Silvia G Priori

List of Publications by Year
in descending order

477	96,143	384 134	243 303
papers	citations	h-index	g-index

509	509	509	47236
all docs	docs citations	times ranked	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	2015 ESC Guidelines for the management of patients with ventricular arrhythmias and the prevention of sudden cardiac death. European Heart Journal, 2015, 36, 2793-2867.	1.0	3,187
3	Universal Definition of Myocardial Infarction. Circulation, 2007, 116, 2634-2653.	1.6	2,755
4	ESC Guidelines for the diagnosis and treatment of acute and chronic heart failure 2008: The Task Force for the Diagnosis and Treatment of Acute and Chronic Heart Failure 2008 of the European Society of Cardiology. Developed in collaboration with the Heart Failure Association of the ESC (HFA) and endorsed by the European Society of Intensive Care Medicine (ESICM). European Heart Journal, 2008, 29, 2388-2442.	1.0	2,656
5	European guidelines on cardiovascular disease prevention in clinical practice: executive summary: Fourth Joint Task Force of the European Society of Cardiology and Other Societies on Cardiovascular Disease Prevention in Clinical Practice (Constituted by representatives of nine societies and by invited) Tj ETQq1 1 0.784314 rgBT /Ov	1.0	2,331
6	From Vulnerable Plaque to Vulnerable Patient. Circulation, 2003, 108, 1664-1672.	1.6	2,308
7	ACC/AHA/ESC 2006 Guidelines for the Management of Patients With Atrial Fibrillation. Circulation, 2006, 114, e257-354.	1.6	2,120
8	Guidelines on the management of valvular heart disease: The Task Force on the Management of Valvular Heart Disease of the European Society of Cardiology. European Heart Journal, 2006, 28, 230-268.	1.0	1,802
9	ESC Guidelines on the management of cardiovascular diseases during pregnancy: The Task Force on the Management of Cardiovascular Diseases during Pregnancy of the European Society of Cardiology (ESC). European Heart Journal, 2011, 32, 3147-3197.	1.0	1,694
10	Genotype-Phenotype Correlation in the Long-QT Syndrome. Circulation, 2001, 103, 89-95.	1.6	1,641
11	HRS/EHRA/APHRS Expert Consensus Statement on the Diagnosis and Management of Patients with Inherited Primary Arrhythmia Syndromes. Heart Rhythm, 2013, 10, 1932-1963.	0.3	1,587
12	From Vulnerable Plaque to Vulnerable Patient. Circulation, 2003, 108, 1772-1778.	1.6	1,562
13	American College of Cardiology/European Society of Cardiology Clinical Expert Consensus Document on Hypertrophic Cardiomyopathy. Journal of the American College of Cardiology, 2003, 42, 1687-1713.	1.2	1,444
14	CaV1.2 Calcium Channel Dysfunction Causes a Multisystem Disorder Including Arrhythmia and Autism. Cell, 2004, 119, 19-31.	13.5	1,403
15	Risk Stratification in the Long-QT Syndrome. New England Journal of Medicine, 2003, 348, 1866-1874.	13.9	1,314
16	Mutations in the Cardiac Ryanodine Receptor Gene (hRyR2) Underlie Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation, 2001, 103, 196-200.	1.6	1,291
17	ACC/AHA/ESC 2006 Guidelines for Management of Patients With Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death. Journal of the American College of Cardiology, 2006, 48, e247-e346.	1.2	1,280
18	Guidelines on the management of stable angina pectoris: executive summary: The Task Force on the Management of Stable Angina Pectoris of the European Society of Cardiology. European Heart Journal, 2006, 27, 1341-1381.	1.0	1,192

#	Article	IF	Citations
19	Spectrum of Mutations in Long-QT Syndrome Genes. Circulation, 2000, 102, 1178-1185.	1.6	1,157
20	Guidelines on diabetes, pre-diabetes, and cardiovascular diseases: executive summary: The Task Force on Diabetes and Cardiovascular Diseases of the European Society of Cardiology (ESC) and of the European Association for the Study of Diabetes (EASD). European Heart Journal, 2006, 28, 88-136.	1.0	1,144
21	Executive summary of the guidelines on the diagnosis and treatment of acute heart failure: The Task Force on Acute Heart Failure of the European Society of Cardiology. European Heart Journal, 2005, 26, 384-416.	1.0	1,114
22	Clinical and Molecular Characterization of Patients With Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation, 2002, 106, 69-74.	1.6	1,103
23	ACC/AHA/ESC 2006 Guidelines for the Management of Patients With Atrial Fibrillationâ€"Executive Summary. Journal of the American College of Cardiology, 2006, 48, 854-906.	1.2	1,044
24	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
25	ACC/AHA/ESC 2006 Guidelines for Management of Patients With Ventricular Arrhythmias and the Prevention of Sudden Cardiac Death. Circulation, 2006, 114, e385-484.	1.6	1,031
26	HRS/EHRA Expert Consensus Statement on the State of Genetic Testing for the Channelopathies and Cardiomyopathies. Heart Rhythm, 2011, 8, 1308-1339.	0.3	995
27	Natural History of Brugada Syndrome. Circulation, 2002, 105, 1342-1347.	1.6	984
28	Fourth Joint Task Force of the European Society of Cardiology and other Societies on Cardiovascular Disease Prevention in Clinical Practice (constituted by representatives of nine societies and by invited) Tj ETQq0 0 0 rgBT /Overlock 10 T	3.1	975
29	Guidelines on diagnosis and treatment of pulmonary arterial hypertension. The Task Force on Diagnosis and Treatment of Pulmonary Arterial Hypertension of the European Society of Cardiology. European Heart Journal, 2004, 25, 2243-2278.	1.0	903
30	Recommendations for competitive sports participation in athletes with cardiovascular disease: A consensus document from the Study Group of Sports Cardiology of the Working Group of Cardiac Rehabilitation and Exercise Physiology and the Working Group of Myocardial and Pericardial Diseases of the European Society of Cardiology. European Heart Journal, 2005, 26, 1422-1445.	1.0	860
31	Low Penetrance in the Long-QT Syndrome. Circulation, 1999, 99, 529-533.	1.6	783
32	Effectiveness and Limitations of Î²-Blocker Therapy in Congenital Long-QT Syndrome. Circulation, 2000, 101, 616-623.	1.6	783
33	Proposed Diagnostic Criteria for the Brugada Syndrome. Circulation, 2002, 106, 2514-2519.	1.6	779
34	2011 ACCF/AHA/HRS Focused Updates Incorporated Into the ACC/AHA/ESC 2006 Guidelines for the Management of Patients With Atrial Fibrillation. Circulation, 2011, 123, e269-367.	1.6	747
35	Influence of the Genotype on the Clinical Course of the Long-QT Syndrome. New England Journal of Medicine, 1998, 339, 960-965.	13.9	728
36	Reduction of hospitalizations for myocardial infarction in Italy in the COVID-19 era. European Heart Journal, 2020, 41, 2083-2088.	1.0	716

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37	Task Force on Sudden Cardiac Death of the European Society of Cardiology. European Heart Journal, 2001, 22, 1374-1450.	1.0	699
38	HRS/EHRA Expert Consensus Statement on the State of Genetic Testing for the Channelopathies and Cardiomyopathies: This document was developed as a partnership between the Heart Rhythm Society (HRS) and the European Heart Rhythm Association (EHRA). Europace, 2011, 13, 1077-1109.	0.7	699
39	Long QT Syndrome Patients With Mutations of the SCN5A and HERG Genes Have Differential Responses to Na ⁺ Channel Blockade and to Increases in Heart Rate. Circulation, 1995, 92, 3381-3386.	1.6	689
40	FKBP12.6 Deficiency and Defective Calcium Release Channel (Ryanodine Receptor) Function Linked to Exercise-Induced Sudden Cardiac Death. Cell, 2003, 113, 829-840.	13.5	683
41	ACC/AHA/ESC 2006 guidelines for the management of patients with atrial fibrillation: full text: A report of the American College of Cardiology/American Heart Association Task Force on practice guidelines and the European Society of Cardiology Committee for Practice Guidelines (Writing) Tj ETQq1 1 0.784314 rgBT /Overlock 10 Developed in collaboration with the European Heart Rhythm Association and the Heart Rhythm	0.7	661
42	ACC/AHA/ESC guidelines for the management of patients with supraventricular arrhythmiasâ^—â^—This document does not cover atrial fibrillation; atrial fibrillation is covered in the ACC/AHA/ESC guidelines on the management of patients with atrial fibrillation found on the ACC, AHA, and ESC Web sites.â€"executive summary. Journal of the American College of Cardiology, 2003, 42, 1493-1531.	1.2	660
43	ACC/AHA/ESC Guidelines for the Management of Patients With Supraventricular Arrhythmiasâ€"Executive Summary. Circulation, 2003, 108, 1871-1909.	1.6	651
44	2015 ESC Guidelines for the management of patients with ventricular arrhythmias and the prevention of sudden cardiac death. Europace, 2015, 17, euv319.	0.7	635
45	ACC/AHA/ESC 2006 guidelines for the management of patients with atrial fibrillationâ€"executive summary. European Heart Journal, 2006, 27, 1979-2030.	1.0	612
46	Guidelines on Prevention, Diagnosis and Treatment of Infective Endocarditis Executive Summary The Task Force on Infective Endocarditis of the European Society of Cardiology. European Heart Journal, 2004, 25, 267-276.	1.0	606
47	Left Cardiac Sympathetic Denervation in the Management of High-Risk Patients Affected by the Long-QT Syndrome. Circulation, 2004, 109, 1826-1833.	1.6	600
48	A Novel Form of Short QT Syndrome (SQT3) Is Caused by a Mutation in the KCNJ2 Gene. Circulation Research, 2005, 96, 800-807.	2.0	575
49	ACC/AHA/ESC 2006 guidelines for management of patients with ventricular arrhythmias and the prevention of sudden cardiac death: A report of the American College of Cardiology/American Heart Association Task Force and the European Society of Cardiology Committee for Practice Guidelines (Writing Committee to Develop Guidelines for Management of Patients With Ventricular Arrhythmias) Tj ETQq1 1 0.784314 rgBT /Overl Rhythm Association and the Heart R. Europace, 2006, 8, 746-837.	0.7	546
50	Association of Long QT Syndrome Loci and Cardiac Events Among Patients Treated With Î²-Blockers. JAMA - Journal of the American Medical Association, 2004, 292, 1341.	3.8	538
51	Risk Stratification in Brugada Syndrome. Journal of the American College of Cardiology, 2012, 59, 37-45.	1.2	523
52	Executive summary: HRS/EHRA/APHRS expert consensus statement on the diagnosis and management of patients with inherited primary arrhythmia syndromes. Europace, 2013, 15, 1389-1406.	0.7	494
53	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy. Heart Rhythm, 2019, 16, e301-e372.	0.3	494
54	Clinical and Genetic Heterogeneity of Right Bundle Branch Block and ST-Segment Elevation Syndrome. Circulation, 2000, 102, 2509-2515.	1.6	490

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55	A common polymorphism associated with antibiotic-induced cardiac arrhythmia. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 10613-10618.	3.3	466
56	Expert consensus document on ?-adrenergic receptor blockersThe Task Force on Beta-Blockers of the European Society of Cardiology. European Heart Journal, 2004, 25, 1341-1362.	1.0	465
57	ACC/AHA/ESC 2006 guidelines for management of patients with ventricular arrhythmias and the prevention of sudden cardiac death--executive summary: A report of the American College of Cardiology/American Heart Association Task Force and the European Society of Cardiology Committee for Practice Guidelines (Writing Committee to Develop Guidelines for Management of Patients with) Tj ETQq1 1 0.784314 rgBT /Over	1.0	454
58	with the European Heart Rhythm Associat. European Heart Journal, 2006, 27, 2099-2140. Age- and Sex-Related Differences in Clinical Manifestations in Patients With Congenital Long-QT Syndrome. Circulation, 1998, 97, 2237-2244.	1.6	451
59	Management of Grown Up Congenital Heart Disease. European Heart Journal, 2003, 24, 1035-1084.	1.0	446
60	Genetic Testing in the Long QT Syndrome. JAMA - Journal of the American Medical Association, 2005, 294, 2975.	3.8	413
61	Spectrum of ST-Tâ€“Wave Patterns and Repolarization Parameters in Congenital Long-QT Syndrome. Circulation, 2000, 102, 2849-2855.	1.6	409
62	Inherited Dysfunction of Sarcoplasmic Reticulum Ca ²⁺ Handling and Arrhythmogenesis. Circulation Research, 2011, 108, 871-883.	2.0	396
63	Increased Risk of Arrhythmic Events in Long-QT Syndrome With Mutations in the Pore Region of the Human Ether-a-go-goâ€“Related Gene Potassium Channel. Circulation, 2002, 105, 794-799.	1.6	370
64	Long QT Syndrome in Adults. Journal of the American College of Cardiology, 2007, 49, 329-337.	1.2	369
65	Guidelines on management (diagnosis and treatment) of syncope-update 2004. Executive Summary. European Heart Journal, 2004, 25, 2054-2072.	1.0	360
66	A Network of Macrophages Supports Mitochondrial Homeostasis in the Heart. Cell, 2020, 183, 94-109.e23.	13.5	360
67	Dispersion of the QT interval. A marker of therapeutic efficacy in the idiopathic long QT syndrome.. Circulation, 1994, 89, 1681-1689.	1.6	356
68	Nav1.5 E1053K mutation causing Brugada syndrome blocks binding to ankyrin-G and expression of Nav1.5 on the surface of cardiomyocytes. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 17533-17538.	3.3	349
69	Fourth Joint Task Force of the European Society of Cardiology and Other Societies on Cardiovascular Disease Prevention in Clinical Practice (Constituted by representatives of nine societies and by invited) Tj ETQq1 1 0.784314 rgBT /Over	8.7	348
70	Dilated cardiomyopathy. Nature Reviews Disease Primers, 2019, 5, 32.	18.1	347
71	Drugs and Brugada syndrome patients: Review of the literature, recommendations, and an up-to-date website (www.brugadadrugs.org). Heart Rhythm, 2009, 6, 1335-1341.	0.3	342
72	A Molecular Link between the Sudden Infant Death Syndrome and the Long-QT Syndrome. New England Journal of Medicine, 2000, 343, 262-267.	13.9	340

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73	Outcome parameters for trials in atrial fibrillation: executive summary: Recommendations from a consensus conference organized by the German Atrial Fibrillation Competence NETwork (AFNET) and the European Heart Rhythm Association (EHRA). European Heart Journal, 2007, 28, 2803-2817.	1.0	335
74	Expert Consensus Document on the Use of Antiplatelet Agents The Task Force on the Use of Antiplatelet Agents in Patients with Atherosclerotic Cardiovascular Disease of the European Society of Cardiology. European Heart Journal, 2004, 25, 166-181.	1.0	334
75	Cardiovascular diseases in women: a statement from the policy conference of the European Society of Cardiology. European Heart Journal, 2006, 27, 994-1005.	1.0	321
76	Cardiac Histological Substrate in Patients With Clinical Phenotype of Brugada Syndrome. Circulation, 2005, 112, 3680-3687.	1.6	317
77	Cardiac sodium channel mutations in patients with long QT syndrome, an inherited cardiac arrhythmia. Human Molecular Genetics, 1995, 4, 1603-1607.	1.4	316
78	Drug-Induced Torsades de Pointes and Implications for Drug Development. Journal of Cardiovascular Electrophysiology, 2004, 15, 475-495.	0.8	314
79	Evidence for a Cardiac Ion Channel Mutation Underlying Drug-Induced QT Prolongation and Life-Threatening Arrhythmias. Journal of Cardiovascular Electrophysiology, 2000, 11, 691-696.	0.8	312
80	Missense Mutations in Plakophilin-2 Cause Sodium Current Deficit and Associate With a Brugada Syndrome Phenotype. Circulation, 2014, 129, 1092-1103.	1.6	305
81	A cardiac arrhythmia syndrome caused by loss of ankyrin-B function. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 9137-9142.	3.3	301
82	Long QT Syndrome and Pregnancy. Journal of the American College of Cardiology, 2007, 49, 1092-1098.	1.2	299
83	2010 Focused Update of ESC Guidelines on device therapy in heart failure. Europace, 2010, 12, 1526-1536.	0.7	297
84	Genetic and Molecular Basis of Cardiac Arrhythmias: Impact on Clinical Management Parts I and II. Circulation, 1999, 99, 518-528.	1.6	295
85	Arrhythmogenesis in Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation Research, 2006, 99, 292-298.	2.0	293
86	European guidelines on cardiovascular disease prevention in clinical practice: Executive summary. Atherosclerosis, 2007, 194, 1-45.	0.4	292
87	HRS/EHRA Expert Consensus on the Monitoring of Cardiovascular Implantable Electronic Devices (CIEDs): Description of Techniques, Indications, Personnel, Frequency and Ethical Considerations. Heart Rhythm, 2008, 5, 907-925.	0.3	279
88	High Efficacy of Î²-Blockers in Long-QT Syndrome Type 1. Circulation, 2009, 119, 215-221.	1.6	274
89	Risk for Life-Threatening Cardiac Events in Patients With Genotype-Confirmed Long-QT Syndrome and Normal-Range Corrected QT Intervals. Journal of the American College of Cardiology, 2011, 57, 51-59.	1.2	268
90	Who Are the Long-QT Syndrome Patients Who Receive an Implantable Cardioverter-Defibrillator and What Happens to Them?. Circulation, 2010, 122, 1272-1282.	1.6	261

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91	Sodium channel mutations and arrhythmias. Nature Reviews Cardiology, 2009, 6, 337-348.	6.1	260
92	Risk of Aborted Cardiac Arrest or Sudden Cardiac Death During Adolescence in the Long-QT Syndrome. JAMA - Journal of the American Medical Association, 2006, 296, 1249.	3.8	258
93	Modulating effects of age and gender on the clinical course of long QT syndrome by genotype. Journal of the American College of Cardiology, 2003, 42, 103-109.	1.2	257
94	Risk Factors for Aborted Cardiac Arrest and Sudden Cardiac Death in Children With the Congenital Long-QT Syndrome. Circulation, 2008, 117, 2184-2191.	1.6	255
95	Outcome parameters for trials in atrial fibrillation: Recommendations from a consensus conference organized by the German Atrial Fibrillation Competence NETwork and the European Heart Rhythm Association. Europace, 2007, 9, 1006-1023.	0.7	254
96	Arrhythmogenic Mechanisms in a Mouse Model of Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation Research, 2007, 101, 1039-1048.	2.0	252
97	Bidirectional Ventricular Tachycardia and Fibrillation Elicited in a Knock-In Mouse Model Carrier of a Mutation in the Cardiac Ryanodine Receptor. Circulation Research, 2005, 96, e77-82.	2.0	247
98	Brugada syndrome and sudden cardiac death in children. Lancet, The, 2000, 355, 808-809.	6.3	244
99	The Elusive Link Between LQT3 and Brugada Syndrome. Circulation, 2000, 102, 945-947.	1.6	243
100	Desmoplakin Cardiomyopathy, a Fibrotic and Inflammatory Form of Cardiomyopathy Distinct From Typical Dilated or Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation, 2020, 141, 1872-1884.	1.6	229
101	Magnetic resonance imaging in individuals with cardiovascular implantable electronic devices. Europace, 2008, 10, 336-346.	0.7	221
102	Comparison of clinical and genetic variables of cardiac events associated with loud noise versus swimming among subjects with the long QT syndrome. American Journal of Cardiology, 1999, 84, 876-879.	0.7	219
103	Update on Expert Consensus on the Monitoring of Cardiovascular Implantable Electronic Devices (CIEDs): Description of Techniques, Indications, Personnel, Frequency and Ethical Considerations: Developed in partnership with the Heart Rhythm Society (HRS) and the European Heart Rhythm Association (EHRA); and in collaboration with the American College of Cardiology (ACC), the American Heart Association (AHA), the European Society of Cardiology (ESC), the Heart Failure Association of ESC (HFA), and the Heart Fail. Europace, 2008, 10, 707-725.	0.7	215
104	The European cardiac resynchronization therapy survey. European Heart Journal, 2009, 30, 2450-2460.	1.0	215
105	Genetics of Sudden Cardiac Death. Circulation Research, 2015, 116, 1919-1936.	2.0	211
106	Epinephrine unmasks latent mutation carriers with LQT1 form of congenital long-QT syndrome. Journal of the American College of Cardiology, 2003, 41, 633-642.	1.2	201
107	Programmed Ventricular Stimulation for Risk Stratification in the Brugada Syndrome. Circulation, 2016, 133, 622-630.	1.6	201
108	Gating Properties of SCN5A Mutations and the Response to Mexiletine in Long-QT Syndrome Type 3 Patients. Circulation, 2007, 116, 1137-1144.	1.6	194

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109	Novel Insight Into the Natural History of Short QT Syndrome. Journal of the American College of Cardiology, 2014, 63, 1300-1308.	1.2	191
110	Gene-Specific Therapy With Mexiletine Reduces Arrhythmic Events in Patients With Long QT Syndrome Type 3. Journal of the American College of Cardiology, 2016, 67, 1053-1058.	1.2	191
111	Clinical Phenotype and Functional Characterization of CASQ2 Mutations Associated With Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation, 2006, 114, 1012-1019.	1.6	189
112	Involvement of the cardiac ryanodine receptor/calcium release channel in catecholaminergic polymorphic ventricular tachycardia. Journal of Cellular Physiology, 2002, 190, 1-6.	2.0	182
113	Inherited Brugada and Long QT-3 Syndrome Mutations of a Single Residue of the Cardiac Sodium Channel Confer Distinct Channel and Clinical Phenotypes. Journal of Biological Chemistry, 2001, 276, 30623-30630.	1.6	181
114	Pre-participation cardiovascular evaluation for athletic participants to prevent sudden death: Position paper from the EHRA and the EACPR, branches of the ESC. Endorsed by APHRS, HRS, and SOLAECE. European Journal of Preventive Cardiology, 2017, 24, 41-69.	0.8	181
115	Protective effect of vagal stimulation on reperfusion arrhythmias in cats.. Circulation Research, 1987, 61, 429-435.	2.0	179
116	Abnormal Interactions of Calsequestrin With the Ryanodine Receptor Calcium Release Channel Complex Linked to Exercise-Induced Sudden Cardiac Death. Circulation Research, 2006, 98, 1151-1158.	2.0	179
117	Long QT syndrome, Brugada syndrome, and conduction system disease are linked to a single sodium channel mutation. Journal of Clinical Investigation, 2002, 110, 1201-1209.	3.9	172
118	Cellular Dysfunction of LQT5-MinK Mutants: Abnormalities of IKs, IKr and Trafficking in Long QT Syndrome. Human Molecular Genetics, 1999, 8, 1499-1507.	1.4	170
119	Acute heart failure congestion and perfusion statusÂâ€“Âimpact of the clinical classification on inâ€hospital and longâ€term outcomes; insights from the ESCâ€EORPâ€HFA Heart Failure Longâ€Term Registry. European Journal of Heart Failure, 2019, 21, 1338-1352.	2.9	170
120	Polymorphisms in the NOS1APGene Modulate QT Interval Duration and Risk of Arrhythmias in the Long QT Syndrome. Journal of the American College of Cardiology, 2010, 55, 2745-2752.	1.2	163
121	Delayed afterdepolarizations elicited in vivo by left stellate ganglion stimulation.. Circulation, 1988, 78, 178-185.	1.6	159
122	Executive Summary: HRS/EHRA/APHRS Expert Consensus Statement on the Diagnosis and Management of Patients with Inherited Primary Arrhythmia Syndromes. Heart Rhythm, 2013, 10, e85-e108.	0.3	159
123	Abnormal Calcium Signaling and Sudden Cardiac Death Associated With Mutation of Calsequestrin. Circulation Research, 2004, 94, 471-477.	2.0	158
124	Inherited calcium channelopathies in the pathophysiology of arrhythmias. Nature Reviews Cardiology, 2012, 9, 561-575.	6.1	158
125	Molecular diagnosis in a child with sudden infant death syndrome. Lancet, The, 2001, 358, 1342-1343.	6.3	157
126	Diagnostic value of epinephrine test for genotyping LQT1, LQT2, and LQT3 forms of congenital long QT syndrome. Heart Rhythm, 2004, 1, 276-283.	0.3	156

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127	Catecholaminergic Polymorphic Ventricular Tachycardia. Progress in Cardiovascular Diseases, 2008, 51, 23-30.	1.6	156
128	Magnetic resonance investigations in Brugada syndrome reveal unexpectedly high rate of structural abnormalities. European Heart Journal, 2009, 30, 2241-2248.	1.0	156
129	The consensus of the task force of the European Society of Cardiology concerning the clinical investigation of the use of autologous adult stem cells for repair of the heart. European Heart Journal, 2006, 27, 1338-1340.	1.0	155
130	Evaluation of the Spatial Aspects of T-Wave Complexity in the Long-QT Syndrome. Circulation, 1997, 96, 3006-3012.	1.6	151
131	Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American College of Cardiology, 2016, 68, 2540-2550.	1.2	148
132	Differential Response to Na ⁺ Channel Blockade, Î²-Adrenergic Stimulation, and Rapid Pacing in a Cellular Model Mimicking the SCN5A and HERG Defects Present in the Long-QT Syndrome. Circulation Research, 1996, 78, 1009-1015.	2.0	148
133	Recommendations for participation in competitive sport and leisure-time physical activity in individuals with cardiomyopathies, myocarditis and pericarditis. European Journal of Cardiovascular Prevention and Rehabilitation, 2006, 13, 876-885.	3.1	146
134	A Recessive Variant of the Romano-Ward Long-QT Syndrome?. Circulation, 1998, 97, 2420-2425.	1.6	139
135	Increased Ca ²⁺ Sensitivity of the Ryanodine Receptor Mutant RyR2 ^{R4496C} Underlies Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation Research, 2009, 104, 201-209.	2.0	137
136	Screening for Sudden Cardiac Death in the Young. Circulation, 2011, 123, 1911-1918.	1.6	137
137	Interplay Between Genetic Substrate, QTcÂDuration, and Arrhythmia Risk in Patients With Long QT Syndrome. Journal of the American College of Cardiology, 2018, 71, 1663-1671.	1.2	137
138	Update of the guidelines on sudden cardiac death of the European Society of Cardiology. European Heart Journal, 2003, 24, 13-15.	1.0	135
139	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy: Executive summary. Heart Rhythm, 2019, 16, e373-e407.	0.3	135
140	Long-QT Syndrome After Age 40. Circulation, 2008, 117, 2192-2201.	1.6	134
141	Diagnosis and treatment of catecholaminergic polymorphic ventricular tachycardia. Heart Rhythm, 2007, 4, 675-678.	0.3	133
142	Sudden Cardiac Death and Genetic Ion Channelopathies. Circulation, 2012, 125, 2027-2034.	1.6	133
143	Genetic and Molecular Basis of Cardiac Arrhythmias: Impact on Clinical Management Part III. Circulation, 1999, 99, 674-681.	1.6	131
144	Yield of Genetic Screening in Inherited Cardiac Channelopathies. Circulation: Arrhythmia and Electrophysiology, 2009, 2, 6-15.	2.1	131

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145	Mutation site-specific differences in arrhythmic risk and sensitivity to sympathetic stimulation in the LQT1 form of congenital long QT syndrome. Journal of the American College of Cardiology, 2004, 44, 117-125.	1.2	130
146	KCNJ2 mutation in short QT syndrome 3 results in atrial fibrillation and ventricular proarrhythmia. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 4291-4296.	3.3	130
147	Efficacy of Flecainide in the Treatment of Catecholaminergic Polymorphic Ventricular Tachycardia. JAMA Cardiology, 2017, 2, 759.	3.0	127
148	Unexpected Structural and Functional Consequences of the R33Q Homozygous Mutation in Cardiac Calsequestrin. Circulation Research, 2008, 103, 298-306.	2.0	124
149	Torsade de Pointes. Drugs, 1994, 47, 51-65.	4.9	122
150	Luminal Ca2+ Regulation of Single Cardiac Ryanodine Receptors: Insights Provided by Calsequestrin and its Mutants. Journal of General Physiology, 2008, 131, 325-334.	0.9	122
151	Cost-effectiveness of neonatal ECG screening for the long QT syndrome. European Heart Journal, 2006, 27, 1824-1832.	1.0	121
152	Implementation of device therapy (cardiac resynchronization therapy and implantable cardioverter) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 5 of Heart Failure, 2009, 11, 1143-1151.	2.9	118
153	Sympathetic stimulation produces a greater increase in both transmural and spatial dispersion of repolarization in LQT1 than LQT2 forms of congenital long QT syndrome. Journal of the American College of Cardiology, 2001, 37, 911-919.	1.2	115
154	Novel Arrhythmogenic Mechanism Revealed by a Long-QT Syndrome Mutation in the Cardiac Na+Channel. Circulation Research, 2001, 88, 740-745.	2.0	114
155	Molecular and Electrophysiological Bases of Catecholaminergic Polymorphic Ventricular Tachycardia. Journal of Cardiovascular Electrophysiology, 2007, 18, 791-797.	0.8	113
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