Carla Spazzolini

List of Publications by Year in Descending Order

39 papers	5,948 citations	23 h-index	41 g-index
41 ext. papers	7,002 ext. citations	10.8 avg, IF	4.43 L-index

#	Paper	IF	Citations
39	Left Cardiac Sympathetic Denervation for Long QT Syndrome: 50 YearsSExperience Provides Guidance for Management.. *JACC: Clinical Electrophysiology*, **2022**, 8, 281-294	4.6	2
38	Estimating the Posttest Probability of Long QT Syndrome Diagnosis for Rare Variants. *Circulation Genomic and Precision Medicine*, **2021**, 14, e003289	5.2	0
37	Mutation location and IKs regulation in the arrhythmic risk of long QT syndrome type 1: the importance of the KCNQ1 S6 region. *European Heart Journal*, **2021**, 42, 4743-4755	9.5	4
36	An International Multicenter Evaluation of Type 5 Long QT Syndrome: A Low Penetrant Primary Arrhythmic Condition. *Circulation*, **2020**, 141, 429-439	16.7	15
35	Exercise Training-Induced Repolarization Abnormalities Masquerading as Congenital Long QT Syndrome. *Circulation*, **2020**, 142, 2405-2415	16.7	8
34	Mothers with long QT syndrome are at increased risk for fetal death: findings from a multicenter international study. *American Journal of Obstetrics and Gynecology*, **2020**, 222, 263.e1-263.e11	6.4	18
33	Calmodulin mutations and life-threatening cardiac arrhythmias: insights from the International Calmodulinopathy Registry. *European Heart Journal*, **2019**, 40, 2964-2975	9.5	61
32	From patient-specific induced pluripotent stem cells to clinical translation in long QT syndrome Type 2. *European Heart Journal*, **2019**, 40, 1832-1836	9.5	41
31	The genetics underlying idiopathic ventricular fibrillation: A special role for catecholaminergic polymorphic ventricular tachycardia?. *International Journal of Cardiology*, **2018**, 250, 139-145	3.2	22
30	SCN5A mutations in 442 neonates and children: genotype-phenotype correlation and identification of higher-risk subgroups. *European Heart Journal*, **2018**, 39, 2879-2887	9.5	18
29	Cardiac arrest and Brugada syndrome: Is drug-induced type 1 ECG pattern always a marker of low risk?. *International Journal of Cardiology*, **2018**, 254, 142-145	3.2	9
28	For neonatal ECG screening there is no reason to relinquish old Bazett's correction. *European Heart Journal*, **2018**, 39, 2888-2895	9.5	19
27	Desmoplakin missense and non-missense mutations in arrhythmogenic right ventricular cardiomyopathy: Genotype-phenotype correlation. *International Journal of Cardiology*, **2017**, 249, 268-273	3.2	46
26	Clinical Aspects of Type 3 Long-QT Syndrome: An International Multicenter Study. *Circulation*, **2016**, 134, 872-82	16.7	118
25	Genetic Modifiers for the Long-QT Syndrome: How Important Is the Role of Variants in the 3S Untranslated Region of KCNQ1?. *Circulation: Cardiovascular Genetics*, **2016**, 9, 330-9		16
24	The genetics underlying acquired long QT syndrome: impact for genetic screening. *European Heart Journal*, **2016**, 37, 1456-64	9.5	108
23	Response to Letters Regarding Article, "Clinical Management of Catecholaminergic Polymorphic Ventricular Tachycardia: The Role of Left Cardiac Sympathetic Denervation". *Circulation*, **2016**, 133, e366-7	16.7	2

22	Response by Crotti et al to Letter Regarding Article, "Genetic Modifiers for the Long-QT Syndrome: How Important Is the Role of Variants in the 3SUntranslated Region of KCNQ1?". *Circulation: Cardiovascular Genetics*, **2016**, 9, 581-582		10
21	Clinical Management of Catecholaminergic Polymorphic Ventricular Tachycardia: The Role of Left Cardiac Sympathetic Denervation. *Circulation*, **2015**, 131, 2185-93	16.7	174
20	Propranolol prevents life-threatening arrhythmias in LQT3 transgenic mice: implications for the clinical management of LQT3 patients. *Heart Rhythm*, **2014**, 11, 126-32	6.7	26
19	Tumor necrosis factor-predicts response to cardiac resynchronization therapy in patients with chronic heart failure. *Circulation Journal*, **2014**, 78, 2232-9	2.9	20
18	A comprehensive electrocardiographic, molecular, and echocardiographic study of Brugada syndrome: validation of the 2013 diagnostic criteria. *Heart Rhythm*, **2014**, 11, 1176-83	6.7	25
17	Rat experimental model of myocardial ischemia/reperfusion injury: an ethical approach to set up the analgesic management of acute post-surgical pain. *PLoS ONE*, **2014**, 9, e95913	3.7	11
16	Vagal reflexes following an exercise stress test: a simple clinical tool for gene-specific risk stratification in the long QT syndrome. *Journal of the American College of Cardiology*, **2012**, 60, 2515-24	15.1	42
15	Not all beta-blockers are equal in the management of long QT syndrome types 1 and 2: higher recurrence of events under metoprolol. *Journal of the American College of Cardiology*, **2012**, 60, 2092-9	15.1	168
14	Who are the long-QT syndrome patients who receive an implantable cardioverter-defibrillator and what happens to them?: data from the European Long-QT Syndrome Implantable Cardioverter-Defibrillator (LQTS ICD) Registry. *Circulation*, **2010**, 122, 1272-82	16.7	209
13	Mutation-specific risk in two genetic forms of type 3 long QT syndrome. *American Journal of Cardiology*, **2010**, 105, 210-3	3	22
12	Prevalence of the congenital long-QT syndrome. *Circulation*, **2009**, 120, 1761-7	16.7	649
11	High efficacy of beta-blockers in long-QT syndrome type 1: contribution of noncompliance and QT-prolonging drugs to the occurrence of beta-blocker treatment "failures". *Circulation*, **2009**, 119, 215-21	16.7	235
10	Clinical implications for patients with long QT syndrome who experience a cardiac event during infancy. *Journal of the American College of Cardiology*, **2009**, 54, 832-7	15.1	70
9	All LQT3 patients need an ICD: true or false?. *Heart Rhythm*, **2009**, 6, 113-20	6.7	81
8	Neural control of heart rate is an arrhythmia risk modifier in long QT syndrome. *Journal of the American College of Cardiology*, **2008**, 51, 920-9	15.1	88
7	The common long-QT syndrome mutation KCNQ1/A341V causes unusually severe clinical manifestations in patients with different ethnic backgrounds: toward a mutation-specific risk stratification. *Circulation*, **2007**, 116, 2366-75	16.7	130
6	Cost-effectiveness of neonatal ECG screening for the long QT syndrome. *European Heart Journal*, **2006**, 27, 1824-32	9.5	101
5	The Jervell and Lange-Nielsen syndrome: natural history, molecular basis, and clinical outcome. *Circulation*, **2006**, 113, 783-90	16.7	274

4	Phenotypic variability and unusual clinical severity of congenital long-QT syndrome in a founder population. *Circulation*, **2005**, 112, 2602-10	16.7	150
3	Left cardiac sympathetic denervation in the management of high-risk patients affected by the long-QT syndrome. *Circulation*, **2004**, 109, 1826-33	16.7	503
2	Risk stratification in the long-QT syndrome. *New England Journal of Medicine*, **2003**, 348, 1866-74	59.2	1090
1	Genotype-phenotype correlation in the long-QT syndrome: gene-specific triggers for life-threatening arrhythmias. *Circulation*, **2001**, 103, 89-95	16.7	1363