.	2.2	27

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91	Spectrum of Biventricular Involvement on CMR Among Carriers of ARVD/C-Associated Mutations. JACC: Cardiovascular Imaging, 2015, 8, 863-864.	5.3	25
92	Right Ventricular Strain Predicts Structural Disease Progression in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American Heart Association, 2020, 9, e015016.	3.7	24
93	Subcutaneous Implantable Cardioverterâ€Ðefibrillator in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia: A Transatlantic Experience. Journal of the American Heart Association, 2018, 7, e008782.	3.7	23
94	Exercise restriction is protective for genotype-positive family members of arrhythmogenic right ventricular cardiomyopathy patients. Europace, 2020, 22, 1270-1278.	1.7	23
95	Shared Desmosome Gene Findings in Early and Late Onset Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of Cardiovascular Translational Research, 2010, 3, 663-673.	2.4	21
96	Cadherin 2-Related Arrhythmogenic Cardiomyopathy. Circulation Genomic and Precision Medicine, 2021, 14, e003097.	3.6	21
97	Altered Electrical, Biomolecular, and Immunologic Phenotypes in a Novel Patient-Derived Stem Cell Model of Desmoglein-2 Mutant ARVC. Journal of Clinical Medicine, 2021, 10, 3061.	2.4	21
98	Enrolling Genomics Research Participants through a Clinical Setting: the Impact of Existing Clinical Relationships on Informed Consent and Expectations for Return of Research Results. Journal of Genetic Counseling, 2018, 27, 263-273.	1.6	19
99	Electrocardiographic Features Differentiating Arrhythmogenic RightÂVentricular Cardiomyopathy FromÂan Athleteâ€™s Heart. JACC: Clinical Electrophysiology, 2018, 4, 1613-1625.	3.2	19
100	Left ventricular fibro-fatty replacement in arrhythmogenic right ventricular dysplasia/cardiomyopathy: prevalence, patterns, and association with arrhythmias. Journal of Cardiovascular Magnetic Resonance, 2021, 23, 58.	3.3	19
101	Role of plakophilin-2 expression on exercise-related progression of arrhythmogenic right ventricular cardiomyopathy: a translational study. European Heart Journal, 2022, 43, 1251-1264.	2.2	19
102	Quality of life metrics in arrhythmogenic right ventricular cardiomyopathy patients: The impact of age, shock and sex. International Journal of Cardiology, 2017, 248, 216-220.	1.7	17
103	Performance of the 2015 International Task Force Consensus Statement Risk Stratification Algorithm for Implantable Cardioverter-Defibrillator Placement in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2018, 11, e005593.	4.8	17
104	2020 APHRS/HRS expert consensus statement on the investigation of decedents with sudden unexplained death and patients with sudden cardiac arrest, and of their families. Journal of Arrhythmia, 2021, 37, 481-534.	1.2	17
105	Multimodality Imaging in Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Cardiovascular Imaging, 2022, 15, CIRCIMAGING121013725.	2.6	17
106	Update on Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy (ARVD/C). Current Treatment Options in Cardiovascular Medicine, 2013, 15, 476-487.	0.9	16
107	Fibrofatty Changes: Incidence at Cardiac MR Imaging in Patients with Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Radiology, 2016, 280, 405-412.	7.3	16
108	Minding the Genes: a Multidisciplinary Approach towards Genetic Assessment of Cardiovascular Disease. Journal of Genetic Counseling, 2017, 26, 224-231.	1.6	16

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109	Clinical outcomes of catheter ablation of ventricular tachycardia in patients with arrhythmogenic right ventricular cardiomyopathy: Insights from the Johns Hopkins ARVC Program. Heart Rhythm, 2021, 18, 1369-1376.	0.7	16
110	Psychosocial care and cardiac genetic counseling following sudden cardiac death in the young. Progress in Pediatric Cardiology, 2017, 45, 31-36.	0.4	15
111	Identification of sarcomeric variants in probands with a clinical diagnosis of arrhythmogenic right ventricular cardiomyopathy (ARVC). Journal of Cardiovascular Electrophysiology, 2018, 29, 1004-1009.	1.7	15
112	Arrhythmogenic Right Ventricular Cardiomyopathy: Progress Toward Personalized Management. Annual Review of Medicine, 2019, 70, 1-18.	12.2	15
113	Statistical evaluation of reproducibility of automated ECG measurements: An example from arrhythmogenic right ventricular dysplasia/cardiomyopathy clinic. Biomedical Signal Processing and Control, 2014, 13, 23-30.	5.7	14
114	Safety and Utility of Cardiopulmonary Exercise Testing in Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Journal of the American Heart Association, 2020, 9, e013695.	3.7	14
115	Clinical Findings and Diagnostic Yield of Arrhythmogenic Cardiomyopathy Through Genomic Screening of Pathogenic or Likely Pathogenic Desmosome Gene Variants. Circulation Genomic and Precision Medicine, 2021, 14, e003302.	3.6	14
116	Psychosocial Stress Hastens Disease Progression and Sudden Death in Mice with Arrhythmogenic Cardiomyopathy. Journal of Clinical Medicine, 2020, 9, 3804.	2.4	13
117	Impact of Genetic Variant Reassessment on the Diagnosis of Arrhythmogenic Right Ventricular Cardiomyopathy Based on the 2010 Task Force Criteria. Circulation Genomic and Precision Medicine, 2021, 14, e003047.	3.6	13
118	The genetic architecture of Plakophilin 2 cardiomyopathy. Genetics in Medicine, 2021, 23, 1961-1968.	2.4	13
119	Premature Ventricular Contraction Variability in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of Cardiovascular Electrophysiology, 2015, 26, 53-57.	1.7	12
120	Epicardial Fat Distribution Assessed with Cardiac CT in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Radiology, 2018, 289, 641-648.	7.3	12
121	Arrhythmic outcome of arrhythmogenic right ventricular cardiomyopathy patients without implantable defibrillators. Journal of Cardiovascular Electrophysiology, 2018, 29, 1396-1402.	1.7	12
122	Managing Secondary Genomic Findings Associated With Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation Genomic and Precision Medicine, 2018, 11, e002237.	3.6	11
123	Contemporary and Future Approaches toÂPrecision Medicine in InheritedÂCardiomyopathies. Journal of the American College of Cardiology, 2021, 77, 2551-2572.	2.8	11
124	Misdiagnosis of ARVC leading to inappropriate ICD implant and subsequent ICD removal â€“ lessons learned. Journal of Cardiovascular Electrophysiology, 2019, 30, 2020-2026.	1.7	10
125	Arrhythmogenic right ventricular cardiomyopathy: evidence for progression increases. European Heart Journal, 2020, 41, 1411-1413.	2.2	10
126	The Value of Cardiac Magnetic Resonance Imaging in Evaluation of Pediatric Patients for Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of the American College of Cardiology, 2015, 66, 873-874.	2.8	9

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127	Return of secondary findings in genomic sequencing: Military implications. Molecular Genetics & Genomic Medicine, 2019, 7, e00483.	1.2	9
128	Anxiety and depression in inherited channelopathy patients with implantable cardioverter-defibrillators. Heart Rhythm O2, 2021, 2, 388-393.	1.7	9
129	Comparing clinical performance of current implantable cardioverter-defibrillator implantation recommendations in arrhythmogenic right ventricular cardiomyopathy. Europace, 2022, 24, 296-305.	1.7	9
130	Heart transplantation strategies in arrhythmogenic right ventricular cardiomyopathy: a tertiary ARVC centre experience. ESC Heart Failure, 2022, 9, 1008-1017.	3.1	9
131	Reader- and Instrument-Dependent Variability in the Electrocardiographic Assessment of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of Cardiovascular Electrophysiology, 2011, 22, 561-568.	1.7	8
132	Global approaches to cardiogenetic evaluation after sudden cardiac death in the young: A survey among health care professionals. Heart Rhythm, 2021, 18, 1637-1644.	0.7	8
133	Association of Premature Ventricular Contraction Burden on Serial Holter Monitoring With Arrhythmic Risk in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy. JAMA Cardiology, 2022, 7, 378.	6.1	8
134	Loss-of-Function FLNC Variants Are Associated With Arrhythmogenic Cardiomyopathy Phenotypes When Identified Through Exome Sequencing of a General Clinical Population. Circulation Genomic and Precision Medicine, 2022, 15, .	3.6	8
135	Efficacy of Catheter Ablation for Atrial Arrhythmias in Patients with Arrhythmogenic Right Ventricular Cardiomyopathyâ€"A Multicenter Study. Journal of Clinical Medicine, 2021, 10, 4962.	2.4	7
136	Arrhythmogenic Right Ventricular Cardiomyopathy in Pediatric Patients: An Important but Underrecognized Clinical Entity. Frontiers in Pediatrics, 2021, 9, 750916.	1.9	7
137	Ability of Patients to Distinguish Among Cardiac Genomic Variant Subclassifications. Circulation Genomic and Precision Medicine, 2018, 11, e001975.	3.6	6
138	Nocturnal Premature Ventricular Contraction Burden as a Marker ofÂDisease Severity inÂArrhythmogenic RightÂVentricular Cardiomyopathy. JACC: Clinical Electrophysiology, 2017, 3, 1607-1608.	3.2	5
139	Risk stratification for ventricular arrhythmias and sudden cardiac death in arrhythmogenic right ventricular cardiomyopathy: an update. Expert Review of Cardiovascular Therapy, 2019, 17, 645-651.	1.5	5
140	Strength of the genetic counselor: patient relationship is associated with extent of increased empowerment in patients with arrhythmogenic cardiomyopathy. Journal of Genetic Counseling, 2022, 31, 388-397.	1.6	5
141	Integrating Exercise Into Personalized Ventricular Arrhythmia Risk Prediction in Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2022, 15, CIRCEP121010221.	4.8	5
142	Influence of Panel Selection on Yield of Clinically Useful Variants in Arrhythmogenic Right Ventricular Cardiomyopathy Families. Circulation Genomic and Precision Medicine, 2020, 13, 548-550.	3.6	4
143	Nature and Nurture in Arrhythmogenic Right Ventricular Cardiomyopathy â€“ A Clinical Perspective. Arrhythmia and Electrophysiology Review, 2015, 4, 156.	2.4	4
144	Genetic Dilated Cardiomyopathy Due to TTN Variants Without Known Familial Disease. Circulation Genomic and Precision Medicine, 2020, 13, e003082.	3.6	4

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145	Arrhythmogenic Right Ventricular Cardiomyopathy Prevalence and Arrhythmic Outcomes in At-Risk Family Members: A Systematic Review and Meta-Analysis. Circulation Genomic and Precision Medicine, 2022, 15, 101161CIRCGEN121003530.	3.6	4
146	Absence of a Primary Role for SCN10A Mutations in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of Cardiovascular Translational Research, 2016, 9, 87-89.	2.4	3
147	Focused revision: An addendum to a National Society of Genetic Counselors (NSGC) practice resource. Journal of Genetic Counseling, 2020, 29, 135-135.	1.6	3
148	Efficacy of catheter ablation for premature ventricular contractions in arrhythmogenic right ventricular cardiomyopathy. Journal of Cardiovascular Electrophysiology, 2021, 32, 1665-1674.	1.7	3
149	Patientâ€™s Guide to Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Circulation, 2014, 130, e89-92.	1.6	2
150	Surgical correction of tricuspid regurgitation in patients with ARVD/C. HeartRhythm Case Reports, 2015, 1, 326-330.	0.4	2
151	Further Progress in Predicting Life-Threatening Arrhythmias in PatientsÂWith Arrhythmogenic RightÂVentricular Cardiomyopathy âˆ—. Journal of the American College of Cardiology, 2016, 68, 2551-2553.	2.8	2
152	No major role for rare plectin variants in arrhythmogenic right ventricular cardiomyopathy. PLoS ONE, 2018, 13, e0203078.	2.5	2
153	Risk Stratification in Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American College of Cardiology, 2020, 75, 2766-2768.	2.8	2
154	Response to Letters Regarding Article, â€œElectrocardiographic Features of Arrhythmogenic Right Ventricular Dysplasiaâ€• Circulation, 2010, 121, .	1.6	1
155	Dual-Organ Transplantation in a WomanÂWith Right Ventricular Failure SecondaryÂto Arrhythmogenic RightÂVentricular Cardiomyopathy. JACC: Case Reports, 2020, 2, 59-63.	0.6	1
156	Abstract 13804: Correlation between Electrocardiographic Features and Local Activation Pattern in Arrhythmogenic Right Ventricular Dysplasia. Circulation, 2014, 130, .	1.6	1
157	Abstract 16584: Abnormal Right Ventricular Strain by Cardiac Magnetic Resonance in Preclinical Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation, 2014, 130, .	1.6	1
158	Cardiac genetic counselor: An important member of your healthcare team. PACE - Pacing and Clinical Electrophysiology, 2018, 41, 1022-1024.	1.2	0
159	The Role of Exercise in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. , 2018, , 299-322.		0
160	Arrhythmogenic right ventricular cardiomyopathy. , 2020, , 375-388.		0
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