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	Clinical characteristics and risk stratification of desmoplakin cardiomyopathy. Europace, 2022, 24, 268-277.	1.7	41
2	Comparing clinical performance of current implantable cardioverter-defibrillator implantation recommendations in arrhythmogenic right ventricular cardiomyopathy. Europace, 2022, 24, 296-305.	1.7	9
3	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
4	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
5	Integrating Exercise Into Personalized Ventricular Arrhythmia Risk Prediction in Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2022, 15, CIRCEP121010221.	4.8	5
6	Multimodality Imaging in Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Cardiovascular Imaging, 2022, 15, CIRCIMAGING121013725.	2.6	17
7	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
8	Heart transplantation strategies in arrhythmogenic right ventricular cardiomyopathy: a tertiary ARVC centre experience. ESC Heart Failure, 2022, 9, 1008-1017.	3.1	9
9	A new prediction model for ventricular arrhythmias in arrhythmogenic right ventricular cardiomyopathy. European Heart Journal, 2022, 43, e1-e9.	2.2	35
10	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
11	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
12	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
13	Keeping balance: Authorâ€™s reply. Europace, 2021, 23, 157-158.	1.7	0
14	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
15	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
16	Sudden Cardiac Death Prediction in Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2021, 14, e008509.	4.8	82
17	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
18	Exercise triggers CAPN1-mediated AIF truncation, inducing myocyte cell death in arrhythmogenic cardiomyopathy. Science Translational Medicine, 2021, 13, .	12.4	46

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19	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
20	Efficacy of catheter ablation for premature ventricular contractions in arrhythmogenic right ventricular cardiomyopathy. Journal of Cardiovascular Electrophysiology, 2021, 32, 1665-1674.	1.7	3
21	Arrhythmogenic Right Ventricular Cardiomyopathy Presenting as Clinical Myocarditis in Women. American Journal of Cardiology, 2021, 145, 128-134.	1.6	38
22	Cadherin 2-Related Arrhythmogenic Cardiomyopathy. Circulation Genomic and Precision Medicine, 2021, 14, e003097.	3.6	21
23	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
24	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
25	Contemporary and Future Approaches toÂPrecision Medicine in InheritedÂCardiomyopathies. Journal of the American College of Cardiology, 2021, 77, 2551-2572.	2.8	11
26	Evidence-Based Assessment of Genes in Dilated Cardiomyopathy. Circulation, 2021, 144, 7-19.	1.6	213
27	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
28	The genetic architecture of Plakophilin 2 cardiomyopathy. Genetics in Medicine, 2021, 23, 1961-1968.	2.4	13
29	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
30	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
31	Anxiety and depression in inherited channelopathy patients with implantable cardioverter-defibrillators. Heart Rhythm O2, 2021, 2, 388-393.	1.7	9
32	Phenotypic Expression, Natural History, and Risk Stratification of Cardiomyopathy Caused by Filamin C Truncating Variants. Circulation, 2021, 144, 1600-1611.	1.6	43
33	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
34	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
35	Arrhythmogenic Right Ventricular Cardiomyopathy in Pediatric Patients: An Important but Underrecognized Clinical Entity. Frontiers in Pediatrics, 2021, 9, 750916.	1.9	7
36	Arrhythmogenic right ventricular cardiomyopathy: evidence for progression increases. European Heart Journal, 2020, 41, 1411-1413.	2.2	10

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37	Arrhythmogenic right ventricular cardiomyopathy. , 2020, , 375-388.		0
38	Psychosocial Stress Hastens Disease Progression and Sudden Death in Mice with Arrhythmogenic Cardiomyopathy. Journal of Clinical Medicine, 2020, 9, 3804.	2.4	13
39	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
40	Risk Stratification in Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American College of Cardiology, 2020, 75, 2766-2768.	2.8	2
41	Exercise restriction is protective for genotype-positive family members of arrhythmogenic right ventricular cardiomyopathy patients. Europace, 2020, 22, 1270-1278.	1.7	23
42	The role of genetics in cardiovascular disease: arrhythmogenic cardiomyopathy. European Heart Journal, 2020, 41, 1393-1400.	2.2	54
43	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
44	FLNC truncations cause arrhythmogenic right ventricular cardiomyopathy. Journal of Medical Genetics, 2020, 57, 254-257.	3.2	43
45	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
46	Dual-Organ Transplantation in a WomanÂWith Right Ventricular Failure SecondaryÂto Arrhythmogenic RightÂVentricular Cardiomyopathy. JACC: Case Reports, 2020, 2, 59-63.	0.6	1
47	Diagnosing arrhythmogenic right ventricular cardiomyopathy by 2010 Task Force Criteria: clinical performance and simplified practical implementation. Europace, 2020, 22, 787-796.	1.7	40
48	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
49	Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. European Heart Journal, 2020, 41, 1414-1429.	2.2	239
50	Genetic Dilated Cardiomyopathy Due to TTN Variants Without Known Familial Disease. Circulation Genomic and Precision Medicine, 2020, 13, e003082.	3.6	4
51	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
52	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
53	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
54	Arrhythmogenic Right Ventricular Cardiomyopathy-Associated Desmosomal Variants Are Rarely De Novo. Circulation Genomic and Precision Medicine, 2019, 12, e002467.	3.6	38

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55	Cardiac sympathectomy for refractory ventricular tachycardia in arrhythmogenic right ventricular cardiomyopathy. Heart Rhythm, 2019, 16, 1003-1010.	0.7	42
56	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy. Heart Rhythm, 2019, 16, e301-e372.	0.7	494
57	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
58	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
59	Return of secondary findings in genomic sequencing: Military implications. Molecular Genetics & Genomic Medicine, 2019, 7, e00483.	1.2	9
60	Arrhythmogenic Right Ventricular Cardiomyopathy: Progress Toward Personalized Management. Annual Review of Medicine, 2019, 70, 1-18.	12.2	15
61	Diagnostic and therapeutic strategies for arrhythmogenic right ventricular dysplasia/cardiomyopathy patient. Europace, 2019, 21, 9-21.	1.7	33
62	Ankyrin-B dysfunction predisposes to arrhythmogenic cardiomyopathy and is amenable to therapy. Journal of Clinical Investigation, 2019, 129, 3171-3184.	8.2	42
63	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
64	Predicting arrhythmic risk in arrhythmogenic right ventricular cardiomyopathy: A systematic review and meta-analysis. Heart Rhythm, 2018, 15, 1097-1107.	0.7	79
65	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
66	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
67	Communication of genetic information to families with inherited rhythm disorders. Heart Rhythm, 2018, 15, 780-786.	0.7	30
68	Subcutaneous Implantable CardioverterâEĐefibrillator in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia: A Transatlantic Experience. Journal of the American Heart Association, 2018, 7, e008782.	3.7	23
69	Electrocardiographic Features Differentiating Arrhythmogenic RightÂVentricular Cardiomyopathy FromÂan Athleteâ€™s Heart. JACC: Clinical Electrophysiology, 2018, 4, 1613-1625.	3.2	19
70	Managing Secondary Genomic Findings Associated With Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation Genomic and Precision Medicine, 2018, 11, e002237.	3.6	11
71	No major role for rare plectin variants in arrhythmogenic right ventricular cardiomyopathy. PLoS ONE, 2018, 13, e0203078.	2.5	2
72	Cardiac genetic counselor: An important member of your healthcare team. PACE - Pacing and Clinical Electrophysiology, 2018, 41, 1022-1024.	1.2	0

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73	The Role of Exercise in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. , 2018, , 299-322.		0
74	Epicardial Fat Distribution Assessed with Cardiac CT in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Radiology, 2018, 289, 641-648.	7.3	12
75	Ability of Patients to Distinguish Among Cardiac Genomic Variant Subclassifications. Circulation Genomic and Precision Medicine, 2018, 11, e001975.	3.6	6
76	Arrhythmic outcome of arrhythmogenic right ventricular cardiomyopathy patients without implantable defibrillators. Journal of Cardiovascular Electrophysiology, 2018, 29, 1396-1402.	1.7	12
77	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
78	Evaluation of Structural Progression in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. JAMA Cardiology, 2017, 2, 293.	6.1	53
79	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
80	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
81	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
82	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
83	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
84	Psychosocial care and cardiac genetic counseling following sudden cardiac death in the young. Progress in Pediatric Cardiology, 2017, 45, 31-36.	0.4	15
85	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
86	Arrhythmogenic cardiomyopathy: pathology, genetics, and concepts in pathogenesis. Cardiovascular Research, 2017, 113, 1521-1531.	3.8	98
87	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
88	Heart Failure Is Common and Under-Recognized in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Circulation: Heart Failure, 2017, 10, .	3.9	53
89	Minding the Genes: a Multidisciplinary Approach towards Genetic Assessment of Cardiovascular Disease. Journal of Genetic Counseling, 2017, 26, 224-231.	1.6	16
90	Nocturnal Premature Ventricular Contraction Burden as a Marker ofÂDisease Severity inÂArrhythmogenic RightÂVentricular Cardiomyopathy. JACC: Clinical Electrophysiology, 2017, 3, 1607-1608.	3.2	5

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91	Abstract 24032: Exercise Instigates Apoptosis-inducing Factor Nuclear Translocation and Myocyte Death in Arrhythmogenic Cardiomyopathy. Circulation, 2017, 136, .	1.6	0
92	Pregnancy course and outcomes in women with arrhythmogenic right ventricular cardiomyopathy. Heart, 2016, 102, 303-312.	2.9	50
93	Further Progress in Predicting Life-Threatening Arrhythmias in PatientsÂWith Arrhythmogenic RightÂVentricular Cardiomyopathy â^—. Journal of the American College of Cardiology, 2016, 68, 2551-2553.	2.8	2
94	Electroanatomic Correlates of Depolarization Abnormalities in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of Cardiovascular Electrophysiology, 2016, 27, 443-452.	1.7	31
95	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
96	Characterizing the Molecular Pathology of Arrhythmogenic Cardiomyopathy in Patient Buccal Mucosa Cells. Circulation: Arrhythmia and Electrophysiology, 2016, 9, e003688.	4.8	35
97	Fibrofatty Changes: Incidence at Cardiac MR Imaging in Patients with Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Radiology, 2016, 280, 405-412.	7.3	16
98	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
99	Safety of American Heart Association-recommended minimum exercise for desmosomal mutation carriers. Heart Rhythm, 2016, 13, 199-207.	0.7	76
100	Approach to family screening in arrhythmogenic right ventricular dysplasia/cardiomyopathy. European Heart Journal, 2016, 37, 755-763.	2.2	68
101	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
102	Surgical correction of tricuspid regurgitation in patients with ARVD/C. HeartRhythm Case Reports, 2015, 1, 326-330.	0.4	2
103	Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy in the PediatricÂPopulation. JACC: Clinical Electrophysiology, 2015, 1, 551-560.	3.2	74
104	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
105	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
106	Premature Ventricular Contraction Variability in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of Cardiovascular Electrophysiology, 2015, 26, 53-57.	1.7	12
107	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
108	Spectrum of Biventricular Involvement on CMR Among Carriers of ARVD/C-Associated Mutations. JACC: Cardiovascular Imaging, 2015, 8, 863-864.	5.3	25

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109	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
110	Nature and Nurture in Arrhythmogenic Right Ventricular Cardiomyopathy â€“ A Clinical Perspective. Arrhythmia and Electrophysiology Review, 2015, 4, 156.	2.4	4
111	Cardiac MR Findings and Potential Diagnostic Pitfalls in Patients Evaluated for Arrhythmogenic Right Ventricular Cardiomyopathy. Radiographics, 2014, 34, 1553-1570.	3.3	52
112	Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy and Cardiac Sarcoidosis. Circulation: Arrhythmia and Electrophysiology, 2014, 7, 230-236.	4.8	112
113	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
114	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
115	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
116	Patientâ€™s Guide to Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Circulation, 2014, 130, e89-92.	1.6	2
117	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
118	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
119	Abstract 13804: Correlation between Electrocardiographic Features and Local Activation Pattern in Arrhythmogenic Right Ventricular Dysplasia. Circulation, 2014, 130, .	1.6	1
120	Abstract 16584: Abnormal Right Ventricular Strain by Cardiac Magnetic Resonance in Preclinical Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation, 2014, 130, .	1.6	1
121	Update on Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy (ARVD/C). Current Treatment Options in Cardiovascular Medicine, 2013, 15, 476-487.	0.9	16
122	Mutationâ€Positive Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy: The Triangle of Dysplasia Displaced. Journal of Cardiovascular Electrophysiology, 2013, 24, 1311-1320.	1.7	148
123	Prevalence of atrial arrhythmias in arrhythmogenic right ventricular dysplasia/cardiomyopathy. Heart Rhythm, 2013, 10, 1661-1668.	0.7	71
124	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
125	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
126	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

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127	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
128	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
129	Risk Stratification in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathyâ€"Associated Desmosomal Mutation Carriers. Circulation: Arrhythmia and Electrophysiology, 2013, 6, 569-578.	4.8	94
130	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
131	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
132	Outcomes of Catheter Ablation of Ventricular Tachycardia in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2012, 5, 499-505.	4.8	175
133	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
134	Cardiac Transplantation in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of the American College of Cardiology, 2012, 59, 289-290.	2.8	76
135	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
136	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
137	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
138	Response to Letters Regarding Article, â€œElectrocardiographic Features of Arrhythmogenic Right Ventricular Dysplasiaâ€• Circulation, 2010, 121, .	1.6	1
139	Ethical and Practical Guidelines for Reporting Genetic Research Results to Study Participants. Circulation: Cardiovascular Genetics, 2010, 3, 574-580.	5.1	328
140	Comprehensive Desmosome Mutation Analysis in North Americans With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Circulation: Cardiovascular Genetics, 2009, 2, 428-435.	5.1	195
141	Electrocardiographic Features of Arrhythmogenic Right Ventricular Dysplasia. Circulation, 2009, 120, 477-487.	1.6	88
142	Morphologic Variants of Familial Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Journal of the American College of Cardiology, 2009, 53, 1289-1299.	2.8	84
143	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
144	Adults With X-Linked Agammaglobulinemia. Medicine (United States), 2008, 87, 253-258.	1.0	38

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145	Abstract 4116: Electrocardiographic Markers of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy: A reappraisal. Circulation, 2008, 118, .	1.6	0
146	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
147	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
148	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
149	Utility of Tissue Doppler and Strain Echocardiography in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Cardiology, 2007, 100, 507-512.	1.6	73
150	DSG2 Mutations Contribute to Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Human Genetics, 2006, 79, 136-142.	6.2	206
151	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
152	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
153	Feasibility and Variability of Three Dimensional Echocardiography in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Cardiology, 2006, 97, 703-709.	1.6	71
154	Recessive arrhythmogenic right ventricular dysplasia due to novel cryptic splice mutation inPKP2. Human Mutation, 2006, 27, 1157-1157.	2.5	77
155	Clinical Features of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy Associated With Mutations in Plakophilin-2. Circulation, 2006, 113, 1641-1649.	1.6	225
156	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
157	Arrhythmogenic Right Ventricular Dysplasia. Circulation, 2005, 112, 3823-3832.	1.6	434
158	Predictors of appropriate implantable defibrillator therapies in patients with arrhythmogenic right ventricular dysplasia. Heart Rhythm, 2005, 2, 1188-1194.	0.7	114
159	Electrocardiographic Features of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy According to Disease Severity. Circulation, 2004, 110, 1527-1534.	1.6	261
160	Findings on magnetic resonance imaging of idiopathic right ventricular outflow tachycardia. American Journal of Cardiology, 2004, 94, 1441-1445.	1.6	61
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