Andrea Nava

List of Publications by Year
in descending order

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88	16,047	38742 50	56724 83
papers	citations	h-index	g-index

89	89	89	6413
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Noninvasive Cardiac Screening in Young Athletes With Ventricular Arrhythmias. American Journal of Cardiology, 2013, 111, 557-562.	1.6	34
2	Desmin Mutations and Arrhythmogenic Right Ventricular Cardiomyopathy. American Journal of Cardiology, 2013, 111, 400-405.	1.6	62
3	Identification of a PKP2 gene deletion in a family with arrhythmogenic right ventricular cardiomyopathy. European Journal of Human Genetics, 2013, 21, 1226-1231.	2.8	39
4	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
5	Mutations in the area composita protein Î±T-catenin are associated with arrhythmogenic right ventricular cardiomyopathy. European Heart Journal, 2013, 34, 201-210.	2.2	175
6	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.	1.6	8
7	Clinical phenotype and diagnosis of arrhythmogenic right ventricular cardiomyopathy in pediatric patients carrying desmosomal gene mutations. Heart Rhythm, 2011, 8, 1686-1695.	0.7	66
8	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, 776-782.	2.8	19
9	Diagnosis of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Circulation, 2010, 121, 1533-1541.	1.6	1,839
10	Diagnosis of arrhythmogenic right ventricular cardiomyopathy/dysplasia: Proposed Modification of the Task Force Criteria. European Heart Journal, 2010, 31, 806-814.	2.2	1,177
11	A Long Lasting Electrocardiographic History. Heart Rhythm, 2010, 7, 1521.	0.7	4
12	Compound and Digenic Heterozygosity Contributes to Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American College of Cardiology, 2010, 55, 587-597.	2.8	282
13	Multiple mutations in desmosomal proteins encoding genes in arrhythmogenic right ventricular cardiomyopathy/dysplasia. Heart Rhythm, 2010, 7, 22-29.	0.7	161
14	Myocyte necrosis underlies progressive myocardial dystrophy in mouse dsg2-related arrhythmogenic right ventricular cardiomyopathy. Journal of Experimental Medicine, 2009, 206, 1787-1802.	8.5	184
15	Electrocardiographic Pattern in Arrhythmogenic Right Ventricular Cardiomyopathy. American Journal of Cardiology, 2009, 103, 1302-1308.	1.6	82
16	Arrhythmogenic right ventricular cardiomyopathy. Lancet, The, 2009, 373, 1289-1300.	13.7	785
17	Comparison of Clinical Features of Arrhythmogenic Right Ventricular Cardiomyopathy in Men Versus Women. American Journal of Cardiology, 2008, 102, 1252-1257.	1.6	81
18	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.	2.2	161

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19	Homozygous SCN5A mutation in Brugada syndrome with monomorphic ventricular tachycardia and structural heart abnormalities. Europace, 2007, 9, 391-397.	1.7	41
20	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	61
21	Introduction: Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia Clarified. , 2007, , 1-5.		2
22	Genotype-Phenotype Correlations. , 2007, , 21-28.		2
23	Pregnancy in women with arrhythmogenic right ventricular cardiomyopathy/dysplasia. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2006, 127, 186-189.	1.1	73
24	Late-onset arrhythmogenic right ventricular cardiomyopathy. Journal of Cardiovascular Medicine, 2006, 7, 74-76.	1.5	8
25	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, 246-249.	1.5	1
26	Long-term follow-up of the signal-averaged ECG in arrhythmogenic right ventricular cardiomyopathy: correlation with arrhythmic events and echocardiographic findings. Europace, 2006, 8, 423-429.	1.7	29
27	Mutations in Desmoglein-2 Gene Are Associated With Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation, 2006, 113, 1171-1179.	1.6	509
28	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.	2.2	219
29	Letter Regarding Article by Norman et al, â€œNovel Mutation in Desmoplakin Causes Arrhythmogenic Left Ventricular Cardiomyopathyâ€• Circulation, 2006, 113, e68; author reply e69.	1.6	5
30	Regulatory mutations in transforming growth factor-?3 gene cause arrhythmogenic right ventricular cardiomyopathy type 1. Cardiovascular Research, 2005, 65, 366-373.	3.8	364
31	Three-Dimensional Electroanatomic Voltage Mapping Increases Accuracy of Diagnosing Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Circulation, 2005, 111, 3042-3050.	1.6	237
32	Letters Regarding Article by Nasir et al, â€œElectrocardiographic Features of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy According to Disease Severity: A Need to Broaden Diagnostic Criteriaâ€• Circulation, 2005, 112, e68; author reply e68-9.	1.6	1
33	Clinical profile of four families with arrhythmogenic right ventricular cardiomyopathy caused by dominant desmoplakin mutations. European Heart Journal, 2005, 26, 1666-1675.	2.2	267
34	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, 761-767.	2.0	58
35	Echocardiographic Findings in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Journal of the American College of Cardiology, 2005, 46, 1962.	2.8	0
36	Arrhythmogenic right ventricular cardiomyopathy: clinical registry and database, evaluation of therapies, pathology registry, DNA banking. European Heart Journal, 2004, 25, 531-534.	2.2	48

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37	Noninvasive Risk Stratification in Arrhythmogenic Right Ventricular Cardiomyopathy. Annals of Noninvasive Electrocardiology, 2003, 8, 161-169.	1.1	28
38	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.	2.8	54
39	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.	2.8	213
40	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.	6.2	570
41	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, 362-364.	1.2	6
42	Heart Rate Variability in Arrhythmogenic Right Ventricular Cardiomyopathy Correlation with Clinical and Prognostic Features. PACE - Pacing and Clinical Electrophysiology, 2002, 25, 1285-1292.	1.2	18
43	Arrhythmogenic Right Ventricular Cardiomyopathy: Current Diagnostic and Management Strategies. Cardiology in Review, 2001, 9, 259-265.	1.4	71
44	Arrhythmogenic Right Ventricular Dysplasia: cardiomyopathy current opinions on diagnostic and therapeutic aspects. Current Opinion in Cardiology, 2001, 16, 8-16.	1.8	17
45	Dispersion of Ventricular Depolarization-Repolarization. Circulation, 2001, 103, 3075-3080.	1.6	158
46	Right Bundle Branch Block, Right Precordial ST-Segment Elevation, and Sudden Death in Young People. Circulation, 2001, 103, 710-717.	1.6	223
47	Arrhythmogenic Right Ventricular Dyspiasia/Cardiomyopathy:. Journal of Cardiovascular Electrophysiology, 2000, 11, 827-832.	1.7	24
48	Familial effort polymorphic ventricular arrhythmias in arrhythmogenic right ventricular cardiomyopathy map to chromosome 1q42-43. American Journal of Cardiology, 2000, 85, 573-579.	1.6	84
49	Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. Circulation, 2000, 101, E101-6.	1.6	272
50	Right Bundle-Branch Block, ST-Segment Elevation, and Sudden Death. Circulation, 2000, 101, E176.	1.6	1
51	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, 766-774.	2.1	49
52	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.	2.8	414
53	Late potentials and ventricular arrhythmias in arrhythmogenic right ventricular cardiomyopathy. American Journal of Cardiology, 1999, 83, 1214-1219.	1.6	106
54	Incidence of Atrial Fibrillation in Patients with Different Mode of Pacing. Long-term Follow-up. PACE - Pacing and Clinical Electrophysiology, 1998, 21, 260-263.	1.2	15

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55	Radiographic Assessment of Atrial Dipole Position in Single Pass Lead VDD and DDD Pacing. PACE - Pacing and Clinical Electrophysiology, 1998, 21, 2240-2245.	1.2	3
56	ARVD4, a New Locus for Arrhythmogenic Right Ventricular Cardiomyopathy, Maps to Chromosome 2 Long Arm. Genomics, 1997, 45, 259-263.	2.9	170
57	Spectrum of Clinicopathologic Manifestations of Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia: A Multicenter Study. Journal of the American College of Cardiology, 1997, 30, 1512-1520.	2.8	884
58	Arrhythmogenic Right Ventricular Cardiomyopathy A Still Underrecognized Clinic Entity. Trends in Cardiovascular Medicine, 1997, 7, 84-90.	4.9	68
59	Arrhythmogenic right ventricular cardiomyopathy: a survey of the investigations at the university of padua. Clinical Cardiology, 1997, 20, 333-336.	1.8	13
60	Familial cardiomyopathy underlies syndrome of right bundle branch block, ST segment elevation and sudden death. Journal of the American College of Cardiology, 1996, 27, 443-448.	2.8	229
61	Endomyocardial biopsy in arrhythmogenic right ventricular cardiomyopathy. American Heart Journal, 1996, 132, 203-206.	2.7	121
62	Upright Tilt Test: Correlation Between Results and Patient Clinical Features. PACE - Pacing and Clinical Electrophysiology, 1996, 19, 1582-1587.	1.2	12
63	Arrhythmogenic Right Ventricular Cardiomyopathy. Circulation, 1996, 94, 983-991.	1.6	724
64	Signal-averaged electrocardiography in familial form of arrhythmogenic right ventricular cardiomyopathy. American Journal of Cardiology, 1995, 75, 1038-1041.	1.6	48
65	Arrhythmogenic right ventricular cardiomyopathy in young versus adult patients: Similarities and differences. Journal of the American College of Cardiology, 1995, 25, 655-664.	2.8	140
66	The gene for arrhythmogenic right ventricular cardiomyopathy maps to chromosome 14q23â€“q24. Human Molecular Genetics, 1994, 3, 959-962.	2.9	326
67	Long-Term Follow-Up of Patients with Single Lead VDD Stimulation. PACE - Pacing and Clinical Electrophysiology, 1994, 17, 1854-1858.	1.2	18
68	QT-interval variability in hypertrophic cardiomyopathy patients with cardiac arrest. International Journal of Cardiology, 1994, 45, 121-127.	1.7	53
69	Comparison of QT dispersion in hypertrophic cardiomyopathy between patients with and without ventricular arrhythmias and sudden death. American Journal of Cardiology, 1993, 72, 973-976.	1.6	358
70	Right ventricular cardiomyopathy in identical and nonidentical young twins. American Heart Journal, 1993, 126, 1187-1193.	2.7	22
71	Right bundle branch block, persistent ST segment elevation and sudden cardiac death. Journal of the American College of Cardiology, 1993, 22, 633.	2.8	31
72	Endomyocardial biopsy in right ventricular cardiomyopathy. International Journal of Cardiology, 1993, 40, 273-282.	1.7	64

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73	Clinical profile of concealed form of arrhythmogenic right ventricular cardiomyopathy presenting with apparently idiopathic ventricular arrhythmias. International Journal of Cardiology, 1992, 35, 195-206.	1.7	60
74	A casual spontaneous mutation as possible cause of the familial form of arrhythmogenic right ventricular cardiomyopathy (arrhythmogenic right ventricular dysplasia). Clinical Cardiology, 1992, 15, 217-219.	1.8	9
75	Spontaneous and induced vasodepressor/vasovagal syncope in hypertrophic cardiomyopathy. Clinical Cardiology, 1992, 15, 387-389.	1.8	6
76	Cardiomyopathy: a necessary revision of the WHO classification. International Journal of Cardiology, 1991, 30, 1-7.	1.7	39
77	LETTERS TO THE EDITOR. PACE - Pacing and Clinical Electrophysiology, 1991, 14, 245-245.	1.2	0
78	Arrhythmia Development in a Young Subject with Right Ventricular Cardiomyopathy. (Right) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 542 T	0.6	0
79	Diagnostic accuracy of right ventriculography in arrhythmogenic right ventricular cardiomyopathy. American Journal of Cardiology, 1990, 66, 741-745.	1.6	103
80	Monomorphic repetitive rhythms originating from the outflow tract in patients with minor forms of right ventricular cardiomyopathy. International Journal of Cardiology, 1990, 27, 211-221.	1.7	11
81	Sudden death in young competitive athletes: clinicopathologic correlations in 22 cases. American Journal of Medicine, 1990, 89, 588-596.	1.5	518
82	Ventricular fibrillation without apparent heart disease: Description of six cases. American Heart Journal, 1989, 118, 1203-1209.	2.7	338
83	Electrovectorcardiographic study of negative T waves on precordial leads in arrhythmogenic right ventricular dysplasia: Relationship with right ventricular volumes. Journal of Electrocardiology, 1988, 21, 239-245.	0.9	75
84	Complex arrhythmias in a patient with predominantly right ventricular cardiomyopathy. International Journal of Cardiology, 1988, 19, 268-271.	1.7	1
85	Familial occurrence of right ventricular dysplasia: A study involving nine families. Journal of the American College of Cardiology, 1988, 12, 1222-1228.	2.8	362
86	Juvenile sudden death and effort ventricular tachycardias in a family with right ventricular cardiomyopathy. International Journal of Cardiology, 1988, 21, 111-123.	1.7	42
87	Right Ventricular Cardiomyopathy and Sudden Death in Young People. New England Journal of Medicine, 1988, 318, 129-133.	27.0	1,490
88	A polymorphic form of familial arrhythmogenic right ventricular dysplasia. American Journal of Cardiology, 1987, 59, 1405-1409.	1.6	93