Elijah R Behr

List of Publications by Year
in descending order

241	12,664	25034	29157
papers	citations	57 h-index	104 g-index

248	248	248	10402
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	ESC guidance for the diagnosis and management of cardiovascular disease during the COVID-19 pandemic: part 2â€"care pathways, treatment, and follow-up. European Heart Journal, 2022, 43, 1059-1103.	2.2	111
2	Investigation on Sudden Unexpected Death in the Young (SUDY) in Europe: results of the European Heart Rhythm Association Survey. Europace, 2022, 24, 331-339.	1.7	23
3	Role of subcutaneous implantable loop recorder for the diagnosis of arrhythmias in Brugada syndrome: A United Kingdom single-center experience. Heart Rhythm, 2022, 19, 70-78.	0.7	20
4	The prevalence of left and right bundle branch block morphology ventricular tachycardia amongst patients with arrhythmogenic cardiomyopathy and sustained ventricular tachycardia: insights from the European Survey on Arrhythmogenic Cardiomyopathy. Europace, 2022, 24, 285-295.	1.7	7
5	Electrocardiogram screening programme in detecting sudden cardiac disease in the young: cost efficiency and diagnostic yieldâ€"Authorsâ€™ reply. Europace, 2022, 24, 524-525.	1.7	0
6	European Society of Cardiology guidance for the diagnosis and management of cardiovascular disease during the COVID-19 pandemic: part 1â€"epidemiology, pathophysiology, and diagnosis. European Heart Journal, 2022, 43, 1033-1058.	2.2	80
7	Primary systemic sclerosis heart involvement: A systematic literature review and preliminary data-driven, consensus-based WSF/HFA definition. Journal of Scleroderma and Related Disorders, 2022, 7, 24-32.	1.7	25
8	Analysis of buccal mucosa as a prognostic tool in children with arrhythmogenic cardiomyopathy. Progress in Pediatric Cardiology, 2022, 64, 101458.	0.4	3
9	Mechanism of the effects of sodium channel blockade on the arrhythmogenic substrate of Brugada syndrome. Heart Rhythm, 2022, 19, 407-416.	0.7	17
10	Efficacy and Safety of Appropriate Shocks and Antitachycardia Pacing in Transvenous and Subcutaneous Implantable Defibrillators: Analysis of All Appropriate Therapy in the PRAETORIAN Trial. Circulation, 2022, 145, 321-329.	1.6	28
11	Association of Sexual Intercourse With Sudden Cardiac Death in Young Individuals in the United Kingdom. JAMA Cardiology, 2022, 7, 358.	6.1	6
12	Interpretation and actionability of genetic variants in cardiomyopathies: a position statement from the European Society of Cardiology Council on cardiovascular genomics. European Heart Journal, 2022, 43, 1901-1916.	2.2	32
13	Genetics of sudden cardiac death. Current Opinion in Cardiology, 2022, 37, 212-218.	1.8	4
14	An International Multicenter Cohort Study on Î²-Blockers for the Treatment of Symptomatic Children With Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation, 2022, 145, 333-344.	1.6	28
15	Rare Variation in Drug Metabolism and Long QT Genes and the Genetic Susceptibility to Acquired Long QT Syndrome. Circulation Genomic and Precision Medicine, 2022, 15, CIRCGEN121003391.	3.6	7
16	European Society of Cardiology guidance for the diagnosis and management of cardiovascular disease during the COVID-19 pandemic: part 1â€"epidemiology, pathophysiology, and diagnosis. Cardiovascular Research, 2022, 118, 1385-1412.	3.8	27
17	Exome Sequencing Highlights a Potential Role for Concealed Cardiomyopathies in Youthful Sudden Cardiac Death. Circulation Genomic and Precision Medicine, 2022, 15, CIRCGEN121003497.	3.6	15
18	Genome-wide association analyses identify new Brugada syndrome risk loci and highlight a new mechanism of sodium channel regulation in disease susceptibility. Nature Genetics, 2022, 54, 232-239.	21.4	55

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19	Brugada Syndrome. JACC: Clinical Electrophysiology, 2022, 8, 386-405.	3.2	26
20	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. Europace, 2022, 24, 1307-1367.	1.7	108
21	Explaining the unexplained: applying genetic testing after cardiac arrest and sudden death. European Heart Journal, 2022, 43, 3082-3084.	2.2	3
22	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. Heart Rhythm, 2022, 19, e1-e60.	0.7	78
23	European Heart Rhythm Association (<scp>EHRA</scp>)/Heart Rhythm Society (<scp>HRS</scp>)/Asia Pacific Heart Rhythm Society (<scp>APHRS</scp>)/Latin American Heart Rhythm Society (<scp>LAHRS</scp>) Expert Consensus Statement on the state of genetic testing for cardiac diseases. Journal of Arrhythmia, 2022, 38, 491-553.	1.2	24
24	ESC guidance for the diagnosis and management of cardiovascular disease during the COVID-19 pandemic: part 2â€"care pathways, treatment, and follow-up. Cardiovascular Research, 2022, 118, 1618-1666.	3.8	32
25	Mitral valve abnormalities in decedents of sudden cardiac death due to hypertrophic cardiomyopathy and idiopathic left ventricular hypertrophy. Heart Rhythm, 2022, 19, 1684-1685.	0.7	2
26	Contact force sensing in ablation of ventricular arrhythmias using a 56-hole open-irrigation catheter: a propensity-matched analysis. Journal of Interventional Cardiac Electrophysiology, 2021, 60, 543-553.	1.3	6
27	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
28	Non-invasive detection of exercise-induced cardiac conduction abnormalities in sudden cardiac death survivors in the inherited cardiac conditions. Europace, 2021, 23, 305-312.	1.7	8
29	Brugada syndrome and arrhythmogenic cardiomyopathy: overlapping disorders of the connexome?. Europace, 2021, 23, 653-664.	1.7	16
30	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
31	Enhancing rare variant interpretation in inherited arrhythmias through quantitative analysis of consortium disease cohorts and population controls. Genetics in Medicine, 2021, 23, 47-58.	2.4	57
32	Brugada syndrome and reduced right ventricular outflow tract conduction reserve: a final common pathway?. European Heart Journal, 2021, 42, 1073-1081.	2.2	56
33	Cardiac channelopathies: diagnosis and contemporary management. Heart, 2021, 107, 1092-1099.	2.9	4
34	Sex-Related Differences in Cardiac Channelopathies. Circulation, 2021, 143, 739-752.	1.6	23
35	Diagnostic yield and financial implications of a nationwide electrocardiographic screening programme to detect cardiac disease in the young. Europace, 2021, 23, 1295-1301.	1.7	15
36	Familial Evaluation in Idiopathic Ventricular Fibrillation. Circulation: Arrhythmia and Electrophysiology, 2021, 14, e009089.	4.8	15

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37	Cadherin 2-Related Arrhythmogenic Cardiomyopathy. Circulation Genomic and Precision Medicine, 2021, 14, e003097.	3.6	21
38	Importance of Dedicated Units for the Management of Patients With Inherited Arrhythmia Syndromes. Circulation Genomic and Precision Medicine, 2021, 14, e003313.	3.6	7
39	Genetics and genomics of arrhythmic risk: current and future strategies to prevent sudden cardiac death. Nature Reviews Cardiology, 2021, 18, 774-784.	13.7	15
40	Management of Congenital Long-QT Syndrome: Commentary From the Experts. Circulation: Arrhythmia and Electrophysiology, 2021, 14, e009726.	4.8	5
41	Genotype-Phenotype Correlation of SCN5A Genotype in Patients With Brugada Syndrome and Arrhythmic Events: Insights From the SABRUS in 392 Probands. Circulation Genomic and Precision Medicine, 2021, 14, e003222.	3.6	7
42	Application of artificial intelligence to the electrocardiogram. European Heart Journal, 2021, 42, 4717-4730.	2.2	96
43	The genomic architecture of the Brugada syndrome. Heart Rhythm, 2021, 18, 1707-1708.	0.7	3
44	Biventricular Myocardial Fibrosis and Sudden Death in Patients With BrugadaÂSyndrome. Journal of the American College of Cardiology, 2021, 78, 1511-1521.	2.8	18
45	J-Wave Syndromes, SCN5A, and CardiacÂConduction Reserve. Journal of the American College of Cardiology, 2021, 78, 1618-1620.	2.8	4
46	The European Cardiac Arrhythmia Genetics (ECGen) Focus Group. European Heart Journal, 2021, , .	2.2	0
47	Confirmation of Cause of Death Via Comprehensive Autopsy and Whole Exome Molecular Sequencing in People With Epilepsy and Sudden Unexpected Death. Journal of the American Heart Association, 2021, 10, e021170.	3.7	9
48	Hourly variability in outflow tract ectopy as a predictor of its site of origin. Journal of Cardiovascular Electrophysiology, 2021, , .	1.7	2
49	Sudden Death in Female Athletes: Insights From a Large Regional Registry in the United Kingdom. Circulation, 2021, 144, 1827-1829.	1.6	6
50	Cardiac arrest as first presentation of arrhythmogenic left ventricular cardiomyopathy due to Filamin C mutation: a case report. European Heart Journal - Case Reports, 2021, 5, ytab422.	0.6	2
51	The yield of postmortem genetic testing in sudden death cases with structural findings at autopsy. European Journal of Human Genetics, 2020, 28, 17-22.	2.8	38
52	Accuracy of the 2017 international recommendations for clinicians who interpret adolescent athletesâ€™ ECGs: a cohort study of 11 168 British white and black soccer players. British Journal of Sports Medicine, 2020, 54, 739-745.	6.7	41
53	Channelopathies in clinical medicineâ€”cardiac arrhythmias. , 2020, , 133-152.		0
54	Prevalence and Phenotypic Correlations of Calmodulinopathy-Causative CALM1-3 Variants Detected in a Multicenter Molecular Autopsy Cohort of Sudden Unexplained Death Victims. Circulation Genomic and Precision Medicine, 2020, 13, e003032.	3.6	3

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55	Response to eLetter: Fascinating helpful article, but how typical were the patients with DCM and what does this tell us?. Heart, 2020, 106, 1532.2-1533.	2.9	0
56	Subcutaneous or Transvenous Defibrillator Therapy. New England Journal of Medicine, 2020, 383, 526-536.	27.0	278
57	J-Wave Syndromes. JACC: Clinical Electrophysiology, 2020, 6, 1862-1863.	3.2	1
58	SCN5A Mutation Type and a Genetic Risk Score Associate Variably With Brugada Syndrome Phenotype in SCN5A Families. Circulation Genomic and Precision Medicine, 2020, 13, e002911.	3.6	41
59	Morphometric characterization of collagen and fat in normal ventricular myocardium. Cardiovascular Pathology, 2020, 48, 107224.	1.6	17
60	Transethnic Genome-Wide Association Study Provides Insights in the Genetic Architecture and Heritability of Long QT Syndrome. Circulation, 2020, 142, 324-338.	1.6	83
61	An International Multicenter Evaluation of Type 5 Long QT Syndrome. Circulation, 2020, 141, 429-439.	1.6	39
62	Continued misuse of orphan drug legislation: a life-threatening risk for mexiletine. European Heart Journal, 2020, 41, 614-617.	2.2	15
63	Differentiation between athleteâ€™s heart and dilated cardiomyopathy in athletic individuals. Heart, 2020, 106, 1059-1065.	2.9	47
64	SARS-CoV-2, COVID-19, and inherited arrhythmia syndromes. Heart Rhythm, 2020, 17, 1456-1462.	0.7	162
65	Triadin Knockout Syndrome Is Absent in a Multi-Center Molecular Autopsy Cohort of Sudden Infant Death Syndrome and Sudden Unexplained Death in the Young and Is Extremely Rare in the General Population. Circulation Genomic and Precision Medicine, 2020, 13, e002731.	3.6	4
66	Diagnostic yield of hypertrophic cardiomyopathy in first-degree relatives of decedents with idiopathic left ventricular hypertrophy. Europace, 2020, 22, 632-642.	1.7	20
67	Diagnosis, family screening, and treatment of inherited arrhythmogenic diseases in Europe: results of the European Heart Rhythm Association Survey. Europace, 2020, 22, 1904-1910.	1.7	11
68	Brugada Syndrome. , 2020, , 25-39.		0
69	Long QT Syndrome. , 2020, , 193-217.		1
70	Noncardiac genetic predisposition in sudden infant death syndrome. Genetics in Medicine, 2019, 21, 641-649.	2.4	9
71	Type 8 long QT syndrome: pathogenic variants in CACNA1C-encoded Cav1.2 cluster in STAC protein binding site. Europace, 2019, 21, 1725-1732.	1.7	15
72	Evaluation After Sudden Death in the Young. Circulation: Arrhythmia and Electrophysiology, 2019, 12, e007453.	4.8	19

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73	Next-generation sequencing using microfluidic PCR enrichment for molecular autopsy. BMC Cardiovascular Disorders, 2019, 19, 174.	1.7	7
74	Genetic susceptibility and the Brugada syndrome. European Heart Journal, 2019, 40, 3094-3096.	2.2	3
75	Ethnic differences in patients with Brugada syndrome and arrhythmic events: New insights from Survey on Arrhythmic Events in Brugada Syndrome. Heart Rhythm, 2019, 16, 1468-1474.	0.7	22
76	Mapping and Ablation of Ventricular Fibrillation Associated With Early Repolarization Syndrome. Circulation, 2019, 140, 1477-1490.	1.6	80
77	Anomalous Coronary Artery Origin and Sudden Cardiac Death. JACC: Clinical Electrophysiology, 2019, 5, 516-522.	3.2	58
78	Sudden Death and Left Ventricular Involvement in Arrhythmogenic Cardiomyopathy. Circulation, 2019, 139, 1786-1797.	1.6	122
79	Calmodulin mutations and life-threatening cardiac arrhythmias: insights from the International Calmodulinopathy Registry. European Heart Journal, 2019, 40, 2964-2975.	2.2	116
80	Implantable cardioverter-defibrillators in previously undiagnosed patients with catecholaminergic polymorphic ventricular tachycardia resuscitated from sudden cardiac arrest. European Heart Journal, 2019, 40, 2953-2961.	2.2	96
81	Genotypeâ€“phenotype association in patients with SCN4A mutation â€“ Authors' reply. Lancet, The, 2019, 393, 2301-2302.	13.7	0
82	National registry for sudden unexpected deaths of infants and children in England: why do we need one and do families want one?. Archives of Disease in Childhood, 2019, 104, 989-993.	1.9	3
83	Inherited cardiomyopathies. BMJ: British Medical Journal, 2019, 365, l1570.	2.3	12
84	Risk score for the exclusion of arrhythmic events in arrhythmogenic right ventricular cardiomyopathy at first presentation. International Journal of Cardiology, 2019, 290, 100-105.	1.7	8
85	Myocardial Inflammation in Brugada Syndrome. Journal of the American College of Cardiology, 2019, 73, 1369-1370.	2.8	0
86	Characterization and Management of Arrhythmic Events in Young Patients With Brugada Syndrome. Journal of the American College of Cardiology, 2019, 73, 1756-1765.	2.8	53
87	Comparison of Ajmaline and Procainamide Provocation Tests in the Diagnosis of Brugada Syndrome. JACC: Clinical Electrophysiology, 2019, 5, 504-512.	3.2	32
88	Sudden Death Can Be the First Manifestation of Hypertrophic Cardiomyopathy. JACC: Clinical Electrophysiology, 2019, 5, 252-254.	3.2	20
89	Response by Sheikh et al to Letter Regarding Article, â€œDiagnostic Yield of Genetic Testing in Young Athletes With T-Wave Inversionâ€. Circulation, 2019, 139, 996-997.	1.6	3
90	Electrocardiographic differentiation between â€˜benign T-wave inversionâ€™ and arrhythmogenic right ventricular cardiomyopathy. Europace, 2019, 21, 332-338.	1.7	36

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91	Time-to-first appropriate shock in patients implanted prophylactically with an implantable cardioverter-defibrillator: data from the Survey on Arrhythmic Events in BRUgada Syndrome (SABRUS). Europace, 2019, 21, 796-802.	1.7	16
92	The narrow-sense and common single nucleotide polymorphism heritability of early repolarization. International Journal of Cardiology, 2019, 279, 135-140.	1.7	7
93	KCND3 potassium channel gene variant confers susceptibility to electrocardiographic early repolarization pattern. JCI Insight, 2019, 4, .	5.0	15
94	The Role of Medical Therapy in Idiopathic Ventricular Fibrillation. European Journal of Arrhythmia & Electrophysiology, 2019, 5, 87.	0.2	1
95	Dysfunction of NaV1.4, a skeletal muscle voltage-gated sodium channel, in sudden infant death syndrome: a case-control study. Lancet, The, 2018, 391, 1483-1492.	13.7	63
96	Fever-related arrhythmic events in the multicenter Survey on Arrhythmic Events in Brugada Syndrome. Heart Rhythm, 2018, 15, 1394-1401.	0.7	71
97	Next-generation sequencing of AV nodal reentrant tachycardia patients identifies broad spectrum of variants in ion channel genes. European Journal of Human Genetics, 2018, 26, 660-668.	2.8	12
98	Profile of patients with Brugada syndrome presenting with their first documented arrhythmic event: Data from the Survey on Arrhythmic Events in BRUgada Syndrome (SABRUS). Heart Rhythm, 2018, 15, 716-724.	0.7	57
99	Obesity and sudden cardiac death in the young: Clinical and pathological insights from a large national registry. European Journal of Preventive Cardiology, 2018, 25, 395-401.	1.8	58
100	Proof of concept study of a novel pacemapping algorithm as a basis to guide ablation of ventricular arrhythmias. Europace, 2018, 20, 1647-1656.	1.7	5
101	Cardiac Genetic Predisposition in SuddenÂInfant Death Syndrome. Journal of the American College of Cardiology, 2018, 71, 1217-1227.	2.8	66
102	The Diagnostic Yield of Brugada Syndrome After Sudden Death WithÂNormal Autopsy. Journal of the American College of Cardiology, 2018, 71, 1204-1214.	2.8	84
103	24â€…Anomalous coronary artery origin and sudden cardiac death. data from a large regional registry. , 2018, , .		0
104	Exome-Wide Rare Variant Analyses in Sudden Infant Death Syndrome. Journal of Pediatrics, 2018, 203, 423-428.e11.	1.8	17
105	Diagnostic Yield of Genetic Testing in Young Athletes With T-Wave Inversion. Circulation, 2018, 138, 1184-1194.	1.6	43
106	Lack of genotype-phenotype correlation in Brugada Syndrome and Sudden Arrhythmic Death Syndrome families with reported pathogenic SCN1B variants. Heart Rhythm, 2018, 15, 1051-1057.	0.7	15
107	SCN5A mutations in 442 neonates and children: genotypeâ€“phenotype correlation and identification of higher-risk subgroups. European Heart Journal, 2018, 39, 2879-2887.	2.2	33
108	Gender differences in patients with Brugada syndrome and arrhythmic events: Data from a survey on arrhythmic events in 678 patients. Heart Rhythm, 2018, 15, 1457-1465.	0.7	65

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109	Importance of Variant Interpretation in Whole-Exome Molecular Autopsy. Circulation, 2018, 137, 2705-2715.	1.6	36
110	Genetics and Genomics of Sudden Unexplained Cardiac Death. , 2018, , 755-779.		0
111	25â€...Sudden cardiac death in elderly patients with hypertrophic cardiomyopathy. data from a large pathology registry. , 2018, , .		0
112	Common Genetic Variant Risk Score Is Associated With Drug-Induced QT Prolongation and Torsade de Pointes Risk. Circulation, 2017, 135, 1300-1310.	1.6	96
113	Fever vs drug: Battling with the Brugada syndrome substrate. Heart Rhythm, 2017, 14, 518-519.	0.7	2
114	Genetic testing in idiopathic ventricular fibrillation: Searching for a needle in a haystack?. Heart Rhythm, 2017, 14, 1041-1042.	0.7	0
115	Sudden Cardiac Death. JACC: Clinical Electrophysiology, 2017, 3, 473-481.	3.2	13
116	Utility of Post-Mortem Genetic Testing in Cases of Sudden Arrhythmic Death Syndrome. Journal of the American College of Cardiology, 2017, 69, 2134-2145.	2.8	219
117	Late gadolinium enhancement in Brugada syndrome: A marker for subtle underlying cardiomyopathy?. Heart Rhythm, 2017, 14, 583-589.	0.7	38
118	Sudden Cardiac Death inÂPre-Excitation and Wolff-Parkinson-White. Journal of the American College of Cardiology, 2017, 69, 1644-1645.	2.8	17
119	Investigation of the family of sudden cardiac death victims. Progress in Pediatric Cardiology, 2017, 45, 25-29.	0.4	1
120	Relationship Between Distance and Change in Surface ECG Morphology During Pacemapping as a Guide to Ablation of Ventricular Arrhythmias. Circulation: Arrhythmia and Electrophysiology, 2017, 10, .	4.8	12
121	Anterior T-Wave Inversion in Young WhiteÂAthletes and Nonathletes. Journal of the American College of Cardiology, 2017, 69, 1-9.	2.8	91
122	Drugs and life-threatening ventricular arrhythmia risk: results from the DARE study cohort. BMJ Open, 2017, 7, e016627.	1.9	30
123	Sudden infant death syndrome and inherited cardiac conditions. Nature Reviews Cardiology, 2017, 14, 715-726.	13.7	36
124	121â€...Left ventricular morphology in elite athletes with extreme anthropometry. Heart, 2017, 103, A91.1-A91.	2.9	0
125	Loss-of-Function KCNE2 Variants. Circulation: Arrhythmia and Electrophysiology, 2017, 10, .	4.8	31
126	Surviving Sudden Death. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	0

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127	Reply. Journal of the American College of Cardiology, 2017, 70, 297-298.	2.8	0
128	Sudden death and cardiac arrest without phenotype: the utility of genetic testing. Trends in Cardiovascular Medicine, 2017, 27, 207-213.	4.9	14
129	Age of First Arrhythmic Event in Brugada Syndrome. Circulation: Arrhythmia and Electrophysiology, 2017, 10, .	4.8	57
130	Mutations in HYAL2, Encoding Hyaluronidase 2, Cause a Syndrome of Orofacial Clefting and Cor Triatriatum Sinister in Humans and Mice. PLoS Genetics, 2017, 13, e1006470.	3.5	20
131	Next-Generation Sequencing in Post-mortem Genetic Testing of Young Sudden Cardiac Death Cases. Frontiers in Cardiovascular Medicine, 2016, 3, 13.	2.4	20
132	Comparison of hypertrophic cardiomyopathy in Afro-Caribbean versus white patients in the UK. Heart, 2016, 102, 1797-1804.	2.9	52
133	New Insights Into the Genetic Basis of Inherited Arrhythmia Syndromes. Circulation: Cardiovascular Genetics, 2016, 9, 569-577.	5.1	45
134	Reply. Journal of the American College of Cardiology, 2016, 67, 1658-1659.	2.8	1
135	Opening Pandora's Box â€" incidental genetic findings. Nature Reviews Cardiology, 2016, 13, 187-188.	13.7	0
136	Evaluation of the Achieve Mapping Catheter in cryoablation for atrial fibrillation: a prospective randomized trial. Journal of Interventional Cardiac Electrophysiology, 2016, 45, 179-187.	1.3	7
137	Etiology of Sudden Death in Sports. Journal of the American College of Cardiology, 2016, 67, 2108-2115.	2.8	399
138	Pharmacological treatment of acquired QT prolongation and torsades de pointes. British Journal of Clinical Pharmacology, 2016, 81, 420-427.	2.4	105
139	Reply. Journal of the American College of Cardiology, 2016, 68, 2126.	2.8	0
140	66â€...Sudden Death in Wolf-Parkinson-White. Description of Post-mortem Pathological Findings and Clinical Correlates in 19 Cases: Abstract 66 Table 1. Heart, 2016, 102, A48-A49.	2.9	0
141	The Prevalence and Significance of the Early Repolarization Pattern in Sudden Arrhythmic Death Syndrome Families. Circulation: Arrhythmia and Electrophysiology, 2016, 9, .	4.8	19
142	New approaches to predicting the risk of sudden death. Clinical Medicine, 2016, 16, 283-283.	1.9	9
143	Sudden unexpected death in epilepsy genetics: Molecular diagnostics and prevention. Epilepsia, 2016, 57, 17-25.	5.1	74
144	The ventricular ectopic QRS interval (VEQSI): Diagnosis of arrhythmogenic right ventricular cardiomyopathy in patients with incomplete disease expression. Heart Rhythm, 2016, 13, 1504-1512.	0.7	13

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145	Impact of clinical and genetic findings on the management of young patients with Brugada syndrome. Heart Rhythm, 2016, 13, 1274-1282.	0.7	89
146	Novel electrocardiographic criteria for the diagnosis of arrhythmogenic right ventricular cardiomyopathy. Europace, 2016, 18, 1420-1426.	1.7	11
147	The prevalence and significance of a short QT interval in 18â€...825 low-risk individuals including athletes. British Journal of Sports Medicine, 2016, 50, 124-129.	6.7	74
148	Sudden cardiac arrest in sports â€“ need for uniform registration: A Position Paper from the Sport Cardiology Section of the European Association for Cardiovascular Prevention and Rehabilitation. European Journal of Preventive Cardiology, 2016, 23, 657-667.	1.8	78
149	The role of genetic testing in unexplained sudden death. Translational Research, 2016, 168, 59-73.	5.0	33
150	Long QT Syndrome. , 2016, , 155-173.		2
151	Reply. Journal of the American College of Cardiology, 2015, 66, 2471-2472.	2.8	0
152	Faulty cardiac repolarization reserve in alternating hemiplegia of childhood broadens the phenotype. Brain, 2015, 138, 2859-2874.	7.6	30
153	Author's reply. Europace, 2015, 17, 1739-40.	1.7	0
154	QRSâ€“STâ€“T triangulation with repolarization shortening as a precursor of sustained ventricular tachycardia during acute myocardial ischemia. Journal of Arrhythmia, 2015, 31, 118-120.	1.2	1
155	Role of common and rare variants in SCN10A: results from the Brugada syndrome QRS locus gene discovery collaborative study. Cardiovascular Research, 2015, 106, 520-529.	3.8	108
156	VERP in Brugada syndrome â€" Very effective risk predictor?. International Journal of Cardiology, 2015, 184, 270-271.	1.7	3
157	Clinical Differentiation Between Physiological Remodeling and Arrhythmogenic Right Ventricular Cardiomyopathy in Athletes With Marked Electrocardiographic Repolarization Anomalies. Journal of the American College of Cardiology, 2015, 65, 2702-2711.	2.8	98
158	Computed bipolar precordial leads for improved P wave detection. Journal of Electrocardiology, 2015, 48, 188-189.	0.9	0
159	Fibrosis, Connexin-43, and Conduction Abnormalities in the Brugada Syndrome. Journal of the American College of Cardiology, 2015, 66, 1976-1986.	2.8	315
160	Electrocardiographic methods for diagnosis and risk stratification in the Brugada syndrome. Journal of the Saudi Heart Association, 2015, 27, 96-108.	0.4	19
161	Sudden Unexplained Death â€“ Treating the Family. Arrhythmia and Electrophysiology Review, 2014, 3, 156.	2.4	7
162	Burden of Sudden Cardiac Death in Persons Aged 1 to 49 Years. Circulation: Arrhythmia and Electrophysiology, 2014, 7, 205-211.	4.8	142

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163	Clinical Characteristics and Circumstances of Death in the Sudden Arrhythmic Death Syndrome. Circulation: Arrhythmia and Electrophysiology, 2014, 7, 1078-1083.	4.8	61
164	MicroRNAs in cardiac arrhythmia: DNA sequence variation of MiR-1 and MiR-133A in long QT syndrome. Scandinavian Journal of Clinical and Laboratory Investigation, 2014, 74, 485-491.	1.2	11
165	Executive Summary: HRS/EHRA/APHRS Expert Consensus Statement on the Diagnosis and Management of Patients with Inherited Primary Arrhythmia Syndromes. Journal of Arrhythmia, 2014, 30, 29-47.	1.2	16
166	HRS/EHRA/APHRS Expert Consensus Statement on the Diagnosis and Management of Patients with Inherited Primary Arrhythmia Syndromes. Journal of Arrhythmia, 2014, 30, 1-28.	1.2	49
167	Clinical utility of computed electrocardiographic leads. Journal of Electrocardiology, 2014, 47, 281-287.	0.9	4
168	The genetics of proâ€arrhythmic adverse drug reactions. British Journal of Clinical Pharmacology, 2014, 77, 618-625.	2.4	7
169	Genetic association study of QT interval highlights role for calcium signaling pathways in myocardial repolarization. Nature Genetics, 2014, 46, 826-836.	21.4	281
170	Cardiac Evaluation of Pediatric Relatives in Sudden Arrhythmic Death Syndrome. Circulation: Arrhythmia and Electrophysiology, 2014, 7, 800-806.	4.8	33
171	Novel Calmodulin Mutations Associated With Congenital Arrhythmia Susceptibility. Circulation: Cardiovascular Genetics, 2014, 7, 466-474.	5.1	165
172	Sudden unexplained death in infants and children: the role of undiagnosed inherited cardiac conditions. Europace, 2014, 16, 1706-1713.	1.7	34
173	The importance of specialist cardiac histopathological examination in the investigation of young sudden cardiac deaths. Europace, 2014, 16, 899-907.	1.7	104
174	Common variants at SCN5A-SCN10A and HEY2 are associated with Brugada syndrome, a rare disease with high risk of sudden cardiac death. Nature Genetics, 2013, 45, 1044-1049.	21.4	467
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