Henrik K Jensen

List of Publications by Citations

138 papers	3,525 citations	29 h-index	56 g-index
150 ext. papers	4,299 ext. citations	4.5 avg, IF	4.8 L-index

#	Paper	IF	Citations
138	Tissue Doppler imaging predicts improved systolic performance and reversed left ventricular remodeling during long-term cardiac resynchronization therapy. *Journal of the American College of Cardiology*, **2002**, 40, 723-30	15.1	499
137	Lipoprotein(a), PCSK9 Inhibition, and Cardiovascular Risk. *Circulation*, **2019**, 139, 1483-1492	16.7	288
136	A sequence variation: 713-8delC in the transforming growth factor-beta 1 gene has higher prevalence in osteoporotic women than in normal women and is associated with very low bone mass in osteoporotic women and increased bone turnover in both osteoporotic and normal women. *Bone*, **1997**, 20, 289-94	4.7	194
135	The relationship of molecular genetic to clinical diagnosis of familial hypercholesterolemia in a Danish population. *Atherosclerosis*, **2005**, 180, 155-60	3.1	134
134	His or para-His pacing preserves left ventricular function in atrioventricular block: a double-blind, randomized, crossover study. *Europace*, **2014**, 16, 1189-96	3.9	101
133	Cardiac magnetic resonance and electroanatomical mapping of acute and chronic atrial ablation injury: a histological validation study. *European Heart Journal*, **2014**, 35, 1486-95	9.5	89
132	Ablation of atrial tachycardia after surgery for congenital and acquired heart disease using an electroanatomic mapping system: Which circuits to expect in which substrate?. *Heart Rhythm*, **2005**, 2, 64-72	6.7	87
131	Risk of arrhythmia induced by psychotropic medications: a proposal for clinical management. *European Heart Journal*, **2014**, 35, 1306-15	9.5	73
130	Multimodality imaging-guided left ventricular lead placement in cardiac resynchronization therapy: a randomized controlled trial. *European Journal of Heart Failure*, **2016**, 18, 1365-1374	12.3	70
129	Left ventricular lead performance in cardiac resynchronization therapy: impact of lead localization and complications. *PACE - Pacing and Clinical Electrophysiology*, **2005**, 28, 483-8	1.6	69
128	Mutations in the potassium channel subunit KCNE1 are associated with early-onset familial atrial fibrillation. *BMC Medical Genetics*, **2012**, 13, 24	2.1	67
127	Impact of acute biventricular pacing on left ventricular performance and volumes in patients with severe heart failure. A tissue doppler and three-dimensional echocardiographic study. *Cardiology*, **2001**, 95, 173-82	1.6	67
126	Catheter ablation of arrhythmias in ebstein's anomaly: a multicenter study. *Journal of Cardiovascular Electrophysiology*, **2011**, 22, 1391-6	2.7	65
125	The prevalence of mutations in KCNQ1, KCNH2, and SCN5A in an unselected national cohort of young sudden unexplained death cases. *Journal of Cardiovascular Electrophysiology*, **2012**, 23, 1092-8	2.7	57
124	The diagnostic performance of imaging methods in ARVC using the 2010 Task Force criteria. *European Heart Journal Cardiovascular Imaging*, **2014**, 15, 1219-25	4.1	56
123	Transcatheter left atrial appendage occlusion in patients with atrial fibrillation and a high bleeding risk using aspirin alone for post-implant antithrombotic therapy. *EuroIntervention*, **2017**, 12, 2075-2082	3.1	56
122	Amiodarone significantly decreases atrial fibrillation in patients undergoing surgery for lung cancer. *Annals of Thoracic Surgery*, **2012**, 94, 339-44; discussion 345-6	2.7	50

121	Biventricular pacing preserves left ventricular performance in patients with high-grade atrio-ventricular block: a randomized comparison with DDD(R) pacing in 50 consecutive patients. *Europace*, **2008**, 10, 314-20	3.9	50
120	T(peak)T(end) interval in long QT syndrome. *Journal of Electrocardiology*, **2008**, 41, 603-8	1.4	48
119	Loss-of-activity-mutation in the cardiac chloride-bicarbonate exchanger AE3 causes short QT syndrome. *Nature Communications*, **2017**, 8, 1696	17.4	46
118	Three dimensional echocardiography documents haemodynamic improvement by biventricular pacing in patients with severe heart failure. *British Heart Journal*, **2001**, 85, 514-20		45
117	His and para-His pacing in AV block: feasibility and electrocardiographic findings. *Journal of Interventional Cardiac Electrophysiology*, **2011**, 31, 255-62	2.4	43
116	Identifying drug-induced repolarization abnormalities from distinct ECG patterns in congenital long QT syndrome: a study of sotalol effects on T-wave morphology. *Drug Safety*, **2009**, 32, 599-611	5.1	43
115	The TMEM43 Newfoundland mutation p.S358L causing ARVC-5 was imported from Europe and increases the stiffness of the cell nucleus. *European Heart Journal*, **2015**, 36, 872-81	9.5	40
114	Phenotypic variation in patients heterozygous for familial defective apolipoprotein B (FDB) in three European countries. *Arteriosclerosis, Thrombosis, and Vascular Biology*, **1997**, 17, 741-7	9.4	40
113	The platelet polymorphism PlA2 is a genetic risk factor for myocardial infarction. *Journal of Internal Medicine*, **2004**, 255, 637-44	10.8	40
112	Spectrum of LDL receptor gene mutations in Denmark: implications for molecular diagnostic strategy in heterozygous familial hypercholesterolemia. *Atherosclerosis*, **1999**, 146, 337-44	3.1	35
111	Superior transseptal approach to mitral valve is associated with a higher need for pacemaker implantation than the left atrial approach. *Annals of Thoracic Surgery*, **2007**, 83, 77-82	2.7	31
110	No genetic linkage or molecular evidence for involvement of the PCSK9, ARH or CYP7A1 genes in the Familial Hypercholesterolemia phenotype in a sample of Danish families without pathogenic mutations in the LDL receptor and apoB genes. *Atherosclerosis*, **2004**, 177, 415-22	3.1	29
109	A common W556S mutation in the LDL receptor gene of Danish patients with familial hypercholesterolemia encodes a transport-defective protein. *Atherosclerosis*, **1997**, 131, 67-72	3.1	28
108	Mutated desmoglein-2 proteins are incorporated into desmosomes and exhibit dominant-negative effects in arrhythmogenic right ventricular cardiomyopathy. *Human Mutation*, **2013**, 34, 697-705	4.7	27
107	DDD(R)-pacing, but not AAI(R)-pacing induces left ventricular desynchronization in patients with sick sinus syndrome: tissue-Doppler and 3D echocardiographic evaluation in a randomized controlled comparison. *Europace*, **2008**, 10, 127-33	3.9	27
106	Truncating plakophilin-2 mutations in arrhythmogenic cardiomyopathy are associated with protein haploinsufficiency in both myocardium and epidermis. *Circulation: Cardiovascular Genetics*, **2014**, 7, 230-40		26
105	An Exploratory Analysis of Proprotein Convertase Subtilisin/Kexin Type 9 Inhibition and Aortic Stenosis in the FOURIER Trial. *JAMA Cardiology*, **2020**, 5, 709-713	16.2	25
104	Radiofrequency ablation of accessory pathways in patients with the Wolff-Parkinson-White syndrome: the long-term mortality and risk of atrial fibrillation. *Europace*, **2015**, 17, 117-22	3.9	25

103	Identification of recurrent and novel mutations in the LDL receptor gene in Spanish patients with familial hypercholesterolemia. *Human Mutation*, **1998**, 11, 413-413	4.7	25
102	Atrial incision affects the incidence of atrial tachycardia after mitral valve surgery. *Annals of Thoracic Surgery*, **2006**, 81, 509-13	2.7	25
101	The Trp23-Stop and Trp66-Gly mutations in the LDL receptor gene: common causes of familial hypercholesterolemia in Denmark. *Atherosclerosis*, **1996**, 120, 57-65	3.1	25
100	Protein expression studies of desmoplakin mutations in cardiomyopathy patients reveal different molecular disease mechanisms. *Clinical Genetics*, **2013**, 84, 20-30	4	24
99	Molecular autopsy in young sudden cardiac death victims with suspected cardiomyopathy. *Forensic Science International*, **2012**, 219, 33-8	2.6	24
98	Adverse effect of right ventricular pacing prevented by biventricular pacing during long-term follow-up: a randomized comparison. *European Journal of Echocardiography*, **2011**, 12, 767-72		24
97	Characterization of a disease-causing Glu119-Lys mutation in the low-density lipoprotein receptor gene in two Danish families with heterozygous familial hypercholesterolemia. *Human Mutation*, **1994**, 4, 102-13	4.7	24
96	Low disease prevalence and inappropriate implantable cardioverter defibrillator shock rate in Brugada syndrome: a nationwide study. *Europace*, **2012**, 14, 1025-9	3.9	23
95	Regional myocardial perfusion during chronic biventricular pacing and after acute change of the pacing mode in patients with congestive heart failure and bundle branch block treated with an atrioventricular sequential biventricular pacemaker. *European Journal of Heart Failure*, **2003**, 5, 179-86	12.3	23
94	Long QT interval in Turner syndrome--a high prevalence of LQTS gene mutations. *PLoS ONE*, **2013**, 8, e69614	3.7	23
93	Postmortem genetic testing of the ryanodine receptor 2 (RYR2) gene in a cohort of sudden unexplained death cases. *International Journal of Legal Medicine*, **2013**, 127, 139-44	3.1	22
92	Two mutations in the same low-density lipoprotein receptor allele act in synergy to reduce receptor function in heterozygous familial hypercholesterolemia. *Human Mutation*, **1997**, 9, 437-44	4.7	22
91	Sodium current and potassium transient outward current genes in Brugada syndrome: screening and bioinformatics. *Canadian Journal of Cardiology*, **2012**, 28, 196-200	3.8	19
90	Incidence of the apolipoprotein B-3500 mutation in Denmark. *Clinica Chimica Acta*, **1994**, 230, 101-4	6.2	18
89	Cardiac ryanodine receptor calcium release deficiency syndrome. *Science Translational Medicine*, **2021**, 13,	17.5	18
88	Prophylactic cavotricuspid isthmus block during atrial fibrillation ablation in patients without atrial flutter: a randomised controlled trial. *Heart*, **2009**, 95, 994-9	5.1	17
87	The superior transseptal surgical approach to mitral valve creates slow conduction. *PACE - Pacing and Clinical Electrophysiology*, **2006**, 29, 719-26	1.6	17
86	Long-term proarrhythmic pharmacotherapy among patients with congenital long QT syndrome and risk of arrhythmia and mortality. *European Heart Journal*, **2019**, 40, 3110-3117	9.5	16

85	Cascade screening in families with inherited cardiac diseases driven by cardiologists: feasibility and nationwide outcome in long QT syndrome. *Cardiology*, **2013**, 126, 131-7	1.6	16
84	Electrically vs. imaging-guided left ventricular lead placement in cardiac resynchronization therapy: a randomized controlled trial. *Europace*, **2019**, 21, 1369-1377	3.9	15
83	Multifocal atrial and ventricular premature contractions with an increased risk of dilated cardiomyopathy caused by a Na1.5 gain-of-function mutation (G213D). *International Journal of Cardiology*, **2018**, 257, 160-167	3.2	14
82	Heart transplantation in arrhythmogenic right ventricular cardiomyopathy - Experience from the Nordic ARVC Registry. *International Journal of Cardiology*, **2018**, 250, 201-206	3.2	14
81	Management of patients with Arrhythmogenic Right Ventricular Cardiomyopathy in the Nordic countries. *Scandinavian Cardiovascular Journal*, **2015**, 49, 299-307	2	14
80	Coronary artery disease-associated genetic variants and biomarkers of inflammation. *PLoS ONE*, **2017**, 12, e0180365	3.7	14
79	Adding the implantable cardioverter-defibrillator to cardiac resynchronization therapy is associated with improved long-term survival in ischaemic, but not in non-ischaemic cardiomyopathy. *Europace*, **2016**, 18, 413-9	3.9	13
78	Nationwide experience of catecholaminergic polymorphic ventricular tachycardia caused by RyR2 mutations. *Heart*, **2017**, 103, 901-909	5.1	13
77	A genetic risk score predicts cardiovascular events in patients with stable coronary artery disease. *International Journal of Cardiology*, **2017**, 241, 411-416	3.2	12
76	LDL-receptor gene mutations and the hypocholesterolemic response to statin therapy. *Clinical Genetics*, **2001**, 59, 397-405	4	12
75	Association of coronary heart disease with age-adjusted aortocoronary calcification in patients with familial hypercholesterolaemia. *Journal of Internal Medicine*, **2000**, 247, 479-84	10.8	12
74	P1A1/A2 polymorphism of platelet glycoprotein IIIa and risk of acute coronary syndromes in heterozygous familial hypercholesterolemia. *Atherosclerosis*, **1999**, 143, 99-104	3.1	12
73	Radiofrequency ablation lesions in low-, intermediate-, and normal-voltage myocardium: an in vivo study in a porcine heart model. *Europace*, **2019**, 21, 1919-1927	3.9	11
72	Mutations in Danish patients with long QT syndrome and the identification of a large founder family with p.F29L in KCNH2. *BMC Medical Genetics*, **2014**, 15, 31	2.1	11
71	Sudden cardiac death in young adults: environmental risk factors and genetic aspects of premature atherosclerosis. *Journal of Forensic Sciences*, **2012**, 57, 658-62	1.8	11
70	Flow cytometry with a monoclonal antibody to the low density lipoprotein receptor compared with gene mutation detection in diagnosis of heterozygous familial hypercholesterolemia. *Clinical Chemistry*, **1998**, 44, 966-972	5.5	11
69	Prevention of atrial flutter with cryoablation may be proarrhythmogenic. *Annals of Thoracic Surgery* , **2007**, 83, 1717-23	2.7	10
68	LDL receptor mutations and ApoB mutations are not risk factors for ischemic cerebrovascular disease of the young, but lipids and lipoproteins are. *European Journal of Neurology*, **1999**, 6, 691-6	6	10

67	Flow cytometric assessment of LDL receptor activity in peripheral blood mononuclear cells compared to gene mutation detection in diagnosis of heterozygous familial hypercholesterolemia. *Cytometry*, **1999**, 36, 52-9		10
66	Allele-specific measurement of low-density lipoprotein receptor transcript levels. *Human Mutation*, **1996**, 8, 126-33	4.7	10
65	Coronary Plaque Burden and Adverse Plaque Characteristics Are Increased in Healthy Relatives of Patients With Early‐Onset Coronary Artery Disease. *JACC: Cardiovascular Imaging*, **2017**, 10, 1128-1135	8.4	9
64	Genetic Risk of Coronary Artery Disease, Features of Atherosclerosis, and Coronary Plaque Burden. *Journal of the American Heart Association*, **2020**, 9, e014795	6	9
63	Long-Term Cardiovascular Risk in Heterozygous Familial Hypercholesterolemia Relatives Identified by Cascade Screening. *Journal of the American Heart Association*, **2017**, 6,	6	9
62	PatientsSattitudes toward detection of heterozygous familial hypercholesterolemia. *Archives of Internal Medicine*, **1997**, 157, 553-560		9
61	A 45-SNP genetic risk score is increased in early-onset coronary artery disease but independent of familial disease clustering. *Atherosclerosis*, **2017**, 257, 172-178	3.1	8
60	Outcome of clinical management in relatives of sudden cardiac death victims. *International Journal of Cardiology*, **2018**, 262, 45-50	3.2	8
59	Aetiologies and temporal trends of atrioventricular block in young patients: a 20-year nationwide study. *Europace*, **2019**, 21, 1710-1716	3.9	8
58	LDL receptor mutation genotype and vascular disease phenotype in heterozygous familial hypercholesterolaemia. *Clinical Genetics*, **2002**, 61, 408-15	4	8
57	Is the knowledge of contact force beneficial in pulmonary vein antrum isolation?. *Scandinavian Cardiovascular Journal*, **2017**, 51, 129-137	2	7
56	Identification of a novel mutation in exon 13 of the LDL receptor gene causing familial hypercholesterolemia in two Spanish families. *Biochimica Et Biophysica Acta - Molecular Basis of Disease*, **1996**, 1316, 1-4	6.9	7
55	Targeted molecular genetic testing in young sudden cardiac death victims from Western Denmark. *International Journal of Legal Medicine*, **2020**, 134, 111-121	3.1	7
54	The QTc interval and risk of cardiac events in bulimia nervosa: A long-term follow-up study. *International Journal of Eating Disorders*, **2018**, 51, 1331-1338	6.3	7
53	Incomplete thromboxane inhibition with 100 mg of intravenous acetylsalicylic acid in patients with acute ST elevation myocardial infarction: a placebo-controlled pilot trial. *Thrombosis Research*, **2001** , 104, 175-80	8.2	6
52	Long-term outcomes in young patients with atrioventricular block of unknown aetiology. *European Heart Journal*, **2021**, 42, 2060-2068	9.5	6
51	Primary Prevention of Sudden Cardiac Death With Implantable Cardioverter-Defibrillator Therapy in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy. *American Journal of Cardiology* , **2019**, 123, 1156-1162	3	5
50	Outcome after catheter ablation for left atrial flutter. *Scandinavian Cardiovascular Journal*, **2019**, 53, 133-140	2	5

49	A randomized trial of contact force in atrial flutter ablation. *Europace*, **2020**, 22, 947-955	3.9	5
48	Functional characterization of two low density lipoprotein receptor gene mutations by fluorescence flow cytometric assessment of receptor activity in stimulated human T-lymphocytes. *Clinical Genetics*, **2000**, 57, 110-5	4	5
47	Cardiovascular risk factor control is insufficient in young patients with coronary artery disease. *Vascular Health and Risk Management*, **2016**, 12, 219-27	4.4	5
46	Atrial fibrillation as a clinical characteristic of arrhythmogenic right ventricular cardiomyopathy: Experience from the Nordic ARVC Registry. *International Journal of Cardiology*, **2020**, 298, 39-43	3.2	5
45	The ABO locus is associated with increased platelet aggregation in patients with stable coronary artery disease. *International Journal of Cardiology*, **2019**, 286, 152-158	3.2	4
44	Flow cytometric assessment of effects of fluvastatin on low-density lipoprotein receptor activity in stimulated T-lymphocytes from patients with heterozygous familial hypercholesterolemia. *Journal of Clinical Pharmacology*, **2000**, 40, 421-9	2.9	4
43	Appropriate use of genetics in a young patient with atrioventricular block and family history of sudden cardiac death. *HeartRhythm Case Reports*, **2019**, 5, 169-172	1	3
42	Implantable cardioverter-defibrillator therapy and device-related complications in young patients with inherited cardiomyopathies or channelopathies: a 17-year cohort study. *Europace*, **2018**, 20, 1849-1855	3.9	3
41	The SH2B3 and KCNK5 loci may be implicated in regulation of platelet count, volume, and maturity. *Thrombosis Research*, **2017**, 158, 86-92	8.2	3
40	A frequent HhaI polymorphism in intron 9 of the low density lipoprotein receptor gene detected by the denaturing gradient gel electrophoresis technique. *Clinical Genetics*, **1995**, 48, 221-2	4	3
39	A robust method for quantification of IKr-related T-wave morphology abnormalities **2007**,		3
38	Flow Cytometric Assessment of LDL Ligand Function for Detection of Heterozygous Familial Defective Apolipoprotein B-100. *Clinical Chemistry*, **2000**, 46, 224-233	5.5	3
37	Linking genotype to aorto-coronary atherosclerosis: a model using familial hypercholesterolemia and aorto-coronary calcification. *Annals of Human Genetics*, **1999**, 63, 511-520	2.2	3
36	Comparison of the effects of xamoterol, atenolol and propranolol on breathlessness, fatigue and plasma electrolytes during exercise in healthy volunteers. *European Journal of Clinical Pharmacology*, **1991**, 41, 51-5	2.8	3
35	Pharmacokinetics and dynamic effects of diltiazem in the isolated guinea-pig heart. *Basic and Clinical Pharmacology and Toxicology*, **1988**, 62, 166-71		3
34	Clinical and Functional Characterization of Ryanodine Receptor 2 Variants Implicated in Calcium-Release Deficiency Syndrome. *JAMA Cardiology*, **2021**,	16.2	3
33	Incidence, Predictors, and Success of Ventricular Tachycardia Catheter Ablation in Arrhythmogenic Right Ventricular Cardiomyopathy (from the Nordic ARVC Registry). *American Journal of Cardiology*, **2020**, 125, 803-811	3	3
32	Pregnancies, ventricular arrhythmias, and substrate progression in women with arrhythmogenic right ventricular cardiomyopathy in the Nordic ARVC Registry. *Europace*, **2020**, 22, 1873-1879	3.9	3

31	Absence of ECG Task Force Criteria does not rule out structural changes in genotype positive ARVC patients. *International Journal of Cardiology*, **2020**, 317, 152-158	3.2	2
30	Voltage and pace-capture mapping of linear ablation lesions overestimates chronic ablation gap size. *Europace*, **2018**, 20, 2028-2035	3.9	2
29	Failure of ICD therapy in lethal arrhythmogenic right ventricular cardiomyopathy type 5 caused by the p.Ser358Leu mutation. *HeartRhythm Case Reports*, **2016**, 2, 217-222	1	2
28	Ablation for idiopathic ventricular arrhythmia - with and without arrhythmia-induced cardiomyopathy. *Scandinavian Cardiovascular Journal*, **2014**, 48, 130-7	2	2
27	Catheter ablation for ventricular tachycardia in ischaemic heart disease; acute success and long-term outcome. *Scandinavian Cardiovascular Journal*, **2014**, 48, 27-34	2	2
26	Intronic mutations at splice junctions in the low-density lipoprotein receptor gene. *Molecular and Cellular Probes*, **1999**, 13, 257-60	3.3	2
25	Polygenic Risk Score-Enhanced Risk Stratification of Coronary Artery Disease in Patients With Stable Chest Pain. *Circulation Genomic and Precision Medicine*, **2021**, 14, e003298	5.2	2
24	Discontinuation of oral anticoagulation and risk of stroke and death after ablation for typical atrial flutter: A nation-wide Danish cohort study. *International Journal of Cardiology*, **2021**, 333, 110-116	3.2	2
23	Complications of implantable cardioverter-defibrillator treatment in arrhythmogenic right ventricular cardiomyopathy. *Europace*, **2021**,	3.9	2
22	Severity of congenital long QT syndrome disease manifestation and risk of depression, anxiety, and mortality: a nationwide study. *Europace*, **2021**,	3.9	2
21	Recurrence after pulmonary vein isolation is associated with low contact force. *Scandinavian Cardiovascular Journal*, **2018**, 52, 28-33	2	2
20	Rapid Exclusion of COVID Infection With the Artificial Intelligence Electrocardiogram. *Mayo Clinic Proceedings*, **2021**, 96, 2081-2094	6.4	2
19	Human RyR2 (Ryanodine Receptor 2) Loss-of-Function Mutations: Clinical Phenotypes and In Vitro Characterization. *Circulation: Arrhythmia and Electrophysiology*, **2021**, 14, e010013	6.4	2
18	Phenotypic characterization of late onset cardiac amyloidosis caused by the transthyretin mutation TTRA45S, p.(Ala65Ser). *Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis*, **2017**, 24, 70-71	2.7	1
17	The ABO Locus is Associated with Increased Fibrin Network Formation in Patients with Stable Coronary Artery Disease. *Thrombosis and Haemostasis*, **2020**, 120, 1248-1256	7	1
16	Left ventricular regional remodeling and lead position during cardiac resynchronization therapy. *Heart Rhythm*, **2018**, 15, 1542-1549	6.7	1
15	Variable primary phenotypic manifestations in a rare familial form of Wolff-Parkinson-White syndrome and hypertrophic cardiomyopathy. *Europace*, **2016**, 18, 397	3.9	1
14	Cardiac magnetic resonance characteristics in young survivors of aborted sudden cardiac death. *European Journal of Radiology*, **2018**, 105, 141-147	4.7	1

13	Radiofrequency ablation of atrial fibrillation: effectiveness and safety in 102 consecutive patients. *Heart*, **2005**, 91, 1611-2	5.1	1
12	Long-term outcomes in a randomized controlled trial of multimodality imaging-guided left ventricular lead placement in cardiac resynchronization therapy.. *Europace*, **2022**,	3.9	1
11	Repeat pulmonary vein isolation in patients with atrial fibrillation: low ablation index is associated with increased risk of recurrent arrhythmia. *Scandinavian Cardiovascular Journal*, **2021**, 55, 29-34	2	1
10	Genotype-phenotype correlation in arrhythmogenic right ventricular cardiomyopathy-risk of arrhythmias and heart failure. *Journal of Medical Genetics*, **2021**,	5.8	1
9	Genetic Variant Score and Arrhythmogenic Right Ventricular Cardiomyopathy Phenotype in Plakophilin-2 Mutation Carriers. *Cardiology*, **2021**, 146, 763-771	1.6	1
8	Two mutations in the same low-density lipoprotein receptor allele act in synergy to reduce receptor function in heterozygous familial hypercholesterolemia **1997**, 9, 437		1
7	Identification of recurrent and novel mutations in the LDL receptor gene in Spanish patients with familial hypercholesterolemia **1998**, 11, 413		1
6	Recurrent atrial arrhythmia in a randomised controlled trial comparing contact force-guided and contact force-blinded ablation for typical atrial flutter.. *Journal of Interventional Cardiac Electrophysiology*, **2022**, 1	2.4	0
5	Aortic Dissections in the Population-Based Danish National Patient Registry from 1996-2016: A Validation Study.. *Clinical Epidemiology*, **2022**, 14, 51-58	5.9	0
4	Electrocardiographic Findings, Arrhythmias, and Left Ventricular Involvement in Familial ST-Depression Syndrome.. *Circulation: Arrhythmia and Electrophysiology*, **2022**, 101161CIRCEP121010688	6.4	0
3	Reply to: TFC ECG in arrhythmogenic cardiomyopathy: Inadequate mixture of criteria?. *International Journal of Cardiology*, **2021**, 323, 203	3.2	
2	Cardiac computed tomography-verified right ventricular lead position and outcomes in cardiac resynchronization therapy.. *Journal of Interventional Cardiac Electrophysiology*, **2022**, 1	2.4	
1	Diagnostic Yield of Genetic Testing in Young Patients With Atrioventricular Block of Unknown Cause.. *Journal of the American Heart Association*, **2022**, e025643	6	