Alex HÃ¸rby Christensen

List of Publications by Year
in descending order

29 papers	220 citations	1162889 8 h-index	1125617 13 g-index
31 all docs	31 docs citations	31 times ranked	363 citing authors

#	Article	IF	Citations
1	Complications of implantable cardioverter-defibrillator treatment in arrhythmogenic right ventricular cardiomyopathy. Europace, 2022, 24, 306-312.	0.7	12
2	Genotypeâ€“phenotype correlation in arrhythmogenic right ventricular cardiomyopathyâ€”risk of arrhythmias and heart failure. Journal of Medical Genetics, 2022, 59, 858-864.	1.5	13
3	Cardiotoxicity in metastatic melanoma patients treated with BRAF and MEK inhibitors in a real-world setting. Acta OncolÃ³gica, 2022, 61, 45-51.	0.8	3
4	Electrocardiographic Findings, Arrhythmias, and Left Ventricular Involvement in Familial ST-Depression Syndrome. Circulation: Arrhythmia and Electrophysiology, 2022, , 101161CIRCEP121010688.	2.1	5
5	Classification of Left and Right Coronary Arteries in Coronary Angiographies Using Deep Learning. Electronics (Switzerland), 2022, 11, 2087.	1.8	2
6	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.	1.5	14
7	Long QT syndrome type 1 and 2 patients respond differently to arrhythmic triggers: The TriQarr inÂvivo study. Heart Rhythm, 2021, 18, 241-249.	0.3	6
8	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.	0.7	9
9	Precordial ECG Amplitudes in the Days After Birth: Electrocardiographic Changes During Transition from Fetal to Neonatal Circulation. Pediatric Cardiology, 2021, 42, 832-839.	0.6	4
10	The Evolution of the Neonatal QRS Axis during the First Four Weeks of Life. Neonatology, 2021, 118, 155-162.	0.9	4
11	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.	1.2	4
12	Reassessment of Gene-Elusive Familial Dilated Cardiomyopathy Leading to the Discovery of a Homozygous AARS2 Variantâ€”The Importance of Regular Reassessment of Genetic Findings. Neurology International, 2021, 11, 122-128.	0.2	0
13	Effect of moderate potassium-elevating treatment in long QT syndrome: the TriQarr Potassium Study. Open Heart, 2021, 8, e001670.	0.9	2
14	Gestational Age and Neonatal Electrocardiograms. Pediatrics, 2021, 148, .	1.0	6
15	Cardiogenetic screening amongst families of sudden cardiac death victims: Authorsâ€™ reply. Europace, 2020, 22, 1754-1755.	0.7	0
16	Diagnostic findings and follow-up outcomes in relatives to young non-autopsied sudden death victims. International Journal of Cardiology, 2020, 318, 61-66.	0.8	4
17	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.	0.3	6
18	Diagnostic yield in victims of sudden cardiac death and their relatives. Europace, 2020, 22, 964-971.	0.7	18

#	Article	IF	Citations
19	Non-diagnostic autopsy findings in sudden unexplained death victims. BMC Cardiovascular Disorders, 2020, 20, 58.	0.7	12
20	Screening relatives in arrhythmogenic right ventricular cardiomyopathy: yield of imaging and electrical investigations. European Heart Journal Cardiovascular Imaging, 2019, 21, 175-182.	0.5	7
21	Rare non-coding Desmoglein-2 variant contributes to Arrhythmogenic right ventricular cardiomyopathy. Journal of Molecular and Cellular Cardiology, 2019, 131, 164-170.	0.9	7
22	A Novel <i>SCN5A</i> Variant Associated with Abnormal Repolarization, Atrial Fibrillation, and Reversible Cardiomyopathy. Cardiology, 2018, 140, 8-13.	0.6	3
23	Citalopram and the KCNE1 D85N variant: a case report on the implications of a genetic modifier. European Heart Journal - Case Reports, 2018, 2, yty106.	0.3	2
24	Functional Promoter Variant in Desmocollin-2 Contributes to Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Cardiovascular Genetics, 2016, 9, 384-387.	5.1	3
25	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.	1.4	5
26	Mutation analysis of the candidate genes SCN1B-4B, FHL1, and LMNA in patients with arrhythmogenic right ventricular cardiomyopathy. Applied & Translational Genomics, 2012, 1, 44-46.	2.1	3
27	Screening of Three Novel Candidate Genes in Arrhythmogenic Right Ventricular Cardiomyopathy. Genetic Testing and Molecular Biomarkers, 2011, 15, 267-271.	0.3	18
28	Missense Variants in <i>Plakophilin-2</i> in Arrhythmogenic Right Ventricular Cardiomyopathy Patients â€“ Disease-Causing or Innocent Bystanders?. Cardiology, 2010, 115, 148-154.	0.6	44
29	Myocarditis Mimicking Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American College of Cardiology, 2009, 54, 663-664.	1.2	3