Andrea Nava

List of Publications by Year in Descending Order

87 papers	13,251 citations	49 h-index	89 g-index
89 ext. papers	14,781 ext. citations	6.7 avg, IF	5.28 L-index

#	Paper	IF	Citations
87	Noninvasive cardiac screening in young athletes with ventricular arrhythmias. *American Journal of Cardiology*, **2013**, 111, 557-62	3	30
86	Desmin mutations and arrhythmogenic right ventricular cardiomyopathy. *American Journal of Cardiology*, **2013**, 111, 400-5	3	51
85	Identification of a PKP2 gene deletion in a family with arrhythmogenic right ventricular cardiomyopathy. *European Journal of Human Genetics*, **2013**, 21, 1226-31	5.3	33
84	Compound and digenic heterozygosity predicts lifetime arrhythmic outcome and sudden cardiac death in desmosomal gene-related arrhythmogenic right ventricular cardiomyopathy. *Circulation: Cardiovascular Genetics*, **2013**, 6, 533-42		168
83	Mutations in the area composita protein □-catenin are associated with arrhythmogenic right ventricular cardiomyopathy. *European Heart Journal*, **2013**, 34, 201-10	9.5	142
82	Follow-up with exercise test of effort-induced ventricular arrhythmias linked to ryanodine receptor type 2 gene mutations. *American Journal of Cardiology*, **2012**, 109, 1015-9	3	7
81	Clinical phenotype and diagnosis of arrhythmogenic right ventricular cardiomyopathy in pediatric patients carrying desmosomal gene mutations. *Heart Rhythm*, **2011**, 8, 1686-95	6.7	56
80	The p.A897KfsX4 frameshift variation in desmocollin-2 is not a causative mutation in arrhythmogenic right ventricular cardiomyopathy. *European Journal of Human Genetics*, **2010**, 18, 776-82	5.3	17
79	Diagnosis of arrhythmogenic right ventricular cardiomyopathy/dysplasia: proposed modification of the task force criteria. *Circulation*, **2010**, 121, 1533-41	16.7	1341
78	Diagnosis of arrhythmogenic right ventricular cardiomyopathy/dysplasia: proposed modification of the Task Force Criteria. *European Heart Journal*, **2010**, 31, 806-14	9.5	889
77	A long lasting electrocardiographic history. *Heart Rhythm*, **2010**, 7, 1521	6.7	1
76	Compound and digenic heterozygosity contributes to arrhythmogenic right ventricular cardiomyopathy. *Journal of the American College of Cardiology*, **2010**, 55, 587-97	15.1	242
75	Multiple mutations in desmosomal proteins encoding genes in arrhythmogenic right ventricular cardiomyopathy/dysplasia. *Heart Rhythm*, **2010**, 7, 22-9	6.7	143
74	Myocyte necrosis underlies progressive myocardial dystrophy in mouse dsg2-related arrhythmogenic right ventricular cardiomyopathy. *Journal of Experimental Medicine*, **2009**, 206, 1787-802	16.6	140
73	Electrocardiographic pattern in arrhythmogenic right ventricular cardiomyopathy. *American Journal of Cardiology*, **2009**, 103, 1302-8	3	68
72	Arrhythmogenic right ventricular cardiomyopathy. *Lancet, The*, **2009**, 373, 1289-300	40	622
71	Comparison of clinical features of arrhythmogenic right ventricular cardiomyopathy in men versus women. *American Journal of Cardiology*, **2008**, 102, 1252-7	3	68

70	Quantitative assessment of endomyocardial biopsy in arrhythmogenic right ventricular cardiomyopathy/dysplasia: an in vitro validation of diagnostic criteria. *European Heart Journal*, **2008**, 29, 2760-71	9.5	128
69	Missense mutations in desmocollin-2 N-terminus, associated with arrhythmogenic right ventricular cardiomyopathy, affect intracellular localization of desmocollin-2 in vitro. *BMC Medical Genetics*, **2007**, 8, 65	2.1	51
68	Homozygous SCN5A mutation in Brugada syndrome with monomorphic ventricular tachycardia and structural heart abnormalities. *Europace*, **2007**, 9, 391-7	3.9	35
67	Introduction: Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia Clarified **2007**, 1-5		1
66	Genotype-Phenotype Correlations **2007**, 21-28		1
65	Long-term follow-up of the signal-averaged ECG in arrhythmogenic right ventricular cardiomyopathy: correlation with arrhythmic events and echocardiographic findings. *Europace*, **2006**, 8, 423-9	3.9	26
64	Mutations in desmoglein-2 gene are associated with arrhythmogenic right ventricular cardiomyopathy. *Circulation*, **2006**, 113, 1171-9	16.7	450
63	Ultrastructural evidence of intercalated disc remodelling in arrhythmogenic right ventricular cardiomyopathy: an electron microscopy investigation on endomyocardial biopsies. *European Heart Journal*, **2006**, 27, 1847-54	9.5	183
62	Letter regarding article by Norman et al, "novel mutation in desmoplakin causes arrhythmogenic left ventricular cardiomyopathy". *Circulation*, **2006**, 113, e68; author reply e69	16.7	3
61	Pregnancy in women with arrhythmogenic right ventricular cardiomyopathy/dysplasia. *European Journal of Obstetrics, Gynecology and Reproductive Biology*, **2006**, 127, 186-9	2.4	61
60	Late-onset arrhythmogenic right ventricular cardiomyopathy. *Journal of Cardiovascular Medicine*, **2006**, 7, 74-6	1.9	8
59	Arrhythmogenic right ventricular cardiomyopathy is a life-threatening disease at high risk for cardiac arrest during effort. Minor forms are as dangerous as major forms?. *Journal of Cardiovascular Medicine*, **2006**, 7, 246-9	1.9	1
58	Clinical profile of four families with arrhythmogenic right ventricular cardiomyopathy caused by dominant desmoplakin mutations. *European Heart Journal*, **2005**, 26, 1666-75	9.5	229
57	Juvenile sudden death in a family with polymorphic ventricular arrhythmias caused by a novel RyR2 gene mutation: evidence of specific morphological substrates. *Human Pathology*, **2005**, 36, 761-7	3.7	50
56	Echocardiographic findings in patients with arrhythmogenic right ventricular cardiomyopathy/dysplasia. *Journal of the American College of Cardiology*, **2005**, 46, 1962; author reply 1962-3	15.1	
55	Regulatory mutations in transforming growth factor-beta3 gene cause arrhythmogenic right ventricular cardiomyopathy type 1. *Cardiovascular Research*, **2005**, 65, 366-73	9.9	315
54	Three-dimensional electroanatomic voltage mapping increases accuracy of diagnosing arrhythmogenic right ventricular cardiomyopathy/dysplasia. *Circulation*, **2005**, 111, 3042-50	16.7	200
53	Letter regarding article by Nasir et al, "Electrocardiographic features of arrhythmogenic right ventricular dysplasia/cardiomyopathy according to disease severity: a need to broaden diagnostic criteria". *Circulation*, **2005**, 112, e68; author reply e68-9	16.7	

52	Arrhythmogenic right ventricular cardiomyopathy: clinical registry and database, evaluation of therapies, pathology registry, DNA banking. *European Heart Journal*, **2004**, 25, 531-4	9.5	44
51	Noninvasive risk stratification in arrhythmogenic right ventricular cardiomyopathy. *Annals of Noninvasive Electrocardiology*, **2003**, 8, 161-9	1.5	24
50	Arrhythmogenic right ventricular cardiomyopathy type 1 (ARVD1): confirmation of locus assignment and mutation screening of four candidate genes. *European Journal of Human Genetics*, **2003**, 11, 69-76	5.3	50
49	Signal-averaged electrocardiographic parameter progression as a marker of increased electrical instability in two cases with an overt form of arrhythmogenic right ventricular cardiomyopathy. *PACE - Pacing and Clinical Electrophysiology*, **2002**, 25, 362-4	1.6	6
48	Heart rate variability in arrhythmogenic right ventricular cardiomyopathy correlation with clinical and prognostic features. *PACE - Pacing and Clinical Electrophysiology*, **2002**, 25, 1285-92	1.6	15
47	Screening for ryanodine receptor type 2 mutations in families with effort-induced polymorphic ventricular arrhythmias and sudden death: early diagnosis of asymptomatic carriers. *Journal of the American College of Cardiology*, **2002**, 40, 341-9	15.1	188
46	Mutation in human desmoplakin domain binding to plakoglobin causes a dominant form of arrhythmogenic right ventricular cardiomyopathy. *American Journal of Human Genetics*, **2002**, 71, 1200-6	11	493
45	Dispersion of ventricular depolarization-repolarization: a noninvasive marker for risk stratification in arrhythmogenic right ventricular cardiomyopathy. *Circulation*, **2001**, 103, 3075-80	16.7	142
44	Right bundle branch block, right precordial st-segment elevation, and sudden death in young people. *Circulation*, **2001**, 103, 710-7	16.7	186
43	Arrhythmogenic right ventricular cardiomyopathy: current diagnostic and management strategies. *Cardiology in Review*, **2001**, 9, 259-65	3.2	60
42	Arrhythmogenic right ventricular dysplasia: cardiomyopathy current opinions on diagnostic and therapeutic aspects. *Current Opinion in Cardiology*, **2001**, 16, 8-16	2.1	14
41	Arrhythmogenic right ventricular dysplasia/cardiomyopathy: need for an international registry. European Society of Cardiology and the Scientific Council on Cardiomyopathies of the World Heart Federation. *Journal of Cardiovascular Electrophysiology*, **2000**, 11, 827-32	2.7	22
40	Familial effort polymorphic ventricular arrhythmias in arrhythmogenic right ventricular cardiomyopathy map to chromosome 1q42-43. *American Journal of Cardiology*, **2000**, 85, 573-9	3	67
39	Arrhythmogenic right ventricular dysplasia/cardiomyopathy: need for an international registry. Study Group on Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy of the Working Groups on Myocardial and Pericardial Disease and Arrhythmias of the European Society of Cardiology and of the Scientific Council on Cardiomyopathies of the World Heart Federation. *Circulation*, **2000**, 101, E101-6	16.7	208
38	Right bundle-branch block, ST-segment elevation, and sudden death. *Circulation*, **2000**, 101, E176	16.7	1
37	Characterization of C14orf4, a novel intronless human gene containing a polyglutamine repeat, mapped to the ARVD1 critical region. *Biochemical and Biophysical Research Communications*, **2000**, 278, 766-74	3.4	42
36	Clinical profile and long-term follow-up of 37 families with arrhythmogenic right ventricular cardiomyopathy. *Journal of the American College of Cardiology*, **2000**, 36, 2226-33	15.1	355
35	Late potentials and ventricular arrhythmias in arrhythmogenic right ventricular cardiomyopathy. *American Journal of Cardiology*, **1999**, 83, 1214-9	3	96

34	Incidence of atrial fibrillation in patients with different mode of pacing. Long-term follow-up. *PACE - Pacing and Clinical Electrophysiology*, **1998**, 21, 260-3	1.6	13
33	Radiographic assessment of atrial dipole position in single pass lead VDD and DDD pacing. The Multicenter Study Group. *PACE - Pacing and Clinical Electrophysiology*, **1998**, 21, 2240-5	1.6	3
32	ARVD4, a new locus for arrhythmogenic right ventricular cardiomyopathy, maps to chromosome 2 long arm. *Genomics*, **1997**, 45, 259-63	4.3	144
31	Spectrum of clinicopathologic manifestations of arrhythmogenic right ventricular cardiomyopathy/dysplasia: a multicenter study. *Journal of the American College of Cardiology*, **1997**, 30, 1512-20	15.1	750
30	Arrhythmogenic right ventricular cardiomyopathy a still underrecognized clinic entity. *Trends in Cardiovascular Medicine*, **1997**, 7, 84-90	6.9	58
29	Arrhythmogenic right ventricular cardiomyopathy: a survey of the investigations at the University of Padua. *Clinical Cardiology*, **1997**, 20, 333-6	3.3	11
28	Familial cardiomyopathy underlies syndrome of right bundle branch block, ST segment elevation and sudden death. *Journal of the American College of Cardiology*, **1996**, 27, 443-8	15.1	189
27	Upright tilt test: correlation between results and patient clinical features. *PACE - Pacing and Clinical Electrophysiology*, **1996**, 19, 1582-7	1.6	11
26	Arrhythmogenic right ventricular cardiomyopathy. Dysplasia, dystrophy, or myocarditis?. *Circulation* , **1996**, 94, 983-91	16.7	581
25	Arrhythmogenic right ventricular cardiomyopathy in young versus adult patients: similarities and differences. *Journal of the American College of Cardiology*, **1995**, 25, 655-64	15.1	121
24	Signal-averaged electrocardiography in familial form of arrhythmogenic right ventricular cardiomyopathy. *American Journal of Cardiology*, **1995**, 75, 1038-41	3	44
23	The gene for arrhythmogenic right ventricular cardiomyopathy maps to chromosome 14q23-q24. *Human Molecular Genetics*, **1994**, 3, 959-62	5.6	280
22	Long-term follow-up of patients with single lead VDD stimulation. *PACE - Pacing and Clinical Electrophysiology*, **1994**, 17, 1854-8	1.6	13
21	QT-interval variability in hypertrophic cardiomyopathy patients with cardiac arrest. *International Journal of Cardiology*, **1994**, 45, 121-7	3.2	45
20	Right ventricular cardiomyopathy in identical and nonidentical young twins. *American Heart Journal*, **1993**, 126, 1187-93	4.9	21
19	Right bundle branch block, persistent ST segment elevation and sudden cardiac death. *Journal of the American College of Cardiology*, **1993**, 22, 633	15.1	12
18	Endomyocardial biopsy in right ventricular cardiomyopathy. *International Journal of Cardiology*, **1993**, 40, 273-82	3.2	60
17	Comparison of QT dispersion in hypertrophic cardiomyopathy between patients with and without ventricular arrhythmias and sudden death. *American Journal of Cardiology*, **1993**, 72, 973-6	3	320

16	Clinical profile of concealed form of arrhythmogenic right ventricular cardiomyopathy presenting with apparently idiopathic ventricular arrhythmias. *International Journal of Cardiology*, **1992**, 35, 195-206; discussion 207-9	3.2	51
15	A casual spontaneous mutation as possible cause of the familial form of arrhythmogenic right ventricular cardiomyopathy (arrhythmogenic right ventricular dysplasia). *Clinical Cardiology*, **1992**, 15, 217-9	3.3	9
14	Spontaneous and induced vasodepressor/vasovagal syncope in hypertrophic cardiomyopathy. *Clinical Cardiology*, **1992**, 15, 387-9	3.3	4
13	Electrophysiological features and the clinical follow-up of patients affected by ventricular tachycardias. *PACE - Pacing and Clinical Electrophysiology*, **1991**, 14, 245	1.6	
12	Cardiomyopathy: a necessary revision of the WHO classification. *International Journal of Cardiology*, **1991**, 30, 1-7	3.2	34
11	Arrhythmia development in a young subject with right ventricular cardiomyopathy (right ventricular dysplasia). *International Heart Journal*, **1991**, 32, 403-8		
10	Diagnostic accuracy of right ventriculography in arrhythmogenic right ventricular cardiomyopathy. *American Journal of Cardiology*, **1990**, 66, 741-5	3	93
9	Monomorphic repetitive rhythms originating from the outflow tract in patients with minor forms of right ventricular cardiomyopathy. *International Journal of Cardiology*, **1990**, 27, 211-21	3.2	8
8	Sudden death in young competitive athletes: clinicopathologic correlations in 22 cases. *American Journal of Medicine*, **1990**, 89, 588-96	2.4	436
7	Ventricular fibrillation without apparent heart disease: description of six cases. *American Heart Journal*, **1989**, 118, 1203-9	4.9	269
6	Electrovectorcardiographic study of negative T waves on precordial leads in arrhythmogenic right ventricular dysplasia: relationship with right ventricular volumes. *Journal of Electrocardiology*, **1988**, 21, 239-45	1.4	69
5	Complex arrhythmias in a patient with predominantly right ventricular cardiomyopathy. *International Journal of Cardiology*, **1988**, 19, 268-71	3.2	1
4	Familial occurrence of right ventricular dysplasia: a study involving nine families. *Journal of the American College of Cardiology*, **1988**, 12, 1222-8	15.1	325
3	Juvenile sudden death and effort ventricular tachycardias in a family with right ventricular cardiomyopathy. *International Journal of Cardiology*, **1988**, 21, 111-26	3.2	40
2	Right ventricular cardiomyopathy and sudden death in young people. *New England Journal of Medicine*, **1988**, 318, 129-33	59.2	1266
1	A polymorphic form of familial arrhythmogenic right ventricular dysplasia. *American Journal of Cardiology*, **1987**, 59, 1405-9	3	86