Henning Bundgaard

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2664481/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	2014 ESC Guidelines on diagnosis and management of hypertrophic cardiomyopathy. European Heart Journal, 2014, 35, 2733-2779.	2.2	3,469
2	Partial Oral versus Intravenous Antibiotic Treatment of Endocarditis. New England Journal of Medicine, 2019, 380, 415-424.	27.0	502
3	Risk of COVID-19 in health-care workers in Denmark: an observational cohort study. Lancet Infectious Diseases, The, 2020, 20, 1401-1408.	9.1	357
4	Nationwide study of sudden cardiac death in persons aged 1–35 years. European Heart Journal, 2011, 32, 983-990.	2.2	303
5	Effectiveness of Adding a Mask Recommendation to Other Public Health Measures to Prevent SARS-CoV-2 Infection in Danish Mask Wearers. Annals of Internal Medicine, 2021, 174, 335-343.	3.9	279
6	Risk of post-pregnancy hypertension in women with a history of hypertensive disorders of pregnancy: nationwide cohort study. BMJ: British Medical Journal, 2017, 358, j3078.	2.3	195
7	Simvastatin Effects on Skeletal Muscle. Journal of the American College of Cardiology, 2013, 61, 44-53.	2.8	156
8	Long-term clinical outcome after alcohol septal ablation for obstructive hypertrophic cardiomyopathy: results from the Euro-ASA registry. European Heart Journal, 2016, 37, 1517-1523.	2.2	148
9	Cardiac manifestations of myotonic dystrophy type 1. International Journal of Cardiology, 2012, 160, 82-88.	1.7	146
10	Burden of Sudden Cardiac Death in Persons Aged 1 to 49 Years. Circulation: Arrhythmia and Electrophysiology, 2014, 7, 205-211.	4.8	142
11	Echocardiographic Strain Imaging to Assess Early and Late Consequences of Sarcomere Mutations in Hypertrophic Cardiomyopathy. Circulation: Cardiovascular Genetics, 2009, 2, 314-321.	5.1	140
12	Sudden cardiac death in children (1-18 years): symptoms and causes of death in a nationwide setting. European Heart Journal, 2014, 35, 868-875.	2.2	134
13	The Na, K-ATPase in the failing human heart. Cardiovascular Research, 2003, 57, 913-920.	3.8	130
14	Penetrance of Hypertrophic Cardiomyopathy in Children and Adolescents. Circulation, 2013, 127, 48-54.	1.6	121
15	Modeling of waning immunity after SARS-CoV-2 vaccination and influencing factors. Nature Communications, 2022, 13, 1614.	12.8	117
16	Diagnostic yield, interpretation, and clinical utility of mutation screening of sarcomere encoding genes in Danish hypertrophic cardiomyopathy patients and relatives. Human Mutation, 2009, 30, 363-370.	2.5	110
17	Association Between Hypertensive Disorders of Pregnancy and Later Risk of Cardiomyopathy. JAMA - Journal of the American Medical Association, 2016, 315, 1026.	7.4	106
18	Efficacy and safety of the angiotensin II receptor blocker losartan for hypertrophic cardiomyopathy: the INHERIT randomised, double-blind, placebo-controlled trial. Lancet Diabetes and Endocrinology,the, 2015, 3, 123-131.	11.4	104

#	Article	IF	CITATIONS
19	Risk of arrhythmia induced by psychotropic medications: a proposal for clinical management. European Heart Journal, 2014, 35, 1306-1315.	2.2	103
20	Long COVID symptoms in SARS-CoV-2-positive adolescents and matched controls (LongCOVIDKidsDK): a national, cross-sectional study. The Lancet Child and Adolescent Health, 2022, 6, 240-248.	5.6	93
21	<i>Enterococcus faecalis</i> Infective Endocarditis. Circulation, 2013, 127, 1810-1817.	1.6	92
22	Incidence of infective endocarditis among patients considered at high risk. European Heart Journal, 2018, 39, 623-629.	2.2	89
23	Long-Term Outcome of Percutaneous Transluminal Septal Myocardial Ablation in Hypertrophic Obstructive Cardiomyopathy. Circulation: Cardiovascular Interventions, 2011, 4, 256-265.	3.9	88
24	Risk prediction of ventricular arrhythmias and myocardial function in Lamin A/C mutation positive subjects. Europace, 2014, 16, 563-571.	1.7	88
25	Prosthetic Valve Endocarditis After Transcatheter Aortic Valve Implantation. Circulation: Cardiovascular Interventions, 2015, 8, .	3.9	88
26	Loss-of-activity-mutation in the cardiac chloride-bicarbonate exchanger AE3 causes short QT syndrome. Nature Communications, 2017, 8, 1696.	12.8	88
27	Alcohol septal ablation in patients with hypertrophic obstructive cardiomyopathy: low incidence of sudden cardiac death and reduced risk profile. Heart, 2013, 99, 1012-1017.	2.9	87
28	Pregnancy Loss and Later Risk of Atherosclerotic Disease. Circulation, 2013, 127, 1775-1782.	1.6	86
29	The role of sarcomere gene mutations in patients with idiopathic dilated cardiomyopathy. European Journal of Human Genetics, 2009, 17, 1241-1249.	2.8	79
30	β ₃ Adrenergic Stimulation of the Cardiac Na ⁺ -K ⁺ Pump by Reversal of an Inhibitory Oxidative Modification. Circulation, 2010, 122, 2699-2708.	1.6	79
31	Electrocardiographic Features of Sarcomere Mutation Carriers With and Without Clinically Overt Hypertrophic Cardiomyopathy. American Journal of Cardiology, 2011, 108, 1606-1613.	1.6	77
32	Echocardiographic abnormalities and predictors of mortality in hospitalized COVIDâ€19 patients: the ECHOVIDâ€19 study. ESC Heart Failure, 2020, 7, 4189-4197.	3.1	77
33	Fabry disease mimicking hypertrophic cardiomyopathy: genetic screening needed for establishing the diagnosis in women. European Journal of Heart Failure, 2010, 12, 535-540.	7.1	75
34	Association Between Fetal Congenital Heart Defects and Maternal Risk of Hypertensive Disorders of Pregnancy in the Same Pregnancy and Across Pregnancies. Circulation, 2017, 136, 39-48.	1.6	73
35	Prevalence and Progression of Late Gadolinium Enhancement in Children and Adolescents With Hypertrophic Cardiomyopathy. Circulation, 2018, 138, 782-792.	1.6	72
36	Hyperaldosteronemia in Rabbits Inhibits the Cardiac Sarcolemmal Na ⁺ -K ⁺ Pump. Circulation Research, 2000, 86, 37-42.	4.5	71

#	Article	IF	CITATIONS
37	The diagnostic performance of imaging methods in ARVC using the 2010 Task Force criteria. European Heart Journal Cardiovascular Imaging, 2014, 15, 1219-1225.	1.2	70
38	Diagnostic yield of molecular autopsy in patients with sudden arrhythmic death syndrome using targeted exome sequencing. Europace, 2016, 18, 888-896.	1.7	69
39	Genetic investigation of 100 heart genes in sudden unexplained death victims in a forensic setting. European Journal of Human Genetics, 2016, 24, 1797-1802.	2.8	65
40	Genetic and phenotypic characterization of mutations in myosin-binding protein C (MYBPC3) in 81 families with familial hypertrophic cardiomyopathy: total or partial haploinsufficiency. European Journal of Human Genetics, 2004, 12, 673-677.	2.8	64
41	Anoctamin 5 muscular dystrophy in Denmark: prevalence, genotypes, phenotypes, cardiac findings, and muscle protein expression. Journal of Neurology, 2013, 260, 2084-2093.	3.6	63
42	Prevalence of Bicuspid Aortic Valve and Associated Aortopathy in Newborns in Copenhagen, Denmark. JAMA - Journal of the American Medical Association, 2021, 325, 561.	7.4	62
43	Endotoxemia stimulates skeletal muscle Na ⁺ -K ⁺ -ATPase and raises blood lactate under aerobic conditions in humans. American Journal of Physiology - Heart and Circulatory Physiology, 2003, 284, H1028-H1034.	3.2	61
44	Long-Term Adverse Cardiac Outcomes in Patients With Sarcoidosis. Journal of the American College of Cardiology, 2020, 76, 767-777.	2.8	61
45	Genetic variability in the absorption of dietary sterols affects the risk of coronary artery disease. European Heart Journal, 2020, 41, 2618-2628.	2.2	61
46	Wide spectrum of desmosomal mutations in Danish patients with arrhythmogenic right ventricular cardiomyopathy. Journal of Medical Genetics, 2010, 47, 736-744.	3.2	58
47	Cardiac involvement in myotonic dystrophy: a nationwide cohort study. European Heart Journal, 2014, 35, 2158-2164.	2.2	56
48	Incidence and factors associated with infective endocarditis in patients undergoing left-sided heart valve replacement. European Heart Journal, 2018, 39, 2668-2675.	2.2	55
49	Risk of cardiovascular disease in family members of young sudden cardiac death victims. European Heart Journal, 2013, 34, 503-511.	2.2	54
50	Temporal Release of High-Sensitivity Cardiac Troponin T and I and Copeptin After Brief Induced Coronary Artery Balloon Occlusion in Humans. Circulation, 2021, 143, 1095-1104.	1.6	54
51	The firstâ€inâ€man randomized trial of a beta3 adrenoceptor agonist in chronic heart failure: the <scp>BEATâ€HF</scp> trial. European Journal of Heart Failure, 2017, 19, 566-575.	7.1	53
52	Infective endocarditis following percutaneous pulmonary valve replacement: Diagnostic challenges and application of intra-cardiac echocardiography. International Journal of Cardiology, 2013, 169, 425-429.	1.7	52
53	Outcomes of Alcohol Septal Ablation inÂYoungerÂPatients With Obstructive HypertrophicÂCardiomyopathy. JACC: Cardiovascular Interventions, 2017, 10, 1134-1143.	2.9	52
54	Long-Term Outcomes of Partial Oral Treatment of Endocarditis. New England Journal of Medicine, 2019, 380, 1373-1374.	27.0	51

#	Article	IF	CITATIONS
55	Valsartan in early-stage hypertrophic cardiomyopathy: a randomized phase 2 trial. Nature Medicine, 2021, 27, 1818-1824.	30.7	51
56	Next-generation sequencing of 34 genes in sudden unexplained death victims in forensics and in patients with channelopathic cardiac diseases. International Journal of Legal Medicine, 2015, 129, 793-800.	2.2	49
57	Next-generation sequencing of 100 candidate genes in young victims of suspected sudden cardiac death with structural abnormalities of the heart. International Journal of Legal Medicine, 2016, 130, 91-102.	2.2	47
58	Clinical and genetic characteristics of cardiac actin gene mutations in hypertrophic cardiomyopathy. Journal of Medical Genetics, 2004, 41, 10e-10.	3.2	46
59	The nitric oxide donor sodium nitroprusside stimulates the Na+-K+pump in isolated rabbit cardiac myocytes. Journal of Physiology, 2005, 565, 815-825.	2.9	46
60	Effect of Institutional Experience on Outcomes of Alcohol Septal Ablation for Hypertrophic Obstructive Cardiomyopathy. Canadian Journal of Cardiology, 2018, 34, 16-22.	1.7	45
61	High prevalence of cardiac involvement in patients with myotonic dystrophy type 1: A cross-sectional study. International Journal of Cardiology, 2014, 174, 31-36.	1.7	44
62	Gender differences in sudden cardiac death in the young-a nationwide study. BMC Cardiovascular Disorders, 2017, 17, 19.	1.7	44
63	Hyperbaric oxygen therapy augments tobramycin efficacy in experimental Staphylococcus aureus endocarditis. International Journal of Antimicrobial Agents, 2017, 50, 406-412.	2.5	44
64	Myocardial fibrosis in patients with myotonic dystrophy type 1: a cardiovascular magnetic resonance study. Journal of Cardiovascular Magnetic Resonance, 2014, 16, 59.	3.3	43
65	Predictors of coronary in-stent restenosis: importance of angiotensin-converting enzyme gene polymorphism and treatment with angiotensin-converting enzyme inhibitors. Journal of the American College of Cardiology, 2001, 38, 1434-1439.	2.8	42
66	One third of Danish hypertrophic cardiomyopathy patients have mutations in MYH7 rod region. European Journal of Human Genetics, 2005, 13, 161-165.	2.8	42
67	Prevalence of infective endocarditis in patients with positive blood cultures: a Danish nationwide study. European Heart Journal, 2019, 40, 3237-3244.	2.2	40
68	Recovery of cardiac function following <scp>COVID</scp> â€19–Â <scp>ECHOVID</scp> â€19: a prospective longitudinal cohort study. European Journal of Heart Failure, 2021, 23, 1903-1912.	7.1	40
69	Outcome of clinical versus genetic family screening in hypertrophic cardiomyopathy with focus on cardiac β-myosin gene mutations. Cardiovascular Research, 2003, 57, 347-357.	3.8	38
70	Mitochondrial Haplogroups Modify the Risk of Developing Hypertrophic Cardiomyopathy in a Danish Population. PLoS ONE, 2013, 8, e71904.	2.5	38
71	Partial oral treatment of endocarditis. American Heart Journal, 2013, 165, 116-122.	2.7	37
72	Human myocardial Na,K-ATPase concentration in heart failure. Molecular and Cellular Biochemistry, 1996, 163-164, 277-283.	3.1	36

#	Article	IF	CITATIONS
73	Natriuretic peptides stimulate the cardiac sodium pump via NPR-C-coupled NOS activation. American Journal of Physiology - Cell Physiology, 2008, 294, C1067-C1073.	4.6	36
74	Progression of cardiac involvement in patients with limb-girdle type 2 and Becker muscular dystrophies: A 9-year follow-up study. International Journal of Cardiology, 2015, 182, 403-411.	1.7	36
75	Severe intracellular magnesium and potassium depletion in patients after treatment with cisplatin. British Journal of Cancer, 2003, 89, 1633-1637.	6.4	35
76	Temporal changes in the incidence of infective endocarditis in Denmark 1997–2017: A nationwide study. International Journal of Cardiology, 2021, 326, 145-152.	1.7	35
77	Short- and long-term outcomes of alcohol septal ablation for hypertrophic obstructive cardiomyopathy in patients with mild left ventricular hypertrophy: a propensity score matching analysis. European Heart Journal, 2019, 40, 1681-1687.	2.2	33
78	Hypertensive disorders of pregnancy and peripartum cardiomyopathy: A nationwide cohort study. PLoS ONE, 2019, 14, e0211857.	2.5	33
79	Humoral response to two doses of BNT162b2 vaccination in people with HIV. Journal of Internal Medicine, 2022, 291, 513-518.	6.0	33
80	Copenhagen Baby Heart Study: a population study of newborns with prenatal inclusion. European Journal of Epidemiology, 2019, 34, 79-90.	5.7	32
81	Anti-biofilm Approach in Infective Endocarditis Exposes New Treatment Strategies for Improved Outcome. Frontiers in Cell and Developmental Biology, 2021, 9, 643335.	3.7	32
82	Multifocal atrial and ventricular premature contractions with an increased risk of dilated cardiomyopathy caused by a Na v 1.5 gain-of-function mutation (G213D). International Journal of Cardiology, 2018, 257, 160-167.	1.7	31
83	Intrinsic mitral valve alterations in hypertrophic cardiomyopathy sarcomere mutation carriers. European Heart Journal Cardiovascular Imaging, 2018, 19, 1109-1116.	1.2	31
84	The val606met mutation in the cardiac beta-myosin heavy chain gene in patients with familial hypertrophic cardiomyopathy is associated with a high risk of sudden death at young age. American Journal of Cardiology, 2001, 87, 1315-1317.	1.6	30
85	Genome-wide association study identifies 18 novel loci associated with left atrial volume and function. European Heart Journal, 2021, 42, 4523-4534.	2.2	30
86	Analyses of more than 60,000 exomes questions the role of numerous genes previously associated with dilated cardiomyopathy. Molecular Genetics & amp; Genomic Medicine, 2016, 4, 617-623.	1.2	29
87	Functional effects of losartan in hypertrophic cardiomyopathy—a randomised clinical trial. Heart, 2016, 102, 285-291.	2.9	29
88	Outcome of Alcohol Septal Ablation in Mildly Symptomatic Patients With Hypertrophic Obstructive Cardiomyopathy: A Longâ€Term Followâ€Up Study Based on the Euroâ€Alcohol Septal Ablation Registry. Journal of the American Heart Association, 2017, 6, .	3.7	29
89	Incidence of infective endocarditis in patients considered at moderate risk. European Heart Journal, 2019, 40, 1355-1361.	2.2	29
90	<i>MT YB</i> mutations in hypertrophic cardiomyopathy. Molecular Genetics & Genomic Medicine, 2013, 1, 54-65.	1.2	28

#	Article	IF	CITATIONS
91	Validation of the HCM Risk-SCD model in patients with hypertrophic cardiomyopathy following alcohol septal ablation. Europace, 2018, 20, f198-f203.	1.7	28
92	Long-term proarrhythmic pharmacotherapy among patients with congenital long QT syndrome and risk of arrhythmia and mortality. European Heart Journal, 2019, 40, 3110-3117.	2.2	28
93	Comparison of Valsalva manoeuvre and exercise in echocardiographic evaluation of left ventricular outflow tract obstruction in hypertrophic cardiomyopathy. European Journal of Echocardiography, 2010, 11, 763-769.	2.3	27
94	Low disease prevalence and inappropriate implantable cardioverter defibrillator shock rate in Brugada syndrome: a nationwide study. Europace, 2012, 14, 1025-1029.	1.7	27
95	Genetic insight into sick sinus syndrome. European Heart Journal, 2021, 42, 1959-1971.	2.2	27
96	The role of Lamin A/C mutations in Danish patients with idiopathic dilated cardiomyopathy. European Journal of Heart Failure, 2009, 11, 1031-1035.	7.1	26
97	K+ supplementation increases muscle [Na+-K+-ATPase] and improves extrarenal K+homeostasis in rats. Journal of Applied Physiology, 1997, 82, 1136-1144.	2.5	25
98	Q Fever in Greenland. Emerging Infectious Diseases, 2010, 16, 511-513.	4.3	25
99	Cardiac symptoms before sudden cardiac death caused by coronary artery disease: a nationwide study among young Danish people. Heart, 2013, 99, 938-943.	2.9	25
100	Heart transplantation in arrhythmogenic right ventricular cardiomyopathy — Experience from the Nordic ARVC Registry. International Journal of Cardiology, 2018, 250, 201-206.	1.7	25
101	Aetiologies and temporal trends of atrioventricular block in young patients: a 20-year nationwide study. Europace, 2019, 21, 1710-1716.	1.7	25
102	The use of beta3-adrenergic receptor agonists in the treatment of heart failure. Current Opinion in Investigational Drugs, 2009, 10, 955-62.	2.3	25
103	Redox-dependent regulation of the Na+–K+ pump: New twists to an old target for treatment of heart failure. Journal of Molecular and Cellular Cardiology, 2013, 61, 94-101.	1.9	24
104	Long-term outcomes in young patients with atrioventricular block of unknown aetiology. European Heart Journal, 2021, 42, 2060-2068.	2.2	24
105	Five-Year Outcomes of the Partial Oral Treatment of Endocarditis (POET) Trial. New England Journal of Medicine, 2022, 386, 601-602.	27.0	24
106	Ocular, bulbar, limb, and cardiopulmonary involvement in oculopharyngeal muscular dystrophy. Acta Neurologica Scandinavica, 2014, 130, 125-130.	2.1	23
107	A Detailed Family History of Myocardial Infarction and Risk of Myocardial Infarction – A Nationwide Cohort Study. PLoS ONE, 2015, 10, e0125896.	2.5	23
108	Data Resource Profile: The Copenhagen Hospital Biobank (CHB). International Journal of Epidemiology, 2021, 50, 719-720e.	1.9	23

#	Article	IF	CITATIONS
109	Decline in Antibody Concentration 6 Months After Two Doses of SARS-CoV-2 BNT162b2 Vaccine in Solid Organ Transplant Recipients and Healthy Controls. Frontiers in Immunology, 2022, 13, 832501.	4.8	23
110	Effect of Nitroglycerin in Patients with Increased Pulmonary Vascular Resistance Undergoing Cardiac Transplantation. Scandinavian Cardiovascular Journal, 1997, 31, 339-342.	1.2	22
111	Sick Sinus Syndrome, Progressive Cardiac Conduction Disease, Atrial Flutter and Ventricular Tachycardia Caused by a Novel <i>SCN5A</i> Mutation. Cardiology, 2010, 115, 311-316.	1.4	21
112	Infective endocarditis following transcatheter aortic valve replacement—: Diagnostic and management challenges. Catheterization and Cardiovascular Interventions, 2013, 81, 623-627.	1.7	21
113	Clinical utility of 18F-FDG positron emission tomography/computed tomography scan vs. 99mTc-HMPAO white blood cell single-photon emission computed tomography in extra-cardiac work-up of infective endocarditis. International Journal of Cardiovascular Imaging, 2017, 33, 751-760.	1.5	21
114	Self-Collected versus Healthcare Worker-Collected Swabs in the Diagnosis of Severe Acute Respiratory Syndrome Coronavirus 2. Diagnostics, 2020, 10, 678.	2.6	21
115	β-Adrenergic Regulation of the Cardiac Na+-K+ ATPase Mediated by Oxidative Signaling. Trends in Cardiovascular Medicine, 2012, 22, 83-87.	4.9	20
116	Nationwide experience of catecholaminergic polymorphic ventricular tachycardia caused by RyR2 mutations. Heart, 2017, 103, 901-909.	2.9	20
117	Functional Effects of Receptor-Binding Domain Mutations of SARS-CoV-2 B.1.351 and P.1 Variants. Frontiers in Immunology, 2021, 12, 757197.	4.8	20
118	Lipoprotein(a) Levels at Birth and in Early Childhood: The COMPARE Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, 324-335.	3.6	20
119	Potassium depletion increases potassium clearance capacity in skeletal muscles in vivo during acute repletion. American Journal of Physiology - Cell Physiology, 2002, 283, C1163-C1170.	4.6	19
120	Risk of Cardiomyopathy in Younger Persons With a Family History of Death from Cardiomyopathy. Circulation, 2015, 132, 1013-1019.	1.6	19
121	High readmission rates and mental distress after infective endocarditis — Results from the national population-based CopenHeart IE survey. International Journal of Cardiology, 2017, 235, 133-140.	1.7	19
122	The D313Y variant in the <i>GLA</i> gene – no evidence of a pathogenic role in Fabry disease. Scandinavian Journal of Clinical and Laboratory Investigation, 2017, 77, 617-621.	1.2	19
123	Strategy for clinical evaluation and screening of sudden cardiac death relatives. Fundamental and Clinical Pharmacology, 2010, 24, 619-635.	1.9	18
124	Linezolid as rescue treatment for left-sided infective endocarditis: an observational, retrospective, multicenter study. European Journal of Clinical Microbiology and Infectious Diseases, 2012, 31, 2567-2574.	2.9	18
125	Management of patients with Arrhythmogenic Right Ventricular Cardiomyopathy in the Nordic countries. Scandinavian Cardiovascular Journal, 2015, 49, 299-307.	1.2	18
126	Adjunctive dabigatran therapy improves outcome of experimental left-sided Staphylococcus aureus endocarditis. PLoS ONE, 2019, 14, e0215333.	2.5	18

#	Article	IF	CITATIONS
127	Diagnostic yield in victims of sudden cardiac death and their relatives. Europace, 2020, 22, 964-971.	1.7	18
128	Cascade Screening in Families with Inherited Cardiac Diseases Driven by Cardiologists: Feasibility and Nationwide Outcome in Long QT Syndrome. Cardiology, 2013, 126, 131-137.	1.4	17
129	A comparison of genetic findings in sudden cardiac death victims and cardiac patients: the importance of phenotypic classification. Europace, 2015, 17, 350-357.	1.7	17
130	A Novel Familial Cardiac Arrhythmia Syndrome with Widespread ST-Segment Depression. New England Journal of Medicine, 2018, 379, 1780-1781.	27.0	17
131	Differences in mortality in patients undergoing surgery for infective endocarditis according to age and valvular surgery. BMC Infectious Diseases, 2020, 20, 705.	2.9	17
132	Antibodyâ€dependent neutralizing capacity of the SARS oVâ€2 vaccine BNT162b2 with and without previous COVIDâ€19 priming. Journal of Internal Medicine, 2021, 290, 1272-1274.	6.0	17
133	Familial Hypertrophic Cardiomyopathy Associated with a Novel Missense Mutation Affecting the ATP-binding Region of the Cardiac Beta-myosin Heavy Chain. Journal of Molecular and Cellular Cardiology, 1999, 31, 745-750.	1.9	16
134	Echocardiographic Findings Suggestive of Infective Endocarditis in Asymptomatic Danish Injection Drug Users Attending Urban Injection Facilities. American Journal of Cardiology, 2014, 114, 100-104.	1.6	16
135	Human genetic variation in GLS2 is associated with development of complicated Staphylococcus aureus bacteremia. PLoS Genetics, 2018, 14, e1007667.	3.5	16
136	Is the alphaâ€galactosidase A variant <scp>p.Asp313Tyr</scp> (<scp>p.D313Y</scp>) pathogenic for Fabry disease? A systematic review. Journal of Inherited Metabolic Disease, 2020, 43, 922-933.	3.6	16
137	Cardiac Myotonic Dystrophy Mimicking Arrhythmogenic Right Ventricular Cardiomyopathy in a Young Sudden Cardiac Death Victim. Circulation: Arrhythmia and Electrophysiology, 2008, 1, 317-320.	4.8	15
138	Deep sedation as temporary bridge to definitive treatment of ventricular arrhythmia storm. European Heart Journal: Acute Cardiovascular Care, 2020, 9, 657-664.	1.0	15
139	Cohort Profile: The Copenhagen Baby Heart Study (CBHS). International Journal of Epidemiology, 2022, 50, 1778-1779m.	1.9	15
140	Testing Denmark: a Danish Nationwide Surveillance Study of COVID-19. Microbiology Spectrum, 2021, 9, e0133021.	3.0	15
141	Myocardial Na,Kâ€ATPase and Digoxin Therapy in Human Heart Failure. Annals of the New York Academy of Sciences, 2003, 986, 702-707.	3.8	14
142	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	14
143	Echocardiographic and clinical findings in patients with Fabry disease during long-term enzyme replacement therapy: a nationwide Danish cohort study. Scandinavian Cardiovascular Journal, 2017, 51, 207-216.	1.2	14
144	Temporal changes in infective endocarditis guidelines during the last 12 years: High-level evidence needed. American Heart Journal, 2017, 193, 70-75.	2.7	14

#	Article	IF	CITATIONS
145	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.	1.7	14
146	Dilated cardiomyopathy caused by truncating titin variants: long-term outcomes, arrhythmias, response to treatment and sex differences. Journal of Medical Genetics, 2021, 58, 832-841.	3.2	14
147	Influence of Septal Thickness on the Clinical Outcome After Alcohol Septal Alation in Hypertrophic Cardiomyopathy. Circulation: Cardiovascular Interventions, 2016, 9, .	3.9	13
148	Repeatability and Reproducibility of Neonatal Echocardiography: The Copenhagen Baby Heart Study. Journal of the American Society of Echocardiography, 2019, 32, 895-905.e2.	2.8	13
149	Genotype–phenotype correlation in arrhythmogenic right ventricular cardiomyopathy—risk of arrhythmias and heart failure. Journal of Medical Genetics, 2022, 59, 858-864.	3.2	13
150	Potassium depletion improves myocardial potassium uptake in vivo. American Journal of Physiology - Cell Physiology, 2004, 287, C135-C141.	4.6	12
151	Survival and sudden cardiac death after septal ablation for hypertrophic obstructive cardiomyopathy. Scandinavian Cardiovascular Journal, 2011, 45, 153-160.	1.2	12
152	Low efficacy of tobramycin in experimental Staphylococcus aureus endocarditis. European Journal of Clinical Microbiology and Infectious Diseases, 2015, 34, 2349-2357.	2.9	12
153	Association between pregnancy losses in women and risk of atherosclerotic disease in their relatives: a nationwide cohort study. European Heart Journal, 2016, 37, 900-907.	2.2	12
154	Cardiac symptoms before sudden cardiac death caused by hypertrophic cardiomyopathy: a nationwide study among the young in Denmark. Europace, 2016, 18, euv403.	1.7	12
155	Infective endocarditis in patients who have undergone transcatheter aortic valve implantation: a review. Clinical Microbiology and Infection, 2020, 26, 999-1007.	6.0	12
156	Complications of implantable cardioverter-defibrillator treatment in arrhythmogenic right ventricular cardiomyopathy. Europace, 2022, 24, 306-312.	1.7	12
157	Non-diagnostic autopsy findings in sudden unexplained death victims. BMC Cardiovascular Disorders, 2020, 20, 58.	1.7	12
158	Seroprevalence of SARS-CoV-2 antibodies in social housing areas in Denmark. BMC Infectious Diseases, 2022, 22, 143.	2.9	12
159	Temporal trends of mortality in patients with infective endocarditis: a nationwide study. European Heart Journal Quality of Care & Clinical Outcomes, 2022, 9, 24-33.	4.0	12
160	Detection of microbial diversity in endocarditis using cultivation-independent molecular techniques. Scandinavian Journal of Infectious Diseases, 2011, 43, 857-869.	1.5	11
161	A randomised clinical trial of comprehensive cardiac rehabilitation versus usual care for patients treated for infective endocarditis—the CopenHeartIEtrial protocol. BMJ Open, 2012, 2, e001929.	1.9	11
162	Private Mitochondrial DNA Variants in Danish Patients with Hypertrophic Cardiomyopathy. PLoS ONE, 2015, 10, e0124540.	2.5	11

#	Article	IF	CITATIONS
163	Long QT syndrome is associated with an increased burden of diabetes, psychiatric and neurological comorbidities: a nationwide cohort study. Open Heart, 2019, 6, e001161.	2.3	11
164	Alcohol septal ablation in patients with severe septal hypertrophy. Heart, 2020, 106, 462-466.	2.9	11
165	Risk Factors for Being Seronegative following SARS-CoV-2 Infection in a Large Cohort of Health Care Workers in Denmark. Microbiology Spectrum, 2021, 9, e0090421.	3.0	11
166	Differences in investigations of sudden unexpected deaths in young people in a nationwide setting. International Journal of Legal Medicine, 2012, 126, 223-229.	2.2	10
167	Atrioventricular conduction after alcohol septal ablation for obstructive hypertrophic cardiomyopathy. Journal of Cardiovascular Medicine, 2014, 15, 214-221.	1.5	10
168	Low procedure-related mortality achieved with alcohol septal ablation in European patients. International Journal of Cardiology, 2016, 209, 194-195.	1.7	10
169	CIED infection with either pocket or systemic infection presentation – complete device removal and long-term antibiotic treatment; long-term outcome. Scandinavian Cardiovascular Journal, 2016, 50, 52-57.	1.2	10
170	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.	1.6	10
171	Congenital myopathies are mainly associated with a mild cardiac phenotype. Journal of Neurology, 2019, 266, 1367-1375.	3.6	10
172	Baseline Characteristics of the VANISH Cohort. Circulation: Heart Failure, 2019, 12, e006231.	3.9	10
173	Ticagrelor and the risk of <i>Staphylococcus aureus</i> bacteraemia and other infections. European Heart Journal - Cardiovascular Pharmacotherapy, 2022, 8, 13-19.	3.0	10
174	Pregnancies, ventricular arrhythmias, and substrate progression in women with arrhythmogenic right ventricular cardiomyopathy in the Nordic ARVC Registry. Europace, 2020, 22, 1873-1879.	1.7	10
175	Prevention of sudden cardiac death in hypertrophic cardiomyopathy: Risk assessment using left atrial diameter predicted from left atrial volume. Clinical Cardiology, 2020, 43, 581-586.	1.8	10
176	Heart Rate Recovery After Exercise Is Associated With Arrhythmic Events in Patients With Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation: Arrhythmia and Electrophysiology, 2020, 13, e007471.	4.8	10
177	Clinical usefulness of FDG-PET/CT for identification of abnormal extra-cardiac foci in patients with infective endocarditis. International Journal of Cardiovascular Imaging, 2020, 36, 939-946.	1.5	10
178	Association of Variants Near the Bradykinin Receptor B2 Gene With Angioedema in Patients Taking ACEÂInhibitors. Journal of the American College of Cardiology, 2021, 78, 696-709.	2.8	10
179	Myocardial Impairment and AcuteÂRespiratory Distress Syndrome inÂHospitalized Patients With COVID-19. JACC: Cardiovascular Imaging, 2020, 13, 2474-2476.	5.3	10
180	SARS-CoV-2 antibody prevalence among homeless people and shelter workers in Denmark: a nationwide cross-sectional study. BMC Public Health, 2022, 22, .	2.9	10

#	Article	IF	CITATIONS
181	β3 -Adrenoceptor agonist stimulation of the Na+ ,K+ -pump in rat skeletal muscle is mediated by β2 - rather than β3 -adrenoceptors. British Journal of Pharmacology, 2006, 149, 635-646.	5.4	9
182	The KCNE genes in hypertrophic cardiomyopathy: a candidate gene study. Journal of Negative Results in BioMedicine, 2011, 10, 12.	1.4	9
183	Cardiomyopathy in Friedreich Ataxia. Circulation, 2012, 125, 1591-1593.	1.6	9
184	Reappraisal of variants previously linked with sudden infant death syndrome: results from three population-based cohorts. European Journal of Human Genetics, 2019, 27, 1427-1435.	2.8	9
185	Nursing Home Admission and Initiation of Domiciliary Care Following Infective Endocarditis. Global Heart, 2020, 14, 41.	2.3	9
186	Long-term outcome of repeated septal reduction therapy after alcohol septal ablation for hypertrophic obstructive cardiomyopathy: insight from the Euro-ASA registry. Archives of Medical Science, 2020, 16, 1239-1242.	0.9	9
187	Dabigatran and the Risk of <i>Staphylococcus aureus</i> Bacteremia: A Nationwide Cohort Study. Clinical Infectious Diseases, 2021, 73, 480-486.	5.8	9
188	Recurrent infective endocarditis versus first-time infective endocarditis after heart valve surgery. Clinical Research in Cardiology, 2020, 109, 1342-1351.	3.3	9
189	Defining the normal QT interval in newborns: the natural history and reference values for the first 4 weeks of life. Europace, 2021, 23, 278-286.	1.7	9
190	Alcohol dose in septal ablation for hypertrophic obstructive cardiomyopathy. International Journal of Cardiology, 2021, 333, 127-132.	1.7	9
191	Novel human <i>inÂvitro</i> vegetation simulation model for infective endocarditis. Apmis, 2021, 129, 653-662.	2.0	9
192	Seroprevalence of SARS-CoV-2 antibodies and reduced risk of reinfection through 6Âmonths: a Danish observational cohort study of 44Â000 healthcare workers. Clinical Microbiology and Infection, 2022, 28, 710-717.	6.0	9
193	Chronic K-supplementation Decreases Myocardial [Na,K-ATPase] and Net K-uptake Capacity in Rodents. Journal of Molecular and Cellular Cardiology, 1998, 30, 2037-2046.	1.9	8
194	Cardiac implantable electronic device and associated risk of infective endocarditis in patients undergoing aortic valve replacement. Europace, 2018, 20, e164-e170.	1.7	8
195	Long-term Risk of Hemorrhagic Stroke in Patients With Infective Endocarditis: A Danish Nationwide Cohort Study. Clinical Infectious Diseases, 2019, 68, 668-675.	5.8	8
196	Clinical characteristics and risk factors of arrhythmia during followâ€up of patients with idiopathic ventricular fibrillation. Journal of Cardiovascular Electrophysiology, 2020, 31, 2677-2686.	1.7	8
197	Risk for infective endocarditis in bacteremia with Gram positive cocci. Infection, 2020, 48, 905-912.	4.7	8
198	Targeting Cardiac Myocyte Na ⁺ -K ⁺ Pump Function With β3 Adrenergic Agonist in Rabbit Model of Severe Congestive Heart Failure. Circulation: Heart Failure, 2020, 13, e006753.	3.9	8

#	Article	IF	CITATIONS
199	Cardiac Involvement in Women With Pathogenic Dystrophin Gene Variants. Frontiers in Neurology, 2021, 12, 707838.	2.4	8
200	Face masks for the prevention of COVID-19 - Rationale and design of the randomised controlled trial DANMASK-19. Danish Medical Journal, 2020, 67, .	0.5	8
201	Mutation detection by cleavase in combination with capillary electrophoresis analysis: Application to mutations causing hypertrophic cardiomyopathy and long-QT syndrome*. Molecular Diagnosis and Therapy, 1998, 3, 105-111.	1.1	7
202	Adult-onset familial hypertrophic cardiomyopathy caused by a novel mutation, R694C, in the MYH7 gene. Clinical Genetics, 1999, 56, 244-246.	2.0	7
203	Screening relatives in arrhythmogenic right ventricular cardiomyopathy: yield of imaging and electrical investigations. European Heart Journal Cardiovascular Imaging, 2019, 21, 175-182.	1.2	7
204	Rare non-coding Desmoglein-2 variant contributes to Arrhythmogenic right ventricular cardiomyopathy. Journal of Molecular and Cellular Cardiology, 2019, 131, 164-170.	1.9	7
205	Risk of stroke subsequent to infective endocarditis: A nationwide study. American Heart Journal, 2019, 212, 144-151.	2.7	7
206	Effect of impaired cardiac conduction after alcohol septal ablation on clinical outcomes: insights from the Euro-ASA registry. European Heart Journal Quality of Care & Clinical Outcomes, 2019, 5, 252-258.	4.0	7
207	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.	1.6	7
208	Severity of congenital long QT syndrome disease manifestation and risk of depression, anxiety, and mortality: a nationwide study. Europace, 2022, 24, 620-629.	1.7	7
209	Sex-Related Differences in Outcomes of Alcohol Septal Ablation for Hypertrophic Obstructive Cardiowascular Interventions, 2021, 14, 1390-1392.	2.9	7
210	Lung Ultrasound Findings Associated With COVID-19 ARDS, ICU Admission, and All-Cause Mortality. Respiratory Care, 2022, 67, 66-75.	1.6	7
211	Incidence of Positive Severe Acute Respiratory Syndrome Coronavirus Polymerase Chain Reaction After Coronavirus Disease 2019 Vaccination With up to 8 Months of Follow-up: Real-life Data From the Capital Region of Denmark. Clinical Infectious Diseases, 2022, 75, e675-e682.	5.8	7
212	Aortic Dissections in the Population-Based Danish National Patient Registry from 1996–2016: A Validation Study. Clinical Epidemiology, 2022, Volume 14, 51-58.	3.0	7
213	Potential Advances of Adjunctive Hyperbaric Oxygen Therapy in Infective Endocarditis. Frontiers in Cellular and Infection Microbiology, 2022, 12, 805964.	3.9	7
214	Lung ultrasound findings following COVID-19 hospitalization: A prospective longitudinal cohort study. Respiratory Medicine, 2022, 197, 106826.	2.9	7
215	Cohort profile: Copenhagen Hospital Biobank - Cardiovascular Disease Cohort (CHB-CVDC): Construction of a large-scale genetic cohort to facilitate a better understanding of heart diseases. BMJ Open, 2021, 11, e049709.	1.9	7
216	Antibody responses and risk factors associated with impaired immunological outcomes following two doses of BNT162b2 COVID-19 vaccination in patients with chronic pulmonary diseases. BMJ Open Respiratory Research, 2022, 9, e001268.	3.0	7

#	Article	IF	CITATIONS
217	Echocardiographic evaluation of pre-diagnostic development in young relatives genetically predisposed to hypertrophic cardiomyopathy. International Journal of Cardiovascular Imaging, 2015, 31, 1511-1518.	1.5	6
218	Clinical evaluation of unselected cardiac arrest survivors in a tertiary center over a 1-year period (the LAZARUZ study). Journal of Electrocardiology, 2016, 49, 707-713.	0.9	6
219	Cardiac magnetic resonance imaging after ventricular tachyarrhythmias increases diagnostic precision and reduces the need for family screening for inherited cardiac disease. Europace, 2016, 18, euv446.	1.7	6
220	Increased Risk of Ischemic Stroke After Treatment of Infective Endocarditis: A Danish, Nationwide, Propensity Score–Matched Cohort Study. Clinical Infectious Diseases, 2019, 70, 1186-1192.	5.8	6
221	External validation of the HANDOC score – high sensitivity to identify patients with non-beta-haemolytic streptococcal endocarditis. Infectious Diseases, 2020, 52, 54-57.	2.8	6
222	Patients with Unstable Atherosclerosis Have More Echolucent Carotid Plaques Compared with Stable Atherosclerotic Patients: A 3-D Ultrasound Study. Ultrasound in Medicine and Biology, 2020, 46, 2164-2172.	1.5	6
223	Diagnostic yield and long-term outcome of nonischemic sudden cardiac arrest survivors and their relatives: Results from a tertiary referral center. Heart Rhythm, 2020, 17, 1679-1686.	0.7	6
224	Long QT syndrome type 1 and 2 patients respond differently to arrhythmic triggers: The TriQarr inÂvivo study. Heart Rhythm, 2021, 18, 241-249.	0.7	6
225	Personalized medicine and preventive health care: juxtaposing health policy and clinical practice. Critical Public Health, 2021, 31, 327-337.	2.4	6
226	Long-term prognosis following hospitalization for acute myocarditis – a matched nationwide cohort study. Scandinavian Cardiovascular Journal, 2021, 55, 264-269.	1.2	6
227	Gestational Age and Neonatal Electrocardiograms. Pediatrics, 2021, 148, .	2.1	6
228	Acute symptoms in SARS-CoV-2 positive adolescents aged 15–18 years – Results from a Danish national cross-sectional survey study. Lancet Regional Health - Europe, The, 2022, 16, 100354.	5.6	6
229	High incidence of discrepancies in new Siemens assay– A comparison of cardiac troponin I assays. Clinical Chemistry and Laboratory Medicine, 2022, .	2.3	6
230	Dynamics of a <i>Staphylococcus aureus</i> infective endocarditis simulation model. Apmis, 2022, 130, 515-523.	2.0	6
231	Reduction of cerebral cortical [3H]ouabain binding site (Na+,K+-ATPase) density in dementia as evaluated in fresh human cerebral cortical biopsies. Cognitive Brain Research, 1996, 4, 281-287.	3.0	5
232	Outcome of Septal Myectomy in Patients with Hypertrophic Obstructive Cardiomyopathy. Scandinavian Cardiovascular Journal, 2000, 34, 564-569.	1.2	5
233	Potassium dynamics are attenuated in hyperkalemia and a determinant of QT adaptation in exercising hemodialysis patients. Journal of Applied Physiology, 2013, 115, 498-504.	2.5	5
234	Left ventricular volume predicts exercise capacity in hypertrophic cardiomyopathy. International Journal of Cardiology, 2016, 203, 676-678.	1.7	5

#	Article	IF	CITATIONS
235	Plakophilin-2 c.419C>T and risk of heart failure and arrhythmias in the general population. European Journal of Human Genetics, 2016, 24, 732-738.	2.8	5
236	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.	0.4	5
237	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.	1.7	5
238	Evolution of P-wave indices during long-term follow-up as markers of atrial substrate progression in arrhythmogenic right ventricular cardiomyopathy. Europace, 2021, 23, i29-i37.	1.7	5
239	Danish citizens' preferences for at-home oropharyngeal/nasal SARS-CoV-2 specimen collection. International Journal of Infectious Diseases, 2021, 109, 195-198.	3.3	5
240	Electrocardiographic Findings, Arrhythmias, and Left Ventricular Involvement in Familial ST-Depression Syndrome. Circulation: Arrhythmia and Electrophysiology, 2022, , 101161CIRCEP121010688.	4.8	5
241	Polygenic risk score for ACE-inhibitor-associated cough based on the discovery of new genetic loci. European Heart Journal, 2022, 43, 4707-4718.	2.2	5
242	Skeletal Muscle Na,K-ATPase Concentration Changes and Intramuscular and Extrarenal K Homeostasis in Animals and Humans. Annals of the New York Academy of Sciences, 1997, 834, 648-650.	3.8	4
243	Late potentials and their correlation with ventricular structure in patients with ventricular arrhythmias. PACE - Pacing and Clinical Electrophysiology, 2017, 40, 1466-1471.	1.2	4
244	Diagnostic findings and follow-up outcomes in relatives to young non-autopsied sudden death victims. International Journal of Cardiology, 2020, 318, 61-66.	1.7	4
245	Precordial ECG Amplitudes in the Days After Birth: Electrocardiographic Changes During Transition from Fetal to Neonatal Circulation. Pediatric Cardiology, 2021, 42, 832-839.	1.3	4
246	The Evolution of the Neonatal QRS Axis during the First Four Weeks of Life. Neonatology, 2021, 118, 155-162.	2.0	4
247	Plasma levels of glucagon but not GLP-1 are elevated in response to inflammation in humans. Endocrine Connections, 2021, 10, 205-213.	1.9	4
248	Prenatal cardiac biometries and flow assessments in fetuses at 20 weeks with a bicuspid aortic valve compared to healthy controls: a multicenter, cohort study. Ultrasound in Obstetrics and Gynecology, 2021, 58, 846-852.	1.7	4
249	Natural History and Clinical Characteristics of the First 10 Danish Families With Familial ST-Depression Syndrome. Journal of the American College of Cardiology, 2021, 77, 2617-2619.	2.8	4
250	Accelerated treatment of endocarditis—The POET II trial: Rationale and design of a randomized controlled trial. American Heart Journal, 2020, 227, 40-46.	2.7	4
251	Severity of anaemia and association with all-cause mortality in patients with medically managed left-sided endocarditis. Heart, 2022, 108, 882-888.	2.9	4
252	Effect of Influenza Vaccination on Risk of Coronavirus Disease 2019: A Prospective Cohort Study of 46â€000 Healthcare Workers. Journal of Infectious Diseases, 2022, 226, 6-10.	4.0	4

#	Article	IF	CITATIONS
253	Proteomic Characterization of Atherosclerotic Lesions In Situ Using Percutaneous Coronary Intervention Angioplasty Balloons—Brief Report. Arteriosclerosis, Thrombosis, and Vascular Biology, 2022, 42, 857-864.	2.4	4
254	Prevalence of Left Ventricular Noncompaction in Newborns. Circulation: Cardiovascular Imaging, 2022, 15, .	2.6	4
255	Association of Common and Rare Genetic Variation in the 3â€Hydroxyâ€3â€Methylglutaryl Coenzyme A Reductase Gene and Cataract Risk. Journal of the American Heart Association, 2022, 11, .	3.7	4
256	Time-dependent response of both ventricles after septal ablation: Implications for biventricular support after left ventricular assist device placement. Journal of Thoracic and Cardiovascular Surgery, 2007, 134, 579-586.	0.8	3
257	Functional Promoter Variant in <i>Desmocollin-2</i> Contributes to Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Cardiovascular Genetics, 2016, 9, 384-387.	5.1	3
258	In-hospital metabolite changes in infective endocarditis—a longitudinal 1H NMR-based study. European Journal of Clinical Microbiology and Infectious Diseases, 2019, 38, 1553-1560.	2.9	3
259	Evolutionary dissection of mtDNA hg H: a susceptibility factor for hypertrophic cardiomyopathy. Mitochondrial DNA Part A: DNA Mapping, Sequencing, and Analysis, 2020, 31, 238-244.	0.7	3
260	Comprehensive cardiac rehabilitation for patients following infective endocarditis : r esults of the randomized CopenHeartIE trial . European Journal of Cardiovascular Nursing, 2022, 21, 261-270.	0.9	3
261	Genetic Variant Score and Arrhythmogenic Right Ventricular Cardiomyopathy Phenotype in Plakophilin-2 Mutation Carriers. Cardiology, 2021, 146, 763-771.	1.4	3
262	The impact of partial-oral endocarditis treatment on anxiety and depression in the POET trial. Journal of Psychosomatic Research, 2022, 154, 110718.	2.6	3
263	Patient-prioritized primary endpoints in clinical trials. Scandinavian Cardiovascular Journal, 2022, 56, 4-5.	1.2	3
264	Familial risk of atrioventricular block in first-degree relatives. Heart, 2022, 108, 1194-1199.	2.9	3
265	Sudden unexplained death versusÂnonautopsied possible sudden cardiac death: Findings in relatives. Journal of Cardiovascular Electrophysiology, 2022, 33, 254-261.	1.7	3
266	β-blocker adherence among patients with congenital long QT syndrome: a nationwide study. European Heart Journal Quality of Care & Clinical Outcomes, 2022, 9, 76-84.	4.0	3
267	Between observer variation is not eliminated by standardised analysis of dobutamine-atropine stress echocardiography. International Journal of Cardiovascular Imaging, 2002, 18, 169-179.	0.6	2
268	Expanding the cerebrovascular phenotype of the p.R258H variant in ACTA2 related hereditary thoracic aortic disease (HTAD). Journal of the Neurological Sciences, 2020, 415, 116897.	0.6	2
269	Specificity and cross-reactivity of a test for anti-SARS-CoV-2 antibodies – Authors' reply. Lancet Infectious Diseases, The, 2021, 21, e119.	9.1	2
270	Effect of moderate potassium-elevating treatment in long QT syndrome: the TriQarr Potassium Study. Open Heart, 2021, 8, e001670.	2.3	2

#	Article	IF	CITATIONS
271	Regulation of Myocardial and Skeletal Muscle Na,K-ATPase in Diabetes Mellitus in Humans and Animals. Advances in Experimental Medicine and Biology, 2001, 498, 319-322.	1.6	2
272	Digoxin affects potassium homeostasis during exercise in patients with heart failure. Cardiovascular Research, 1995, 29, 506-511.	3.8	2
273	Receptor Occupancy with Digoxin vs. Receptor Occupancy with a Putative Endogenous Digitalislike Factor. Hypertension Research, 2000, 23, S39-S43.	2.7	2
274	Diagnostic Yield of Genetic Testing in Young Patients With Atrioventricular Block of Unknown Cause. Journal of the American Heart Association, 2022, 11, e025643.	3.7	2
275	Classification of Left and Right Coronary Arteries in Coronary Angiographies Using Deep Learning. Electronics (Switzerland), 2022, 11, 2087.	3.1	2
276	Regulation of Myocardial Na,K-ATPase Concentration in Experimental and Human Heart Disease. Annals of the New York Academy of Sciences, 1997, 834, 676-679.	3.8	1
277	Post-mortem investigation of young deceased individuals with ischemic heart disease—outcome of supplementary genetic testing for dyslipidemia. International Journal of Legal Medicine, 2016, 130, 947-948.	2.2	1
278	The D313Y variant in the GLA gene - no evidence of a pathogenic role in Fabry disease in 2 Danish families. Molecular Genetics and Metabolism, 2018, 123, S44.	1.1	1
279	Lung ultrasound findings in hospitalized COVID-19 patients in relation to venous thromboembolic events: the ECHOVID-19 study. Journal of Ultrasound, 2021, , 1.	1.3	1
280	Mid-regional pro-atrial natriuretic peptide levels before and after hemodialysis predict long-term prognosis. Clinical Biochemistry, 2021, 94, 20-26.	1.9	1
281	Effectiveness of Adding a Mask Recommendation to Other Public Health Measures. Annals of Internal Medicine, 2021, 174, 1194-1195.	3.9	1
282	Pacemaker implantation in a patient with brugada and sick sinus syndrome. World Journal of Cardiology, 2013, 5, 65.	1.5	1
283	Coagulation parameters in the newborn and infant– the Copenhagen Baby Heart and COMPARE studies. Clinical Chemistry and Laboratory Medicine, 2021, .	2.3	1
284	Effect of Loss-of-Function Genetic Variants in <i>PCSK9</i> on Glycemic Traits, Neurocognitive Impairment, and Hepatobiliary Function. Diabetes Care, 2022, 45, 251-254.	8.6	1
285	Clinical Implications of <i>SCN10A</i> Loss-of-Function Variants in 169 610 Exomes Representing the General Population. Circulation Genomic and Precision Medicine, 2022, 15, CIRCGEN121003574.	3.6	1
286	Expression of Na,K-ATPase in hearts of diabetic and sew-starving rats: A comparative sudy. Journal of Molecular and Cellular Cardiology, 2001, 33, A138.	1.9	0
287	Muscular K learance Capacity <i>in Vivo</i> Must Be Evaluated on the Basis of K and Na,Kâ€ATPase Concentrations. Annals of the New York Academy of Sciences, 2003, 986, 623-624.	3.8	0
288	Exercise echocardiography in hypertrophic cardiomyopathy: reply. European Journal of Echocardiography, 2010, 11, 730-731.	2.3	0

#	Article	IF	CITATIONS
289	MT-CYB mutations in hypertrophic cardiomyopathy. Molecular Genetics & Genomic Medicine, 2013, 1, 187-187.	1.2	0
290	Response letter to "Cardiac involvement in myotonic dystrophy type 1 — Do not forget the loop recorder!― International Journal of Cardiology, 2013, 168, 1541.	1.7	0
291	Reply. Journal of the American College of Cardiology, 2013, 61, 2393.	2.8	0
292	Cardiac fibrosis in myotonic dystrophy type 1; an early marker of cardiac involvement. European Heart Journal, 2013, 34, P2987-P2987.	2.2	0
293	Nationwide study on sudden cardiac death in Danes aged 1-49 years. European Heart Journal, 2013, 34, 1741-1741.	2.2	Ο
294	Cardiac symptoms before sudden cardiac death caused by coronary artery disease: a nationwide study among young Danes. European Heart Journal, 2013, 34, P1359-P1359.	2.2	0
295	MULTIMODALITY IMAGING TO ASSESS DETERMINANTS OF EXERCISE CAPACITY IN HYPERTROPHIC CARDIOMYOPATHY. Journal of the American College of Cardiology, 2015, 65, A978.	2.8	0
296	Presymptomatic diagnosis of Fabry's disease: a case report. Journal of Medical Case Reports, 2016, 10, 330.	0.8	0
297	Cardiac magnetic resonance imaging provides more than a diagnosis. Europace, 2017, 19, euw253.	1.7	0
298	Association Between Hypertensive Disorders of Pregnancy and Later Risk of Cardiomyopathy. Obstetrical and Gynecological Survey, 2016, 71, 387-389.	0.4	0
299	Characterization of a Na V 1.5 Gain-of-Function Mutation (G213D) causing Multifocal Atrial and Ventricular Premature Ectopies and an Increased Risk of Dilated Cardiomyopathy. Biophysical Journal, 2017, 112, 104a.	0.5	0
300	RISK OF INFECTIVE ENDOCARDITIS ASSOCIATED WITH CONCOMITANT PACEMAKER AMONG PATIENTS UNDERGOING AORTIC VALVE REPLACEMENT: A NATIONWIDE STUDY. Journal of the American College of Cardiology, 2017, 69, 1959.	2.8	0
301	DUCHENNE MUSCULAR DYSTROPHY - GENETICS. Neuromuscular Disorders, 2018, 28, S98.	0.6	0
302	A comparison of cholesterol levels in umbilical cord blood and in neonatal blood - The compare study. Atherosclerosis, 2018, 275, e92.	0.8	0
303	TCTAP A-078 Short- and Long-term Outcome of Alcohol Septal Ablation for Hypertrophic Obstructive Cardiomyopathy in Patients with Mild Left Ventricular Hypertrophy: A Propensity Score Matching Analysis. Journal of the American College of Cardiology, 2019, 73, S40.	2.8	0
304	Corrigendum to: Incidence of infective endocarditis in patients considered at moderate risk. European Heart Journal, 2019, 40, 1361-1361.	2.2	0
305	Cardiogenetic screening amongst families of sudden cardiac death victims: Authors' reply. Europace, 2020, 22, 1754-1755.	1.7	0
306	Reply to: TFC ECG in arrhythmogenic cardiomyopathy: Inadequate mixture of criteria?. International Journal of Cardiology, 2021, 323, 203.	1.7	0

#	Article	IF	CITATIONS
307	Improval of outcome in patients with endocarditis. European Heart Journal: Acute Cardiovascular Care, 2021, 10, 367-368.	1.0	0
308	Response by Hasselbalch et al to Letter Regarding Article, "Temporal Release of High-Sensitivity Cardiac Troponin T and I and Copeptin After Brief Induced Coronary Artery Balloon Occlusion in Humans― Circulation, 2021, 144, e169-e170.	1.6	0
309	Infective endocarditis caused by Bartonella quintana in Greenland. JMM Case Reports, 2014, 1, .	1.3	0
310	Rare variants in HCN4 identified in the general population are associated with complete atrioventricular (AV) block, 1. degree AV block and bundle branch block, results from 50.000 exomes. European Heart Journal, 2020, 41, .	2.2	0
311	Echolucent Carotid Plaques Becomes More Echogenic over Time – A 3D Ultrasound Study. Annals of Vascular Surgery, 2022, , .	0.9	0
312	Self-assessed health status and associated mortality in endocarditis: secondary findings from the POET trial. Quality of Life Research, 2022, , 1.	3.1	0
313	Abstract 13222: Genetic Variants Close to <i>NKX2-5</i> and <i>MYH6</i> Are Associated With AV Nodal Reentry Tachycardia in First Genome-Wide Association Study. Circulation, 2021, 144, .	1.6	0
314	Abstract 12229: Left Ventricular Non-Compaction in Childhood: Echocardiographic Follow-Up and Prevalence in First-Degree Relatives. Circulation, 2021, 144, .	1.6	0
315	Abstract 12519: Electrocardiographic Findings and Arrhythmias in Patients With Familial ST-Depression Syndrome. Circulation, 2021, 144, .	1.6	0
316	Abstract 12265: The Impact of Gestational Age on the Neonatal Electrocardiogram. Circulation, 2021, 144, .	1.6	0
317	Clinical Characteristics, Incidences, and Mortality Rates for Aortic Dissections Type A and B: A Nationwide Danish Population-Based Cohort Study from 1996 to 2016, Aorta, 2022,	0.5	0