

Dawood Darbar

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174
papers

10,174
citations

51
h-index

99
g-index

201
ext. papers

12,586
ext. citations

8.3
avg, IF

5.55
L-index

#	Paper	IF	Citations
174	2012 ACCF/AHA/HRS focused update incorporated into the ACCF/AHA/HRS 2008 guidelines for device-based therapy of cardiac rhythm abnormalities: a report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines and the Heart Rhythm Society. <i>Journal of the American College of Cardiology</i> , 2013 , 61, e6-75	15.1	602
173	Meta-analysis identifies six new susceptibility loci for atrial fibrillation. <i>Nature Genetics</i> , 2012 , 44, 670-5	36.3	429
172	2012 ACCF/AHA/HRS focused update incorporated into the ACCF/AHA/HRS 2008 guidelines for device-based therapy of cardiac rhythm abnormalities: a report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines and the Heart Rhythm Society. <i>Circulation</i> , 2012 , 127, e283-356	16.7	393
171	Prevention of atrial fibrillation: report from a national heart, lung, and blood institute workshop. <i>Circulation</i> , 2009 , 119, 606-18	16.7	378
170	A sequence variant in ZFX3 on 16q22 associates with atrial fibrillation and ischemic stroke. <i>Nature Genetics</i> , 2009 , 41, 876-8	36.3	365
169	Common variants in KCNN3 are associated with lone atrial fibrillation. <i>Nature Genetics</i> , 2010 , 42, 240-4	36.3	362
168	Metabolic syndrome and risk of development of atrial fibrillation: the Niigata preventive medicine study. <i>Circulation</i> , 2008 , 117, 1255-60	16.7	308
167	Drug-induced long QT syndrome. <i>Pharmacological Reviews</i> , 2010 , 62, 760-81	22.5	296
166	Multi-ethnic genome-wide association study for atrial fibrillation. <i>Nature Genetics</i> , 2018 , 50, 1225-1233	36.3	277
165	2012 ACCF/AHA/HRS focused update of the 2008 guidelines for device-based therapy of cardiac rhythm abnormalities: a report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines. <i>Journal of the American College of Cardiology</i> , 2012 , 60, 1297-313	15.1	273
164	Familial atrial fibrillation is a genetically heterogeneous disorder. <i>Journal of the American College of Cardiology</i> , 2003 , 41, 2185-92	15.1	273
163	Sequencing of 53,831 diverse genomes from the NHLBI TOPMed Program. <i>Nature</i> , 2021 , 590, 290-299	50.4	268
162	2012 ACCF/AHA/HRS focused update of the 2008 guidelines for device-based therapy of cardiac rhythm abnormalities: a report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines and the Heart Rhythm Society. [corrected]. <i>Circulation</i> , 2012 , 126, 1784-800	16.7	250
161	Cardiac sodium channel (SCN5A) variants associated with atrial fibrillation. <i>Circulation</i> , 2008 , 117, 1927-35	36.7	245
160	Atrial natriuretic peptide frameshift mutation in familial atrial fibrillation. <i>New England Journal of Medicine</i> , 2008 , 359, 158-65	59.2	236
159	A rare variant in MYH6 is associated with high risk of sick sinus syndrome. <i>Nature Genetics</i> , 2011 , 43, 316-20	30.3	228
158	Identification of heart rate-associated loci and their effects on cardiac conduction and rhythm disorders. <i>Nature Genetics</i> , 2013 , 45, 621-31	36.3	219

157	Role of inflammation and oxidative stress in atrial fibrillation. <i>Heart Rhythm</i> , 2010 , 7, 438-44	6.7	213
156	Large-scale analyses of common and rare variants identify 12 new loci associated with atrial fibrillation. <i>Nature Genetics</i> , 2017 , 49, 946-952	36.3	176
155	2014 AATS guidelines for the prevention and management of perioperative atrial fibrillation and flutter for thoracic surgical procedures. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2014 , 148, e153-93 ⁵	15.5	170
154	Mutations in sodium channel α - and β -subunits associated with atrial fibrillation. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2009 , 2, 268-75	6.4	169
153	Large scale replication and meta-analysis of variants on chromosome 4q25 associated with atrial fibrillation. <i>European Heart Journal</i> , 2009 , 30, 813-9	9.5	165
152	Integrating genetic, transcriptional, and functional analyses to identify 5 novel genes for atrial fibrillation. <i>Circulation</i> , 2014 , 130, 1225-35	16.7	143
151	Inherited causes of clonal haematopoiesis in 97,691 whole genomes. <i>Nature</i> , 2020 , 586, 763-768	50.4	127
150	Diagnostic value of B-type natriuretic peptide concentrations in patients with acute myocardial infarction. <i>American Journal of Cardiology</i> , 1996 , 78, 284-7	3	107
149	Relation of the severity of obstructive sleep apnea in response to anti-arrhythmic drugs in patients with atrial fibrillation or atrial flutter. <i>American Journal of Cardiology</i> , 2012 , 110, 369-72	3	103
148	The APPLE score: a novel and simple score for the prediction of rhythm outcomes after catheter ablation of atrial fibrillation. <i>Clinical Research in Cardiology</i> , 2015 , 104, 871-6	6.1	96
147	Symptomatic response to antiarrhythmic drug therapy is modulated by a common single nucleotide polymorphism in atrial fibrillation. <i>Journal of the American College of Cardiology</i> , 2012 , 60, 539-45	15.1	93
146	Role of common and rare variants in SCN10A: results from the Brugada syndrome QRS locus gene discovery collaborative study. <i>Cardiovascular Research</i> , 2015 , 106, 520-9	9.9	86
145	A KCNJ8 mutation associated with early repolarization and atrial fibrillation. <i>Europace</i> , 2012 , 14, 1428-32 ⁹	3.9	84
144	Unmasking of brugada syndrome by lithium. <i>Circulation</i> , 2005 , 112, 1527-31	16.7	84
143	Variation in the 4q25 chromosomal locus predicts atrial fibrillation after coronary artery bypass graft surgery. <i>Circulation: Cardiovascular Genetics</i> , 2009 , 2, 499-506		79
142	Atrial Fibrillation Is an Independent Predictor of Mortality in Critically Ill Patients. <i>Critical Care Medicine</i> , 2015 , 43, 2104-11	1.4	77
141	Polymorphism modulates symptomatic response to antiarrhythmic drug therapy in patients with lone atrial fibrillation. <i>Heart Rhythm</i> , 2007 , 4, 743-9	6.7	77
140	Augmented potassium current is a shared phenotype for two genetic defects associated with familial atrial fibrillation. <i>Journal of Molecular and Cellular Cardiology</i> , 2010 , 48, 181-90	5.8	76

139	Exaggerated QT prolongation after cardioversion of atrial fibrillation. <i>Journal of the American College of Cardiology</i> , 1999 , 34, 396-401	15.1	76
138	Association Between Titin Loss-of-Function Variants and Early-Onset Atrial Fibrillation. <i>JAMA - Journal of the American Medical Association</i> , 2018 , 320, 2354-2364	27.4	75
137	Voltage-gated sodium channels: biophysics, pharmacology, and related channelopathies. <i>Frontiers in Pharmacology</i> , 2012 , 3, 124	5.6	72
136	Common genetic variants and response to atrial fibrillation ablation. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2015 , 8, 296-302	6.4	70
135	Sensitivity and specificity of QTc dispersion for identification of risk of cardiac death in patients with peripheral vascular disease. <i>BMJ: British Medical Journal</i> , 1996 , 312, 874-8; discussion 878-9		70
134	Common genetic polymorphism at 4q25 locus predicts atrial fibrillation recurrence after successful cardioversion. <i>Heart Rhythm</i> , 2013 , 10, 849-55	6.7	67
133	Chromosome 4q25 variants are genetic modifiers of rare ion channel mutations associated with familial atrial fibrillation. <i>Journal of the American College of Cardiology</i> , 2012 , 60, 1173-81	15.1	65
132	A common β -adrenergic receptor polymorphism predicts favorable response to rate-control therapy in atrial fibrillation. <i>Journal of the American College of Cardiology</i> , 2012 , 59, 49-56	15.1	64
131	P wave signal-averaged electrocardiography to identify risk for atrial fibrillation. <i>PACE - Pacing and Clinical Electrophysiology</i> , 2002 , 25, 1447-53	1.6	63
130	Genetic mechanisms of atrial fibrillation: impact on response to treatment. <i>Nature Reviews Cardiology</i> , 2013 , 10, 317-29	14.8	60
129	Novel KCNA5 mutation implicates tyrosine kinase signaling in human atrial fibrillation. <i>Heart Rhythm</i> , 2010 , 7, 1246-52	6.7	58
128	Genetic Risk Prediction of Atrial Fibrillation. <i>Circulation</i> , 2017 , 135, 1311-1320	16.7	56
127	Effect of Peripheral Arterial Disease in Patients Undergoing Percutaneous Coronary Intervention With Intracoronary Stents. <i>Mayo Clinic Proceedings</i> , 2004 , 79, 1113-1118	6.4	53
126	A clinical prediction model to estimate risk for 30-day adverse events in emergency department patients with symptomatic atrial fibrillation. <i>Annals of Emergency Medicine</i> , 2011 , 57, 1-12	2.1	51
125	Prolonged signal-averaged P-wave duration as an intermediate phenotype for familial atrial fibrillation. <i>Journal of the American College of Cardiology</i> , 2008 , 51, 1083-9	15.1	51
124	Relation of morbid obesity and female gender to risk of procedural complications in patients undergoing atrial fibrillation ablation. <i>American Journal of Cardiology</i> , 2013 , 111, 368-73	3	48
123	2014 AATS guidelines for the prevention and management of perioperative atrial fibrillation and flutter for thoracic surgical procedures. Executive summary. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2014 , 148, 772-91	1.5	47
122	Ictal asystole and ictal syncope: insights into clinical management. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2015 , 8, 159-64	6.4	46

121	SCN10A/Nav1.8 modulation of peak and late sodium currents in patients with early onset atrial fibrillation. <i>Cardiovascular Research</i> , 2014 , 104, 355-63	9.9	46
120	A Missense Variant in PLEC Increases Risk of Atrial Fibrillation. <i>Journal of the American College of Cardiology</i> , 2017 , 70, 2157-2168	15.1	43
119	Effect of omega-three polyunsaturated fatty acids on inflammation, oxidative stress, and recurrence of atrial fibrillation. <i>American Journal of Cardiology</i> , 2015 , 115, 196-201	3	42
118	Epinephrine-induced changes in serum potassium and cardiac repolarization and effects of pretreatment with propranolol and diltiazem. <i>American Journal of Cardiology</i> , 1996 , 77, 1351-5	3	42
117	Suppression of spontaneous ca elevations prevents atrial fibrillation in caldesmon 2-null hearts. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2014 , 7, 313-20	6.4	40
116	Differential impact of race and risk factors on incidence of atrial fibrillation. <i>American Heart Journal</i> , 2011 , 162, 31-7	4.9	39
115	Persistent atrial fibrillation is associated with reduced risk of torsades de pointes in patients with drug-induced long QT syndrome. <i>Journal of the American College of Cardiology</i> , 2008 , 51, 836-42	15.1	38
114	Arrhythmogenic right ventricular cardiomyopathy due to a novel plakophilin 2 mutation: wide spectrum of disease in mutation carriers within a family. <i>Heart Rhythm</i> , 2006 , 3, 939-44	6.7	37
113	Race-specific impact of atrial fibrillation risk factors in blacks and whites in the southern community cohort study. <i>American Journal of Cardiology</i> , 2012 , 110, 1637-42	3	35
112	Functional modeling in zebrafish demonstrates that the atrial-fibrillation-associated gene GREM2 regulates cardiac laterality, cardiomyocyte differentiation and atrial rhythm. <i>DMM Disease Models and Mechanisms</i> , 2013 , 6, 332-41	4.1	35
111	Tremor-induced ECG artifact mimicking ventricular tachycardia. <i>Circulation</i> , 2000 , 102, 1337-8	16.7	35
110	Characterization of genome-wide association-identified variants for atrial fibrillation in African Americans. <i>PLoS ONE</i> , 2012 , 7, e32338	3.7	33
109	Dietary salt increases first-pass elimination of oral quinidine. <i>Clinical Pharmacology and Therapeutics</i> , 1997 , 61, 292-300	6.1	32
108	Evaluation of a Prediction Model for the Development of Atrial Fibrillation in a Repository of Electronic Medical Records. <i>JAMA Cardiology</i> , 2016 , 1, 1007-1013	16.2	32
107	Electrophysiologic Characterization of Calcium Handling in Human Induced Pluripotent Stem Cell-Derived Atrial Cardiomyocytes. <i>Stem Cell Reports</i> , 2018 , 10, 1867-1878	8	31
106	Whole-exome sequencing in familial atrial fibrillation. <i>European Heart Journal</i> , 2014 , 35, 2477-83	9.5	30
105	2012 ACCF/AHA/HRS focused update of the 2008 guidelines for device-based therapy of cardiac rhythm abnormalities: a report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2012 , 144, e127-45	1.5	29
104	Modulation by dietary salt of verapamil disposition in humans. <i>Circulation</i> , 1998 , 98, 2702-8	16.7	28

103	Symptomatic burden as an endpoint to evaluate interventions in patients with atrial fibrillation. <i>Heart Rhythm</i> , 2005 , 2, 544-9	6.7	27
102	Prediction of late cardiac events by dipyridamole thallium scintigraphy in patients with intermittent claudication and occult coronary artery disease. <i>American Journal of Cardiology</i> , 1996 , 78, 736-40	3	25
101	Genetic and clinical risk prediction model for postoperative atrial fibrillation. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2015 , 8, 25-31	6.4	24
100	Genetics of atrial fibrillation: rare mutations, common polymorphisms, and clinical relevance. <i>Heart Rhythm</i> , 2008 , 5, 483-6	6.7	23
99	Risk factors for bradycardia requiring pacemaker implantation in patients with atrial fibrillation. <i>American Journal of Cardiology</i> , 2012 , 110, 1315-21	3	22
98	Coding variants in and increase risk of atrial fibrillation. <i>Communications Biology</i> , 2018 , 1, 68	6.7	21
97	Pharmacogenetics of antiarrhythmic therapy. <i>Expert Opinion on Pharmacotherapy</i> , 2006 , 7, 1583-90	4	21
96	The AFFORD clinical decision aid to identify emergency department patients with atrial fibrillation at low risk for 30-day adverse events. <i>American Journal of Cardiology</i> , 2015 , 115, 763-70	3	20
95	Selective targeting of gain-of-function KCNQ1 mutations predisposing to atrial fibrillation. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2013 , 6, 960-6	6.4	20
94	ACE I/D polymorphism associated with abnormal atrial and atrioventricular conduction in lone atrial fibrillation and structural heart disease: implications for electrical remodeling. <i>Heart Rhythm</i> , 2009 , 6, 1327-32	6.7	20
93	A rate-independent method of assessing QT-RR slope following conversion of atrial fibrillation. <i>Journal of Cardiovascular Electrophysiology</i> , 2007 , 18, 636-41	2.7	20
92	Variants in the SCN5A Promoter Associated With Various Arrhythmia Phenotypes. <i>Journal of the American Heart Association</i> , 2016 , 5,	6	18
91	Localization of the origin of arrhythmias for ablation: from Electrocardiography to advanced endocardial mapping systems. <i>Journal of Cardiovascular Electrophysiology</i> , 2001 , 12, 1309-25	2.7	18
90	Common SCN10A variants modulate PR interval and heart rate response during atrial fibrillation. <i>Europace</i> , 2014 , 16, 485-90	3.9	17
89	Future of antiarrhythmic drugs. <i>Current Opinion in Cardiology</i> , 2006 , 21, 361-7	2.1	17
88	Ion Channel and Structural Remodeling in Obesity-Mediated Atrial Fibrillation. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2020 , 13, e008296	6.4	17
87	Effect of peripheral arterial disease in patients undergoing percutaneous coronary intervention with intracoronary stents. <i>Mayo Clinic Proceedings</i> , 2004 , 79, 1113-8	6.4	16
86	Clinical use of and future perspectives on antiarrhythmic drugs. <i>European Journal of Clinical Pharmacology</i> , 2008 , 64, 1139-46	2.8	15

85	Candidate gene approach to identifying rare genetic variants associated with lone atrial fibrillation. <i>Heart Rhythm</i> , 2014 , 11, 46-52	6.7	14
84	The "missing" link in atrial fibrillation heritability. <i>Journal of Electrocardiology</i> , 2011 , 44, 641-4	1.4	14
83	Genomics, heart failure and sudden cardiac death. <i>Heart Failure Reviews</i> , 2010 , 15, 229-38	5	14
82	Implantable cardioverter-defibrillator malfunction due to mechanical failure of the header connection. <i>Journal of Cardiovascular Electrophysiology</i> , 2004 , 15, 1095-9	2.7	14
81	Electrophysiologic and molecular mechanisms of a frameshift NPPA mutation linked with familial atrial fibrillation. <i>Journal of Molecular and Cellular Cardiology</i> , 2019 , 132, 24-35	5.8	13
80	A common variant on chromosome 4q25 is associated with prolonged PR interval in subjects with and without atrial fibrillation. <i>American Journal of Cardiology</i> , 2014 , 113, 309-13	3	13
79	The Role of Pharmacogenetics in Atrial Fibrillation Therapeutics: Is Personalized Therapy in Sight?. <i>Journal of Cardiovascular Pharmacology</i> , 2016 , 67, 9-18	3.1	13
78	Association Between Family History and Early-Onset Atrial Fibrillation Across Racial and Ethnic Groups. <i>JAMA Network Open</i> , 2018 , 1, e182497	10.4	13
77	Genetic Susceptibility for Atrial Fibrillation in Patients Undergoing Atrial Fibrillation Ablation. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2020 , 13, e007676	6.4	12
76	Evaluating the HATCH score for predicting progression to sustained atrial fibrillation in ED patients with new atrial fibrillation. <i>American Journal of Emergency Medicine</i> , 2013 , 31, 792-7	2.9	12
75	A genotype-dependent intermediate ECG phenotype in patients with persistent lone atrial fibrillation genotype ECG-phenotype correlation in atrial fibrillation. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2009 , 2, 24-8	6.4	12
74	Sympathetic activation enhances QT prolongation by quinidine. <i>Journal of Cardiovascular Electrophysiology</i> , 2001 , 12, 9-14	2.7	12
73	A genome-wide association study to identify genomic modulators of rate control therapy in patients with atrial fibrillation. <i>American Journal of Cardiology</i> , 2014 , 114, 593-600	3	11
72	Improved understanding of the pathophysiology of atrial fibrillation through the lens of discrete pathological pathways. <i>Global Cardiology Science & Practice</i> , 2014 , 2014, 24-36	0.7	11
71	Association of Rare Genetic Variants and Early-Onset Atrial Fibrillation in Ethnic Minority Individuals. <i>JAMA Cardiology</i> , 2021 , 6, 811-819	16.2	11
70	Prevalence and predictors of atrial fibrillation among patients undergoing bariatric surgery. <i>Obesity Surgery</i> , 2014 , 24, 611-6	3.7	10
69	Sotalol-induced torsades de pointes precipitated during treatment with oseltamivir for H1N1 influenza. <i>Heart Rhythm</i> , 2010 , 7, 1454-7	6.7	10
68	On the relationship among QT interval, atrial fibrillation, and torsade de pointes. <i>Europace</i> , 2007 , 9 Suppl 4, iv1-3	3.9	10

67	Examining rare and low-frequency genetic variants previously associated with lone or familial forms of atrial fibrillation in an electronic medical record system: a cautionary note. <i>Circulation: Cardiovascular Genetics</i> , 2015 , 8, 58-63		9
66	Proarrhythmic and Torsadogenic Effects of Potassium Channel Blockers in Patients. <i>Cardiac Electrophysiology Clinics</i> , 2016 , 8, 481-93	1.4	9
65	Atrial Fibrillation and Variants. <i>Cardiac Electrophysiology Clinics</i> , 2014 , 6, 741-748	1.4	9
64	Severity of obstructive sleep apnea influences the effect of genotype on response to anti-arrhythmic drug therapy for atrial fibrillation. <i>Journal of Clinical Sleep Medicine</i> , 2014 , 10, 503-7	3.1	9
63	Atrial fibrillation and flutter outcomes and risk determination (AFFORD): design and rationale. <i>Journal of Cardiology</i> , 2011 , 58, 124-30	3	9
62	Early-Onset Atrial Fibrillation and the Prevalence of Rare Variants in Cardiomyopathy and Arrhythmia Genes. <i>JAMA Cardiology</i> , 2021 ,	16.2	9
61	Genetic modulation of atrial fibrillation risk in a Hispanic/Latino cohort. <i>PLoS ONE</i> , 2018 , 13, e0194480	3.7	8
60	Investigating the Genetic Architecture of the PR Interval Using Clinical Phenotypes. <i>Circulation: Cardiovascular Genetics</i> , 2017 , 10,		7
59	Is it time to develop a "pathogenicity" score to distinguish long QT syndrome causing mutations from "background" genetic noise?. <i>Heart Rhythm</i> , 2009 , 6, 1304-5	6.7	7
58	Images in cardiovascular medicine. Himalayan T waves in the congenital long-QT syndrome. <i>Circulation</i> , 2005 , 111, e161	16.7	6
57	Lone AF - Etiologic Factors and Genetic Insights into Pathophysiology. <i>Journal of Atrial Fibrillation</i> , 2010 , 3, 236	0.8	6
56	Association Between Obesity-Mediated Atrial Fibrillation and Therapy With Sodium Channel Blocker Antiarrhythmic Drugs. <i>JAMA Cardiology</i> , 2020 , 5, 57-64	16.2	6
55	Human induced pluripotent stem cell-derived atrial cardiomyocytes carrying an SCN5A mutation identify nitric oxide signaling as a mediator of atrial fibrillation. <i>Stem Cell Reports</i> , 2021 , 16, 1542-1554	8	6
54	Relation of Body Mass Index to Symptom Burden in Patients with Atrial Fibrillation. <i>American Journal of Cardiology</i> , 2018 , 122, 235-241	3	6
53	Assessing the contribution of rare variants to complex trait heritability from whole-genome sequence data.. <i>Nature Genetics</i> , 2022 ,	36.3	6
52	Genotype influence in responses to therapy for atrial fibrillation. <i>Expert Review of Cardiovascular Therapy</i> , 2016 , 14, 1119-31	2.5	5
51	Genome-wide assessment for genetic variants associated with ventricular dysfunction after primary coronary artery bypass graft surgery. <i>PLoS ONE</i> , 2011 , 6, e24593	3.7	5
50	Arrhythmia pharmacogenomics: methodological considerations. <i>Current Pharmaceutical Design</i> , 2009 , 15, 3734-41	3.3	5

49	Recurrence of atrial tachyarrhythmias in implantable cardioverter-defibrillator recipients. <i>PACE - Pacing and Clinical Electrophysiology</i> , 2005 , 28, 1047-51	1.6	5
48	Relation of Obstructive Sleep Apnea and a Common Variant at Chromosome 4q25 to Atrial Fibrillation. <i>American Journal of Cardiology</i> , 2017 , 119, 1387-1391	3	4
47	Impact of traditional risk factors for the outcomes of atrial fibrillation across race and ethnicity and sex groups. <i>IJC Heart and Vasculature</i> , 2020 , 28, 100538	2.4	4
46	Rare variants in genes encoding the cardiac sodium channel and associated compounds and their impact on outcome of catheter ablation of atrial fibrillation. <i>PLoS ONE</i> , 2017 , 12, e0183690	3.7	4
45	Triggers for cardiac events in patients with type 2 long QT syndrome. <i>Heart Rhythm</i> , 2010 , 7, 1806-7	6.7	4
44	Cardiac sodium channel variants: action players with many faces. <i>Heart Rhythm</i> , 2008 , 5, 1441-3	6.7	4
43	Congenital long QT syndrome aggravated by salt-wasting nephropathy. <i>Heart Rhythm</i> , 2005 , 2, 304-6	6.7	4
42	Heart failure: a diagnostic and therapeutic dilemma in elderly patients. <i>Age and Ageing</i> , 1998 , 27, 539-43		4
41	Can qualitative echocardiography be used to select patients for angiotensin-converting enzyme inhibitors following acute myocardial infarction?. <i>European Heart Journal</i> , 1996 , 17, 1783-6	9.5	4
40	Loss of quinidine gluconate injection in a polyvinyl chloride infusion system. <i>American Journal of Health-System Pharmacy</i> , 1996 , 53, 655-8	2.2	4
39	Common genetic variants associated with obesity in an African-American and Hispanic/Latino population. <i>PLoS ONE</i> , 2021 , 16, e0250697	3.7	4
38	Measurement of diffuse ventricular fibrosis with myocardial T1 in patients with atrial fibrillation. <i>Journal of Arrhythmia</i> , 2016 , 32, 51-6	1.5	4
37	Whole Genome Sequencing Identifies CRISPLD2 as a Lung Function Gene in Children With Asthma. <i>Chest</i> , 2019 , 156, 1068-1079	5.3	3
36	Multi-Ethnic Genome-Wide Association Study of Decomposed Cardioelectric Phenotypes Illustrates Strategies to Identify and Characterize Evidence of Shared Genetic Effects for Complex Traits. <i>Circulation Genomic and Precision Medicine</i> , 2020 , 13, e002680	5.2	3
35	Association of atrial fibrillation risk alleles and response to acute rate control therapy. <i>American Journal of Emergency Medicine</i> , 2016 , 34, 735-40	2.9	3
34	GWAS of QRS duration identifies new loci specific to Hispanic/Latino populations. <i>PLoS ONE</i> , 2019 , 14, e0217796	3.7	3
33	Attitudes of physicians in the treatment of congestive heart failure in older adults. <i>Journal of the American Geriatrics Society</i> , 1995 , 43, 943-4	5.6	3
32	Abstract 4099: Genetic and Clinical Predictors of Response to Rate Control Therapy in Patients with Atrial Fibrillation. <i>Circulation</i> , 2008 , 118,	16.7	3

31	Pathogenic mutations perturb calmodulin regulation of Na1.8 channel. <i>Biochemical and Biophysical Research Communications</i> , 2020 , 533, 168-174	3.4	3
30	Mendelian randomization supports bidirectional causality between telomere length and clonal hematopoiesis of indeterminate potential.. <i>Science Advances</i> , 2022 , 8, eabl6579	14.3	3
29	Standard Antiarrhythmic Drugs 2014 , 1095-1110		2
28	P1-84. <i>Heart Rhythm</i> , 2006 , 3, S135-S136	6.7	2
27	Abstract 356: Loss of Function Mutations of Sodium Channel Beta-1 and Beta-2 Subunits Associated with Atrial Fibrillation and ST-segment Elevation. <i>Circulation</i> , 2007 , 116,	16.7	2
26	Clinical and Genetic Contributors to New-Onset Atrial Fibrillation in Critically Ill Adults. <i>Critical Care Medicine</i> , 2020 , 48, 22-30	1.4	2
25	Bioengineering approaches to mature induced pluripotent stem cell-derived atrial cardiomyocytes to model atrial fibrillation. <i>Experimental Biology and Medicine</i> , 2021 , 246, 1816-1828	3.7	2
24	Common genetic variation in circadian clock genes are associated with cardiovascular risk factors in an African American and Hispanic/Latino cohort. <i>IJC Heart and Vasculature</i> , 2021 , 34, 100808	2.4	2
23	A new paradigm for predicting risk of Torsades de Pointes during drug development: Commentary on: "Improved prediction of drug-induced Torsades de Pointes through simulations of dynamics and machine learning algorithms". <i>Clinical Pharmacology and Therapeutics</i> , 2016 , 100, 324-6	6.1	2
22	Mutant ANP induces mitochondrial and ion channel remodeling in a human iPSC-derived atrial fibrillation model.. <i>JCI Insight</i> , 2022 , 7,	9.9	2
21	Standard Antiarrhythmic Drugs 2018 , 1062-1075		1
20	Congenital right coronary artery aneurysm causing myocardial infarction, pseudoaneurysm formation, and right atrial compression. <i>Journal of the American Society of Echocardiography</i> , 2002 , 15, 736-8	5.8	1
19	Localized aortic dissection: unusual features by transesophageal echocardiography. <i>Journal of the American Society of Echocardiography</i> , 2000 , 13, 1130-4	5.8	1
18	Genetic determinants of telomere length from 109,122 ancestrally diverse whole-genome sequences in TOPMed.. <i>Cell Genomics</i> , 2022 , 2, 100084-100084		1
17	Atrial Fibrillation Susceptibility Alleles on Chromosome 4q25 Modulate Response to Catheter Ablation. <i>Journal of Atrial Fibrillation</i> , 2010 , 3, 272	0.8	1
16	Genetic and Molecular Basis of Arrhythmias 2011 , 65-86		1
15	Mutations in RPL3L and MYZAP increase risk of atrial fibrillation		1
14	Atrial Fibrillation Genetic Risk Differentiates Cardioembolic Stroke from other Stroke Subtypes		1

13	Genetics of atrial fibrillation-practical applications for clinical management: if not now, when and how?. <i>Cardiovascular Research</i> , 2021 , 117, 1718-1731	9.9	1
12	Atrial Fibrillation in Inherited Channelopathies. <i>Cardiac Electrophysiology Clinics</i> , 2021 , 13, 155-163	1.4	0
11	Assessment of the Framingham risk factors among ED patients with newly diagnosed atrial fibrillation. <i>American Journal of Emergency Medicine</i> , 2012 , 30, 151-7	2.9	
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