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	Clinical profile and long-term follow-up of a cohort of patients with desmoplakin cardiomyopathy. Heart Rhythm, 2022, 19, 1315-1324.	0.7	22
2	Hypertrophic Cardiomyopathy and Primary Restrictive Cardiomyopathy: Similarities, Differences and Phenocopies. Journal of Clinical Medicine, 2021, 10, 1954.	2.4	16
3	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
4	Evolving Diagnostic Criteria for Arrhythmogenic Cardiomyopathy. Journal of the American Heart Association, 2021, 10, e021987.	3.7	60
5	Circulating miR-185-5p as a Potential Biomarker for Arrhythmogenic Right Ventricular Cardiomyopathy. Cells, 2021, 10, 2578.	4.1	5
6	â€˜Hot phaseâ€™ clinical presentation in arrhythmogenic cardiomyopathy. Europace, 2021, 23, 907-917.	1.7	67
7	Recent Advances in CRISPR/Cas9-Based Genome Editing Tools for Cardiac Diseases. International Journal of Molecular Sciences, 2021, 22, 10985.	4.1	5
8	Arrhythmogenic Cardiomyopathy. European Heart Journal, 2020, 41, 4457-4462.	2.2	12
9	Modeling Cardiovascular Diseases with hiPSC-Derived Cardiomyocytes in 2D and 3D Cultures. International Journal of Molecular Sciences, 2020, 21, 3404.	4.1	46
10	Diagnosis of arrhythmogenic cardiomyopathy: The Padua criteria. International Journal of Cardiology, 2020, 319, 106-114.	1.7	283
11	Pathogenic Potential of Hic1-Expressing Cardiac Stromal Progenitors. Cell Stem Cell, 2020, 26, 205-220.e8.	11.1	60
12	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
13	Arrhythmogenic right ventricular cardiomyopathy: evaluation of the current diagnostic criteria and differential diagnosis. European Heart Journal, 2020, 41, 1414-1429.	2.2	239
14	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
15	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
16	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
17	Large Genomic Rearrangements of Desmosomal Genes in Italian Arrhythmogenic Cardiomyopathy Patients. Circulation: Arrhythmia and Electrophysiology, 2017, 10, .	4.8	35
18	Co-inheritance of mutations associated with arrhythmogenic cardiomyopathy and hypertrophic cardiomyopathy. European Journal of Human Genetics, 2017, 25, 1165-1169.	2.8	10

#	Article	IF	Citations
19	Wnt/Î²-catenin pathway in arrhythmogenic cardiomyopathy. Oncotarget, 2017, 8, 60640-60655.	1.8	46
20	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
21	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
22	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
23	The novel S59P mutation in the TNFRSF1A gene identified in an adult onset TNF receptor associated periodic syndrome (TRAPS) constitutively activates NF-Î¹ºB pathway. Arthritis Research and Therapy, 2015, 17, 93.	3.5	43
24	Arrhythmogenic cardiomyopathy: a disease of intercalated discs. Cell and Tissue Research, 2015, 360, 491-500.	2.9	41
25	Homozygous Desmocollin-2 Mutations and Arrhythmogenic Cardiomyopathy. American Journal of Cardiology, 2015, 116, 1245-1251.	1.6	38
26	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
27	Intercalated Discs and Arrhythmogenic Cardiomyopathy. Circulation: Cardiovascular Genetics, 2014, 7, 930-940.	5.1	41
28	Comparison of Clinical Features of Arrhythmogenic Right Ventricular Cardiomyopathy in Men Versus Women. American Journal of Cardiology, 2008, 102, 1252-1257.	1.6	81
29	Genetic bases of arrhythmogenic right ventricular cardiomyopathy. Heart International, 2006, 2, 17.	1.4	13
30	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
31	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
32	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