

# Barbara Bauce

## List of Publications by Year in descending order

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Version: 2024-02-01

68  
papers

8,186  
citations

126708

33  
h-index

106150

65  
g-index

69  
all docs

69  
docs citations

69  
times ranked

5146  
citing authors

| #  | ARTICLE   | IF  | CITATIONS |
|----|---|-----|-----------|
| 1  | Clinical application of CMR in cardiomyopathies: evolving concepts and techniques. <i>Heart Failure Reviews</i> , 2023, 28, 77-95.  | 1.7 | 19        |
| 2  | Cardiac magnetic resonance imaging of arrhythmogenic cardiomyopathy: evolving diagnostic perspectives. <i>European Radiology</i> , 2023, 33, 270-282.   | 2.3 | 12        |
| 3  | Filamin-C variant-associated cardiomyopathy: A pooled analysis of individual patient data to evaluate the clinical profile and risk of sudden cardiac death. <i>Heart Rhythm</i> , 2022, 19, 235-243.   | 0.3 | 33        |
| 4  | Role of Cardiac Magnetic Resonance Imaging in the Evaluation of Athletes with Premature Ventricular Beats. <i>Journal of Clinical Medicine</i> , 2022, 11, 426.   | 1.0 | 11        |
| 5  | The 2020 "Padua Criteria" for Diagnosis and Phenotype Characterization of Arrhythmogenic Cardiomyopathy in Clinical Practice. <i>Journal of Clinical Medicine</i> , 2022, 11, 279.  | 1.0 | 9         |
| 6  | Strength of clinical indication and therapeutic impact of the implantable cardioverter defibrillator in patients with hypertrophic cardiomyopathy. <i>International Journal of Cardiology</i> , 2022, 353, 62-67.   | 0.8 | 2         |
| 7  | Thyroid dysfunction on the heart: clinical effects, prognostic impact and management strategies. <i>Monaldi Archives for Chest Disease</i> , 2022, , .  | 0.3 | 1         |
| 8  | Diagnosis and Management of Rare Cardiomyopathies in Adult and Paediatric Patients. A Position Paper of the Italian Society of Cardiology (SIC) and Italian Society of Paediatric Cardiology (SICP). <i>International Journal of Cardiology</i> , 2022, 357, 55-71. | 0.8 | 36        |
| 9  | Clinical profile and long-term follow-up of a cohort of patients with desmoplakin cardiomyopathy. <i>Heart Rhythm</i> , 2022, 19, 1315-1324.  | 0.3 | 22        |
| 10 | Importance of genotype for risk stratification in arrhythmogenic right ventricular cardiomyopathy using the 2019 ARVC risk calculator. <i>European Heart Journal</i> , 2022, 43, 3053-3067.   | 1.0 | 41        |
| 11 | Arrhythmic Mitral Valve Prolapse in the Young: A Rare but Concerning Entity. <i>Diagnostics</i> , 2022, 12, 1519.   | 1.3 | 0         |
| 12 | Added Value of CCTA-Derived Features to Predict MACEs in Stable Patients Undergoing Coronary Computed Tomography. <i>Diagnostics</i> , 2022, 12, 1446.  | 1.3 | 4         |
| 13 | Ventricular arrhythmias in mitral valve prolapse: new explanations for an old problem. <i>Heart</i> , 2021, 107, 353-354.   | 1.2 | 5         |
| 14 | Reply to "signal averaged electrocardiogram findings among right ventricular arrhythmogenic cardiomyopathy (ARVC) patients: Do they have a place in ARVC management?" <i>International Journal of Cardiology</i> , 2021, 327, 155.                                  | 0.8 | 1         |
| 15 | Right Ventricular Cardiomyopathies. , 2021, , 267-288.  |     | 0         |
| 16 | Differential diagnosis of arrhythmogenic cardiomyopathy: phenocopies versus disease variants. <i>Minerva Medica</i> , 2021, 112, 269-280.   | 0.3 | 13        |
| 17 | Role of Exercise as a Modulating Factor in Arrhythmogenic Cardiomyopathy. <i>Current Cardiology Reports</i> , 2021, 23, 57.   | 1.3 | 17        |
| 18 | Arrhythmogenic Left Ventricular Cardiomyopathy: Genotype-Phenotype Correlations and New Diagnostic Criteria. <i>Journal of Clinical Medicine</i> , 2021, 10, 2212.  | 1.0 | 18        |

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|----|--|-----|-----------|
| 19 | Arrhythmogenic Cardiomyopathyâ€™ Current Treatment and Future Options. <i>Journal of Clinical Medicine</i> , 2021, 10, 2750.   | 1.0 | 10        |
| 20 | The genetic architecture of Plakophilin 2 cardiomyopathy. <i>Genetics in Medicine</i> , 2021, 23, 1961-1968.   | 1.1 | 13        |
| 21 | Novel pathogenic role for galectin-3 in early disease stages of arrhythmogenic cardiomyopathy. <i>Heart Rhythm</i> , 2021, 18, 1394-1403.  | 0.3 | 8         |
| 22 | Myocardial Tissue Characterization in Arrhythmogenic Cardiomyopathy. <i>JACC: Cardiovascular Imaging</i> , 2021, 14, 1675-1678.  | 2.3 | 13        |
| 23 | Evolving Diagnostic Criteria for Arrhythmogenic Cardiomyopathy. <i>Journal of the American Heart Association</i> , 2021, 10, e021987.  | 1.6 | 60        |
| 24 | â€™Hot phaseâ€™™ clinical presentation in arrhythmogenic cardiomyopathy. <i>Europace</i> , 2021, 23, 907-917.  | 0.7 | 67        |
| 25 | Arrhythmogenic Cardiomyopathy. <i>European Heart Journal</i> , 2020, 41, 4457-4462.  | 1.0 | 12        |
| 26 | Diagnosis of arrhythmogenic cardiomyopathy: The Padua criteria. <i>International Journal of Cardiology</i> , 2020, 319, 106-114.   | 0.8 | 283       |
| 27 | Arrhythmogenic Right Ventricular Cardiomyopathy: Characterization of Left Ventricular Phenotype and Differential Diagnosis With Dilated Cardiomyopathy. <i>Journal of the American Heart Association</i> , 2020, 9, e014628.                           | 1.6 | 92        |
| 28 | A microRNA Expression Profile as Non-Invasive Biomarker in a Large Arrhythmogenic Cardiomyopathy Cohort. <i>International Journal of Molecular Sciences</i> , 2020, 21, 1536.  | 1.8 | 21        |
| 29 | Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. <i>European Heart Journal</i> , 2020, 41, 1414-1429.  | 1.0 | 239       |
| 30 | Management of nonischemic-dilated cardiomyopathies in clinical practice: a position paper of the working group on myocardial and pericardial diseases of Italian Society of Cardiology. <i>Journal of Cardiovascular Medicine</i> , 2020, 21, 927-943. | 0.6 | 5         |
| 31 | Definition and treatment of arrhythmogenic cardiomyopathy: an updated expert panel report. <i>European Journal of Heart Failure</i> , 2019, 21, 955-964.   | 2.9 | 84        |
| 32 | Predictive value of exercise testing in athletes with ventricular ectopy evaluated by cardiac magnetic resonance. <i>Heart Rhythm</i> , 2019, 16, 239-248.   | 0.3 | 45        |
| 33 | Relationship Between Electrocardiographic Findings and Cardiac Magnetic Resonance Phenotypes in Arrhythmogenic Cardiomyopathy. <i>Journal of the American Heart Association</i> , 2018, 7, e009855.  | 1.6 | 58        |
| 34 | Large Genomic Rearrangements of Desmosomal Genes in Italian Arrhythmogenic Cardiomyopathy Patients. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2017, 10, .   | 2.1 | 35        |
| 35 | Co-inheritance of mutations associated with arrhythmogenic cardiomyopathy and hypertrophic cardiomyopathy. <i>European Journal of Human Genetics</i> , 2017, 25, 1165-1169.  | 1.4 | 10        |
| 36 | Anchoring Vignettes in EQ-5D-5L Questionnaire: Validation of a New Instrument. <i>Open Nursing Journal</i> , 2017, 11, 144-156.  | 0.2 | 3         |

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|----|---|-----|-----------|
| 37 | Arrhythmogenic Right Ventricular Cardiomyopathy: Risk Stratification and Indications for Defibrillator Therapy. <i>Current Cardiology Reports</i> , 2016, 18, 57.   | 1.3 | 20        |
| 38 | Morphofunctional Abnormalities of Mitral Annulus and Arrhythmic Mitral Valve Prolapse. <i>Circulation: Cardiovascular Imaging</i> , 2016, 9, e005030.   | 1.3 | 226       |
| 39 | Nonischemic Left Ventricular Scar as a Substrate of Life-Threatening Ventricular Arrhythmias and Sudden Cardiac Death in Competitive Athletes. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2016, 9, .                              | 2.1 | 216       |
| 40 | Relationship Between Arrhythmogenic Right Ventricular Cardiomyopathy and Brugada Syndrome. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2016, 9, e003631.   | 2.1 | 78        |
| 41 | Phenotypic expression is a prerequisite for malignant arrhythmic events and sudden cardiac death in arrhythmogenic right ventricular cardiomyopathy. <i>Europace</i> , 2016, 18, 1086-1094.   | 0.7 | 50        |
| 42 | Treatment of arrhythmogenic right ventricular cardiomyopathy/dysplasia: an international task force consensus statement. <i>European Heart Journal</i> , 2015, 36, ehv162.  | 1.0 | 171       |
| 43 | Arrhythmic Mitral Valve Prolapse and Sudden Cardiac Death. <i>Circulation</i> , 2015, 132, 556-566.   | 1.6 | 422       |
| 44 | Is Internet use associated with anxiety in patients with and at risk for cardiomyopathy?. <i>American Heart Journal</i> , 2015, 170, 87-95.e4.  | 1.2 | 16        |
| 45 | Nonischemic Left Ventricular Scar. <i>Circulation</i> , 2014, 130, e180-2.  | 1.6 | 22        |
| 46 | Noninvasive Cardiac Screening in Young Athletes With Ventricular Arrhythmias. <i>American Journal of Cardiology</i> , 2013, 111, 557-562.   | 0.7 | 34        |
| 47 | Compound and Digenic Heterozygosity Predicts Lifetime Arrhythmic Outcome and Sudden Cardiac Death in Desmosomal Gene-Related Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2013, 6, 533-542. | 5.1 | 209       |
| 48 | Imaging Study of Ventricular Scar in Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2012, 5, 91-100.   | 2.1 | 154       |
| 49 | Prevalence of Cardiomyopathy in Italian Asymptomatic Children With Electrocardiographic T-Wave Inversion at Preparticipation Screening. <i>Circulation</i> , 2012, 125, 529-538.  | 1.6 | 144       |
| 50 | Clinical phenotype and diagnosis of arrhythmogenic right ventricular cardiomyopathy in pediatric patients carrying desmosomal gene mutations. <i>Heart Rhythm</i> , 2012, 9, e11-e12.   | 0.3 | 9         |
| 51 | Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2012, 5, 1233-1246.   | 2.1 | 90        |
| 52 | Follow-Up with Exercise Test of Effort-Induced Ventricular Arrhythmias Linked to Ryanodine Receptor Type 2 Gene Mutations. <i>American Journal of Cardiology</i> , 2012, 109, 1015-1019.  | 0.7 | 8         |
| 53 | Clinical phenotype and diagnosis of arrhythmogenic right ventricular cardiomyopathy in pediatric patients carrying desmosomal gene mutations. <i>Heart Rhythm</i> , 2011, 8, 1686-1695.   | 0.3 | 66        |
| 54 | The p.A897KfsX4 frameshift variation in desmocollin-2 is not a causative mutation in arrhythmogenic right ventricular cardiomyopathy. <i>European Journal of Human Genetics</i> , 2010, 18, 776-782.  | 1.4 | 19        |

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|----|---|-----|-----------|
| 55 | Diagnosis of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. <i>Circulation</i> , 2010, 121, 1533-1541.  | 1.6 | 1,839     |
| 56 | Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Cardiac Electrophysiology Clinics</i> , 2010, 2, 571-586.   | 0.7 | 6         |
| 57 | Compound and Digenic Heterozygosity Contributes to Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2010, 55, 587-597.   | 1.2 | 282       |
| 58 | Multiple mutations in desmosomal proteins encoding genes in arrhythmogenic right ventricular cardiomyopathy/dysplasia. <i>Heart Rhythm</i> , 2010, 7, 22-29.  | 0.3 | 161       |
| 59 | Myocyte necrosis underlies progressive myocardial dystrophy in mouse <i>dsg2</i> -related arrhythmogenic right ventricular cardiomyopathy. <i>Journal of Experimental Medicine</i> , 2009, 206, 1787-1802.                          | 4.2 | 184       |
| 60 | Comparison of Clinical Features of Arrhythmogenic Right Ventricular Cardiomyopathy in Men Versus Women. <i>American Journal of Cardiology</i> , 2008, 102, 1252-1257.   | 0.7 | 81        |
| 61 | Quantitative assessment of endomyocardial biopsy in arrhythmogenic right ventricular cardiomyopathy/dysplasia: an in vitro validation of diagnostic criteria. <i>European Heart Journal</i> , 2008, 29, 2760-2771.                  | 1.0 | 161       |
| 62 | Mutations in Desmoglein-2 Gene Are Associated With Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Circulation</i> , 2006, 113, 1171-1179.  | 1.6 | 509       |
| 63 | Ultrastructural evidence of intercalated disc remodelling in arrhythmogenic right ventricular cardiomyopathy: an electron microscopy investigation on endomyocardial biopsies. <i>European Heart Journal</i> , 2006, 27, 1847-1854. | 1.0 | 219       |
| 64 | Three-Dimensional Electroanatomic Voltage Mapping Increases Accuracy of Diagnosing Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. <i>Circulation</i> , 2005, 111, 3042-3050.  | 1.6 | 237       |
| 65 | Clinical profile of four families with arrhythmogenic right ventricular cardiomyopathy caused by dominant desmoplakin mutations. <i>European Heart Journal</i> , 2005, 26, 1666-1675.   | 1.0 | 267       |
| 66 | Echocardiographic Findings in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. <i>Journal of the American College of Cardiology</i> , 2005, 46, 1962.   | 1.2 | 0         |
| 67 | Mutation in Human Desmoplakin Domain Binding to Plakoglobin Causes a Dominant Form of Arrhythmogenic Right Ventricular Cardiomyopathy. <i>American Journal of Human Genetics</i> , 2002, 71, 1200-1206.                             | 2.6 | 570       |
| 68 | Clinical profile and long-term follow-up of 37 families with arrhythmogenic right ventricular cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2000, 36, 2226-2233.   | 1.2 | 414       |