

CITATION REPORT

List of articles citing

Compound and digenic heterozygosity predicts lifetime arrhythmic outcome and sudden cardiac death in desmosomal gene-related arrhythmogenic right ventricular cardiomyopathy

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#	Paper	IF	Citations
195	Arrhythmogenic right ventricular cardiomyopathy: growing evidence for complex inheritance. <i>Circulation: Cardiovascular Genetics</i> , 2013 , 6, 525-7		10
194	Arrhythmogenic right ventricular dysplasia/cardiomyopathy type 1: a light on molecular mechanisms. 2013 , 2013, 460805		1
193	Arrhythmogenic ventricular cardiomyopathy: A paradigm shift from right to biventricular disease. 2014 , 6, 154-74		37
192	Causality in genetics: the gradient of genetic effects and back to Koch's postulates of causality. 2014 , 114, e18-21		22
191	The usual suspects in sudden cardiac death of the young: a focus on inherited arrhythmogenic diseases. 2014 , 12, 499-519		31
190	When rare illuminates common: how cardiocutaneous syndromes transformed our perspective on arrhythmogenic cardiomyopathy. 2014 , 21, 3-11		6
189	Intercalated discs and arrhythmogenic cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2014 , 7, 930-40		32
188	Genetic testing in cardiovascular diseases. 2014 , 29, 235-40		24
187	Arrhythmogene rechtsventrikuläre Kardiomyopathie. 2014 , 8, 85-98		2
186	Arrhythmogenic right ventricular cardiomyopathy (ARVC): cardiovascular magnetic resonance update. 2014 , 16, 50		90
185	Desmosomes in the heart: a review of clinical and mechanistic analyses. 2014 , 21, 109-28		34
184	Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy in the Pediatric Population: Clinical Characterization and Comparison With Adult-Onset Disease. <i>JACC: Clinical Electrophysiology</i> , 2015 , 1, 551-560	4.6	46
183	Sudden cardiac death in the young: the bogeyman. 2015 , 25, 408-23		5
182	Electrocardiographic features of disease progression in arrhythmogenic right ventricular cardiomyopathy/dysplasia. 2015 , 15, 4		20
181	Arrhythmogenic right ventricular cardiomyopathy and sports activity. <i>European Heart Journal</i> , 2015 , 36, 1708-10	9.5	27
180	The ARVD/C genetic variants database: 2014 update. 2015 , 36, 403-10		61
179	The research venture in arrhythmogenic right ventricular cardiomyopathy: a paradigm of translational medicine. <i>European Heart Journal</i> , 2015 , 36, 837-46	9.5	39

178	Impact of genotype on clinical course in arrhythmogenic right ventricular dysplasia/cardiomyopathy-associated mutation carriers. <i>European Heart Journal</i> , 2015 , 36, 847-55	9.5	238
177	Arrhythmogenic right ventricular dysplasia/cardiomyopathy: clinical challenges in a changing disease spectrum. <i>Trends in Cardiovascular Medicine</i> , 2015 , 25, 191-8	6.9	2
176	Arrhythmogenic cardiomyopathy: a disease of intercalated discs. 2015 , 360, 491-500		31
175	Comprehensive analysis of desmosomal gene mutations in Han Chinese patients with arrhythmogenic right ventricular cardiomyopathy. 2015 , 58, 258-65		10
174	Treatment of arrhythmogenic right ventricular cardiomyopathy/dysplasia: an international task force consensus statement. <i>European Heart Journal</i> , 2015 , 36, 3227-37	9.5	135
173	Desmoglein 2-Dependent Arrhythmogenic Cardiomyopathy Is Caused by a Loss of Adhesive Function. <i>Circulation: Cardiovascular Genetics</i> , 2015 , 8, 553-63		41
172	Arrhythmogenic cardiomyopathy in a patient with a rare loss-of-function KCNQ1 mutation. <i>Journal of the American Heart Association</i> , 2015 , 4, e001526	6	18
171	2015 ESC Guidelines for the management of patients with ventricular arrhythmias and the prevention of sudden cardiac death: The Task Force for the Management of Patients with Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death of the European Society of Cardiology (ESC) Endorsed by: Association for European Paediatric and Congenital Cardiology (AEPC). <i>Europace</i> , 2015 , 17, 1601-87	3.9	426
170	Homozygous Desmocollin-2 Mutations and Arrhythmogenic Cardiomyopathy. 2015 , 116, 1245-51		28
169	Oxidized low-density lipoprotein attenuated desmoglein 1 and desmocollin 2 expression via LOX-1/Ca(2+)/PKC-β signal in human umbilical vein endothelial cells. 2015 , 468, 380-6		12
168	Treatment of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia: An International Task Force Consensus Statement. 2015 , 132, 441-53		199
167	Determined to Fail--the Role of Genetic Mechanisms in Heart Failure. 2015 , 12, 333-8		7
166	2015 ESC Guidelines for the management of patients with ventricular arrhythmias and the prevention of sudden cardiac death: The Task Force for the Management of Patients with Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death of the European Society of Cardiology (ESC). Endorsed by: Association for European Paediatric and Congenital Cardiology	9.5	2187
165	Clinical interpretation of genetic variants in arrhythmogenic right ventricular cardiomyopathy. 2015 , 104, 288-303		9
164	Association of Phenotype and Genotype in the Diagnosis and Prognosis of ARVC/D in the Adult Population. 2016 , 89-103		
163	Arrhythmogenic Right Ventricular Dysplasia in Neuromuscular Disorders. 2016 , 10, 173-180		4
162	Arrhythmogenic right-ventricular cardiomyopathy: molecular genetics into clinical practice in the era of next generation sequencing. 2016 , 17, 399-407		14
161	Arrhythmogenic Right Ventricular Cardiomyopathy: Risk Stratification and Indications for Defibrillator Therapy. 2016 , 18, 57		20

160	Molecular disturbance underlies to arrhythmogenic cardiomyopathy induced by transgene content, age and exercise in a truncated PKP2 mouse model. 2016 , 25, 3676-3688		15
159	Electrocardiographic differentiation of idiopathic right ventricular outflow tract ectopy from early arrhythmogenic right ventricular cardiomyopathy. <i>Europace</i> , 2017 , 19, 622-628	3.9	16
158	The effects of transfection reagent polyethyleneimine (PEI) and non-targeting control siRNAs on global gene expression in human aortic smooth muscle cells. 2016 , 17, 20		12
157	Arrhythmogenic cardiomyopathy. 2016 , 11, 33		80
156	Potentially Lethal Ventricular Arrhythmias and Heart Failure in Arrhythmogenic Right Ventricular Cardiomyopathy: What Are the Differences Between Men and Women?. <i>JACC: Clinical Electrophysiology</i> , 2016 , 2, 546-555	4.6	23
155	Approach to family screening in arrhythmogenic right ventricular dysplasia/cardiomyopathy. <i>European Heart Journal</i> , 2016 , 37, 755-63	9.5	56
154	Arrhythmic risk assessment in genotyped families with arrhythmogenic right ventricular cardiomyopathy. <i>Europace</i> , 2016 , 18, 610-6	3.9	34
153	Phenotypic expression is a prerequisite for malignant arrhythmic events and sudden cardiac death in arrhythmogenic right ventricular cardiomyopathy. <i>Europace</i> , 2016 , 18, 1086-94	3.9	39
152	Multilevel analyses of SCN5A mutations in arrhythmogenic right ventricular dysplasia/cardiomyopathy suggest non-canonical mechanisms for disease pathogenesis. 2017 , 113, 102-111		111
151	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
150	Arrhythmogenic Right Ventricular Cardiomyopathy. 2017 , 376, 1489-90		44
149	Genetic and epigenetic regulation of arrhythmogenic cardiomyopathy. 2017 , 1863, 2064-2069		5
148	Implantable Cardioverter-Defibrillator Therapy in Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy: Predictors of Appropriate Therapy, Outcomes, and Complications. <i>Journal of the American Heart Association</i> , 2017 , 6,	6	47
147	Sex hormones affect outcome in arrhythmogenic right ventricular cardiomyopathy/dysplasia: from a stem cell derived cardiomyocyte-based model to clinical biomarkers of disease outcome. <i>European Heart Journal</i> , 2017 , 38, 1498-1508	9.5	76
146	Targeted Next-Generation Sequencing of 51 Genes Involved in Primary Electrical Disease. 2017 , 19, 445-459		9
145	Molecular mechanisms in the pathogenesis of arrhythmogenic cardiomyopathy. <i>Cardiovascular Pathology</i> , 2017 , 28, 51-58	3.8	2
144	Arrhythmogenic Right Ventricular Cardiomyopathy. 2017 , 376, 61-72		302
143	Unexpected sudden death in pregnancy: arrhythmogenic right ventricular cardiomyopathy/dysplasia: a case report. 2017 , 2, 161-163		2

142	Large Genomic Rearrangements of Desmosomal Genes in Italian Arrhythmogenic Cardiomyopathy Patients. 2017 , 10,		24
141	Arrhythmogenic cardiomyopathy: pathology, genetics, and concepts in pathogenesis. 2017 , 113, 1521-1531		58
140	Unique genetic background and outcome of non-Caucasian Japanese probands with arrhythmogenic right ventricular dysplasia/cardiomyopathy. 2017 , 5, 639-651		10
139	Arrhythmogenic Cardiomyopathy. 2017 , 121, 784-802		167
138	Right ventricular arrhythmogenic cardiomyopathy: genetic and MR for modern clinical diagnosis. 2017 , 18 Suppl 1, e157-e160		
137	Ventricular Arrhythmias and Sudden Cardiac Death. 2017 , 9, 693-708		11
136	Sudden Cardiac Death in Genetic Cardiomyopathies. 2017 , 9, 581-603		10
135	Sudden arrhythmic death and cardiomyopathies in the young: a molecular and pathology overview. 2017 , 23, 486-498		
134	A Novel Mutation in Junctional Plakoglobin Causing Lethal Congenital Epidermolysis Bullosa. 2017 , 191, 266-269.e1		4
133	Risk Stratification in Arrhythmogenic Right Ventricular Cardiomyopathy. 2017 , 136, 2068-2082		100
132	Co-inheritance of mutations associated with arrhythmogenic cardiomyopathy and hypertrophic cardiomyopathy. 2017 , 25, 1165-1169		7
131	High proportion of genetic cases in patients with advanced cardiomyopathy including a novel homozygous Plakophilin 2-gene mutation. <i>PLoS ONE</i> , 2017 , 12, e0189489	3.7	20
130	Case reports of two pedigrees with recessive arrhythmogenic right ventricular cardiomyopathy associated with homozygous Thr335Ala variant in DSG2. 2017 , 18, 86		11
129	Implantable Cardioverter-Defibrillators in Inherited Arrhythmia Syndromes. 2017 , 566-578		
128	Cardiac Channelopathies: Recognition, Treatment, Management. 2018 , 29, 43-57		2
127	Filamin C: A New Arrhythmogenic Cardiomyopathy-Causing Gene?. <i>JACC: Clinical Electrophysiology</i> , 2018 , 4, 515-517	4.6	3
126	Diagnosis of Arrhythmogenic Right Ventricular Cardiomyopathy: Progress and Pitfalls. <i>Heart Lung and Circulation</i> , 2018 , 27, 1310-1317	1.8	10
125	Targeted next generation sequencing in a young population with suspected inherited malignant cardiac arrhythmias. 2018 , 26, 303-313		6

124	Arrhythmogenic right ventricular cardiomyopathy. <i>Journal of Arrhythmia</i> , 2018 , 34, 356-368	1.5	4
123	Diagnostic Criteria, Genetics, and Molecular Basis of Arrhythmogenic Cardiomyopathy. 2018 , 14, 201-213		21
122	Genetic basis of arrhythmogenic cardiomyopathy. 2018 , 33, 276-281		14
121	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 Rhythm Society. <i>Heart Rhythm</i> , 2018 , 15, e103-e108	6.7	264
120	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. <i>Heart Rhythm</i> , 2018 , 15, e72-e100	6.7	151
119	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 Rhythm Society. <i>Journal of the American College of Cardiology</i> , 2018 , 72, e147-e232	15.1	180
118	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 Rhythm Society. <i>Journal of the American College of Cardiology</i> , 2018 , 72, e147-e232		173
117	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 , 138, e272-e391		307
116	Catecholaminergic polymorphic ventricular tachycardia patients with multiple genetic variants in the PACES CPVT Registry. <i>PLoS ONE</i> , 2018 , 13, e0205925	3.7	22
115	Arrhythmogenic Cardiomyopathy. 2018 , 631-639		
114	Compound and heterozygous mutations of DSG2 identified by Whole Exome Sequencing in arrhythmogenic right ventricular cardiomyopathy/dysplasia with ventricular tachycardia. <i>Journal of Electrocardiology</i> , 2018 , 51, 837-843	1.4	3
113	Arrhythmogenic Right Ventricular Cardiomyopathy. 2018 , 291-296		
112	Clinical Diagnosis, Imaging, and Genetics of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia: JACC State-of-the-Art Review. <i>Journal of the American College of Cardiology</i> , 2018 , 72, 784-804	15.1	105
111	Role of right ventricular involvement in acute myocarditis, assessed by cardiac magnetic resonance. <i>International Journal of Cardiology</i> , 2018 , 271, 359-365	3.2	14
110	Multiparametric Approach to Arrhythmogenic Cardiomyopathy: Clinical, Instrumental, and Lifestyle Indications. 2018 , 3, 35		
109	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. <i>Journal of the American College of Cardiology</i> , 2018 , 72, e91-e220	15.1	411
108	Atrial involvement in arrhythmogenic right ventricular cardiomyopathy patients referred for ventricular arrhythmias ablation. <i>Journal of Cardiovascular Electrophysiology</i> , 2018 , 29, 1388-1395	2.7	3
107	Primary Myocardial Fibrosis as an Alternative Phenotype Pathway of Inherited Cardiac Structural Disorders. 2018 , 137, 2716-2726		22

106	A founder homozygous DSG2 variant in East Asia results in ARVC with full penetrance and heart failure phenotype. <i>International Journal of Cardiology</i> , 2019 , 274, 263-270	3.2	17
105	Update on cardiomyopathies and sudden cardiac death. 2019 , 4, 202-210		5
104	A common indel polymorphism of the Desmoglein-2 (DSG2) is associated with sudden cardiac death in Chinese populations. 2019 , 301, 382-387		8
103	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy. <i>Heart Rhythm</i> , 2019 , 16, e301-e372	6.7	247
102	Recessive variants in plakophilin-2 contributes to early-onset arrhythmogenic cardiomyopathy with severe heart failure. <i>Europace</i> , 2019 , 21, 970-977	3.9	2
101	Arrhythmogenic Right Ventricular Cardiomyopathy. 2019 , 410-419.e3		
100	Large next-generation sequencing gene panels in genetic heart disease: yield of pathogenic variants and variants of unknown significance. 2019 , 27, 304-309		8
99	Managing uncertainty in inherited cardiac pathologies-an international multidisciplinary survey. 2019 , 27, 1178-1185		3
98	Heart failure in arrhythmogenic cardiomyopathy: is phenotypic variability just a matter of genetics?. 2019 , 21, 801-802		1
97	Arrhythmogenic cardiomyopathies (ACs): diagnosis, risk stratification and management. <i>Heart</i> , 2019 , 105, 1117-1128	5.1	11
96	High risk of heart failure associated with desmoglein-2 mutations compared to plakophilin-2 mutations in arrhythmogenic right ventricular cardiomyopathy/dysplasia. 2019 , 21, 792-800		20
95	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. <i>International Journal of Cardiology</i> , 2019 , 279, 79-83	3.2	5
94	Predictors of Adverse Outcomes in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy: A Meta-Analysis of Observational Studies. <i>Cardiology in Review</i> , 2019 , 27, 189-197	3.2	2
93	Sudden death in mild hypertrophic cardiomyopathy with compound DSG2/DSC2/MYH6 mutations: Revisiting phenotype after genetic assessment in a master runner athlete. <i>Journal of Electrocardiology</i> , 2019 , 53, 95-99	1.4	7
92	The complex molecular genetics of arrhythmogenic cardiomyopathy. <i>International Journal of Cardiology</i> , 2019 , 284, 59-60	3.2	1
91	Arrhythmogenic Right Ventricular Cardiomyopathy: Progress Toward Personalized Management. <i>Annual Review of Medicine</i> , 2019 , 70, 1-18	17.4	9
90	Arrhythmogenic cardiomyopathy: what blood can reveal?. <i>Heart Rhythm</i> , 2019 , 16, 470-477	6.7	10
89	Diagnostic and therapeutic strategies for arrhythmogenic right ventricular dysplasia/cardiomyopathy patient. <i>Europace</i> , 2019 , 21, 9-21	3.9	19

88	The merits of the ICD for inherited heart rhythm disorders: A critical re-appraisal. <i>Trends in Cardiovascular Medicine</i> , 2020 , 30, 415-421	6.9	1
87	Postmortem diagnosis of left dominant arrhythmogenic cardiomyopathy: the importance of a multidisciplinary network for sudden death victims. "HIC mors gaudet succurere vitae". <i>Cardiovascular Pathology</i> , 2020 , 44, 107157	3.8	1
86	Exercise and Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Heart Lung and Circulation</i> , 2020 , 29, 547-555	1.8	11
85	Arrhythmogenic right ventricular cardiomyopathy. 2020 , 375-388		
84	Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia: Mechanisms and Management. <i>Research Reports in Clinical Cardiology</i> , 2020 , Volume 11, 19-29	0.1	1
83	The Hidden Fragility in the Heart of the Athletes: A Review of Genetic Biomarkers. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	7
82	Genotype-Phenotype Correlation: A Triple DNA Mutational Event in a Boy Entering Sport Conveys an Additional Pathogenicity Risk. <i>Genes</i> , 2020 , 11,	4.2	10
81	Arrhythmogenic Cardiomyopathy: Molecular Insights for Improved Therapeutic Design. <i>Journal of Cardiovascular Development and Disease</i> , 2020 , 7,	4.2	6
80	Plasma testosterone and arrhythmic events in male patients with arrhythmogenic right ventricular cardiomyopathy. <i>ESC Heart Failure</i> , 2020 , 7, 1547-1559	3.7	6
79	Arrhythmogenic right ventricular cardiomyopathy : Evolving from unique clinical features to a complex pathophysiological concept. <i>Herz</i> , 2020 , 45, 243-251	2.6	2
78	The role of genetics in cardiovascular disease: arrhythmogenic cardiomyopathy. <i>European Heart Journal</i> , 2020 , 41, 1393-1400	9.5	30
77	Natural History of Arrhythmogenic Cardiomyopathy. <i>Journal of Clinical Medicine</i> , 2020 , 9,	5.1	19
76	Arrhythmogenic Right Ventricular Cardiomyopathy: Characterization of Left Ventricular Phenotype and Differential Diagnosis With Dilated Cardiomyopathy. <i>Journal of the American Heart Association</i> , 2020 , 9, e014628	6	43
75	Advantages and Perils of Clinical Whole-Exome and Whole-Genome Sequencing in Cardiomyopathy. <i>Cardiovascular Drugs and Therapy</i> , 2020 , 34, 241-253	3.9	13
74	Binding of the periplakin linker requires vimentin acidic residues D176 and E187. <i>Communications Biology</i> , 2020 , 3, 83	6.7	5
73	The evolution of gene-guided management of inherited arrhythmia syndromes: Peering beyond monogenic paradigms towards comprehensive genomic risk scores. <i>Journal of Cardiovascular Electrophysiology</i> , 2020 , 31, 2998-3008	2.7	1
72	Clinical predictors of left ventricular involvement in arrhythmogenic right ventricular cardiomyopathy. <i>American Heart Journal</i> , 2020 , 223, 34-43	4.9	6
71	Emerging concepts in arrhythmogenic dilated cardiomyopathy. <i>Heart Failure Reviews</i> , 2021 , 26, 1219-1239		8

70	Meta-analysis of cardiomyopathy-associated variants in troponin genes identifies loci and intragenic hot spots that are associated with worse clinical outcomes. <i>Journal of Molecular and Cellular Cardiology</i> , 2020 , 142, 118-125	5.8	14
69	2020 ESC Guidelines on sports cardiology and exercise in patients with cardiovascular disease. <i>European Heart Journal</i> , 2021 , 42, 17-96	9.5	264
68	Sex Differences in Right Ventricular Dysfunction: Insights From the Bench to Bedside. <i>Frontiers in Physiology</i> , 2020 , 11, 623129	4.6	4
67	Right Ventricular Cardiomyopathies. 2021 , 267-288		
66	Genetics of Cardiomyopathy: Clinical and Mechanistic Implications for Heart Failure. <i>Korean Circulation Journal</i> , 2021 , 51, 797-836	2.2	3
65	Risk Stratification in Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Arrhythmia and Electrophysiology Review</i> , 2021 , 10, 26-32	3.2	1
64	Update on the Diagnostic Pitfalls of Autopsy and Post-Mortem Genetic Testing in Cardiomyopathies. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	5
63	Genetics of Cardiomyopathy.		
62	Arrhythmogenic Left Ventricular Cardiomyopathy: Genotype-Phenotype Correlations and New Diagnostic Criteria. <i>Journal of Clinical Medicine</i> , 2021 , 10,	5.1	5
61	Contemporary and Future Approaches to 'Precision Medicine in Inherited' Cardiomyopathies: JACC Focus Seminar 3/5. <i>Journal of the American College of Cardiology</i> , 2021 , 77, 2551-2572	15.1	4
60	Pregnancy in arrhythmogenic cardiomyopathy. <i>Herzschrittmachertherapie Und Elektrophysiologie</i> , 2021 , 32, 186-198	0.8	0
59	Guía ESC 2020 sobre cardiología del deporte y el ejercicio en pacientes con enfermedad cardiovascular. <i>Revista Espanola De Cardiologia</i> , 2021 , 74, 545.e1-545.e73	1.5	2
58	Arrhythmogenic Cardiomyopathy-Current Treatment and Future Options. <i>Journal of Clinical Medicine</i> , 2021 , 10,	5.1	1
57	Pathogenic variants in plakophilin-2 gene (PKP2) are associated with better survival in arrhythmogenic right ventricular cardiomyopathy. <i>Journal of Applied Genetics</i> , 2021 , 62, 613-620	2.5	2
56	Arrhythmic risk stratification in arrhythmogenic cardiomyopathy: new predictors for left-sided variants?. <i>European Heart Journal</i> , 2021 , 42, 2851-2853	9.5	4
55	Arrhythmogenic left ventricular cardiomyopathy. <i>Heart</i> , 2021 ,	5.1	7
54	Genotype-phenotype correlation in arrhythmogenic right ventricular cardiomyopathy-risk of arrhythmias and heart failure. <i>Journal of Medical Genetics</i> , 2021 ,	5.8	1
53	Interpretation of Incidental Genetic Findings Localizing to Genes Associated With Cardiac Channelopathies and Cardiomyopathies. <i>Circulation Genomic and Precision Medicine</i> , 2021 , 14, e003200	5.2	2

52	Filamin-C variant-associated cardiomyopathy: A pooled analysis of individual patient data to evaluate the clinical profile and risk of sudden cardiac death. <i>Heart Rhythm</i> , 2021 ,	6.7	4
51	Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. <i>European Heart Journal</i> , 2020 , 41, 1414-1429	9.5	110
50	Contribution of exome sequencing for genetic diagnostic in arrhythmogenic right ventricular cardiomyopathy/dysplasia. <i>PLoS ONE</i> , 2017 , 12, e0181840	3.7	11
49	A novel mutation and intrafamilial phenotypic variability in ARVC/D. <i>Medical Journal of the Islamic Republic of Iran</i> , 2018 , 32, 5	1.1	4
48	Nature and Nurture in Arrhythmogenic Right Ventricular Cardiomyopathy - A Clinical Perspective. <i>Arrhythmia and Electrophysiology Review</i> , 2015 , 4, 156-62	3.2	2
47	Arrhythmogenic Cardiomyopathy: Electrical and Structural Phenotypes. <i>Arrhythmia and Electrophysiology Review</i> , 2016 , 5, 90-101	3.2	41
46	MicroRNA-130a Regulation of Desmocollin 2 in a Novel Model of Arrhythmogenic Cardiomyopathy. <i>MicroRNA (Sharjah, United Arab Emirates)</i> , 2017 , 6, 143-150	2.9	9
45	Practical Aspects in Genetic Testing for Cardiomyopathies and Channelopathies. <i>Clinical Biochemist Reviews</i> , 2019 , 40, 187-200	7.3	4
44	Circulation and Contacts in Sixteenth Century New Cartography: Spain, Portugal and Italy. <i>Culture & History Digital Journal</i> , 2021 , 10, e015	0.2	
43	Arrhythmogenic Right Ventricular Cardiomyopathy: Usefulness of Imaging in Prognostic Stratification and Choice of Treatment. 2014 , 173-181		
42	Genetics: Genotype/Phenotype Correlations in Cardiomyopathies. 2014 , 13-24		
41	Phenotypic Expression and Genetics of J Wave Syndrome in the Early Stage of Arrhythmogenic Right Ventricular Cardiomyopathy. 2016 , 259-280		
40	Arrhythmogenic right ventricular dysplasia / THE DARK VOYAGE OF THE FEARLESS EAGLE /Case presentation and review of the literature. <i>Journal of Heart and Cardiology</i> , 2016 , 2, 1-6		
39	Diagnostic Evaluation of Children with Known or Suspected ARVC/D. 2016 , 105-113		
38	Arrhythmogenic Cardiomyopathy. 2016 , 91-111		
37	Inherited Cardiac Muscle Disorders: Arrhythmogenic Right Ventricular Cardiomyopathy. 2018 , 367-388		
36	PKP2 and DSG2 genetic variations in Latvian arrhythmogenic right ventricular dysplasia/cardiomyopathy registry patients. <i>Anatolian Journal of Cardiology</i> , 2018 , 20, 296-302	0.8	1
35	Genetic Testing for Inheritable Cardiac Channelopathies. <i>Cardiac and Vascular Biology</i> , 2018 , 323-358	0.2	

34	[Molecular genetic basis of sudden cardiac death in the young with cardiomyopathy of various origins]. <i>Sudebno-Meditsinskaya Ekspertisa</i> , 2019 , 62, 48-53		0
33	Risk stratification in families with history of idiopathic ventricular fibrillation. <i>HeartRhythm Case Reports</i> , 2020 , 6, 386-389	1	
32	Approach to inherited arrhythmias in pregnancy. <i>International Journal of Cardiology Congenital Heart Disease</i> , 2021 , 100264	0.7	
31	Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Contemporary Cardiology</i> , 2020 , 791-810	0.1	
30	Arrhythmogenic Cardiomyopathy. 2020 , 99-114		
29	Genetic and Molecular Basis of Cardiac Arrhythmias. <i>Contemporary Cardiology</i> , 2020 , 75-96	0.1	
28	Clinical and genetic features of arrhythmogenic cardiomyopathy: diagnosis, management and the heart failure perspective.. <i>Progress in Pediatric Cardiology</i> , 2021 , 63, 101459-101459	0.4	0
27	Arrhythmogenic Right Ventricular Cardiomyopathy - 4 Swedish families with an associated PKP2 c.2146-1G>C variant. <i>American Journal of Cardiovascular Disease</i> , 2016 , 6, 55-65	0.9	5
26	Risk of sports-related sudden cardiac death in women. <i>European Heart Journal</i> , 2021 ,	9.5	1
25	Arrhythmogenic Right Ventricular Cardiomyopathy in Pediatric Patients: An Important but Underrecognized Clinical Entity.. <i>Frontiers in Pediatrics</i> , 2021 , 9, 750916	3.4	2
24	Cardiovascular Characteristics of Patients with Genetic Variation in Desmoplakin ().. <i>Neurology International</i> , 2022 , 12, 24-36	0	2
23	Modeling reduced contractility and impaired desmosome assembly due to plakophilin-2 deficiency using isogenic iPS cell-derived cardiomyocytes.. <i>Stem Cell Reports</i> , 2022 ,	8	1
22	Genomic and Non-Genomic Regulatory Mechanisms of the Cardiac Sodium Channel in Cardiac Arrhythmias.. <i>International Journal of Molecular Sciences</i> , 2022 , 23,	6.3	3
21	Arrhythmogenic cardiomyopathy in children according to "Padua criteria": Single pediatric center experience.. <i>International Journal of Cardiology</i> , 2022 ,	3.2	3
20	Recent Non-Invasive Parameters to Identify Subjects at High Risk of Sudden Cardiac Death.. <i>Journal of Clinical Medicine</i> , 2022 , 11,	5.1	0
19	Diagnosis and management of rare cardiomyopathies in adult and paediatric patients.. <i>International Journal of Cardiology</i> , 2022 ,	3.2	2
18	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.. <i>Europace</i> , 2022 ,	3.9	5
17	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.. <i>Heart Rhythm</i> , 2022 ,	6.7	6

16	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. <i>Journal of Arrhythmia</i> ,	1.5	2
15	Ventricular arrhythmia management in patients with genetic cardiomyopathies.. <i>Heart Rhythm O2</i> , 2021 , 2, 819-831	1.5	
14	Histopathological Features and Protein Markers of Arrhythmogenic Cardiomyopathy.. <i>Frontiers in Cardiovascular Medicine</i> , 2021 , 8, 746321	5.4	1
13	Arrhythmogenic Right Ventricular Cardiomyopathy.. <i>JACC: Clinical Electrophysiology</i> , 2022 , 8, 533-553	4.6	1
12	OUP accepted manuscript. <i>European Heart Journal</i> ,	9.5	1
11	Molecular genetic testing in athletes: Why and when a position statement from the Italian society of sports cardiology. <i>International Journal of Cardiology</i> , 2022 ,	3.2	0
10	Implantable defibrillators in primary prevention of genetic arrhythmias. A shocking choice?. <i>European Heart Journal</i> ,	9.5	0
9	The value of genetic testing in the diagnosis and risk stratification of arrhythmogenic right ventricular cardiomyopathy. <i>Heart Rhythm</i> , 2022 ,	6.7	
8	Sex Differences in Cardiomyopathy. <i>Current Cardiovascular Risk Reports</i> ,	0.9	
7	2022 ESC Guidelines for the management of patients with ventricular arrhythmias and the prevention of sudden cardiac death.		37
6	Genetics and Genomics of Congenital and Acquired Cardiovascular Disease. 2021 , 1-41		0
5	Varied Presentation of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy (ARVD/C): A Case Series. 2023 ,		0
4	DSP-Related Cardiomyopathy as a Distinct Clinical Entity? Emerging Evidence from an Italian Cohort. 2023 , 24, 2490		1
3	Review: Sex-related differences in the treatment of cardiac arrhythmia. 2023 , 244, 108388		0
2	Disease modeling of desmosome-related cardiomyopathy using induced pluripotent stem cell-derived cardiomyocytes. 15, 71-82		0
1	Cheek-Pro-Heart: What Can the Buccal Mucosa Do for Arrhythmogenic Cardiomyopathy?. 2023 , 11, 1207		0