Cristina Basso

List of Publications by Year in descending order

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352 papers	42,702 citations	3149 92 h-index	2274 200 g-index
364 all docs	364 docs citations	364 times ranked	21125 citing authors

#	Article	IF	Citations
1	2014 ESC Guidelines on diagnosis and management of hypertrophic cardiomyopathy. European Heart Journal, 2014, 35, 2733-2779.	1.0	3,469
2	Current state of knowledge on aetiology, diagnosis, management, and therapy of myocarditis: a position statement of the European Society of Cardiology Working Group on Myocardial and Pericardial Diseases. European Heart Journal, 2013, 34, 2636-2648.	1.0	2,436
3	Diagnosis of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Circulation, 2010, 121, 1533-1541.	1.6	1,839
4	Trends in Sudden Cardiovascular Death in Young Competitive Athletes After Implementation of a Preparticipation Screening Program. JAMA - Journal of the American Medical Association, 2006, 296, 1593.	3.8	1,265
5	Diagnosis of arrhythmogenic right ventricular cardiomyopathy/dysplasia: Proposed Modification of the Task Force Criteria. European Heart Journal, 2010, 31, 806-814.	1.0	1,177
6	Does sports activity enhance the risk of sudden death in adolescents and young adults?. Journal of the American College of Cardiology, 2003, 42, 1959-1963.	1.2	1,133
7	Clinical profile of congenital coronary artery anomalies with origin from the wrong aortic sinus leading to sudden death in young competitive athletes. Journal of the American College of Cardiology, 2000, 35, 1493-1501.	1.2	1,046
8	Cardiovascular pre-participation screening of young competitive athletes for prevention of sudden death: proposal for a common European protocol. European Heart Journal, 2005, 26, 516-524.	1.0	1,037
9	Screening for Hypertrophic Cardiomyopathy in Young Athletes. New England Journal of Medicine, 1998, 339, 364-369.	13.9	890
10	Spectrum of Clinicopathologic Manifestations of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia: A Multicenter Study. Journal of the American College of Cardiology, 1997, 30, 1512-1520.	1.2	884
11	2020 ESC Guidelines on sports cardiology and exercise in patients with cardiovascular disease. European Heart Journal, 2021, 42, 17-96.	1.0	830
12	Arrhythmogenic right ventricular cardiomyopathy. Lancet, The, 2009, 373, 1289-1300.	6.3	785
13	Recommendations for interpretation of 12-lead electrocardiogram in the athlete. European Heart Journal, 2010, 31, 243-259.	1.0	730
14	Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation, 1996, 94, 983-991.	1.6	724
15	Identification of mutations in the cardiac ryanodine receptor gene in families affected with arrhythmogenic right ventricular cardiomyopathy type 2 (ARVD2). Human Molecular Genetics, 2001, 10, 189-194.	1.4	709
16	Mutation in Human Desmoplakin Domain Binding to Plakoglobin Causes a Dominant Form of Arrhythmogenic Right Ventricular Cardiomyopathy. American Journal of Human Genetics, 2002, 71, 1200-1206.	2.6	570
17	Implantable Cardioverter-Defibrillator Therapy for Prevention of Sudden Death in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Circulation, 2003, 108, 3084-3091.	1.6	539
18	Mutations in Desmoglein-2 Gene Are Associated With Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation, 2006, 113, 1171-1179.	1.6	509

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19	Anomalous origin of coronary arteries and risk of sudden death: A study based on an autopsy population of congenital heart disease. Human Pathology, 1998, 29, 689-695.	1.1	480
20	Diagnosis and treatment of cardiac amyloidosis: a position statement of the ESC Working Group on Myocardial and Pericardial Diseases. European Heart Journal, 2021, 42, 1554-1568.	1.0	434
21	A New Diagnostic Test for Arrhythmogenic Right Ventricular Cardiomyopathy. New England Journal of Medicine, 2009, 360, 1075-1084.	13.9	424
22	2011 Consensus statement on endomyocardial biopsy from the Association for European Cardiovascular Pathology and the Society for Cardiovascular Pathology. Cardiovascular Pathology, 2012, 21, 245-274.	0.7	423
23	Arrhythmic Mitral Valve Prolapse and Sudden Cardiac Death. Circulation, 2015, 132, 556-566.	1.6	422
24	Acute myocarditis presenting as a reverse Tako-Tsubo syndrome in a patient with SARS-CoV-2 respiratory infection. European Heart Journal, 2020, 41, 1861-1862.	1.0	415
25	Clinical profile and long-term follow-up of 37 families with arrhythmogenic right ventricular cardiomyopathy. Journal of the American College of Cardiology, 2000, 36, 2226-2233.	1.2	414
26	Genetic counselling and testing in cardiomyopathies: a position statement of the European Society of Cardiology Working Group on Myocardial and Pericardial Diseases. European Heart Journal, 2010, 31, 2715-2726.	1.0	408
27	Hypertrophic cardiomyopathy and sudden death in the young: Pathologic evidence of myocardial ischemia. Human Pathology, 2000, 31, 988-998.	1.1	374
28	Regulatory mutations in transforming growth factor-?3 gene cause arrhythmogenic right ventricular cardiomyopathy type 1. Cardiovascular Research, 2005, 65, 366-373.	1.8	364
29	Guidelines for autopsy investigation of sudden cardiac death: 2017 update from the Association for European Cardiovascular Pathology. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2017, 471, 691-705.	1.4	357
30	Treatment of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Circulation, 2015, 132, 441-453.	1.6	356
31	Management of Acute Myocarditis and Chronic Inflammatory Cardiomyopathy. Circulation: Heart Failure, 2020, 13, e007405.	1.6	353
32	Pathological features of COVID-19-associated myocardial injury: a multicentre cardiovascular pathology study. European Heart Journal, 2020, 41, 3827-3835.	1.0	350
33	Guidelines for autopsy investigation of sudden cardiac death. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2008, 452, 11-18.	1.4	349
34	Remodeling of myocyte gap junctions in arrhythmogenic right ventricular cardiomyopathy due to a deletion in plakoglobin (Naxos disease). Heart Rhythm, 2004, 1, 3-11.	0.3	309
35	Arrhythmogenic Cardiomyopathy. Circulation Research, 2017, 121, 784-802.	2.0	294
36	Diagnosis of arrhythmogenic cardiomyopathy: The Padua criteria. International Journal of Cardiology, 2020, 319, 106-114.	0.8	283

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37	Compound and Digenic Heterozygosity Contributes to Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American College of Cardiology, 2010, 55, 587-597.	1.2	282
38	An echocardiographic survey of primary school children for bicuspid aortic valve. American Journal of Cardiology, 2004, 93, 661-663.	0.7	274
39	Prophylactic Implantable Defibrillator in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia and No Prior Ventricular Fibrillation or Sustained Ventricular Tachycardia. Circulation, 2010, 122, 1144-1152.	1.6	272
40	Clinical profile of four families with arrhythmogenic right ventricular cardiomyopathy caused by dominant desmoplakin mutations. European Heart Journal, 2005, 26, 1666-1675.	1.0	267
41	Outcomes in Athletes with Marked ECG Repolarization Abnormalities. New England Journal of Medicine, 2008, 358, 152-161.	13.9	266
42	Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. European Heart Journal, 2020, 41, 1414-1429.	1.0	239
43	Consensus statement on surgical pathology of the aorta from the Society for Cardiovascular Pathology and the Association for European Cardiovascular Pathology: I. Inflammatory diseases. Cardiovascular Pathology, 2015, 24, 267-278.	0.7	238
44	Three-Dimensional Electroanatomic Voltage Mapping Increases Accuracy of Diagnosing Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Circulation, 2005, 111, 3042-3050.	1.6	237
45	Postmortem diagnosis in sudden cardiac death victims: macroscopic, microscopic and molecular findings. Cardiovascular Research, 2001, 50, 290-300.	1.8	231
46	Morphofunctional Abnormalities of Mitral Annulus and Arrhythmic Mitral Valve Prolapse. Circulation: Cardiovascular Imaging, 2016, 9, e005030.	1.3	226
47	European Association of Preventive Cardiology (EAPC) and European Association of Cardiovascular Imaging (EACVI) joint position statement: recommendations for the indication and interpretation of cardiovascular imaging in the evaluation of the athleteâ€™s heart. European Heart Journal, 2018, 39, 1949-1969.	1.0	224
48	Ultrastructural evidence of intercalated disc remodelling in arrhythmogenic right ventricular cardiomyopathy: an electron microscopy investigation on endomyocardial biopsies. European Heart Journal, 2006, 27, 1847-1854.	1.0	219
49	Nonischemic Left Ventricular Scar as a Substrate of Life-Threatening Ventricular Arrhythmias and Sudden Cardiac Death in Competitive Athletes. Circulation: Arrhythmia and Electrophysiology, 2016, 9, .	2.1	216
50	Screening for ryanodine receptor type 2 mutations in families with effort-induced polymorphic ventricular arrhythmias and sudden death. Journal of the American College of Cardiology, 2002, 40, 341-349.	1.2	213
51	Compound and Digenic Heterozygosity Predicts Lifetime Arrhythmic Outcome and Sudden Cardiac Death in Desmosomal Geneâ€“Related Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Cardiovascular Genetics, 2013, 6, 533-542.	5.1	209
52	Consensus statement on surgical pathology of the aorta from the Society for Cardiovascular Pathology and the Association For European Cardiovascular Pathology: II. Noninflammatory degenerative diseases â€” nomenclature and diagnostic criteria. Cardiovascular Pathology, 2016, 25, 247-257.	0.7	208
53	Aortic elasticity and size in bicuspid aortic valve syndrome. European Heart Journal, 2008, 29, 472-479.	1.0	202
54	Pathophysiology of arrhythmogenic cardiomyopathy. Nature Reviews Cardiology, 2012, 9, 223-233.	6.1	201

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55	Clinicopathological profiles of progressive heart failure in hypertrophic cardiomyopathy. European Heart Journal, 2010, 31, 2111-2123.	1.0	190
56	Myocyte necrosis underlies progressive myocardial dystrophy in mouse dsg2-related arrhythmogenic right ventricular cardiomyopathy. Journal of Experimental Medicine, 2009, 206, 1787-1802.	4.2	184
57	Intercalated disc abnormalities, reduced Na+ current density, and conduction slowing in desmoglein-2 mutant mice prior to cardiomyopathic changes. Cardiovascular Research, 2012, 95, 409-418.	1.8	180
58	Mutations in the area composita protein Î±T-catenin are associated with arrhythmogenic right ventricular cardiomyopathy. European Heart Journal, 2013, 34, 201-210.	1.0	175
59	Treatment of arrhythmogenic right ventricular cardiomyopathy/dysplasia: an international task force consensus statement. European Heart Journal, 2015, 36, ehv162.	1.0	171
60	Frequency of Bicuspid Aortic Valve in Young Male Conscripts by Echocardiogram. American Journal of Cardiology, 2005, 96, 718-721.	0.7	168
61	Three-Dimensional Electroanatomical Voltage Mapping and Histologic Evaluation of Myocardial Substrate in Right Ventricular Outflow Tract Tachycardia. Journal of the American College of Cardiology, 2008, 51, 731-739.	1.2	168
62	Quantitative assessment of endomyocardial biopsy in arrhythmogenic right ventricular cardiomyopathy/dysplasia: an in vitro validation of diagnostic criteria. European Heart Journal, 2008, 29, 2760-2771.	1.0	161
63	Multiple mutations in desmosomal proteins encoding genes in arrhythmogenic right ventricular cardiomyopathy/dysplasia. Heart Rhythm, 2010, 7, 22-29.	0.3	161
64	Altered Desmosomal Proteins in Granulomatous Myocarditis and Potential Pathogenic Links to Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2011, 4, 743-752.	2.1	161
65	Dispersion of Ventricular Depolarization-Repolarization. Circulation, 2001, 103, 3075-3080.	1.6	158
66	Incidence, Predictors, and Outcome of Conduction Disorders After Transcatheter Self-Expandable Aortic Valve Implantation. American Journal of Cardiology, 2011, 107, 747-754.	0.7	156
67	Imaging Study of Ventricular Scar in Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2012, 5, 91-100.	2.1	154
68	Mitral Valve Prolapse, Ventricular Arrhythmias, and Sudden Death. Circulation, 2019, 140, 952-964.	1.6	154
69	Adipositas cordis, fatty infiltration of the right ventricle, and arrhythmogenic right ventricular cardiomyopathy. Just a matter of fat?. Cardiovascular Pathology, 2005, 14, 37-41.	0.7	152
70	Ventricular Arrhythmias in Myocarditis. Journal of the American College of Cardiology, 2020, 75, 1046-1057.	1.2	148
71	Arrhythmogenic right ventricular cardiomyopathy/dysplasia. Orphanet Journal of Rare Diseases, 2007, 2, 45.	1.2	147
72	Prevalence of Cardiomyopathy in Italian Asymptomatic Children With Electrocardiographic T-Wave Inversion at Preparticipation Screening. Circulation, 2012, 125, 529-538.	1.6	144

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73	Congenital coronary artery anomalies: a bridge from embryology to anatomy and pathophysiologyâ€"a position statement of the development, anatomy, and pathology ESC Working Group. Cardiovascular Research, 2016, 109, 204-216.	1.8	143
74	Impact of the presence and amount of myocardial fibrosis by cardiac magnetic resonance on arrhythmic outcome and sudden cardiac death in nonischemic dilated cardiomyopathy. Heart Rhythm, 2014, 11, 856-863.	0.3	142
75	Arrhythmias in myocarditis: State of the art. Heart Rhythm, 2019, 16, 793-801.	0.3	142
76	Cocaine-related sudden death: a prospective investigation in south-west Spain. European Heart Journal, 2010, 31, 318-329.	1.0	140
77	Myocardial bridging, a frequent component of the hypertrophic cardiomyopathy phenotype, lacks systematic association with sudden cardiac death. European Heart Journal, 2009, 30, 1627-1634.	1.0	139
78	Anabolic androgenic steroids abuse and cardiac death in athletes: Morphological and toxicological findings in four fatal cases. Forensic Science International, 2012, 217, e13-e18.	1.3	129
79	Prospective Study of Cardiac Sarcoid Mimicking Arrhythmogenic Right Ventricular Dysplasia. Journal of Cardiovascular Electrophysiology, 2009, 20, 473-476.	0.8	127
80	Comprehensive multi-modality imaging approach in arrhythmogenic cardiomyopathyâ€"an expert consensus document of the European Association of Cardiovascular Imaging. European Heart Journal Cardiovascular Imaging, 2017, 18, 237-253.	0.5	123
81	Endomyocardial biopsy in arrhythmogenic right ventricular cardiomyopathy. American Heart Journal, 1996, 132, 203-206.	1.2	121
82	Molecular biology and clinical management of arrhythmogenic right ventricular cardiomyopathy/dysplasia. Heart, 2011, 97, 530-539.	1.2	120
83	Sudden cardiac death with normal heart:. Cardiovascular Pathology, 2010, 19, 321-325.	0.7	119
84	Prevalence, Characteristics, and Outcomes of COVID-19â€"Associated Acute Myocarditis. Circulation, 2022, 145, 1123-1139.	1.6	118
85	Arrhythmogenic cardiomyopathy. Orphanet Journal of Rare Diseases, 2016, 11, 33.	1.2	116
86	A Novel Circulating Noncoding Small RNA for the Detection of Acute Myocarditis. New England Journal of Medicine, 2021, 384, 2014-2027.	13.9	112
87	Cardiac involvement in patients with Becker muscular dystrophy: new diagnostic and pathophysiological insights by a CMR approach. Journal of Cardiovascular Magnetic Resonance, 2008, 10, 50.	1.6	110
88	Exercise and the Risk of Sudden Cardiac Death. Herz, 2006, 31, 553-558.	0.4	108
89	Heart Failure Association of the ESC, Heart Failure Society of America and Japanese Heart Failure Society Position statement on endomyocardial biopsy. European Journal of Heart Failure, 2021, 23, 854-871.	2.9	105
90	Arrhythmogenic right ventricular cardiomyopathy/dysplasia: is there a role for viruses?. Cardiovascular Pathology, 2006, 15, 11-17.	0.7	102

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91	Classification and histological, immunohistochemical, and molecular diagnosis of inflammatory myocardial disease. Heart Failure Reviews, 2013, 18, 673-681.	1.7	100
92	Arrhythmogenic right ventricular cardiomyopathy: An update. Cardiovascular Pathology, 2001, 10, 109-117.	0.7	99
93	3-Dimensional Echocardiography in Imaging the Tricuspid Valve. JACC: Cardiovascular Imaging, 2019, 12, 500-515.	2.3	99
94	Surgery for Primary Cardiac Tumors in Children. Circulation, 2012, 126, 22-30.	1.6	98
95	Arrhythmogenic cardiomyopathy: pathology, genetics, and concepts in pathogenesis. Cardiovascular Research, 2017, 113, 1521-1531.	1.8	98
96	Recommendations for processing cardiovascular surgical pathology specimens: a consensus statement from the Standards and Definitions Committee of the Society for Cardiovascular Pathology and the Association for European Cardiovascular Pathology. Cardiovascular Pathology, 2012, 21, 2-16.	0.7	95
97	Functional Regurgitation of Atrioventricular Valves and Atrial Fibrillation: An Elusive Pathophysiological Link Deserving Further Attention. Journal of the American Society of Echocardiography, 2020, 33, 42-53.	1.2	94
98	Prognostic Value of Endocardial Voltage Mapping in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Circulation: Arrhythmia and Electrophysiology, 2013, 6, 167-176.	2.1	92
99	Arrhythmogenic Right Ventricular Cardiomyopathy: Characterization of Left Ventricular Phenotype and Differential Diagnosis With Dilated Cardiomyopathy. Journal of the American Heart Association, 2020, 9, e014628.	1.6	92
100	Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2012, 5, 1233-1246.	2.1	90
101	Neonatal Transplantation Confers Maturation of PSC-Derived Cardiomyocytes Conducive to Modeling Cardiomyopathy. Cell Reports, 2017, 18, 571-582.	2.9	90
102	Essay: Sudden death in young athletes. Lancet, The, 2005, 366, S47-S48.	6.3	88
103	Morphologic Validation of Reperfused Hemorrhagic Myocardial Infarction by Cardiovascular Magnetic Resonance. American Journal of Cardiology, 2007, 100, 1322-1327.	0.7	87
104	Postmortem Genetic Testing for Conventional Autopsyâ€“Negative Sudden Unexplained Death. American Journal of Clinical Pathology, 2008, 129, 391-397.	0.4	86
105	Cardiac masses and tumours. Heart, 2016, 102, 1230-1245.	1.2	86
106	Definition and treatment of arrhythmogenic cardiomyopathy: an updated expert panel report. European Journal of Heart Failure, 2019, 21, 955-964.	2.9	84
107	Comparison of Clinical Features of Arrhythmogenic Right Ventricular Cardiomyopathy in Men Versus Women. American Journal of Cardiology, 2008, 102, 1252-1257.	0.7	81
108	European recommendations integrating genetic testing into multidisciplinary management of sudden cardiac death. European Journal of Human Genetics, 2019, 27, 1763-1773.	1.4	78

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109	Diagnosis of myocardial infarction at autopsy: AECVP reappraisal in the light of the current clinical classification. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 476, 179-194.	1.4	78
110	Morphologic spectrum of primary restrictive cardiomyopathy. American Journal of Cardiology, 1997, 80, 1046-1050.	0.7	77
111	Cardiomyopathies: is it time for a molecular classification?. European Heart Journal, 2004, 25, 1772-1775.	1.0	77
112	Arrhythmogenic right ventricular cardiomyopathy/dysplasia on the basis of the revised diagnostic criteria in affected families with desmosomal mutations. European Heart Journal, 2011, 32, 1097-1104.	1.0	77
113	The ARVD/C Genetic Variants Database: 2014 Update. Human Mutation, 2015, 36, 403-410.	1.1	77
114	Clinical presentation and diagnosis of myocarditis. Heart, 2015, 101, 1332-1344.	1.2	77
115	Novel Î±-Actinin 2 Variant Associated With Familial Hypertrophic Cardiomyopathy and Juvenile Atrial Arrhythmias. Circulation: Cardiovascular Genetics, 2014, 7, 741-750.	5.1	74
116	Evidence From Family Studies for Autoimmunity in Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation, 2020, 141, 1238-1248.	1.6	69
117	â€˜Hot phaseâ€™ clinical presentation in arrhythmogenic cardiomyopathy. Europace, 2021, 23, 907-917.	0.7	67
118	Clinical phenotype and diagnosis of arrhythmogenic right ventricular cardiomyopathy in pediatric patients carrying desmosomal gene mutations. Heart Rhythm, 2011, 8, 1686-1695.	0.3	66
119	Defining phenotypes and disease progression in sarcomeric cardiomyopathies: contemporary role of clinical investigations. Cardiovascular Research, 2015, 105, 409-423.	1.8	66
120	Prevention of sudden cardiac death in the young and in athletes: dream or reality?. Cardiovascular Pathology, 2010, 19, 207-217.	0.7	65
121	Right atrial volume is a major determinant of tricuspid annulus area in functional tricuspid regurgitation: a three-dimensional echocardiographic study. European Heart Journal Cardiovascular Imaging, 2021, 22, 660-669.	0.5	65
122	Desmin Mutations and Arrhythmogenic Right Ventricular Cardiomyopathy. American Journal of Cardiology, 2013, 111, 400-405.	0.7	62
123	Significance of Late Gadolinium Enhancement at Right Ventricular Attachment to Ventricular Septum in Patients With Hypertrophic Cardiomyopathy. American Journal of Cardiology, 2015, 116, 436-441.	0.7	62
124	Diagnostic Yield of Electroanatomic Voltage Mapping in Guiding Endomyocardial Biopsies. Circulation, 2020, 142, 1249-1260.	1.6	61
125	Evolving Diagnostic Criteria for Arrhythmogenic Cardiomyopathy. Journal of the American Heart Association, 2021, 10, e021987.	1.6	60
126	Strategies for the prevention of sudden cardiac death during sports. European Journal of Cardiovascular Prevention and Rehabilitation, 2011, 18, 197-208.	3.1	55

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127	Dynamics of neuroeffector coupling at cardiac sympathetic synapses. Journal of Physiology, 2018, 596, 2055-2075.	1.3	55
128	Circulating extracellular vesicles as non-invasive biomarker of rejection in heart transplant. Journal of Heart and Lung Transplantation, 2020, 39, 1136-1148.	0.3	54
129	Myocarditis and Dilated Cardiomyopathy in Athletes: Diagnosis, Management, and Recommendations for Sport Activity. Cardiology Clinics, 2007, 25, 423-429.	0.9	53
130	Phenotypic expression is a prerequisite for malignant arrhythmic events and sudden cardiac death in arrhythmogenic right ventricular cardiomyopathy. Europace, 2016, 18, 1086-1094.	0.7	50
131	Arrhythmogenic right ventricular cardiomyopathy: clinical registry and database, evaluation of therapies, pathology registry, DNA banking. European Heart Journal, 2004, 25, 531-534.	1.0	48
132	Contemporary genetic testing in inherited cardiac disease. Journal of Cardiovascular Medicine, 2018, 19, 1-11.	0.6	48
133	Predictive value of exercise testing in athletes with ventricular ectopy evaluated by cardiac magnetic resonance. Heart Rhythm, 2019, 16, 239-248.	0.3	45
134	Revisiting definition and classification of cardiomyopathies in the era of molecular medicine. European Heart Journal, 2007, 29, 144-146.	1.0	44
135	T-Cellâ€“Mediated Inflammatory Activity in the Stellate Ganglia of Patients With Ion-Channel Disease and Severe Ventricular Arrhythmias. Circulation: Arrhythmia and Electrophysiology, 2014, 7, 224-229.	2.1	43
136	A founderMYBPC3mutation results in HCM with a high risk of sudden death after the fourth decade of life. Journal of Medical Genetics, 2015, 52, 338-347.	1.5	41
137	Importance of genotype for risk stratification in arrhythmogenic right ventricular cardiomyopathy using the 2019 ARVC risk calculator. European Heart Journal, 2022, 43, 3053-3067.	1.0	41
138	Identification of a PKP2 gene deletion in a family with arrhythmogenic right ventricular cardiomyopathy. European Journal of Human Genetics, 2013, 21, 1226-1231.	1.4	39
139	Loss of cardiac Wnt/Î²-catenin signalling in desmoplakin-deficient AC8 zebrafish models is rescuable by genetic and pharmacological intervention. Cardiovascular Research, 2018, 114, 1082-1097.	1.8	39
140	Inflammation as a Predictor of RecurrentÂVentricular Tachycardia After Ablation in Patients With Myocarditis. Journal of the American College of Cardiology, 2020, 76, 1644-1656.	1.2	39
141	Homozygous Desmocollin-2 Mutations and Arrhythmogenic Cardiomyopathy. American Journal of Cardiology, 2015, 116, 1245-1251.	0.7	38
142	Whole-Exome Sequencing Identifies Pathogenic Variants in TJP1 Gene Associated With Arrhythmogenic Cardiomyopathy. Circulation Genomic and Precision Medicine, 2018, 11, e002123.	1.6	38
143	Feasibility of postmortem examination in the era of COVID-19 pandemic: the experience of a Northeast Italy University Hospital. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 477, 341-347.	1.4	38
144	Cardiac hypertrophy at autopsy. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2021, 479, 79-94.	1.4	38

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145	The metamorphosis of myocardial infarction following coronary recanalization. Cardiovascular Pathology, 2010, 19, 22-28.	0.7	37
146	Cardiac sympathetic innervation network shapes the myocardium by locally controlling cardiomyocyte size through the cellular proteolytic machinery. Journal of Physiology, 2019, 597, 3639-3656.	1.3	37
147	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). International Journal of Cardiology, 2022, 357, 55-71.	0.8	36
148	Arrhythmogenic Cardiomyopathy. Circulation: Cardiovascular Genetics, 2011, 4, 318-326.	5.1	35
149	Prevalence and clinical meaning of isolated increase of QRS voltages in hypertrophic cardiomyopathy versus athlete's heart: Relevance to athletic screening. International Journal of Cardiology, 2013, 168, 4494-4497.	0.8	35
150	Comparison of Features of Fatal Versus Nonfatal Cardiac Arrest in Patients With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Cardiology, 2017, 120, 111-117.	0.7	35
151	Large Genomic Rearrangements of Desmosomal Genes in Italian Arrhythmogenic Cardiomyopathy Patients. Circulation: Arrhythmia and Electrophysiology, 2017, 10, .	2.1	35
152	Arrhythmogenic cardiomyopathy: pathogenesis, pro-arrhythmic remodelling, and novel approaches for risk stratification and therapy. Cardiovascular Research, 2020, 116, 1571-1584.	1.8	35
153	Systemic sclerosis myocarditis has unique clinical, histological and prognostic features: a comparative histological analysis. Rheumatology, 2020, 59, 2523-2533.	0.9	35
154	Anomalous origin of coronary arteries from the â€œwrongâ€•sinus in athletes: Diagnosis and management strategies. International Journal of Cardiology, 2018, 252, 13-20.	0.8	33
155	Impact of systemic immune-mediated diseases on clinical features and prognosis of patients with biopsy-proved myocarditis. International Journal of Cardiology, 2019, 280, 110-116.	0.8	33
156	Filamin-C variant-associated cardiomyopathy: AÂpooled analysis of individual patient data to evaluate the clinical profile and risk of sudden cardiac death. Heart Rhythm, 2022, 19, 235-243.	0.3	33
157	Arrhythmogenic right ventricular cardiomyopathy. Herz, 2015, 40, 600-606.	0.4	32
158	Immunosuppressive Therapy and Risk Stratification of Patients With Myocarditis Presenting With Ventricular Arrhythmias. JACC: Clinical Electrophysiology, 2020, 6, 1221-1234.	1.3	32
159	The changing spectrum of arrhythmogenic (right ventricular) cardiomyopathy. Cell and Tissue Research, 2012, 348, 319-323.	1.5	31
160	Long-term follow-up analysis of a highly characterized arrhythmogenic cardiomyopathy cohort with classical and non-classical phenotypesâ€“a real-world assessment of a novel prediction model: does the subtype really matter. Europace, 2020, 22, 797-805.	0.7	31
161	Myocardial Infarction in a Patient with Hypertrophic Cardiomyopathy. New England Journal of Medicine, 2000, 342, 593-594.	13.9	30
162	Liver histopathology in COVID-19 patients: A mono-Institutional series of liver biopsies and autopsy specimens. Pathology Research and Practice, 2021, 221, 153451.	1.0	30

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163	Twenty years of progress and beckoning frontiers in cardiovascular pathology. Cardiovascular Pathology, 2005, 14, 165-169.	0.7	29
164	Characteristics of Patients With Arrhythmogenic Left Ventricular Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2020, 13, e009005.	2.1	29
165	Heart Failure Association, Heart Failure Society of America, and Japanese Heart Failure Society Position Statement on Endomyocardial Biopsy. Journal of Cardiac Failure, 2021, 27, 727-743.	0.7	29
166	Pitfalls in the Diagnosis of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. American Journal of Cardiology, 2010, 105, 1036-1039.	0.7	28
167	Arrhythmogenic Right Ventricular Cardiomyopathy: What's in a Name? From a Congenital Defect (Dysplasia) to a Genetically Determined Cardiomyopathy (Dystrophy). American Journal of Cardiology, 2010, 106, 275-277.	0.7	27
168	Clinical utility gene card for: arrhythmogenic right ventricular cardiomyopathy (ARVC). European Journal of Human Genetics, 2014, 22, 293-293.	1.4	27
169	Diagnostic Criteria, Genetics, and Molecular Basis of Arrhythmogenic Cardiomyopathy. Heart Failure Clinics, 2018, 14, 201-213.	1.0	27
170	TGF-beta1 pathway activation and adherens junction molecular pattern in nonsyndromic mitral valve prolapse. Cardiovascular Pathology, 2015, 24, 359-367.	0.7	25
171	The Immunopathological and Histological Landscape of COVID-19-Mediated Lung Injury. International Journal of Molecular Sciences, 2021, 22, 974.	1.8	25
172	Left atrial myxoma in a child. Cardiovascular Pathology, 2003, 12, 233-236.	0.7	24
173	Syncope as a Warning Symptom of Sudden Cardiac Death in Athletes. Cardiology Clinics, 2015, 33, 423-432.	0.9	23
174	Structurally Normal Hearts Are Uncommonly Associated With Sudden Deaths in Athletes and Young People. Journal of the American College of Cardiology, 2019, 73, 3031-3032.	1.2	23
175	Giant congenital aortic aneurysm with cleft sternum in a neonate: pathological and surgical considerations for optimal management. Cardiovascular Pathology, 2010, 19, 183-186.	0.7	22
176	Nonischemic Left Ventricular Scar. Circulation, 2014, 130, e180-2.	1.6	22
177	Diagnostic value and prognostic implications of early cardiac magnetic resonance in survivors of out-of-hospital cardiac arrest. Heart Rhythm, 2018, 15, 1031-1041.	0.3	22
178	Mitral Annulus Disjunction. Journal of the American College of Cardiology, 2018, 72, 1610-1612.	1.2	22
179	Hybrid FDG-PET/MR or FDG-PET/CT to Detect Disease Activity in Patients With Persisting Arrhythmias After Myocarditis. JACC: Cardiovascular Imaging, 2021, 14, 288-292.	2.3	22
180	Clinical profile and long-term follow-up of a cohort of patients with desmoplakin cardiomyopathy. Heart Rhythm, 2022, 19, 1315-1324.	0.3	22

#	Article	IF	Citations
181	A novel kind of tumor type-characteristic junction: plakophilin-2 as a major protein of adherens junctions in cardiac myxomata. Modern Pathology, 2010, 23, 1429-1437.	2.9	21
182	A microRNA Expression Profile as Non-Invasive Biomarker in a Large Arrhythmogenic Cardiomyopathy Cohort. International Journal of Molecular Sciences, 2020, 21, 1536.	1.8	21
183	Cadherin 2-Related Arrhythmogenic Cardiomyopathy. Circulation Genomic and Precision Medicine, 2021, 14, e003097.	1.6	21
184	Screening of competitive athletes to prevent sudden death. Heart, 2013, 99, 304-306.	1.2	20
185	Morphologic studies of cell endogenous repopulation in decellularized aortic and pulmonary homografts implanted in sheep. Cardiovascular Pathology, 2015, 24, 102-109.	0.7	20
186	Arrhythmogenic Right Ventricular Cardiomyopathy: Risk Stratification and Indications for Defibrillator Therapy. Current Cardiology Reports, 2016, 18, 57.	1.3	20
187	Sudden cardiac death in an Italian competitive athlete: Pre-participation screening and cardiovascular emergency care are both essential. International Journal of Cardiology, 2016, 206, 84-86.	0.8	20
188	Efficacy and safety of mycophenolate mofetil in patients with virus-negative lymphocytic myocarditis: A prospective cohort study. Journal of Autoimmunity, 2020, 106, 102330.	3.0	20
189	Autopsy examination in sudden cardiac death: a current perspective on behalf of the Association for European Cardiovascular Pathology. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2021, 478, 687-693.	1.4	20
190	Clinically Suspected and Biopsy-Proven Myocarditis Temporally Associated with SARS-CoV-2 Infection. Annual Review of Medicine, 2022, 73, 149-166.	5.0	20
191	Predictors of relapse, death or heart transplantation in myocarditis before the introduction of immunosuppression: negative prognostic impact of female gender, fulminant onset, lower ejection fraction and serum autoantibodies. European Journal of Heart Failure, 2022, 24, 1033-1044.	2.9	19
192	Clinical application of CMR in cardiomyopathies: evolving concepts and techniques. Heart Failure Reviews, 2023, 28, 77-95.	1.7	19
193	AECVP and SCVP 2009 Recommendations for Training in Cardiovascular Pathology. Cardiovascular Pathology, 2010, 19, 129-135.	0.7	18
194	Cardiac Magnetic Resonance Features ofÂBiopsy-Proven Endomyocardial Diseases. JACC: Cardiovascular Imaging, 2014, 7, 309-312.	2.3	18
195	The clinico-pathological conference, based upon Giovanni Battista Morgagniâ€™s legacy, remains of fundamental importance even in the era of the vanishing autopsy. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2015, 467, 249-254.	1.4	18
196	Optimized protocol for immunostaining of experimental GFP-expressing and human hearts. Histochemistry and Cell Biology, 2016, 146, 407-419.	0.8	18
197	Anatomy of the cavotricuspid isthmus for radiofrequency ablation in typical atrial flutter. Heart Rhythm, 2019, 16, 1611-1618.	0.3	18
198	Programmed ventricular stimulation in patients with active vs previous arrhythmic myocarditis. Journal of Cardiovascular Electrophysiology, 2020, 31, 692-701.	0.8	18

#	Article	IF	Citations
199	The 2021 WHO Classification of Tumors of the Heart. Journal of Thoracic Oncology, 2022, 17, 510-518.	0.5	18
200	Candida endocarditis complicating transapical aortic valve implantation. European Heart Journal, 2011, 32, 2265-2265.	1.0	17
201	Why and how to support screening strategies to prevent sudden death in athletes. Cell and Tissue Research, 2012, 348, 315-318.	1.5	17
202	Update on cardiomyopathies and sudden cardiac death. Forensic Sciences Research, 2019, 4, 202-210.	0.9	17
203	Serial Versus Single Cardiovascular Screening of Adolescent Athletes. Circulation, 2021, 143, 1729-1731.	1.6	17
204	Sudden coronary death in the young: Evidence of contractile phenotype of smooth muscle cells in the culprit atherosclerotic plaque. International Journal of Cardiology, 2018, 264, 1-6.	0.8	16
205	The Spectrum of COVID-19-Associated Myocarditis: A Patient-Tailored Multidisciplinary Approach. Journal of Clinical Medicine, 2021, 10, 1974.	1.0	16
206	Hypertrophic Cardiomyopathy and Primary Restrictive Cardiomyopathy: Similarities, Differences and Phenocopies. Journal of Clinical Medicine, 2021, 10, 1954.	1.0	16
207	Ventricular arrhythmias in athletes: Role of a comprehensive diagnostic workup. Heart Rhythm, 2022, 19, 90-99.	0.3	16
208	Clinical Applications of FDG-PET Scan in Arrhythmic Myocarditis. JACC: Cardiovascular Imaging, 2022, 15, 1771-1780.	2.3	16
209	Anomalous aortic origin of coronary arteries: Early results on clinical management from an international multicenter study. International Journal of Cardiology, 2019, 291, 189-193.	0.8	15
210	A targeted next-generation gene panel reveals a novel heterozygous nonsense variant in the TP63 gene in patients with arrhythmogenic cardiomyopathy. Heart Rhythm, 2019, 16, 773-780.	0.3	15
211	Sudden Cardiac Death in the Young and Athletes. , 2016, , .		14
212	Endomyocardial Biopsy: The Forgotten Piece in the Arrhythmogenic Cardiomyopathy Puzzle. Journal of the American Heart Association, 2021, 10, e021370.	1.6	14
213	Septal Late Gadolinium Enhancement and Arrhythmic Risk in Genetic and Acquired Non-Ischaemic Cardiomyopathies. Heart Lung and Circulation, 2020, 29, 1356-1365.	0.2	13
214	The genetic architecture of Plakophilin 2 cardiomyopathy. Genetics in Medicine, 2021, 23, 1961-1968.	1.1	13
215	Myocardial Tissue Characterization in Arrhythmogenic Cardiomyopathy. JACC: Cardiovascular Imaging, 2021, 14, 1675-1678.	2.3	13
216	Nerve growth factor transfer from cardiomyocytes to innervating sympathetic neurons activates TrkA receptors at the neuroâ€cardiac junction. Journal of Physiology, 2022, 600, 2853-2875.	1.3	13

#	Article	IF	Citations
217	Pharmacotherapy and Other Therapeutic Modalities for Managing Arrhythmogenic Right Ventricular Cardiomyopathy. Cardiovascular Drugs and Therapy, 2015, 29, 171-177.	1.3	12
218	Story telling of myocarditis. International Journal of Cardiology, 2019, 294, 61-64.	0.8	12
219	How to improve therapy in myocarditis: role of cardiovascular magnetic resonance and of endomyocardial biopsy. European Heart Journal Supplements, 2019, 21, B19-B22.	0.0	12
220	Arrhythmogenic Cardiomyopathy. European Heart Journal, 2020, 41, 4457-4462.	1.0	12
221	Prior myocarditis and ventricular arrhythmias: The importance of scar pattern. Heart Rhythm, 2021, 18, 589-596.	0.3	12
222	Cardiac magnetic resonance imaging of arrhythmogenic cardiomyopathy: evolving diagnostic perspectives. European Radiology, 2023, 33, 270-282.	2.3	12
223	Mechanical dispersion and arrhythmic mitral valve prolapse: substrate and trigger in electrical instability. Heart, 2019, 105, 1053-1054.	1.2	11
224	Recurrent autoimmune myocarditis in a young woman during the coronavirus disease 2019 pandemic. ESC Heart Failure, 2021, 8, 756-760.	1.4	11
225	Immunosuppressive therapy in childhoodâ€onset arrhythmogenic inflammatory cardiomyopathy. PACE - Pacing and Clinical Electrophysiology, 2021, 44, 552-556.	0.5	11
226	Imaging Phenotype vs Genotype in Nonhypertrophic Heritable Cardiomyopathies. Circulation: Cardiovascular Imaging, 2010, 3, 753-765.	1.3	10
227	Relation of Aortic Valve Weight to Severity of Aortic Stenosis. American Journal of Cardiology, 2011, 107, 741-746.	0.7	10
228	Concealed Metastatic Lung Carcinoma Presenting as Acute Coronary Syndrome With Progressive Conduction Abnormalities. Circulation, 2012, 125, e499-502.	1.6	10
229	Morphology of right atrial appendage for permanent atrial pacing and risk of iatrogenic perforation of the aorta by active fixation lead. Heart Rhythm, 2015, 12, 744-750.	0.3	10
230	Cardiovascular medicine in Morgagni's De sedibus: dawn of cardiovascular pathology. Cardiovascular Pathology, 2016, 25, 443-452.	0.7	10
231	Co-inheritance of mutations associated with arrhythmogenic cardiomyopathy and hypertrophic cardiomyopathy. European Journal of Human Genetics, 2017, 25, 1165-1169.	1.4	10
232	Arrhythmogenic Cardiomyopathy Is a Multicellular Disease Affecting Cardiac and Bone Marrow Mesenchymal Stromal Cells. Journal of Clinical Medicine, 2021, 10, 1871.	1.0	10
233	Neuropeptide Y promotes adipogenesis of human cardiac mesenchymal stromal cells in arrhythmogenic cardiomyopathy. International Journal of Cardiology, 2021, 342, 94-102.	0.8	10
234	Renin-angiotensin-aldosterone system inhibition in patients affected by heart failure: efficacy, mechanistic effects and practical use of sacubitril/valsartan. Position Paper of the Italian Society of Cardiology. European Journal of Internal Medicine, 2022, 102, 8-16.	1.0	10

#	Article	IF	Citations
235	ST-Segment Elevation and Sudden Death in the Athlete. Cardiac Electrophysiology Clinics, 2013, 5, 73-84.	0.7	9
236	Left ventricular outflow tract rupture during transcatheter aortic valve implantation: anatomic evidence of the vulnerable area. Cardiovascular Pathology, 2017, 29, 7-10.	0.7	9
237	Right Ventricular Junctional Late Gadolinium Enhancement Correlates With Outcomes in Pulmonary Hypertension. JACC: Cardiovascular Imaging, 2019, 12, 936-938.	2.3	9
238	Functional modulation of atrio-ventricular conduction by enhanced late sodium current and calcium-dependent mechanisms in Scn5a1798insD/+ mice. Europace, 2020, 22, 1579-1589.	0.7	9
239	Right ventricular endomyocardial biopsy in patients with cardiac magnetic resonance showing left ventricular myocarditis. Journal of Cardiovascular Medicine, 2021, 22, 560-566.	0.6	9
240	Prognostic Significance of Feature-Tracking Right Ventricular Global Longitudinal Strain in Non-ischemic Dilated Cardiomyopathy. Frontiers in Cardiovascular Medicine, 2021, 8, 765274.	1.1	9
241	The 2020 â€œPadua Criteriaâ€•for Diagnosis and Phenotype Characterization of Arrhythmogenic Cardiomyopathy in Clinical Practice. Journal of Clinical Medicine, 2022, 11, 279.	1.0	9
242	Autopsy in the era of advanced cardiovascular imaging. European Heart Journal, 2022, 43, 2461-2468.	1.0	9
243	Sudden coronary death--not always atherosclerotic. Heart, 2010, 96, 1084-1085.	1.2	8
244	Follow-Up with Exercise Test of Effort-Induced Ventricular Arrhythmias Linked to Ryanodine Receptor Type 2 Gene Mutations. American Journal of Cardiology, 2012, 109, 1015-1019.	0.7	8
245	Is it time for plakoglobin immune-histochemical diagnostic test for arrhythmogenic cardiomyopathy in the routine pathology practice?. Cardiovascular Pathology, 2013, 22, 312-313.	0.7	8
246	Novel Basic Science Insights to Improve the Management of Heart Failure: Review of the Working Group on Cellular and Molecular Biology of the Heart of the Italian Society of Cardiology. International Journal of Molecular Sciences, 2020, 21, 1192.	1.8	8
247	A standardized postmortem protocol to assess the real burden of sudden infant death syndrome. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 477, 177-183.	1.4	8
248	Novel pathogenic role for galectin-3 in early disease stages of arrhythmogenic cardiomyopathy. Heart Rhythm, 2021, 18, 1394-1403.	0.3	8
249	Additional diagnostic value of cardiac magnetic resonance feature tracking in patients with biopsy-proven arrhythmogenic cardiomyopathy. International Journal of Cardiology, 2021, 339, 203-210.	0.8	8
250	Sudden cardiac death, borderline myocarditis and molecular diagnosis: evidence or assumption?. Medicine, Science and the Law, 2011, 51, 27-29.	0.6	7
251	Aortic and Pulmonary Root Aneurysms in a Child With Loeys-Dietz Syndrome. Annals of Thoracic Surgery, 2016, 101, 1193-1195.	0.7	7
252	Reply. Journal of the American College of Cardiology, 2020, 76, 126-128.	1.2	7

#	Article	IF	Citations
253	Small Vessel Disease: Another Component of the Hypertrophic Cardiomyopathy Phenotype Not Necessarily Associated with Fibrosis. Journal of Clinical Medicine, 2021, 10, 575.	1.0	7
254	Screening professional athletes for cardiovascular diseases at risk of cardiac arrest. European Heart Journal, 2022, 43, 251-254.	1.0	7
255	Cardiac magnetic resonance features of left dominant arrhythmogenic cardiomyopathy: differential diagnosis with myocarditis. International Journal of Cardiovascular Imaging, 2022, 38, 397-405.	0.7	7
256	Spontaneous coronary artery dissection mimicking aortic dissection. Cardiovascular Pathology, 2004, 13, 330-333.	0.7	6
257	Arrhythmogenic Right Ventricular Cardiomyopathy. Cardiac Electrophysiology Clinics, 2010, 2, 571-586.	0.7	6
258	Multifocal Purkinjeâ€“like hamartoma and junctional ectopic tachycardia with a rapidly fatal outcome in a newborn. Heart Rhythm, 2014, 11, 1264-1266.	0.3	6
259	Evidence of aldosterone synthesis in human myocardium in acute myocarditis. International Journal of Cardiology, 2019, 275, 114-119.	0.8	6
260	Cardiac arrest at rest and during sport activity: causes and prevention. European Heart Journal Supplements, 2020, 22, E20-E24.	0.0	6
261	Novel Missense Variant in MYL2 Gene Associated With Hypertrophic Cardiomyopathy Showing High Incidence of Restrictive Physiology. Circulation Genomic and Precision Medicine, 2020, 13, e002824.	1.6	6
262	The three fetal shunts: A story of wrong eponyms. Journal of Anatomy, 2021, 238, 1028-1035.	0.9	6
263	Pathology of the Aorta and Aorta as Homograft. Journal of Cardiovascular Development and Disease, 2021, 8, 76.	0.8	6
264	Arrhythmogenic cardiomyopathy: the ongoing search for mechanism-driven therapies meets extracellular vesicles. European Heart Journal, 2021, 42, 3572-3574.	1.0	6
265	Autopsy and Endomyocardial Biopsy Findings. , 2007, , 29-44.		6
266	Continuous Electrical Monitoring in Patients with Arrhythmic Myocarditis: Insights from a Referral Center. Journal of Clinical Medicine, 2021, 10, 5142.	1.0	6
267	Molecular genetic testing in athletes: Why and when a position statement from the Italian Society of Sports Cardiology. International Journal of Cardiology, 2022, 364, 169-177.	0.8	6
268	Pathobiology of Arrhythmogenic Cardiomyopathy. Cardiac Electrophysiology Clinics, 2011, 3, 193-204.	0.7	5
269	Opacification patterns of cardiac masses using low-mechanical index contrast echocardiography: comparison with histopathological findings. Cardiovascular Pathology, 2017, 30, 72-77.	0.7	5
270	Sequencing of NOTCH1 gene in an Italian population with bicuspid aortic valve: Preliminary results from the GISSI OUTLIERS VAR study. Gene, 2019, 715, 143970.	1.0	5

#	Article	IF	Citations
271	Thyroid dysfunction in adult patients with biopsy-proved myocarditis: Screening and characterization. European Journal of Internal Medicine, 2020, 71, 98-100.	1.0	5
272	The Role of MicroRNAs in Arrhythmogenic Cardiomyopathy: Biomarkers or Innocent Bystanders of Disease Progression?. International Journal of Molecular Sciences, 2020, 21, 6434.	1.8	5
273	Structural valve deterioration and mode of failure of stentless bioprosthetic valves. Cardiovascular Pathology, 2021, 51, 107301.	0.7	5
274	The multiple faces of autoimmune/immuneâ€mediated myocarditis in children: a biopsyâ€proven case series treated with immunosuppressive therapy. ESC Heart Failure, 2021, 8, 1604-1609.	1.4	5
275	Sudden Death and Coronary Artery Anomalies. Frontiers in Cardiovascular Medicine, 2021, 8, 636589.	1.1	5
276	The missing pieces in the puzzle of arrhythmic mitral valve prolapse: Papillary muscles, mitral annulus dysjunction, and myocardial scarring. Heart Rhythm, 2021, 18, 577-578.	0.3	5
277	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. Journal of Cardiovascular Medicine, 2020, 21, 927-943.	0.6	5
278	Multi-Design Differential Expression Profiling of COVID-19 Lung Autopsy Specimens Reveals Significantly Deregulated Inflammatory Pathways and SFTPC Impaired Transcription. Cells, 2022, 11, 1011.	1.8	5
279	In vivo Lesion Index (LSI) validation in percutaneous radiofrequency catheter ablation. Journal of Cardiovascular Electrophysiology, 2022, 33, 874-882.	0.8	5
280	To the Editorâ€”Incidence of Sports-Related Sudden Cardiac Death: The Danish Paradox. Heart Rhythm, 2010, 7, 1917-1918.	0.3	4
281	Hyperacute Valve Thrombosis After Transapical Transcatheter Aortic Valve Replacement in a Patient With PolycythemiaÂVera. JACC: Cardiovascular Interventions, 2016, 9, 1746-1747.	1.1	4
282	â€œPrimaryâ€dilated hearts. International Journal of Cardiology, 2018, 257, 366-370.	0.8	4
283	Filamin A mitral valve disease: prolapse, dystrophy, or dysplasia?. European Heart Journal, 2018, 39, 1278-1280.	1.0	4
284	Cardiac arrest due to acute massive aortic root thrombosis after pericardial bioprosthetic aortic valve replacement. Cardiovascular Pathology, 2019, 41, 8-10.	0.7	4
285	A de novo ryanodine receptor 2 gene variant in a case of sudden cardiac death. International Journal of Legal Medicine, 2020, 134, 619-623.	1.2	4
286	Transcatheter ablation of the atrioventricular junction in refractory atrial fibrillation: A clinicopathological study. International Journal of Cardiology, 2021, 329, 99-104.	0.8	4
287	A rare cause of effusiveâ€“constrictive pericarditis. ESC Heart Failure, 2021, 8, 4313-4317.	1.4	4
288	Autoimmune Myocarditis and Arrhythmogenic Mitral Valve Prolapse: An Unexpected Overlap Syndrome. Journal of Cardiovascular Development and Disease, 2021, 8, 151.	0.8	4

#	Article	IF	Citations
289	Age is a risk factor for maladaptive changes of the pulmonary root in rats exposed to increased pressure loading. Cardiovascular Pathology, 2012, 21, 199-205.	0.7	3
290	Corrado and colleagues reply to Van Brabandt and colleagues. BMJ, The, 2016, 354, i3631.	3.0	3
291	Response to Letters Regarding Article, â€œArrhythmic Mitral Valve Prolapse and Sudden Cardiac Deathâ€. Circulation, 2016, 133, e460.	1.6	3
292	Aortic stenting in the growing sheep causes aortic endothelial dysfunction but not hypertension: Clinical implications for coarctation repair. Congenital Heart Disease, 2017, 12, 74-83.	0.0	3
293	Genetics in cardiovascular diseases. Italian Journal of Medicine, 2019, 13, 137-151.	0.2	3
294	Clinical and Molecular Data Define a Diagnosis of Arrhythmogenic Cardiomyopathy in a Carrier of a Brugada-Syndrome-Associated PKP2 Mutation. Genes, 2020, 11, 571.	1.0	3
295	The fascinating discovery of the electrical system in the heart: A story telling. International Journal of Cardiology, 2020, 317, 81-85.	0.8	3
296	A patient with rapid worsening dyspnoea during Covid-19 pandemic. European Heart Journal, 2021, 42, 717-718.	1.0	3
297	Coronary Arteries: Normal Anatomy With Historical Notes and Embryology of Main Stems. Frontiers in Cardiovascular Medicine, 2021, 8, 649855.	1.1	3
298	Clinicopathological insights from early structural valve deterioration of a surgical and transcatheter valveâ€inâ€valve mitral bioprotheses. Journal of Cardiac Surgery, 2021, 36, 4427-4430.	0.3	3
299	Immune checkpoint inhibitor myocarditis: a call for standardized histopathologic criteria. European Journal of Heart Failure, 2021, 23, 1736-1738.	2.9	3
300	Specific Cardiovascular Diseases and Competitive Sports Participation: Coronary Anomalies and Myocardial Bridging atÂRisk of Sudden Death. , 2020, , 403-421.		3
301	The Northeast Italy, Veneto Region Experience. , 2016, , 171-181.		3
302	Definition, Epidemiology, and Pathophysiologic Mechanisms. , 2016, , 3-20.		3
303	Multiple mycotic aneurysms of the aortic root after aortic valve replacement. Cardiovascular Pathology, 2020, 44, 107152.	0.7	2
304	Recurrent pulmonary embolization of inflammatory myofibroblastic tumor: a case report. Cardiovascular Pathology, 2021, 50, 107270.	0.7	2
305	Atherosclerotic Plaque Healing. New England Journal of Medicine, 2021, 384, 292-294.	13.9	2
306	When self-medication goes wrong: the case of argyria at the Padua Morgagni Museum of Pathology. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2021, , 1.	1.4	2

#	Article	IF	Citations
307	800 years of research at the University of Padua (1222â€“2022): contemporary insights into Sports Cardiology. European Heart Journal, 2022, 43, 1787-1791.	1.0	2
308	Sudden arrhythmic death and the cardiomyopathies: Molecular genetics and pathology. Diagnostic Histopathology, 2010, 16, 31-42.	0.2	1
309	Assessing the Significance of Pathogenic Mutations and Autopsy Findings in the Light of 2010 Arrhythmogenic Right Ventricular Cardiomyopathy Diagnostic Criteria. Circulation: Cardiovascular Genetics, 2012, 5, 384-386.	5.1	1
310	Cardiac Tumors: From Autoptic Observations to Surgical Pathology in the Era of Advanced Cardiac Imaging. , 2013, , 1-22.		1
311	Primary Cardiac Tumors in the Pediatric Age. , 2013, , 59-71.		1
312	Pathologic Substrates of Sudden Cardiac Death During Sports. Cardiac Electrophysiology Clinics, 2013, 5, 1-11.	0.7	1
313	Molecular Pathobiology of Myocarditis. , 2014, , 135-159.		1
314	Arrhythmogenic Cardiomyopathy: History and Pathology. , 2016, , 5-33.		1
315	Are we ready for a â€œtissue identity cardâ€•of inherited cardiomyopathies?. Heart Rhythm, 2017, 14, 1033-1034.	0.3	1
316	Biopsy-Proven Lymphocytic Myocarditis With Heart Failure in a Middle-Aged Female Patient With MixedÂConnective Tissue Disease. JACC: Case Reports, 2019, 1, 171-174.	0.3	1
317	Tullio Terni (1888â€“1946): The life of a neurocardioanatomist with a tragic epilogue. International Journal of Cardiology, 2019, 289, 153-156.	0.8	1
318	Heart failure in arrhythmogenic cardiomyopathy: is phenotypic variability justÂaÂmatter of genetics?. European Journal of Heart Failure, 2019, 21, 801-802.	2.9	1
319	Endomyocardial fibrosis and myocardial infarction leading to diastolic and systolic dysfunction requiring transplantation. Cardiovascular Pathology, 2019, 38, 21-24.	0.7	1
320	Sudden death with massive hemoptysis from aortobronchial fistula. Cardiovascular Pathology, 2020, 44, 107158.	0.7	1
321	Fabrici and the functional power of the image. International Journal of Cardiology, 2020, 316, 252-256.	0.8	1
322	Reply to â€œsignal averaged electrocardiogram findings among right ventricular arrhtyhmogenic cardiomyopathy (ARVC) patients: Do they have a place in ARVC management?â€• International Journal of Cardiology, 2021, 327, 155.	0.8	1
323	Ultra-rapid regrowth of urothelial cardiac metastasis after debulking surgery. Cardiovascular Pathology, 2021, 53, 107334.	0.7	1
324	Isolated Dissection of the Ductus Arteriosus Associated with Sudden Unexpected Intrauterine Death. Journal of Cardiovascular Development and Disease, 2021, 8, 91.	0.8	1

#	Article	IF	Citations
325	How to look at adult congenital left ventricular outpouchings: a step-by-step approach using cardiac magnetic resonance. European Heart Journal Cardiovascular Imaging, 2022, 23, 1001-1005.	0.5	1
326	Pancarditis as the Clinical Presentation of Eosinophilic Granulomatosis with Polyangiitis: A Multimodality Approach to Diagnosis. Neurology International, 2022, 12, 133-141.	0.2	1
327	Sudden Death in Athletes. , 0, , 189-202.		0
328	Arrhythmogenic Cardiomyopathy: AÂHistorical Overview. Cardiac Electrophysiology Clinics, 2011, 3, 179-191.	0.7	0
329	Molecular Pathology and Genetic Testing. , 2016, , 165-169.		0
330	Conduction System Disease. , 2016, , 123-135.		0
331	Ion Channel Disease. , 2016, , 143-147.		0
332	Key role of the post-mortem in sudden cardiac death. Progress in Pediatric Cardiology, 2017, 45, 14-16.	0.2	0
333	Sudden arrhythmic death and cardiomyopathies in the young: a molecular and pathology overview. Diagnostic Histopathology, 2017, 23, 486-498.	0.2	0
334	Reply. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2018, 472, 295-296.	1.4	0
335	Cardiac magnetic resonance imaging in patients with a nonconditional cardiac implantable device: Good-quality images besides safety are essential. Heart Rhythm, 2018, 15, 226-227.	0.3	0
336	Arrhythmogenic Cardiomyopathy. , 2018, , 631-639.		0
337	William Harvey in quarantine and lazaretto when back to Italy at the time of 1630 plague. European Heart Journal, 2021, 42, 4613-4616.	1.0	0
338	An updated approach to sudden cardiac death, the AECVP perspective. International Journal of Legal Medicine, 2021, 135, 1555-1557.	1.2	0
339	Ventricular Arrhythmias in Athletes: Role of a Comprehensive Diagnostic Workup. SSRN Electronic Journal, 0, , .	0.4	0
340	Arrhythmogenic Cardiomyopathy. , 2022, , 57-67.		0
341	Valve Disease. , 2016, , 109-122.		0
342	Inherited Cardiac Muscle Disorders: Arrhythmogenic Right Ventricular Cardiomyopathy. , 2018, , 367-388.		0

#	Article	IF	Citations
343	Interventional Anatomy of Aortic Valve. , 2019, , 11-19.		0
344	Classification, Histopathology, Immunohistology, and Molecular Diagnosis of Myocarditis. , 2020, , 1-17.		0
345	Sudden Death in Athletes: Autoptic Findings. , 2022, , 1-22.		0
346	Echocardiography and Multimodality Cardiac Imaging in COVID-19 Patients. Journal of Cardiovascular Echography, 2020, 30, S18-S24.	0.1	0
347	Guidelines for autopsy investigation of sudden cardiac death. Ceskoslovenska Patologie, 2010, 46, 43-8.	0.1	0
348	TEMPORARY REMOVAL: Sudden death of an athlete due to concealed arrhythmogenic cardiomyopathy: Why preparticipation screening did fail. International Journal of Cardiology, 2022, , .	0.8	0
349	325â€ƒDealing with cardiac amyloidosis diagnosis: keep calm and use the magnifying glasses!. European Heart Journal Supplements, 2021, 23, .	0.0	0
350	332â€ƒClinical and prognostic significance of junctional late gadolinium enhancement in patients with non-ischaemic cardiomyopathy. European Heart Journal Supplements, 2021, 23, .	0.0	0
351	383â€ƒECG in biopsy-proven and clinically suspected myocarditis: morpho-functional correlates and prognostic implications. European Heart Journal Supplements, 2021, 23, .	0.0	0
352	Mechanical and Structural Adaptation of the Pulmonary Root after Ross Operation in a Murine Model. Journal of Clinical Medicine, 2022, 11, 3742.	1.0	0