

Angeliki Asimaki

List of Publications by Citations

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64
papers

3,956
citations

31
h-index

62
g-index

68
ext. papers

4,805
ext. citations

7.9
avg, IF

4.83
L-index

#	Paper	IF	Citations
64	Clinical and genetic characterization of families with arrhythmogenic right ventricular dysplasia/cardiomyopathy provides novel insights into patterns of disease expression. <i>Circulation</i> , 2007 , 115, 1710-20	16.7	401
63	A new diagnostic test for arrhythmogenic right ventricular cardiomyopathy. <i>New England Journal of Medicine</i> , 2009 , 360, 1075-84	59.2	357
62	Arrhythmogenic right ventricular dysplasia/cardiomyopathy associated with mutations in the desmosomal gene desmocollin-2. <i>American Journal of Human Genetics</i> , 2006 , 79, 978-84	11	299
61	Phospholamban R14del mutation in patients diagnosed with dilated cardiomyopathy or arrhythmogenic right ventricular cardiomyopathy: evidence supporting the concept of arrhythmogenic cardiomyopathy. <i>European Journal of Heart Failure</i> , 2012 , 14, 1199-207	12.3	270
60	A novel dominant mutation in plakoglobin causes arrhythmogenic right ventricular cardiomyopathy. <i>American Journal of Human Genetics</i> , 2007 , 81, 964-73	11	184
59	Clinical expression of plakophilin-2 mutations in familial arrhythmogenic right ventricular cardiomyopathy. <i>Circulation</i> , 2006 , 113, 356-64	16.7	167
58	Identification of a new modulator of the intercalated disc in a zebrafish model of arrhythmogenic cardiomyopathy. <i>Science Translational Medicine</i> , 2014 , 6, 240ra74	17.5	165
57	Altered desmosomal proteins in granulomatous myocarditis and potential pathogenic links to arrhythmogenic right ventricular cardiomyopathy. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2011 , 4, 743-52	6.4	122
56	Desmoglein-2 mutations in arrhythmogenic right ventricular cardiomyopathy: a genotype-phenotype characterization of familial disease. <i>European Heart Journal</i> , 2007 , 28, 581-8	9.5	120
55	Electrophysiological abnormalities precede overt structural changes in arrhythmogenic right ventricular cardiomyopathy due to mutations in desmoplakin-A combined murine and human study. <i>European Heart Journal</i> , 2012 , 33, 1942-53	9.5	117
54	Severe cardiac phenotype with right ventricular predominance in a large cohort of patients with a single missense mutation in the DES gene. <i>Heart Rhythm</i> , 2009 , 6, 1574-83	6.7	117
53	Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. <i>European Heart Journal</i> , 2020 , 41, 1414-1429	9.5	110
52	Remodeling of the cardiac sodium channel, connexin43, and plakoglobin at the intercalated disk in patients with arrhythmogenic cardiomyopathy. <i>Heart Rhythm</i> , 2013 , 10, 412-9	6.7	100
51	Diagnosis of arrhythmogenic cardiomyopathy: The Padua criteria. <i>International Journal of Cardiology</i> , 2020 , 319, 106-114	3.2	89
50	Filamin C Truncation Mutations Are Associated With Arrhythmogenic Dilated Cardiomyopathy and Changes in the Cell-Cell Adhesion Structures. <i>JACC: Clinical Electrophysiology</i> , 2018 , 4, 504-514	4.6	84
49	Central role for GSK3 β in the pathogenesis of arrhythmogenic cardiomyopathy. <i>JCI Insight</i> , 2016 , 1,	9.9	84
48	Desmin mutations as a cause of right ventricular heart failure affect the intercalated disks. <i>Heart Rhythm</i> , 2010 , 7, 1058-64	6.7	83

47	Sudden Death and Left Ventricular Involvement in Arrhythmogenic Cardiomyopathy. <i>Circulation</i> , 2019 , 139, 1786-1797	16.7	70
46	Arrhythmogenic right ventricular cardiomyopathy/dysplasia on the basis of the revised diagnostic criteria in affected families with desmosomal mutations. <i>European Heart Journal</i> , 2011 , 32, 1097-104	9.5	64
45	Connexin43 mutation causes heterogeneous gap junction loss and sudden infant death. <i>Circulation</i> , 2012 , 125, 474-81	16.7	59
44	Arrhythmogenic right ventricular cardiomyopathy caused by deletions in plakophilin-2 and plakoglobin (Naxos disease) in families from Greece and Cyprus: genotype-phenotype relations, diagnostic features and prognosis. <i>European Heart Journal</i> , 2006 , 27, 2208-16	9.5	58
43	Therapeutic Modulation of the Immune Response in Arrhythmogenic Cardiomyopathy. <i>Circulation</i> , 2019 , 140, 1491-1505	16.7	57
42	Definition and treatment of arrhythmogenic cardiomyopathy: an updated expert panel report. <i>European Journal of Heart Failure</i> , 2019 , 21, 955-964	12.3	47
41	Electrical coupling and propagation in engineered ventricular myocardium with heterogeneous expression of connexin43. <i>Circulation Research</i> , 2012 , 110, 1445-53	15.7	43
40	Biallelic Truncating Mutations in ALPK3 Cause Severe Pediatric Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2016 , 67, 515-25	15.1	41
39	A novel desmocollin-2 mutation reveals insights into the molecular link between desmosomes and gap junctions. <i>Heart Rhythm</i> , 2011 , 8, 711-8	6.7	40
38	Persistent lone atrial fibrillation: clinicopathologic study of 19 cases. <i>Heart Rhythm</i> , 2014 , 11, 1250-8	6.7	39
37	Mechanistic insights into arrhythmogenic right ventricular cardiomyopathy caused by desmocollin-2 mutations. <i>Cardiovascular Research</i> , 2011 , 90, 77-87	9.9	37
36	Arrhythmogenic right ventricular cardiomyopathy mutations alter shear response without changes in cell-cell adhesion. <i>Cardiovascular Research</i> , 2014 , 104, 280-9	9.9	35
35	Pathogenesis of Arrhythmogenic Cardiomyopathy. <i>Canadian Journal of Cardiology</i> , 2015 , 31, 1313-24	3.8	34
34	Gap junction remodeling in a case of arrhythmogenic right ventricular dysplasia due to plakophilin-2 mutation. <i>Journal of Cardiovascular Electrophysiology</i> , 2008 , 19, 1212-4	2.7	32
33	Novel missense mutations in exon 15 of desmoglein-2: role of the intracellular cadherin segment in arrhythmogenic right ventricular cardiomyopathy?. <i>Heart Rhythm</i> , 2010 , 7, 1446-53	6.7	30
32	Filamin C variants are associated with a distinctive clinical and immunohistochemical arrhythmogenic cardiomyopathy phenotype. <i>International Journal of Cardiology</i> , 2020 , 307, 101-108	3.2	30
31	Disparate effects of different mutations in plakoglobin on cell mechanical behavior. <i>Cytoskeleton</i> , 2008 , 65, 964-78		27
30	Molecular changes in the heart of a severe case of arrhythmogenic right ventricular cardiomyopathy caused by a desmoglein-2 null allele. <i>Cardiovascular Pathology</i> , 2012 , 21, 275-82	3.8	26

29	Arrhythmogenic right ventricular cardiomyopathy: new insights into mechanisms of disease. <i>Cardiovascular Pathology</i> , 2010 , 19, 166-70	3.8	25
28	Characterizing the Molecular Pathology of Arrhythmogenic Cardiomyopathy in Patient Buccal Mucosa Cells. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2016 , 9, e003688	6.4	24
27	Prolonged RV endocardial activation duration: a novel marker of arrhythmogenic right ventricular dysplasia/cardiomyopathy. <i>Heart Rhythm</i> , 2009 , 6, 769-75	6.7	24
26	The role of endomyocardial biopsy in ARVC: looking beyond histology in search of new diagnostic markers. <i>Journal of Cardiovascular Electrophysiology</i> , 2011 , 22, 111-7	2.7	22
25	Unique epidermolytic bullous dermatosis with associated lethal cardiomyopathy related to novel desmoplakin mutations. <i>Journal of Cutaneous Pathology</i> , 2009 , 36, 553-9	1.7	22
24	Expression of cathepsin K and tartrate-resistant acid phosphatase is not confined to osteoclasts but is a general feature of multinucleated giant cells: systematic analysis. <i>Rheumatology</i> , 2013 , 52, 1529-33	3.9	21
23	Remodeling of cell-cell junctions in arrhythmogenic cardiomyopathy. <i>Cell Communication and Adhesion</i> , 2014 , 21, 13-23		18
22	Arrhythmogenic Cardiomyopathy - New Insights into Disease Mechanisms and Drug Discovery. <i>Progress in Pediatric Cardiology</i> , 2014 , 37, 3-7	0.4	16
21	Exercise triggers CAPN1-mediated AIF truncation, inducing myocyte cell death in arrhythmogenic cardiomyopathy. <i>Science Translational Medicine</i> , 2021 , 13,	17.5	15
20	Gap junctions and arrhythmogenic cardiomyopathy. <i>Heart Rhythm</i> , 2012 , 9, 992-5	6.7	13
19	Development of dilated cardiomyopathy and impaired calcium homeostasis with cardiac-specific deletion of ESRR α . <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2017 , 312, H662-H671	5.2	12
18	Arrhythmogenic right ventricular cardiomyopathy: new insights into disease mechanisms and diagnosis. <i>Journal of Investigative Medicine</i> , 2009 , 57, 861-4	2.9	12
17	RNA sequencing-based transcriptome profiling of cardiac tissue implicates novel putative disease mechanisms in FLNC-associated arrhythmogenic cardiomyopathy. <i>International Journal of Cardiology</i> , 2020 , 302, 124-130	3.2	10
16	Distinct molecular signature of phospholamban p.Arg14del arrhythmogenic cardiomyopathy. <i>Cardiovascular Pathology</i> , 2019 , 40, 2-6	3.8	9
15	Morphometric characterization of collagen and fat in normal ventricular myocardium. <i>Cardiovascular Pathology</i> , 2020 , 48, 107224	3.8	8
14	Brugada syndrome and arrhythmogenic cardiomyopathy: overlapping disorders of the connexome?. <i>Europace</i> , 2021 , 23, 653-664	3.9	8
13	Reduced plakoglobin immunoreactivity in arrhythmogenic cardiomyopathy: methodological considerations. <i>Cardiovascular Pathology</i> , 2013 , 22, 314-8	3.8	7
12	The Novel Desmin Variant p.Leu115Ile Is Associated With a Unique Form of Biventricular Arrhythmogenic Cardiomyopathy. <i>Canadian Journal of Cardiology</i> , 2021 , 37, 857-866	3.8	7

11	Homozygous Truncating Variant in PKP2 Causes Hypoplastic Left Heart Syndrome. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e002397	5.2	6
10	Inflammation and Immune Response in Arrhythmogenic Cardiomyopathy: State-of-the-Art Review. <i>Circulation</i> , 2021 , 144, 1646-1655	16.7	5
9	Cardiac sarcoidosis with severe involvement of the right ventricle: a case report. <i>Autopsy and Case Reports</i> , 2015 , 5, 53-63	0.6	5
8	End stage of arrhythmogenic cardiomyopathy with severe involvement of the interventricular septum. <i>Heart Rhythm</i> , 2013 , 10, 283-9	6.7	4
7	Reduced desmoplakin immunofluorescence signal in arrhythmogenic cardiomyopathy with epicardial right ventricular outflow tract tachycardia. <i>HeartRhythm Case Reports</i> , 2019 , 5, 57-62	1	4
6	Desmosomal COP9 regulates proteome degradation in arrhythmogenic right ventricular dysplasia/cardiomyopathy. <i>Journal of Clinical Investigation</i> , 2021 , 131,	15.9	3
5	Left Dominant Arrhythmogenic Cardiomyopathy Caused by a Novel Nonsense Mutation in Desmoplakin. <i>Revista Espanola De Cardiologia (English Ed)</i> , 2011 , 64, 530-534	0.7	1
4	Histopathological Features and Protein Markers of Arrhythmogenic Cardiomyopathy.. <i>Frontiers in Cardiovascular Medicine</i> , 2021 , 8, 746321	5.4	1
3	Clinical and Molecular Aspects of Naxos Disease. <i>Heart Failure Clinics</i> , 2022 , 18, 89-99	3.3	0
2	MY APPROACH to the patient with arrhythmogenic right ventricular cardiomyopathy (ARVC). <i>Trends in Cardiovascular Medicine</i> , 2017 , 27, 293	6.9	
1	Connexin43 Mutation Causes Heterogeneous Gap Junction Loss and Sudden Infant Death. <i>Circulation</i> , 2011 , 1	16.7	