Rachele Adorisio

List of Publications by Year
in descending order

38 papers	691 citations	516561 16 h-index	580701 25 g-index
38 all docs	38 docs citations	38 times ranked	1194 citing authors

#	Article	IF	Citations
1	A current approach to heart failure in Duchenne muscular dystrophy. Heart, 2017, 103, 1770-1779.	1.2	75
2	Abnormal Vasomotor Function of the Epicardial Coronary Arteries in Children Five to Eight Years After Arterial Switch Operation. Journal of the American College of Cardiology, 2005, 46, 1565-1572.	1.2	60
3	Clinical Presentation and Natural History of Hypertrophic Cardiomyopathy in RASopathies. Heart Failure Clinics, 2018, 14, 225-235.	1.0	44
4	Left ventricular assist device in Duchenne Cardiomyopathy: Can we change the natural history of cardiac disease?. International Journal of Cardiology, 2012, 161, e43.	0.8	43
5	Left ventricular assist device as destination therapy in cardiac end-stage dystrophinopathies: Midterm results. Journal of Thoracic and Cardiovascular Surgery, 2017, 153, 669-674.	0.4	41
6	Plastic Bronchitis After Extracardiac Fontan Operation. Annals of Thoracic Surgery, 2012, 94, 860-864.	0.7	39
7	Diagnosis and Management of Rare Cardiomyopathies in Adult and Paediatric Patients. A Position Paper of the Italian Society of Cardiology (SIC) and Italian Society of Paediatric Cardiology (SICP). International Journal of Cardiology, 2022, 357, 55-71.	0.8	36
8	Persistent pulmonary arterial hypertension in the newborn (PPHN): A frequent manifestation of TMEM70 defective patients. Molecular Genetics and Metabolism, 2014, 111, 353-359.	0.5	31
9	Pharmacological treatment of chronic heart failure. Heart Failure Reviews, 2006, 11, 109-123.	1.7	29
10	Clinical Profile of Cardiac Involvement in Danon Disease. Circulation Genomic and Precision Medicine, 2020, 13, e003117.	1.6	29
11	Dystrophin Cardiomyopathies: Clinical Management, Molecular Pathogenesis and Evolution towards Precision Medicine. Journal of Clinical Medicine, 2018, 7, 291.	1.0	24
12	Long-term survival and phenotypic spectrum in heterotaxy syndrome: A 25-year follow-up experience. International Journal of Cardiology, 2018, 268, 100-105.	0.8	24
13	Cardiac Manifestations in Children with SARS-COV-2 Infection: 1-Year Pediatric Multicenter Experience. Children, 2021, 8, 717.	0.6	23
14	Duchenne Dilated Cardiomyopathy: Cardiac Management from Prevention to Advanced Cardiovascular Therapies. Journal of Clinical Medicine, 2020, 9, 3186.	1.0	22
15	The Use of Berlin Heart EXCOR VAD in Children Less than 10 kg: A Single Center Experience. Frontiers in Physiology, 2016, 7, 614.	1.3	19
16	Differences in morbidity and mortality in Down syndrome are related to the type of congenital heart defect. American Journal of Medical Genetics, Part A, 2020, 182, 1342-1350.	0.7	19
17	Heart rate reduction strategy using ivabradine in end-stage Duchenne cardiomyopathy. International Journal of Cardiology, 2019, 280, 99-103.	0.8	17
18	Heterotaxy syndrome with and without spleen: Different infection risk and management. Journal of Allergy and Clinical Immunology, 2017, 139, 1981-1984.e1.	1.5	14

#	Article	IF	Citations
19	Consensus Document of the Italian Association of Hospital Cardiologists (ANMCO), Italian Society of Pediatric Cardiology (SICP), and Italian Society of Gynaecologists and Obstetrics (SIGO): pregnancy and congenital heart diseases. European Heart Journal Supplements, 2017, 19, D256-D292.	0.0	13
20	SOS1 mutations in Noonan syndrome: Cardiomyopathies and not only congenital heart defects! Report of six patients including two novel variants and literature review. American Journal of Medical Genetics, Part A, 2019, 179, 2083-2090.	0.7	10
21	Harmonic Imaging with Levovist for Transthoracic Echocardiographic Reconstruction of Left Ventricle in Patients with Post-Ischemic Left Ventricular Dysfunction and Suboptimal Acoustic Windows. Journal of the American Society of Echocardiography, 2000, 13, 139-145.	1.2	9
22	Cardiovascular Involvement in Pediatric Laminopathies. Report of Six Patients and Literature Revision. Frontiers in Pediatrics, 2020, 8, 374.	0.9	9
23	Coronary plaque composition assessed by intravascular ultrasound virtual histology: Association with longâ€term clinical outcomes after heart transplantation in young adult recipients. Catheterization and Cardiovascular Interventions, 2014, 83, 70-77.	0.7	8
24	First human implantation of a miniaturized axial flow ventricular assist device in a child with end-stage heart failure. Journal of Heart and Lung Transplantation, 2020, 39, 83-87.	0.3	8
25	The use of a numerical model to simulate the cavo-pulmonary assistance in Fontan circulation: a preliminary verification. Journal of Artificial Organs, 2016, 19, 105-113.	0.4	6
26	Myocardial and Arrhythmic Spectrum of Neuromuscular Disorders in Children. Biomolecules, 2021, 11, 1578.	1.8	5
27	Cardiomyopathies in Children and Systemic Disorders When Is It Useful to Look beyond the Heart?. Journal of Cardiovascular Development and Disease, 2022, 9, 47.	0.8	5
28	A New 2D Echocardiographic Approach to Evaluate the Membrane and Valve Movement of the Berlin Heart EXCOR VAD Chamber in Pediatric VAD Patients. Artificial Organs, 2018, 42, 451-456.	1.0	4
29	First evidence of maternally inherited mosaicism in TGFBR1 and subtle primary myocardial changes in Loeys-Dietz syndrome: a case report. BMC Medical Genetics, 2018, 19, 170.	2.1	4
30	Delayed appearance of 3â€methylglutaconic aciduria in neonates with early onset metabolic cardiomyopathies: A potential pitfall for the diagnosis. American Journal of Medical Genetics, Part A, 2020, 182, 64-70.	0.7	4
31	Comment on: â€˜Implantation of a left ventricular assist device to provide long term support for endâ€stage Duchenne muscular dystrophyâ€associated cardiomyopathyâ€™ by Stoller et al.. ESC Heart Failure, 2018, 5, 651-652.	1.4	3
32	Insights from Cardiopulmonary Exercise Testing in Pediatric Patients with Hypertrophic Cardiomyopathy. Biomolecules, 2021, 11, 376.	1.8	3
33	Remember friedreich ataxia even in a toddler with apparently isolated dilated (not hypertrophic!) cardiomyopathy: revisited. Minerva Pediatrics, 2021, , .	0.2	3
34	ICD Outcome in Pediatric Cardiomyopathies. Journal of Cardiovascular Development and Disease, 2022, 9, 33.	0.8	3
35	Real-World Use of Carvedilol in Children With Dilated Cardiomyopathy: Long-Term Effect on Survival and Ventricular Function. Frontiers in Pediatrics, 2022, 10, 845406.	0.9	3
36	Infant miniaturized continuous-flow pumps and permanent support in Pediatrics. Annals of Cardiothoracic Surgery, 2021, 10, 277-280.	0.6	1

#	Article	IF	Citations
37	Cardiomyopathy in Duchenne Muscular Distrophy: Clinical Insights and Therapeutic Implications. , 0, , .		1
38	Persistent myocardial atrophy despite LV reverse remodeling in Duchenne cardiomyopathy treated by LVAD. Pediatric Transplantation, 2021, 25, e13890.	0.5	0