Alessandra Rampazzo

List of Publications by Year
in descending order

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32 papers	2,738 citations	331670 21 h-index	414414 32 g-index
32 all docs	32 docs citations	32 times ranked	2827 citing authors

#	Article	IF	Citations
1	Identification of mutations in the cardiac ryanodine receptor gene in families affected with arrhythmogenic right ventricular cardiomyopathy type 2 (ARVD2). Human Molecular Genetics, 2001, 10, 189-194.	2.9	709
2	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
3	Diagnosis of arrhythmogenic cardiomyopathy: The Padua criteria. International Journal of Cardiology, 2020, 319, 106-114.	1.7	283
4	Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. European Heart Journal, 2020, 41, 1414-1429.	2.2	239
5	Comparison of Clinical Features of Arrhythmogenic Right Ventricular Cardiomyopathy in Men Versus Women. American Journal of Cardiology, 2008, 102, 1252-1257.	1.6	81
6	â€˜Hot phaseâ€™ clinical presentation in arrhythmogenic cardiomyopathy. Europace, 2021, 23, 907-917.	1.7	67
7	Pathogenic Potential of Hic1-Expressing Cardiac Stromal Progenitors. Cell Stem Cell, 2020, 26, 205-220.e8.	11.1	60
8	Evolving Diagnostic Criteria for Arrhythmogenic Cardiomyopathy. Journal of the American Heart Association, 2021, 10, e021987.	3.7	60
9	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
10	Phenotypic expression is a prerequisite for malignant arrhythmic events and sudden cardiac death in arrhythmogenic right ventricular cardiomyopathy. Europace, 2016, 18, 1086-1094.	1.7	50
11	Modeling Cardiovascular Diseases with hiPSC-Derived Cardiomyocytes in 2D and 3D Cultures. International Journal of Molecular Sciences, 2020, 21, 3404.	4.1	46
12	Wnt/Î²-catenin pathway in arrhythmogenic cardiomyopathy. Oncotarget, 2017, 8, 60640-60655.	1.8	46
13	Clinical and Functional Characterization of a Novel Mutation in Lamin A/C Gene in a Multigenerational Family with Arrhythmogenic Cardiac Laminopathy. PLoS ONE, 2015, 10, e0121723.	2.5	43
14	The novel S59P mutation in the TNFRSF1A gene identified in an adult onset TNF receptor associated periodic syndrome (TRAPS) constitutively activates NF-ÎºB pathway. Arthritis Research and Therapy, 2015, 17, 93.	3.5	43
15	Intercalated Discs and Arrhythmogenic Cardiomyopathy. Circulation: Cardiovascular Genetics, 2014, 7, 930-940.	5.1	41
16	A founderMYBPC3mutation results in HCM with a high risk of sudden death after the fourth decade of life. Journal of Medical Genetics, 2015, 52, 338-347.	3.2	41
17	Arrhythmogenic cardiomyopathy: a disease of intercalated discs. Cell and Tissue Research, 2015, 360, 491-500.	2.9	41
18	A novel murine model for arrhythmogenic cardiomyopathy points to a pathogenic role of Wnt signalling and miRNA dysregulation. Cardiovascular Research, 2019, 115, 739-751.	3.8	40

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19	Homozygous Desmocollin-2 Mutations and Arrhythmogenic Cardiomyopathy. American Journal of Cardiology, 2015, 116, 1245-1251.	1.6	38
20	Large Genomic Rearrangements of Desmosomal Genes in Italian Arrhythmogenic Cardiomyopathy Patients. Circulation: Arrhythmia and Electrophysiology, 2017, 10, .	4.8	35
21	Genetics meets epigenetics: Genetic variants that modulate noncoding RNA in cardiovascular diseases. Journal of Molecular and Cellular Cardiology, 2015, 89, 27-34.	1.9	28
22	Clinical profile and long-term follow-up of a cohort of patients with desmoplakin cardiomyopathy. Heart Rhythm, 2022, 19, 1315-1324.	0.7	22
23	Transcriptomic Characterization of a Human In Vitro Model of Arrhythmogenic Cardiomyopathy Under Topological and Mechanical Stimuli. Annals of Biomedical Engineering, 2019, 47, 852-865.	2.5	16
24	Hypertrophic Cardiomyopathy and Primary Restrictive Cardiomyopathy: Similarities, Differences and Phenocopies. Journal of Clinical Medicine, 2021, 10, 1954.	2.4	16
25	A targeted next-generation gene panel reveals a novel heterozygous nonsense variant in the TP63 gene in patients with arrhythmogenic cardiomyopathy. Heart Rhythm, 2019, 16, 773-780.	0.7	15
26	Genetic bases of arrhythmogenic right ventricular cardiomyopathy. Heart International, 2006, 2, 17.	1.4	13
27	Arrhythmogenic Cardiomyopathy. European Heart Journal, 2020, 41, 4457-4462.	2.2	12
28	Co-inheritance of mutations associated with arrhythmogenic cardiomyopathy and hypertrophic cardiomyopathy. European Journal of Human Genetics, 2017, 25, 1165-1169.	2.8	10
29	Novel Missense Variant in MYL2 Gene Associated With Hypertrophic Cardiomyopathy Showing High Incidence of Restrictive Physiology. Circulation Genomic and Precision Medicine, 2020, 13, e002824.	3.6	6
30	Circulating miR-185-5p as a Potential Biomarker for Arrhythmogenic Right Ventricular Cardiomyopathy. Cells, 2021, 10, 2578.	4.1	5
31	Recent Advances in CRISPR/Cas9-Based Genome Editing Tools for Cardiac Diseases. International Journal of Molecular Sciences, 2021, 22, 10985.	4.1	5
32	Pathogenic variants in plakophilin-2 gene (PKP2) are associated with better survival in arrhythmogenic right ventricular cardiomyopathy. Journal of Applied Genetics, 2021, 62, 613-620.	1.9	3