CITATION REPORT List of articles citing

Clinical Presentation, Long-Term Follow-Up, and Outcomes of 1001 Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy Patients and Family Members

DOI: 10.1161/circgenetics.114.001003 Circulation: Cardiovascular Genetics, 2015, 8, 437-46.

Source: https://exaly.com/paper-pdf/62337902/citation-report.pdf

Version: 2024-04-17

This report has been generated based on the citations recorded by exaly.com for the above article. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

#	Paper	IF	Citations
322	Arrhythmogenic right ventricular dysplasia/cardiomyopathy-three decades of progress. 2015 , 79, 901-1	3	37
321	Left Ventricular Involvement in ARVD/C: Is It Time to Readjust Our Views?. 2015 , 8, 1311-2		4
320	Arrhythmogenic Right Ventricular Cardiomyopathy: Toward a Modern Clinical and Genomic Understanding. <i>Circulation: Cardiovascular Genetics</i> , 2015 , 8, 421-4		6
319	The Value of Cardiac Magnetic Resonance Imaging in Evaluation of Pediatric Patients for Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. 2015 , 66, 873-874		7
318	Reply: The Value of Cardiac Magnetic Resonance Imaging in Evaluation of Pediatric Patients for Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. 2015 , 66, 874-875		
317	Arrhythmogenic Right Ventricular Dysplasia in Neuromuscular Disorders. 2016 , 10, 173-180		4
316	Clinical Presentation and Outcomes by Sex in Arrhythmogenic Right Ventricular Cardiomyopathy: Findings from the North American ARVC Registry. 2016 , 27, 555-62		26
315	Fetal arrhythmogenic right ventricular cardiomyopathy with double mutations in TMEM43. 2016 , 58, 409-411		7
314	Arrhythmogenic right-ventricular cardiomyopathy: molecular genetics into clinical practice in the era of next generation sequencing. 2016 , 17, 399-407		14
313	Pregnancy course and outcomes in women with arrhythmogenic right ventricular cardiomyopathy. 2016 , 102, 303-12		35
312	Further Progress in Predicting Life-Threatening Arrhythmias in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy. 2016 , 68, 2551-2553		1
311	Arrhythmogenic Right Ventricular Cardiomyopathy: Clinical Course and Predictors of Arrhythmic Risk. 2016 , 68, 2540-2550		99
310	Genetics and Genomics of Single-Gene Cardiovascular Diseases: Common Hereditary Cardiomyopathies as Prototypes of Single-Gene Disorders. 2016 , 68, 2831-2849		31
309	The genetic background of arrhythmogenic right ventricular cardiomyopathy. 2016, 32, 398-403		36
308	Arrhythmogenic right ventricular cardiomyopathy: implications of next-generation sequencing in appropriate diagnosis. 2017 , 19, 1063-1069		25
307	The ventricular ectopic QRS interval for diagnosis and risk stratification in arrhythmogenic right ventricular dysplasia/cardiomyopathy: Is this the answer?. 2016 , 13, 1513-4		
306	Cardiac Fibro-Adipocyte Progenitors Express Desmosome Proteins and Preferentially Differentiate to Adipocytes Upon Deletion of the Desmoplakin Gene. 2016 , 119, 41-54		57

(2016-2016)

305	Advances in the Diagnosis and Management of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. 2016 , 18, 53	9
304	Clinical characterisation and risk stratification of patients with arrhythmogenic right ventricular dysplasia/cardiomyopathy B0 years of age. 2016 , 24, 740-747	5
303	A 43-Year-Old Man With Daytime Sleepiness and a Heart Murmur. 2016 , 150, e117-e120	
302	Men and Women in Arrhythmogenic Right Ventricular Cardiomyopathy. 2016 , 2, 556-557	1
301	Right Ventricular Imaging and Computer Simulation for Electromechanical Substrate Characterization in Arrhythmogenic Right Ventricular Cardiomyopathy. 2016 , 68, 2185-2197	33
300	The Canadian Arrhythmogenic Right Ventricular Cardiomyopathy Registry: Rationale, Design, and Preliminary Recruitment. 2016 , 32, 1396-1401	4
299	Long-term right ventricular implantable cardioverter-defibrillator lead performance in arrhythmogenic right ventricular cardiomyopathy. 2016 , 13, 1964-70	9
298	Catecholaminergic Polymorphic Ventricular Tachycardia: Activity as Tolerated?. 2016 , 2, 263-265	
297	2015 update on the diagnosis and management of arrhythmogenic right ventricular cardiomyopathy. 2016 , 31, 46-56	20
296	Recent advances in genetic testing and counseling for inherited arrhythmias. 2016 , 32, 389-397	24
295	Arrhythmogenic cardiomyopathy. 2016 , 11, 33	80
294	Mapping and ablation procedures for the treatment of ventricular tachycardia. 2016 , 14, 1071-87	2
293	Right ventricular strain by MR quantitatively identifies regional dysfunction in patients with arrhythmogenic right ventricular cardiomyopathy. 2016 , 43, 1132-9	31
292	Arrhythmogenic right ventricular dysplasia/cardiomyopathy: arrhythmogenesis in the apparently normal heart?. 2016 , 18, 953-4	1
291	Exercise and Inherited Arrhythmias. 2016 , 32, 452-8	14
290	Myocardial expression profiles of candidate molecules in patients with arrhythmogenic right ventricular cardiomyopathy/dysplasia compared to those with dilated cardiomyopathy and healthy controls. 2016 , 13, 731-41	23
289	Evidence Mounts That Severity of Disease Impacts the Prognosis for Patients With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. 2016 , 2, 116-117	
288	Next generation sequencing-based copy number analysis reveals low prevalence of deletions and duplications in 46 genes associated with genetic cardiomyopathies. 2016 , 4, 143-51	27

287	Safety of American Heart Association-recommended minimum exercise for desmosomal mutation carriers. 2016 , 13, 199-207	54
286	Approach to family screening in arrhythmogenic right ventricular dysplasia/cardiomyopathy. 2016 , 37, 755-63	56
285	Comprehensive multi-modality imaging approach in arrhythmogenic cardiomyopathy-an expert consensus document of the European Association of Cardiovascular Imaging. 2017 , 18, 237-253	88
284	Multilevel analyses of SCN5A mutations in arrhythmogenic right ventricular dysplasia/cardiomyopathy suggest non-canonical mechanisms for disease pathogenesis. 2017 , 113, 102-111	111
283	Arrhythmogenic right ventricular dysplasia/cardiomyopathy. 2017, 27, S57-S61	4
282	Genotype-phenotype relationship in patients with arrhythmogenic right ventricular cardiomyopathy caused by desmosomal gene mutations: A systematic review and meta-analysis. 2017 , 7, 41387	28
281	Potential new mechanisms of pro-arrhythmia in arrhythmogenic cardiomyopathy: focus on calcium sensitive pathways. 2017 , 25, 157-169	25
2 80	Editorial commentary: Wolff-Parkinson-White pattern and syndrome: Where do we stand in 2017?. 2017 , 27, 269-270	
279	Identification of Cadherin 2 () Mutations in Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2017 , 10,	94
278	At the Heart of the Pregnancy: What Prenatal and Cardiovascular Genetic Counselors Need to Know about Maternal Heart Disease. 2017 , 26, 669-688	3
277	Comparison of Features of Fatal Versus Nonfatal Cardiac Arrest in Patients With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. 2017 , 120, 111-117	24
276	Electronic health record phenotype in subjects with genetic variants associated with arrhythmogenic right ventricular cardiomyopathy: a study of 30,716 subjects with exome sequencing. 2017 , 19, 1245-1252	33
275	Characterization of the arrhythmogenic substrate in patients with arrhythmogenic right ventricular cardiomyopathy undergoing ventricular tachycardia ablation. 2017 , 19, 1049-1062	24
274	Distinct fibrosis pattern in desmosomal and phospholamban mutation carriers in hereditary cardiomyopathies. 2017 , 14, 1024-1032	39
273	Whole exome sequencing with genomic triangulation implicates CDH2-encoded N-cadherin as a novel pathogenic substrate for arrhythmogenic cardiomyopathy. 2017 , 12, 226-235	35
272	Arrhythmogenic Right Ventricular Cardiomyopathy. 2017 , 376, 61-72	302
271	Arrhythmogenic cardiomyopathy: pathology, genetics, and concepts in pathogenesis. 2017, 113, 1521-1531	58
270	Unique genetic background and outcome of non-Caucasian Japanese probands with arrhythmogenic right ventricular dysplasia/cardiomyopathy. 2017 , 5, 639-651	10

269	Arrhythmogenic Cardiomyopathy. 2017 , 121, 784-802	167
268	: A qualitative study of psychological sequelae to the implantable cardioverter defibrillator as a treatment for the prevention of sudden cardiac death in arrhythmogenic right ventricular cardiomyopathy 2017 , 6, 2048004017698614	
267	Heart Failure Is Common and Under-Recognized in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. 2017 , 10,	38
266	Plakophilin-2 is required for transcription of genes that control calcium cycling and cardiac rhythm. 2017 , 8, 106	94
265	Analysis of 60 706 Exomes Questions the Role of De Novo Variants Previously Implicated in Cardiac Disease. <i>Circulation: Cardiovascular Genetics</i> , 2017 , 10,	3
264	Sudden Cardiac Death in Genetic Cardiomyopathies. 2017 , 9, 581-603	10
263	A Novel Mutation in Junctional Plakoglobin Causing Lethal Congenital Epidermolysis Bullosa. 2017 , 191, 266-269.e1	4
262	Risk Stratification in Arrhythmogenic Right Ventricular Cardiomyopathy. 2017 , 136, 2068-2082	100
261	Miocardiopat\(\text{B} \) III. Miocardiopat\(\text{B} \) restrictiva. Displasia arritmog\(\text{B} \) ica del ventr\(\text{B} \) ulo derecho. Miocardiopat\(\text{B} \) no compactada. 2017 , 12, 2573-2584	
260	Genetic testing in cardiomyopathies: an update on indications and benefits. 2017 , 32, 189-195	3
259	Combination of ECG and Echocardiography for Identification of Arrhythmic Events in Early ARVC. 2017 , 10, 503-513	45
258	Moving From Multimodality Diagnostic Tests Toward Multimodality Risk Stratification in ARVC. 2017 , 10, 514-517	1
257	Minding the Genes: a Multidisciplinary Approach towards Genetic Assessment of Cardiovascular Disease. 2017 , 26, 224-231	14
256	High proportion of genetic cases in patients with advanced cardiomyopathy including a novel homozygous Plakophilin 2-gene mutation. 2017 , 12, e0189489	20
255	Sudden cardiac death: focus on the genetics of channelopathies and cardiomyopathies. 2017, 24, 56	17
254	Performance of the 2015 International Task Force Consensus Statement Risk Stratification Algorithm for Implantable Cardioverter-Defibrillator Placement in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. 2018 , 11, e005593	8
253	Arrhythmic Risk Stratification for Arrhythmogenic Right Ventricular Cardiomyopathy: Should We Ask Who Is at High Risk or Who Is at Low Risk?. 2018 , 11, e006160	
252	Sequencing of Linkage Region on Chromosome 12p11 Identifies as a Candidate Gene for Left Ventricular Mass in Dominican Families. 2018 , 8, 659-668	2

251	Isolated, premature ventricular complex-induced right ventricular dysfunction mimicking arrhythmogenic right ventricular cardiomyopathy. 2018 , 4, 222-226	2
250	Controversies in Brugada syndrome. 2018 , 28, 284-292	5
249	Whole-Exome Sequencing Reveals and Mutations as a Potential Digenic Cause of Left Ventricular Noncompaction. 2018 , 11, e001966	4
248	Identification of sarcomeric variants in probands with a clinical diagnosis of arrhythmogenic right ventricular cardiomyopathy (ARVC). 2018 , 29, 1004-1009	10
247	Arrhythmogenic right ventricular cardiomyopathy. 2018 , 34, 356-368	4
246	Epidemiology and Clinical Aspects of Genetic Cardiomyopathies. 2018, 14, 119-128	22
245	Genetic Evaluation of Cardiomyopathy-A Heart Failure Society of America Practice Guideline. 2018 , 24, 281-302	160
244	Controversies Surrounding Exercise in Genetic Cardiomyopathies. 2018 , 14, 189-200	1
243	Genetic basis of arrhythmogenic cardiomyopathy. 2018 , 33, 276-281	14
242	Clinical genetic testing in pediatric cardiomyopathy: Is bigger better?. 2018 , 93, 33-40	26
241	Lifelong arrhythmic risk stratification in arrhythmogenic right ventricular cardiomyopathy: distribution of events and impact of periodical reassessment. 2018 , 20, f20-f29	9
240	Heart transplantation in arrhythmogenic right ventricular cardiomyopathy - Experience from the Nordic ARVC Registry. 2018 , 250, 201-206	14
239	2017 AHA/ACC/HRS guideline for management of patients with ventricular arrhythmias and the prevention of sudden cardiac death: Executive summary: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Heart	264
238	2017 AHA/ACC/HRS guideline for management of patients with ventricular arrhythmias and the prevention of sudden cardiac death: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Heart Rhythm Society. 2018 ,	151
237	2017 AHA/ACC/HRS Guideline for Management of Patients With Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death: Executive Summary: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Heart	180
236	2017 AHA/ACC/HRS Guideline for Management of Patients With Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death: Executive Summary: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Heart	173
235	2017 AHA/ACC/HRS Guideline for Management of Patients With Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Heart Rhythm Society. 2018 ,	307
	138, e272-e391	

233	Ventricular tachycardia ablation in arrhythmogenic right ventricular cardiomyopathy patients with TMEM43 gene mutations. 2018 , 29, 90-97	6
232	Arrhythmogenic right ventricular dysplasia/cardiomyopathy: an electrocardiogram-based review. 2018 , 20, f3-f12	16
231	Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. 2018, 192-204	
230	Blockade of the Adenosine 2A Receptor Mitigates the Cardiomyopathy Induced by Loss of Plakophilin-2 Expression. 2018 , 9, 1750	7
229	Bioinformatic analysis of a plakophilin-2-dependent transcription network: implications for the mechanisms of arrhythmogenic right ventricular cardiomyopathy in humans and in boxer dogs. 2018 , 20, iii125-iii132	9
228	OBSOLETE: Arrhythmogenic Right Ventricular Cardiomyopathy. 2018,	
227	OBSOLETE: Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. 2018,	1
226	Pleiotropic Phenotypes Associated With PKP2 Variants. 2018 , 5, 184	10
225	Catheter Ablation of Ventricular Tachycardia in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. 2018 , 48, 890-905	2
224	Electrocardiographic Features Differentiating Arrhythmogenic Right Ventricular Cardiomyopathy From an Athlete's Heart. 2018 , 4, 1613-1625	9
223	Additional Genetic Variants in Inherited Dilated Cardiomyopathy: Just Another Brick in the Wall?. 2018 , 11, e002249	1
222	Managing Secondary Genomic Findings Associated With Arrhythmogenic Right Ventricular Cardiomyopathy: Case Studies and Proposal for Clinical Surveillance. 2018 , 11, e002237	11
221	No major role for rare plectin variants in arrhythmogenic right ventricular cardiomyopathy. 2018 , 13, e0203078	2
220	Frequency of genetic variants associated with arrhythmogenic right ventricular cardiomyopathy in the genome aggregation database. 2018 , 26, 1312-1318	20
219	Arrhythmogenic Right Ventricular Cardiomyopathy. 2018, 182-191	
218	Arrhythmogenic Right Ventricular Cardiomyopathy. 2018, 291-296	
217	Right ventricular dysplasia: management and treatment in light of current evidence. 2018 , 8, 101-106	1
216	Clinical Diagnosis, Imaging, and Genetics of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia: JACC State-of-the-Art Review. 2018 , 72, 784-804	105

215	Prediction of Life-Threatening Ventricular Arrhythmia in Patients With Arrhythmogenic Cardiomyopathy: A Primary Prevention Cohort Study. 2018 , 11, 1377-1386	37
214	Primary prevention implantable cardioverter-defibrillator and opportunities for sudden cardiac death risk assessment in non-ischaemic cardiomyopathy. 2018 , 39, 2859-2866	20
213	Absence of a primary role for TTN missense variants in arrhythmogenic cardiomyopathy: From a clinical and pathological perspective. 2018 , 41, 615-622	5
212	2017 AHA/ACC/HRS Guideline for Management of Patients With Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Heart Rhythm Society. 2018 ,	411
211	Atrial involvement in arrhythmogenic right ventricular cardiomyopathy patients referred for ventricular arrhythmias ablation. 2018 , 29, 1388-1395	3
210	Genetic evaluation of cardiomyopathy: a clinical practice resource of the American College of Medical Genetics and Genomics (ACMG). 2018 , 20, 899-909	96
209	Arrhythmic outcome of arrhythmogenic right ventricular cardiomyopathy patients without implantable defibrillators. 2018 , 29, 1396-1402	7
208	Impact of Exercise Restriction on Arrhythmic Risk Among Patients With Arrhythmogenic Right Ventricular Cardiomyopathy. 2018 , 7,	37
207	Prevention of Sudden Cardiac Death in Arrhythmogenic Cardiomyopathy. 2018, 4, 769-770	О
206	A founder homozygous DSG2 variant in East Asia results in ARVC with full penetrance and heart failure phenotype. 2019 , 274, 263-270	17
205	Arrhythmogenic Right Ventricular Cardiomyopathy: A Review of Living and Deceased Probands. 2019 , 28, 1034-1041	4
204	Beyond the One Gene-One Disease Paradigm: Complex Genetics and Pleiotropy in Inheritable Cardiac Disorders. 2019 , 140, 595-610	56
203	For Whom the Bell Tolls: Refining Risk Assessment for Sudden Cardiac Death. 2019 , 21, 106	3
202	Disruption of Ca Homeostasis and Connexin 43 Hemichannel Function in the Right Ventricle Precedes Overt Arrhythmogenic Cardiomyopathy in Plakophilin-2-Deficient Mice. 2019 , 140, 1015-1030	46
201	Phenotypic Manifestations of Arrhythmogenic Cardiomyopathy in Children and Adolescents. 2019 , 74, 346-358	31
200	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy: Executive summary. 2019 , 16, e373-e407	73
199	Plakophilin-2 Haploinsufficiency Causes Calcium Handling Deficits and Modulates the Cardiac Response Towards Stress. 2019 , 20,	23
198	Arrhythmogenic Right Ventricular Cardiomyopathy-Associated Desmosomal Variants Are Rarely De Novo. 2019 , 12, e002467	23

197	Desmoglein 2 mutation provokes skeletal muscle actin expression and accumulation at intercalated discs in murine hearts. 2019 , 132,	3
196	Catecholaminergic Polymorphic (Right) Ventricular Tachycardia?. 2019 , 5, 128-130	
195	Heart failure in patients with arrhythmogenic right ventricular cardiomyopathy: Genetic characteristics. 2019 , 286, 99-103	4
194	From Hypertrophy to Heart Failure: What Is New in Genetic Cardiomyopathies. 2019 , 16, 157-167	7
193	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy. 2019 , 16, e301-e372	247
192	Sudden death related cardiomyopathies - Arrhythmogenic right ventricular cardiomyopathy, arrhythmogenic cardiomyopathy, and exercise-induced cardiomyopathy. 2019 , 62, 217-226	7
191	The Netherlands Arrhythmogenic Cardiomyopathy Registry: design and status update. 2019 , 27, 480-486	13
190	Inherited cardiomyopathies. 2019 , 365, l1570	8
189	Risk score for the exclusion of arrhythmic events in arrhythmogenic right ventricular cardiomyopathy at first presentation. 2019 , 290, 100-105	4
188	Molecular mechanisms of arrhythmogenic cardiomyopathy. 2019 , 16, 519-537	77
187	Arrhythmogenic Right Ventricular Cardiomyopathy. 2019 , 410-419.e3	
186	Genotype-phenotype correlations in ARVC: Toward a precision medicine approach. 2019 , 286, 115-116	O
185	Heart failure in arrhythmogenic cardiomyopathy: is phenotypic variability just a matter of genetics?. 2019 , 21, 801-802	1
184	Arrhythmogenic cardiomyopathies (ACs): diagnosis, risk stratification and management. 2019 , 105, 1117-112	8 11
183	Life-threatening arrhythmic presentation in patients with arrhythmogenic cardiomyopathy before and after entering the genomic era; a two-decade experience from a large volume center. 2019 , 279, 79-83	5
182	Prevalence and Electronic Health Record-Based Phenotype of Loss-of-Function Genetic Variants in Arrhythmogenic Right Ventricular Cardiomyopathy-Associated Genes. 2019 , 12, e002579	20
181	Plakophilin-2 Truncation Variants in Patients Clinically Diagnosed With Catecholaminergic Polymorphic Ventricular Tachycardia and Decedents With Exercise-Associated Autopsy Negative Sudden Unexplained Death in the Young. 2019 , 5, 120-127	24
180	A targeted next-generation gene panel reveals a novel heterozygous nonsense variant in the TP63 gene in patients with arrhythmogenic cardiomyopathy. 2019 , 16, 773-780	7

179	Arrhythmogenic Cardiomyopathy in 2018-2019: ARVC/ALVC or Both?. 2019, 28, 164-177	35
178	Arrhythmogenic Right Ventricular Cardiomyopathy: Progress Toward Personalized Management. 2019 , 70, 1-18	9
177	Diagnostic and therapeutic strategies for arrhythmogenic right ventricular dysplasia/cardiomyopathy patient. 2019 , 21, 9-21	19
176	Arrhythmogenic right ventricular cardiomyopathy: evidence for progression increases. 2020 , 41, 1411-1413	7
175	Distal myopathy induced arrhythmogenic right ventricular cardiomyopathy in a pedigree carrying novel DSG2 null variant. 2020 , 298, 25-31	3
174	Catheter ablation of electrical storm in patients with arrhythmogenic right ventricular cardiomyopathy. 2020 , 17, 41-48	5
173	The merits of the ICD for inherited heart rhythm disorders: A critical re-appraisal. 2020 , 30, 415-421	1
172	Arrhythmogenic right ventricular cardiomyopathy. 2020 , 375-388	
171	Modeling Inherited Cardiomyopathies in Adult Zebrafish for Precision Medicine. 2020, 11, 599244	6
170	The Novel Desmin Variant p.Leu115Ile Is Associated With a Unique Form of Biventricular Arrhythmogenic Cardiomyopathy. 2021 , 37, 857-866	7
169	Syncope in a 14yo female with ventricular tachycardia; an atypical etiology. 2021 , 45, 677.e5-677.e7	0
168	Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia: Mechanisms and Management. 2020 , Volume 11, 19-29	1
167	Arrhythmogenic cardiomyopathy: An in-depth look at molecular mechanisms and clinical correlates. 2021 , 31, 395-402	9
166	Forty-six year old man with palpitations. 2020 , 106, 1235-1282	
165	Potential role of imaging markers in predicting future disease expression of arrhythmogenic cardiomyopathy. 2021 , 17, 647-654	5
164	Established and Emerging Mechanisms in the Pathogenesis of Arrhythmogenic Cardiomyopathy: A Multifaceted Disease. 2020 , 21,	9
163	Arrhythmogenic Right Ventricular Cardiomyopathy: An Exuberant Case Affecting Both Ventricles. 2020 , 13, e010243	1
162	Secondary findings in inherited heart conditions: a genotype-first feasibility study to assess phenotype, behavioural and psychosocial outcomes. 2020 , 28, 1486-1496	5

(2020-2020)

161	State of the Art Review on Genetics and Precision Medicine in Arrhythmogenic Cardiomyopathy. 2020 , 21,	9
160	Characteristics of Patients With Arrhythmogenic Left Ventricular Cardiomyopathy: Combining Genetic and Histopathologic Findings. 2020 , 13, e009005	5
159	Familial Arrhythmogenic Cardiomyopathy: Clinical Determinants of Phenotype Discordance and the Impact of Endurance Sports. 2020 , 9,	1
158	Arrhythmogenic right ventricular cardiomyopathy in patients with biallelic JUP-associated skin fragility. 2020 , 10, 21622	3
157	Development of an algorithm for automatic classification of right ventricle deformation patterns in arrhythmogenic right ventricular cardiomyopathy. 2020 , 37, 698-705	2
156	Desmosome-Dyad Crosstalk: An Arrhythmogenic Axis in Arrhythmogenic Right Ventricular Cardiomyopathy. 2020 , 141, 1494-1497	2
155	Parameter subset reduction for patient-specific modelling of arrhythmogenic cardiomyopathy-related mutation carriers in the CircAdapt model. 2020 , 378, 20190347	4
154	Genotype-Phenotype Correlation: A Triple DNA Mutational Event in a Boy Entering Sport Conveys an Additional Pathogenicity Risk. 2020 , 11,	10
153	Arrhythmogenic Cardiomyopathy: Molecular Insights for Improved Therapeutic Design. 2020, 7,	6
152	Exercise restriction is protective for genotype-positive family members of arrhythmogenic right ventricular cardiomyopathy patients. 2020 , 22, 1270-1278	12
151	Right Heart Failure: Causes and Clinical Epidemiology. 2020 , 38, 175-183	4
150	Absence of ECG Task Force Criteria does not rule out structural changes in genotype positive ARVC patients. 2020 , 317, 152-158	2
149	Arrhythmogenic right ventricular cardiomyopathy: Evolving from unique clinical features to a complex pathophysiological concept. 2020 , 45, 243-251	2
148	The role of genetics in cardiovascular disease: arrhythmogenic cardiomyopathy. 2020 , 41, 1393-1400	30
147	Natural History of Arrhythmogenic Cardiomyopathy. 2020 , 9,	19
146	Integrin I ID Deficiency-Mediated RyR2 Dysfunction Contributes to Catecholamine-Sensitive Ventricular Tachycardia in Arrhythmogenic Right Ventricular Cardiomyopathy. 2020 , 141, 1477-1493	14
145	Emerging concepts in arrhythmogenic dilated cardiomyopathy. 2021 , 26, 1219-1229	8
144	Predicting Heart Failure in Arrhythmogenic Right Ventricular Cardiomyopathy. 2020 , 9, e015702	1

143	Genetic basis and molecular biology of cardiac arrhythmias in cardiomyopathies. 2020, 116, 1600-1619	15
142	Arrhythmogenic cardiomyopathy: pathogenesis, pro-arrhythmic remodelling, and novel approaches for risk stratification and therapy. 2020 , 116, 1571-1584	13
141	Dual-Organ Transplantation in a Woman With Right Ventricular Failure Secondary to Arrhythmogenic Right Ventricular Cardiomyopathy. 2020 , 2, 59-63	
140	Genetic variants of uncertain significance: How to match scientific rigour and standard of proof in sudden cardiac death?. 2020 , 45, 101712	12
139	Right Ventricular Strain Predicts Structural Disease Progression in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy. 2020 , 9, e015016	12
138	Epidemiology of the inherited cardiomyopathies. 2021 , 18, 22-36	32
137	Transcriptomic Coupling of PKP2 With Inflammatory and Immune Pathways Endogenous to Adult Cardiac Myocytes. 2020 , 11, 623190	6
136	Sex Differences in Right Ventricular Dysfunction: Insights From the Bench to Bedside. 2020 , 11, 623129	4
135	Two Novel Variants in Genes of Arrhythmogenic Right Ventricular Cardiomyopathy - a Case Report. 2021 , 28, 127-135	
134	Risk stratification and subclinical phenotyping of dilated and/or arrhythmogenic cardiomyopathy mutation-positive relatives: CVON eDETECT consortium. 2021 , 29, 301-308	
133	Deciphering DSC2 arrhythmogenic cardiomyopathy electrical instability: From ion channels to ECG and tailored drug therapy. 2021 , 11, e319	3
132	Exercise triggers CAPN1-mediated AIF truncation, inducing myocyte cell death in arrhythmogenic cardiomyopathy. 2021 , 13,	15
131	Rare Variants Associated with Arrhythmogenic Cardiomyopathy: Reclassification Five Years Later. 2021 , 11,	3
130	Impact of Genetic Variant Reassessment on the Diagnosis of Arrhythmogenic Right Ventricular Cardiomyopathy Based on the 2010 Task Force Criteria. 2021 , 14, e003047	4
129	Electromechanical substrate characterization in arrhythmogenic cardiomyopathy using imaging-based patient-specific computer simulations. 2021 , 23, i153-i160	3
128	Arrhythmogenic Cardiomyopathy: Mechanisms, Genetics, and Their Clinical Implications. <i>Current Cardiovascular Risk Reports</i> , 2021 , 15, 1	1
127	Risk Stratification in Arrhythmogenic Right Ventricular Cardiomyopathy. 2021 , 10, 26-32	1
126	Update on the Diagnostic Pitfalls of Autopsy and Post-Mortem Genetic Testing in Cardiomyopathies. 2021 , 22,	5

125	Arrhythmogenic Right Ventricular Cardiomyopathy Presenting as Clinical Myocarditis in Women. 2021 , 145, 128-134	12
124	Cadherin 2-Related Arrhythmogenic Cardiomyopathy: Prevalence and Clinical Features. 2021 , 14, e003097	8
123	Clinical Findings and Diagnostic Yield of Arrhythmogenic Cardiomyopathy Through Genomic Screening of Pathogenic or Likely Pathogenic Desmosome Gene Variants. 2021 , 14, e003302	4
122	Genetics of Cardiomyopathy.	
121	Actionable secondary findings in arrhythmogenic right ventricle cardiomyopathy genes: impact and challenge of genetic counseling. 2021 , 11, 637-649	0
120	Clinical Insights Into Heritable Cardiomyopathies. 2021 , 12, 663450	3
119	An actuarial model of arrhythmogenic right ventricular cardiomyopathy and life insurance. 1-21	
118	Arrhythmogenic Left Ventricular Cardiomyopathy: Genotype-Phenotype Correlations and New Diagnostic Criteria. 2021 , 10,	5
117	Peeking Beyond Strain's Peak: Regional Strain Patterns and Dispersion in Arrhythmogenic Right Ventricular Cardiomyopathy. 2021 , 14, 911-914	1
116	Contemporary and Future Approaches to Precision Medicine in Inherited Cardiomyopathies: JACC Focus Seminar 3/5. 2021 , 77, 2551-2572	4
115	Pregnancy in arrhythmogenic cardiomyopathy. 2021 , 32, 186-198	0
114	Right Ventricular Functional Abnormalities in Arrhythmogenic Cardiomyopathy: Association With Life-Threatening Ventricular Arrhythmias. 2021 , 14, 900-910	9
113	State-of-the-art narrative review: multimodality imaging in electrophysiology and cardiac device therapies. 2021 , 11, 881-895	1
112	Diagnostic yield of targeted next generation sequencing in 2002 Dutch cardiomyopathy patients. 2021 , 332, 99-104	2
111	Sporadic and rapidly progressive arrhythmogenic right ventricular cardiomyopathy in a 12-year-old boy who was diagnosed with epilepsy. 2021 , 2021, omab046	0
110	The EP300/TP53 pathway, a suppressor of the Hippo and canonical WNT pathways, is activated in human hearts with arrhythmogenic cardiomyopathy in the absence of overt heart failure. 2021 ,	2
109	Beneficial effect of voluntary physical exercise in Plakophilin2 transgenic mice. 2021 , 16, e0252649	0
108	Pathogenic variants in plakophilin-2 gene (PKP2) are associated with better survival in arrhythmogenic right ventricular cardiomyopathy. 2021 , 62, 613-620	2

107	International Evidence Based Reappraisal of Genes Associated With Arrhythmogenic Right Ventricular Cardiomyopathy Using the Clinical Genome Resource Framework. 2021 , 14, e003273	21
106	Cardiomyopathies: An Overview. 2021 , 22,	6
105	Altered Electrical, Biomolecular, and Immunologic Phenotypes in a Novel Patient-Derived Stem Cell Model of Desmoglein-2 Mutant ARVC. 2021 , 10,	2
104	Anatomical-MRI Correlations in Adults and Children with Arrhythmogenic Right Ventricular Cardiomyopathy. 2021 , 11,	
103	Effect of preload reducing therapy on right ventricular size and function in patients with arrhythmogenic right ventricular cardiomyopathy. 2021 , 18, 1186-1191	3
102	Genotype-phenotype correlation in arrhythmogenic right ventricular cardiomyopathy-risk of arrhythmias and heart failure. 2021 ,	1
101	Outcomes after catheter ablation of ventricular tachycardia without implantable cardioverter-defibrillator in selected patients with arrhythmogenic right ventricular cardiomyopathy. 2021 , 23, 1428-1436	0
100	Peculiar Aspects of Patients with Inherited Arrhythmias during the COVID-19 Pandemic. 2021 , 117, 394-403	О
99	Clinical characteristics and risk stratification of desmoplakin cardiomyopathy. 2021,	6
98	Desmoplakin cardiomyopathy and arrhythmogenic right ventricular cardiomyopathy: two distinct forms of cardiomyopathy?. 2021 ,	
97	Epicardial differentiation drives fibro-fatty remodeling in arrhythmogenic cardiomyopathy. 2021 , 13, eabf2750	3
96	Contribution of concomitant myocarditis to the development of various clinical types of arrhythmogenic right ventricular cardiomyopathy. 2021 , 20, 2781	1
95	Running the Risk: Exercise and Arrhythmogenic Cardiomyopathy 2021, 23, 1	
94	Comparing clinical performance of current implantable cardioverter-defibrillator implantation recommendations in arrhythmogenic right ventricular cardiomyopathy. 2021 ,	1
93	The clinical utility of pediatric cardiomyopathy genetic testing: From diagnosis to a precision medicine-based approach to care. 2021 , 62, 101413-101413	0
92	Uncertainty Quantification of Regional Cardiac Tissue Properties in Arrhythmogenic Cardiomyopathy Using Adaptive Multiple Importance Sampling. 2021 , 12, 738926	1
91	Molecular autopsy and subsequent functional analysis reveal de novo DSG2 mutation as cause of sudden death. 2021 , 64, 104322	
90	Evaluation and monitoring of patients with cardiomyopathies (including myocardial infiltration). 2022 , 77-104.e5	

89	Genetic Restrictive Cardiomyopathy: Causes and Consequences-An Integrative Approach. 2021, 22,	8
88	Inherited cardiomyopathies. 2021 , 277-290	
87	Electrocardiographic features of arrhythmogenic right ventricular cardiomyopathy in school-aged children. 2021 , 36, 863-873	
86	Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. 2020 , 41, 1414-1429	110
85	Appropriate and Inappropriate Implantable Cardioverter Defibrillators Therapies in Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia Patients. 2018 , 9, 204-214	5
84	Nature and Nurture in Arrhythmogenic Right Ventricular Cardiomyopathy - A Clinical Perspective. 2015 , 4, 156-62	2
83	Wnt/Etatenin pathway in arrhythmogenic cardiomyopathy. 2017 , 8, 60640-60655	34
82	Practical Aspects in Genetic Testing for Cardiomyopathies and Channelopathies. 2019 , 40, 187-200	4
81	Circulation and Contacts in Sixteenth Century New Cartography: Spain, Portugal and Italy. 2021 , 10, e015	
80	Heart Failure in Patients with Arrhythmogenic Cardiomyopathy. 2021 , 10,	O
79	How exercise can deteriorate the clinical course of an ARVC patient: a case report. 2021, 5, ytab417	0
78	Diagnostic Evaluation of Children with Known or Suspected ARVC/D. 2016 , 105-113	
77	Arrhythmogenic Cardiomyopathy. 2016 , 91-111	
76	Arrhythmogenic Cardiomyopathy. 2018 , 6-9	
75	Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death Guideline Update. 2020 , 31, 221-227	0
74	SARS-CoV-2 protein Nsp1 alters actomyosin cytoskeleton and phenocopies arrhythmogenic cardiomyopathy-related PKP2 mutant.	
73	Arrhythmogenic right ventricular cardiomyopathy in the pediatric population. 2022, 37, 99-108	4
72	Arrhythmogenic Right Ventricular Cardiomyopathy. 2020 , 791-810	

71	Arrhythmogenic Cardiomyopathy. 2020 , 99-114	
70	Arrhythmogenic Right Ventricular Cardiomyopathy (ARVC). 2020 , 93-123	
69	Genetic and Molecular Basis of Cardiac Arrhythmias. 2020 , 75-96	
68	CLINICAL CASE OF NON-COMPACTION CARDIOMYOPATHY IN A PATIENT WITH PKP2 GENE MUTATION. 2020 , 106-111	
67	Arrhythmogenic right ventricular cardiomyopathy in Boxer dogs: the diagnosis as a link to the human disease. 2017 , 36, 135-150	5
66	Identification of Disrupted Myocardial Calcium Homeostasis as Proarrhythmic Trigger in Arrhythmogenic Cardiomyopathy. 2021 , 12, 732573	
65	In silico Identification of Disrupted Myocardial Calcium Homeostasis as Proarrhythmic Trigger in Arrhythmogenic Cardiomyopathy. 2021 , 12, 732573	1
64	MicroRNAs: From Junk RNA to Life Regulators and Their Role in Cardiovascular Disease. 2021 , 11, 230-254	
63	Clinical and genetic features of arrhythmogenic cardiomyopathy: the electrophysiology perspective. 2021 , 101463	
62	Spectrum of desmosomal gene variations in patients with arrhythmogenic right ventricular cardiomyopathy. 2021 , 26, 4692	О
61	The genetic basis of sudden death in young people - Cardiac and non-cardiac. 2021, 810, 146067	
60	Arrhythmogenic Right Ventricular Cardiomyopathy in Pediatric Patients: An Important but Underrecognized Clinical Entity 2021 , 9, 750916	2
59	Cardiovascular Characteristics of Patients with Genetic Variation in Desmoplakin () 2022, 12, 24-36	2
58	Integrating Exercise Into Personalized Ventricular Arrhythmia Risk Prediction in Arrhythmogenic Right Ventricular Cardiomyopathy 2022 , CIRCEP121010221	О
57	Penetrance and disease expression of (likely) pathogenic variants associated with inherited cardiomyopathies in the general population.	
56	Role of genetic testing in cardiomyopathies: [þrimer for cardiologists 2022 , 14, 29-39	1
55	Arrhythmogenic Right Ventricular Cardiomyopathy in an Older Private Pilot 2022, 93, 111-115	
54	Multimodality Imaging in Arrhythmogenic Right Ventricular Cardiomyopathy 2022 , CIRCIMAGING121013725	5 0

53	Arrhythmogenic Right Ventricular Cardiomyopathy and Differential Diagnosis with Diseases Mimicking Its Phenotypes 2022 , 11,	3
52	Clinical Characteristics and Follow-Up of Pediatric-Onset Arrhythmogenic Right Ventricular Cardiomyopathy 2022 , 8, 306-318	2
51	Atomic Force Microscopy (AFM) Applications in Arrhythmogenic Cardiomyopathy 2022, 23,	2
50	Arrhythmogenic Right Ventricular Cardiomyopathy: Overview and Case Study 2022, 33, 14-22	
49	European Heart Rhythm Association (EHRA)/Heart Rhythm Society (HRS)/Asia Pacific Heart Rhythm Society (APHRS)/Latin American Heart Rhythm Society (LAHRS) Expert Consensus Statement on the state of genetic testing for cardiac diseases 2022 ,	5
48	Elucidating arrhythmogenic right ventricular cardiomyopathy with stem cells 2022,	
47	European Heart Rhythm Association (EHRA)/Heart Rhythm Society (HRS)/Asia Pacific Heart Rhythm Society (APHRS)/Latin American Heart Rhythm Society (LAHRS) Expert Consensus Statement on the state of genetic testing for cardiac diseases 2022 ,	6
46	Practitioners' Confidence and Desires for Education in Cardiovascular and Sudden Cardiac Death Genetics 2022 , e023763	
45	European Heart Rhythm Association (EHRA)/Heart Rhythm Society (HRS)/Asia Pacific Heart Rhythm Society (APHRS)/Latin American Heart Rhythm Society (LAHRS) Expert Consensus Statement on the state of genetic testing for cardiac diseases.	2
44	Myocarditis or inherited disease? - The multifaceted presentation of arrhythmogenic cardiomyopathy 2022 , 827, 146470	0
43	Ventricular arrhythmia management in patients with genetic cardiomyopathies 2021, 2, 819-831	
42	Histopathological Features and Protein Markers of Arrhythmogenic Cardiomyopathy 2021 , 8, 746321	1
41	Heart transplantation strategies in arrhythmogenic right ventricular cardiomyopathy: a tertiary ARVC centre experience 2021 ,	1
40	Pathogenesis, Diagnosis and Risk Stratification in Arrhythmogenic Cardiomyopathy. 2021 , 11, 263-289	
39	The Female Athlete's Heart: Overview and Management of Cardiovascular Diseases 2021, 16, e47	2
38	Fibro-fatty remodelling in arrhythmogenic cardiomyopathy 2022 , 117, 22	O
37	Arrhythmogenic Right Ventricular Cardiomyopathy 2022 , 8, 533-553	1
36	Table_1.XLS. 2018 ,	

New Variant in Placophilin-2 Gene Causing Arrhythmogenic Myocardiopathy. **2022**, 13, 782

34	Arrhythmogenic Right Ventricular Dysplasia (ARVD) With Protein Plakophilin-2 Mutation. 2022,	
33	Exercise Causes Arrhythmogenic Remodeling of Intracellular Calcium Dynamics in Plakophilin-2-Deficient Hearts 2022 ,	2
32	Population Prevalence of Premature Truncating Variants in Plakophilin-2 and Association With Arrhythmogenic Right Ventricular Cardiomyopathy: a UK Biobank Analysis 2022 , 101161CIRCGEN121	003507 ^O
31	Arrhythmogenic Right Ventricular Cardiomyopathy Prevalence and Arrhythmic Outcomes in At-Risk Family Members: A Systematic Review and Meta-Analysis 2022 , 101161CIRCGEN121003530	O
30	Device-measured physical activity, sedentary time, and sleep in patients with arrhythmogenic cardiomyopathy: descriptive values and stability over 30 measurement days.	O
29	Arrhythmias as Presentation of Genetic Cardiomyopathy. 2022, 130, 1698-1722	О
28	The value of genetic testing in the diagnosis and risk stratification of arrhythmogenic right ventricular cardiomyopathy. 2022 ,	
27	Arrhythmic risk prediction in arrhythmogenic right ventricular cardiomyopathy: external validation of the arrhythmogenic right ventricular cardiomyopathy risk calculator.	1
26	Burden of rare variants in arrhythmogenic cardiomyopathy with right dominant form associated genes provides new insights for molecular diagnosis and clinical management	O
25	Family History and Warning Symptoms Precede Sudden Cardiac Death in Arrhythmogenic Right Ventricular Cardiomyopathy (from a Nationwide Study in Sweden). 2022 ,	
24	A rare case of arrhythmogenic right ventricular cardiomyopathy associated with LAMA2 mutation: A case report and literature review. 9,	O
23	Novel Risk Prediction Model to Determine Adverse Heart Failure Outcomes in Arrhythmogenic Right Ventricular Cardiomyopathy. 2022 , 11,	О
22	Sex Differences in Cardiomyopathy. Current Cardiovascular Risk Reports,	0.9
21	Spectrum of Rare and Common Genetic Variants in Arrhythmogenic Cardiomyopathy Patients. 2022 , 12, 1043	
20	Genetic testing for cardiomyopathies when science and health policies join in personalizing cardiovascular prevention.	
19	Loss of Nuclear Envelope Integrity and Increased Oxidant Production Cause DNA Damage in Adult Hearts Deficient in PKP2: A Molecular Substrate of ARVC. 2022 , 146, 851-867	2
18	The Athletel Heart Thallenges and Controversies. 2022 , 80, 1346-1362	O

CITATION REPORT

17	Humanized Dsp ACM Mouse Model Displays Stress-Induced Cardiac Electrical and Structural Phenotypes. 2022 , 11, 3049	1
16	Genotypeβhenotype Correlates in Arrhythmogenic Cardiomyopathies.	Ο
15	Prevalence and Phenotypic Burden of Monogenic Arrhythmias Using Integration of Electronic Health Records With Genetics.	0
14	Genetics and Genomics of Congenital and Acquired Cardiovascular Disease. 2021, 1-41	Ο
13	Arrhythmogenic Cardiomyopathy: A Review of a Rare Case of Biventricular Phenotype. 2022,	O
12	Prevalence and Disease Expression of Pathogenic and Likely Pathogenic Variants Associated With Inherited Cardiomyopathies in the General Population.	O
11	Exercise After Acute Myocarditis. 2023, 41, 107-115	O
10	Clinical characteristics of various arrhythmogenic cardiomyopathy phenotypes in the pediatric population: a systematic review and meta-analysis. 2022 , 27, 5146	O
9	Clinical and echocardiographic evolution of patients with arrhythmogenic cardiomyopathy before heart transplantation.	O
8	Clinical course of arrhythmogenic right ventricular cardiomyopathy with end-stage heart failure and outcome after heart transplantation. 2022 ,	O
7	Modeling incomplete penetrance in arrhythmogenic cardiomyopathy by human induced pluripotent stem cell derived cardiomyocytes. 2023 , 21, 1759-1773	O
6	Catalytic antibodies in arrhythmogenic cardiomyopathy patients cleave desmoglein 2 and N-cadherin and impair cardiomyocyte cohesion.	O
5	A Crossroads Junction That Leads to Heart Failure (Arrhythmogenic Cardiomyopathy): Hope for Future Therapeutics. 2023 , 13, 31-32	О
4	Cardiomyopathies in children: An overview. 2023 ,	O
3	Advances in Ion Channel, Non-Desmosomal Variants and Autophagic Mechanisms Implicated in Arrhythmogenic Cardiomyopathy. 2023 , 45, 2186-2200	О
2	Connexin hemichannels as candidate targets for cardioprotective and anti-arrhythmic treatments. 2023 , 133,	O
1	Disease modeling of desmosome-related cardiomyopathy using induced pluripotent stem cell-derived cardiomyocytes. 15, 71-82	O