

Ramon Brugada

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

147 papers	6,432 citations	31 h-index	79 g-index
168 ext. papers	7,609 ext. citations	5.8 avg, IF	5.33 L-index

#	Paper	IF	Citations
147	Clinical Genetics of Inherited Arrhythmogenic Disease in the Pediatric Population.. <i>Biomedicines</i> , 2022 , 10,	4.8	1
146	Genetic Cardiomyopathy May Overlap Fulminant Myocarditis Clinical Findings.. <i>Circulation: Heart Failure</i> , 2022 , 15, e008443	7.6	
145	Generation of an induced pluripotent stem cell line from a healthy Caucasian male.. <i>Stem Cell Research</i> , 2022 , 60, 102717	1.6	0
144	Rare variants in genes encoding structural myocyte contribute to a thickened ventricular septum in sudden death population without ventricular alterations.. <i>Forensic Science International: Genetics</i> , 2022 , 58, 102688	4.3	0
143	Prevalence of Pathogenic Variants in Cardiomyopathy-Associated Genes in Myocarditis.. <i>Circulation Genomic and Precision Medicine</i> , 2022 , 101161CIRCGEN121003408	5.2	1
142	Microbiota alterations in proline metabolism impact depression.. <i>Cell Metabolism</i> , 2022 , 34, 681-701.e1024.6	24.6	7
141	Analysis of Brugada syndrome loci reveals that fine-mapping clustered GWAS hits enhances the annotation of disease-relevant variants. <i>Cell Reports Medicine</i> , 2021 , 2, 100250	18	0
140	Sudden Death without a Clear Cause after Comprehensive Investigation: An Example of Forensic Approach to Atypical/Uncertain Findings. <i>Diagnostics</i> , 2021 , 11,	3.8	0
139	Early Identification of Prolonged QT Interval for Prevention of Sudden Infant Death. <i>Frontiers in Pediatrics</i> , 2021 , 9, 704580	3.4	1
138	Whole-Brain Dynamics in Aging: Disruptions in Functional Connectivity and the Role of the Rich Club. <i>Cerebral Cortex</i> , 2021 , 31, 2466-2481	5.1	3
137	Enhancing rare variant interpretation in inherited arrhythmias through quantitative analysis of consortium disease cohorts and population controls. <i>Genetics in Medicine</i> , 2021 , 23, 47-58	8.1	13
136	Rare Variants Associated with Arrhythmogenic Cardiomyopathy: Reclassification Five Years Later. <i>Journal of Personalized Medicine</i> , 2021 , 11,	3.6	3
135	Genotype-Phenotype Correlation of Genotype in Patients With Brugada Syndrome and Arrhythmic Events: Insights From the SABRUS in 392 Probands. <i>Circulation Genomic and Precision Medicine</i> , 2021 , 14, e003222	5.2	0
134	Clinical impact of rare variants associated with inherited channelopathies: a 5-year update. <i>Human Genetics</i> , 2021 , 1	6.3	1
133	Obesity-associated deficits in inhibitory control are phenocopied to mice through gut microbiota changes in one-carbon and aromatic amino acids metabolic pathways. <i>Gut</i> , 2021 , 70, 2283-2296	19.2	7
132	Transethnic Genome-Wide Association Study Provides Insights in the Genetic Architecture and Heritability of Long QT Syndrome. <i>Circulation</i> , 2020 , 142, 324-338	16.7	27
131	The arrhythmogenic right ventricular cardiomyopathy in comparison to the athletic heart. <i>Journal of Cardiovascular Electrophysiology</i> , 2020 , 31, 1836-1843	2.7	4

130	The Aging Imageomics Study: rationale, design and baseline characteristics of the study population. <i>Mechanisms of Ageing and Development</i> , 2020 , 189, 111257	5.6	6
129	Genetic Variants as Sudden-Death Risk Markers in Inherited Arrhythmogenic Syndromes: Personalized Genetic Interpretation. <i>Journal of Clinical Medicine</i> , 2020 , 9,	5.1	3
128	Clinical Features and Natural History of PRKAG2 Variant Cardiac Glycogenosis. <i>Journal of the American College of Cardiology</i> , 2020 , 76, 186-197	15.1	16
127	Sudden Cardiac Death and Copy Number Variants: What Do We Know after 10 Years of Genetic Analysis?. <i>Forensic Science International: Genetics</i> , 2020 , 47, 102281	4.3	10
126	Genetic variants of uncertain significance: How to match scientific rigour and standard of proof in sudden cardiac death?. <i>Legal Medicine</i> , 2020 , 45, 101712	1.9	12
125	The role of clinical assessment and electrophysiology study in Brugada syndrome patients with syncope. <i>American Heart Journal</i> , 2020 , 220, 213-223	4.9	5
124	Update on Genetic Basis of Brugada Syndrome: Monogenic, Polygenic or Oligogenic?. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	20
123	Reanalysis and reclassification of rare genetic variants associated with inherited arrhythmogenic syndromes. <i>EBioMedicine</i> , 2020 , 54, 102732	8.8	21
122	Pediatric Malignant Arrhythmias Caused by Rare Homozygous Genetic Variants in : A Comprehensive Interpretation. <i>Frontiers in Pediatrics</i> , 2020 , 8, 601708	3.4	2
121	Sudden infant death as the most severe phenotype caused by genetic modulation in a family with atrial fibrillation. <i>Forensic Science International: Genetics</i> , 2019 , 43, 102159	4.3	0
120	Personalized Interpretation and Clinical Translation of Genetic Variants Associated With Cardiomyopathies. <i>Frontiers in Genetics</i> , 2019 , 10, 450	4.5	4
119	Experimental Models of Brugada syndrome. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	18
118	Update on the Genetic Basis of Sudden Unexpected Death in Epilepsy. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	21
117	Genetic interpretation and clinical translation of minor genes related to Brugada syndrome. <i>Human Mutation</i> , 2019 , 40, 749-764	4.7	17
116	Characterization and Management of Arrhythmic Events in Young Patients With Brugada Syndrome. <i>Journal of the American College of Cardiology</i> , 2019 , 73, 1756-1765	15.1	33
115	Short QT Syndrome: A Comprehensive Genetic Interpretation and Clinical Translation of Rare Variants. <i>Journal of Clinical Medicine</i> , 2019 , 8,	5.1	17
114	Electrocardiogram in Newborns: Beneficial or Not?. <i>Pediatric Cardiology</i> , 2019 , 40, 1320-1321	2.1	
113	Ethnic differences in patients with Brugada syndrome and arrhythmic events: New insights from Survey on Arrhythmic Events in Brugada Syndrome. <i>Heart Rhythm</i> , 2019 , 16, 1468-1474	6.7	14

112	Trafficking and Function of the Voltage-Gated Sodium Channel α Subunit. <i>Biomolecules</i> , 2019 , 9,	5.9	4
111	-Glycosylation of the voltage-gated sodium channel α subunit is required for efficient trafficking of Na _v 1.5/ α to the plasma membrane. <i>Journal of Biological Chemistry</i> , 2019 , 294, 16123-16140	5.4	7
110	Electrocardiographic Assessment and Genetic Analysis in Neonates: a Current Topic of Discussion. <i>Current Cardiology Reviews</i> , 2019 , 15, 30-37	2.4	5
109	Brugada Syndrome: anesthetic considerations and management algorithm. <i>Minerva Anestesiologica</i> , 2019 , 85, 173-188	1.9	5
108	Elucidating the Role of K Channels during In Vitro Capacitation of Boar Spermatozoa: Do SLO1 Channels Play a Crucial Role?. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	6
107	Time-to-first appropriate shock in patients implanted prophylactically with an implantable cardioverter-defibrillator: data from the Survey on Arrhythmic Events in BRUGada Syndrome (SABRUS). <i>Europace</i> , 2019 , 21, 796-802	3.9	7
106	Role of copy number variants in sudden cardiac death and related diseases: genetic analysis and translation into clinical practice. <i>European Journal of Human Genetics</i> , 2018 , 26, 1014-1025	5.3	17
105	Fever-related arrhythmic events in the multicenter Survey on Arrhythmic Events in Brugada Syndrome. <i>Heart Rhythm</i> , 2018 , 15, 1394-1401	6.7	49
104	Profile of patients with Brugada syndrome presenting with their first documented arrhythmic event: Data from the Survey on Arrhythmic Events in BRUGada Syndrome (SABRUS). <i>Heart Rhythm</i> , 2018 , 15, 716-724	6.7	36
103	The long-QT syndrome and exercise practice: The never-ending debate. <i>Journal of Cardiovascular Electrophysiology</i> , 2018 , 29, 489-496	2.7	9
102	Sodium channel current loss of function in induced pluripotent stem cell-derived cardiomyocytes from a Brugada syndrome patient. <i>Journal of Molecular and Cellular Cardiology</i> , 2018 , 114, 10-19	5.8	31
101	Molecular autopsy in a cohort of infants died suddenly at rest. <i>Forensic Science International: Genetics</i> , 2018 , 37, 54-63	4.3	6
100	Plasma microRNAs as biomarkers for Lamin A/C-related dilated cardiomyopathy. <i>Journal of Molecular Medicine</i> , 2018 , 96, 845-856	5.5	22
99	Role of genetic and electrolyte abnormalities in prolonged QTc interval and sudden cardiac death in end-stage renal disease patients. <i>PLoS ONE</i> , 2018 , 13, e0200756	3.7	9
98	Present Status of Brugada Syndrome: JACC State-of-the-Art Review. <i>Journal of the American College of Cardiology</i> , 2018 , 72, 1046-1059	15.1	165
97	Gender differences in patients with Brugada syndrome and arrhythmic events: Data from a survey on arrhythmic events in 678 patients. <i>Heart Rhythm</i> , 2018 , 15, 1457-1465	6.7	36
96	GRAPES: A Versatile Tool for Analyzing Structural Variation From Whole-Genome and Targeted DNA Sequencing Data. <i>FASEB Journal</i> , 2018 , 32, 532.10	0.9	
95	Electroanatomic and Pathologic Right Ventricular Outflow Tract Abnormalities in Patients With Brugada Syndrome. <i>Journal of the American College of Cardiology</i> , 2018 , 72, 2747-2757	15.1	38

94	Recent Advances in Short QT Syndrome. <i>Frontiers in Cardiovascular Medicine</i> , 2018 , 5, 149	5.4	36
93	Genetic analysis in post-mortem samples with micro-ischemic alterations. <i>Forensic Science International</i> , 2017 , 271, 120-125	2.6	1
92	Juvenile myoclonic epilepsy and Brugada type 1 ECG pattern associated with (a novel) plakophilin 2 mutation. <i>Journal of Neurology</i> , 2017 , 264, 792-795	5.5	2
91	Sudden Arrhythmic Death During Exercise: A Post-Mortem Genetic Analysis. <i>Sports Medicine</i> , 2017 , 47, 2101-2115	10.6	10
90	Sudden death due to catecholaminergic polymorphic ventricular tachycardia following negative stress-test outcome: genetics and clinical implications. <i>Forensic Science, Medicine, and Pathology</i> , 2017 , 13, 217-225	1.5	4
89	Electrical Substrate Elimination in 135 Consecutive Patients With Brugada Syndrome. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2017 , 10, e005053	6.4	112
88	Reply to letter to editor: "Genetic basis of dilated cardiomyopathy". <i>International Journal of Cardiology</i> , 2017 , 229, 32	3.2	
87	Brugada Syndrome and Exercise Practice: Current Knowledge, Shortcomings and Open Questions. <i>International Journal of Sports Medicine</i> , 2017 , 38, 573-581	3.6	11
86	Trafficking and localisation to the plasma membrane of Na _v 1.5 promoted by the β subunit is defective due to a β mutation associated with Brugada syndrome. <i>Biology of the Cell</i> , 2017 , 109, 273-291	3.5	9
85	Short QT syndrome in pediatrics. <i>Clinical Research in Cardiology</i> , 2017 , 106, 393-400	6.1	12
84	A novel variant in RyR2 causes familial catecholaminergic polymorphic ventricular tachycardia. <i>Forensic Science International</i> , 2017 , 270, 173-177	2.6	2
83	Patients With Brugada Syndrome and Implanted Cardioverter-Defibrillators: Long-Term Follow-Up. <i>Journal of the American College of Cardiology</i> , 2017 , 70, 1991-2002	15.1	25
82	Incomplete Penetrance and Variable Expressivity: Hallmarks in Channelopathies Associated with Sudden Cardiac Death. <i>Biology</i> , 2017 , 7,	4.9	17
81	Do sodium channel proteolytic fragments regulate sodium channel expression?. <i>Channels</i> , 2017 , 11, 476-481	3.8	2
80	Unmasking the molecular link between arrhythmogenic cardiomyopathy and Brugada syndrome. <i>Nature Reviews Cardiology</i> , 2017 , 14, 744-756	14.8	40
79	Transcriptional regulation of the sodium channel gene (SCN5A) by GATA4 in human heart. <i>Journal of Molecular and Cellular Cardiology</i> , 2017 , 102, 74-82	5.8	20
78	Age of First Arrhythmic Event in Brugada Syndrome: Data From the SABRUS (Survey on Arrhythmic Events in Brugada Syndrome) in 678 Patients. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2017 , 10,	6.4	39
77	Cardiac Channelopathies and Sudden Death: Recent Clinical and Genetic Advances. <i>Biology</i> , 2017 , 6,	4.9	63

76	Additional value of screening for minor genes and copy number variants in hypertrophic cardiomyopathy. <i>PLoS ONE</i> , 2017 , 12, e0181465	3.7	23
75	Targeted next-generation sequencing provides novel clues for associated epilepsy and cardiac conduction disorder/SUDEP. <i>PLoS ONE</i> , 2017 , 12, e0189618	3.7	21
74	Brugada syndrome: clinical and genetic findings. <i>Genetics in Medicine</i> , 2016 , 18, 3-12	8.1	73
73	Genetic basis of dilated cardiomyopathy. <i>International Journal of Cardiology</i> , 2016 , 224, 461-472	3.2	50
72	Proteomic identification of putative biomarkers for early detection of sudden cardiac death in a family with a LMNA gene mutation causing dilated cardiomyopathy. <i>Journal of Proteomics</i> , 2016 , 148, 75-84	3.9	9
71	Sudden infant death syndrome caused by cardiac arrhythmias: only a matter of genes encoding ion channels?. <i>International Journal of Legal Medicine</i> , 2016 , 130, 415-20	3.1	20
70	Genetic investigation of sudden unexpected death in epilepsy cohort by panel target resequencing. <i>International Journal of Legal Medicine</i> , 2016 , 130, 331-9	3.1	39
69	Natural and Undetermined Sudden Death: Value of Post-Mortem Genetic Investigation. <i>PLoS ONE</i> , 2016 , 11, e0167358	3.7	36
68	Familial Dilated Cardiomyopathy Caused by a Novel Frameshift in the BAG3 Gene. <i>PLoS ONE</i> , 2016 , 11, e0158730	3.7	28
67	Large Genomic Imbalances in Brugada Syndrome. <i>PLoS ONE</i> , 2016 , 11, e0163514	3.7	18
66	Spatiotemporal Characteristics of QRS Complexes Enable the Diagnosis of Brugada Syndrome Regardless of the Appearance of a Type 1 ECG. <i>Journal of Cardiovascular Electrophysiology</i> , 2016 , 27, 563-70	2.7	4
65	Genetic basis of atrial fibrillation. <i>Genes and Diseases</i> , 2016 , 3, 257-262	6.6	17
64	Acute, Exercise Dose-Dependent Impairment in Atrial Performance During an Endurance Race: 2D Ultrasound Speckle-Tracking Strain Analysis. <i>JACC: Cardiovascular Imaging</i> , 2016 , 9, 1380-1388	8.4	25
63	Further evidence of the association between LQT syndrome and epilepsy in a family with KCNQ1 pathogenic variant. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2015 , 25, 65-7	3.2	45
62	Clinical and molecular characterization of a cardiac ryanodine receptor founder mutation causing catecholaminergic polymorphic ventricular tachycardia. <i>Heart Rhythm</i> , 2015 , 12, 1636-43	6.7	31
61	Optimized pacing mode for hypertrophic cardiomyopathy: Impact of ECG fusion during pacing. <i>Heart Rhythm</i> , 2015 , 12, 909-16	6.7	8
60	Short QT and atrial fibrillation: A mutation-specific disease. Late follow-up in three unrelated children. <i>Heart Rhythm Case Reports</i> , 2015 , 1, 193-197	1	10
59	A novel mutation in lamin a/c causing familial dilated cardiomyopathy associated with sudden cardiac death. <i>Journal of Cardiac Failure</i> , 2015 , 21, 217-25	3.3	19

58	Genetic analysis, in silico prediction, and family segregation in long QT syndrome. <i>European Journal of Human Genetics</i> , 2015 , 23, 79-85	5.3	12
57	Genetics of channelopathies associated with sudden cardiac death. <i>Global Cardiology Science & Practice</i> , 2015 , 2015, 39	0.7	23
56	Determining the pathogenicity of genetic variants associated with cardiac channelopathies. <i>Scientific Reports</i> , 2015 , 5, 7953	4.9	37
55	Rare Titin (TTN) Variants in Diseases Associated with Sudden Cardiac Death. <i>International Journal of Molecular Sciences</i> , 2015 , 16, 25773-87	6.3	12
54	Genetic Analysis of Arrhythmogenic Diseases in the Era of NGS: The Complexity of Clinical Decision-Making in Brugada Syndrome. <i>PLoS ONE</i> , 2015 , 10, e0133037	3.7	32
53	Comprehensive Genetic Characterization of a Spanish Brugada Syndrome Cohort. <i>PLoS ONE</i> , 2015 , 10, e0132888	3.7	19
52	Dual fatty acid synthase and HER2 signaling blockade shows marked antitumor activity against breast cancer models resistant to anti-HER2 drugs. <i>PLoS ONE</i> , 2015 , 10, e0131241	3.7	37
51	Identification of N-terminal protein acetylation and arginine methylation of the voltage-gated sodium channel in end-stage heart failure human heart. <i>Journal of Molecular and Cellular Cardiology</i> , 2014 , 76, 126-9	5.8	34
50	Common variation in fatty acid metabolic genes and risk of incident sudden cardiac arrest. <i>Heart Rhythm</i> , 2014 , 11, 471-7	6.7	12
49	The role of clinical, genetic and segregation evaluation in sudden infant death. <i>Forensic Science International</i> , 2014 , 242, 9-15	2.6	17
48	Stop-gain mutations in PKP2 are associated with a later age of onset of arrhythmogenic right ventricular cardiomyopathy. <i>PLoS ONE</i> , 2014 , 9, e100560	3.7	16
47	Identification of Genetic Alterations, as Causative Genetic Defects in Long QT Syndrome, Using Next Generation Sequencing Technology. <i>PLoS ONE</i> , 2014 , 9, e114894	3.7	23
46	The smooth muscle-type β subunit potentiates activation by DiBAC4(3) in recombinant BK channels. <i>Channels</i> , 2014 , 8, 95-102	3	1
45	Post-mortem genetic analysis in juvenile cases of sudden cardiac death. <i>Forensic Science International</i> , 2014 , 245, 30-7	2.6	34
44	Subcellular localisation of retromer in post-endocytic pathways of polarised Madin-Darby canine kidney cells. <i>Biology of the Cell</i> , 2014 , 106, 377-93	3.5	12
43	A missense mutation in the sodium channel β subunit reveals SCN2B as a new candidate gene for Brugada syndrome. <i>Human Mutation</i> , 2013 , 34, 961-6	4.7	78
42	Concomitant Brugada-like and short QT electrocardiogram linked to SCN5A mutation. <i>European Journal of Human Genetics</i> , 2012 , 20, 1189-92	5.3	23
41	CONCOMITANT BRUGADA-LIKE AND SHORT QT ELECTROCARDIOGRAM LINKED TO SCN5A MUTATION. <i>Heart</i> , 2012 , 98, E315.3-E316	5.1	1

40	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). <i>Europace</i> , 2011 , 13, 1077-109	3.9	557
39	Sudden death: managing the family, the role of genetics. <i>Heart</i> , 2011 , 97, 676-81	5.1	9
38	Is atrial fibrillation a genetic disease?. <i>Journal of Cardiovascular Electrophysiology</i> , 2005 , 16, 553-6	2.7	24
37	Short QT syndrome and atrial fibrillation caused by mutation in KCNH2. <i>Journal of Cardiovascular Electrophysiology</i> , 2005 , 16, 394-6	2.7	236
36	Sudden death associated with short-QT syndrome linked to mutations in HERG. <i>Circulation</i> , 2004 , 109, 30-5	16.7	685
35	Brugada Syndrome 2001 ,		1
34	Prognostic value of electrophysiologic investigations in Brugada syndrome. <i>Journal of Cardiovascular Electrophysiology</i> , 2001 , 12, 1004-7	2.7	120
33	Single nucleotide polymorphisms and life-threatening arrhythmias: causal or casual?. <i>Journal of Cardiovascular Electrophysiology</i> , 2001 , 12, 1230-1	2.7	2
32	The Brugada syndrome. <i>Cardiovascular Drugs and Therapy</i> , 2001 , 15, 15-7	3.9	2
31	Brugada Syndrome. <i>Circulation</i> , 2001 , 104, 3017-3019	16.7	17
30	Use of intravenous antiarrhythmics to identify concealed Brugada syndrome. <i>Current Controlled Trials in Cardiovascular Medicine</i> , 2000 , 1, 45-47		18
29	Arrhythmia induction by antiarrhythmic drugs. <i>PACE - Pacing and Clinical Electrophysiology</i> , 2000 , 23, 291-2	1.6	51
28	The Brugada Syndrome. <i>Annals of Noninvasive Electrocardiology</i> , 2000 , 5, 88-91	1.5	4
27	The Brugada syndrome. <i>Current Cardiology Reports</i> , 2000 , 2, 507-14	4.2	54
26	Gene therapy for cardiovascular diseases. <i>Expert Opinion on Therapeutic Patents</i> , 2000 , 10, 1385-1393	6.8	2
25	Ionic mechanisms responsible for the electrocardiographic phenotype of the Brugada syndrome are temperature dependent. <i>Circulation Research</i> , 1999 , 85, 803-9	15.7	469
24	Genetics of cardiovascular disease with emphasis on atrial fibrillation. <i>Journal of Interventional Cardiac Electrophysiology</i> , 1999 , 3, 7-13	2.4	8
23	The Brugada Syndrome. <i>Journal of Interventional Cardiac Electrophysiology</i> , 1999 , 3, 202-204		2

22	Genetic basis and molecular mechanism for idiopathic ventricular fibrillation. <i>Nature</i> , 1998 , 392, 293-6	50.4	1455
21	The molecular genetics of arrhythmias and sudden death. <i>Clinical Cardiology</i> , 1998 , 21, 553-60	3.3	18
20	Right bundle-branch block and ST-segment elevation in leads V1 through V3: a marker for sudden death in patients without demonstrable structural heart disease. <i>Circulation</i> , 1998 , 97, 457-60	16.7	592
19	Prognosis in Individuals with Brugada Syndrome184-193		
18	Atrial Tachyarrhythmias in Brugada Syndrome178-183		1
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