

Rachele Adoriso

List of Publications by Year in descending order

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Version: 2024-02-01

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 | ICD Outcome in Pediatric Cardiomyopathies. <i>Journal of Cardiovascular Development and Disease</i> , 2022, 9, 33. | 0.8 | 3 |
| 2 | Cardiomyopathies in Children and Systemic Disorders When Is It Useful to Look beyond the Heart?. <i>Journal of Cardiovascular Development and Disease</i> , 2022, 9, 47. | 0.8 | 5 |
| 3 | 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). <i>International Journal of Cardiology</i> , 2022, 357, 55-71. | 0.8 | 36 |
| 4 | Real-World Use of Carvedilol in Children With Dilated Cardiomyopathy: Long-Term Effect on Survival and Ventricular Function. <i>Frontiers in Pediatrics</i> , 2022, 10, 845406. | 0.9 | 3 |
| 5 | Persistent myocardial atrophy despite LV reverse remodeling in Duchenne cardiomyopathy treated by LVAD. <i>Pediatric Transplantation</i> , 2021, 25, e13890. | 0.5 | 0 |
| 6 | Insights from Cardiopulmonary Exercise Testing in Pediatric Patients with Hypertrophic Cardiomyopathy. <i>Biomolecules</i> , 2021, 11, 376. | 1.8 | 3 |
| 7 | Infant miniaturized continuous-flow pumps and permanent support in Pediatrics. <i>Annals of Cardiothoracic Surgery</i> , 2021, 10, 277-280. | 0.6 | 1 |
| 8 | Cardiac Manifestations in Children with SARS-COV-2 Infection: 1-Year Pediatric Multicenter Experience. <i>Children</i> , 2021, 8, 717. | 0.6 | 23 |
| 9 | Myocardial and Arrhythmic Spectrum of Neuromuscular Disorders in Children. <i>Biomolecules</i> , 2021, 11, 1578. | 1.8 | 5 |
| 10 | Remember friedreich ataxia even in a toddler with apparently isolated dilated (not hypertrophic!) cardiomyopathy: revisited. <i>Minerva Pediatrics</i> , 2021, , . | 0.2 | 3 |
| 11 | First human implantation of a miniaturized axial flow ventricular assist device in a child with end-stage heart failure. <i>Journal of Heart and Lung Transplantation</i> , 2020, 39, 83-87. | 0.3 | 8 |
| 12 | Delayed appearance of 3-oxo-methylglutaconic aciduria in neonates with early onset metabolic cardiomyopathies: A potential pitfall for the diagnosis. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 64-70. | 0.7 | 4 |
| 13 | Duchenne Dilated Cardiomyopathy: Cardiac Management from Prevention to Advanced Cardiovascular Therapies. <i>Journal of Clinical Medicine</i> , 2020, 9, 3186. | 1.0 | 22 |
| 14 | Cardiovascular Involvement in Pediatric Laminopathies. Report of Six Patients and Literature Revision. <i>Frontiers in Pediatrics</i> , 2020, 8, 374. | 0.9 | 9 |
| 15 | Clinical Profile of Cardiac Involvement in Danon Disease. <i>Circulation Genomic and Precision Medicine</i> , 2020, 13, e003117. | 1.6 | 29 |
| 16 | Differences in morbidity and mortality in Down syndrome are related to the type of congenital heart defect. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 1342-1350. | 0.7 | 19 |
| 17 | <i>SOS1</i> mutations in Noonan syndrome: Cardiomyopathies and not only congenital heart defects! Report of six patients including two novel variants and literature review. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 2083-2090. | 0.7 | 10 |
| 18 | Heart rate reduction strategy using ivabradine in end-stage Duchenne cardiomyopathy. <i>International Journal of Cardiology</i> , 2019, 280, 99-103. | 0.8 | 17 |

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 19 | A New 2D Echocardiographic Approach to Evaluate the Membrane and Valve Movement of the Berlin Heart EXCOR VAD Chamber in Pediatric VAD Patients. <i>Artificial Organs</i> , 2018, 42, 451-456. | 1.0 | 4 |
| 20 | Clinical Presentation and Natural History of Hypertrophic Cardiomyopathy in RASopathies. <i>Heart Failure Clinics</i> , 2018, 14, 225-235. | 1.0 | 44 |
| 21 | Dystrophin Cardiomyopathies: Clinical Management, Molecular Pathogenesis and Evolution towards Precision Medicine. <i>Journal of Clinical Medicine</i> , 2018, 7, 291. | 1.0 | 24 |
| 22 | First evidence of maternally inherited mosaicism in TGFBR1 and subtle primary myocardial changes in Loeys-Dietz syndrome: a case report. <i>BMC Medical Genetics</i> , 2018, 19, 170. | 2.1 | 4 |
| 23 | Long-term survival and phenotypic spectrum in heterotaxy syndrome: A 25-year follow-up experience. <i>International Journal of Cardiology</i> , 2018, 268, 100-105. | 0.8 | 24 |
| 24 | Comment on: "Implantation of a left ventricular assist device to provide long term support for end-stage Duchenne muscular dystrophy-associated cardiomyopathy" by Stoller et al. <i>ESC Heart Failure</i> , 2018, 5, 651-652. | 1.4 | 3 |
| 25 | Heterotaxy syndrome with and without spleen: Different infection risk and management. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 1981-1984.e1. | 1.5 | 14 |
| 26 | 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. <i>European Heart Journal Supplements</i> , 2017, 19, D256-D292. | 0.0 | 13 |
| 27 | A current approach to heart failure in Duchenne muscular dystrophy. <i>Heart</i> , 2017, 103, 1770-1779. | 1.2 | 75 |
| 28 | Left ventricular assist device as destination therapy in cardiac end-stage dystrophinopathies: Midterm results. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2017, 153, 669-674. | 0.4 | 41 |
| 29 | The Use of Berlin Heart EXCOR VAD in Children Less than 10 kg: A Single Center Experience. <i>Frontiers in Physiology</i> , 2016, 7, 614. | 1.3 | 19 |
| 30 | The use of a numerical model to simulate the cavo-pulmonary assistance in Fontan circulation: a preliminary verification. <i>Journal of Artificial Organs</i> , 2016, 19, 105-113. | 0.4 | 6 |
| 31 | Coronary plaque composition assessed by intravascular ultrasound virtual histology: Association with long-term clinical outcomes after heart transplantation in young adult recipients. <i>Catheterization and Cardiovascular Interventions</i> , 2014, 83, 70-77. | 0.7 | 8 |
| 32 | Persistent pulmonary arterial hypertension in the newborn (PPHN): A frequent manifestation of TMEM70 defective patients. <i>Molecular Genetics and Metabolism</i> , 2014, 111, 353-359. | 0.5 | 31 |
| 33 | Plastic Bronchitis After Extracardiac Fontan Operation. <i>Annals of Thoracic Surgery</i> , 2012, 94, 860-864. | 0.7 | 39 |
| 34 | Left ventricular assist device in Duchenne Cardiomyopathy: Can we change the natural history of cardiac disease?. <i>International Journal of Cardiology</i> , 2012, 161, e43. | 0.8 | 43 |
| 35 | Pharmacological treatment of chronic heart failure. <i>Heart Failure Reviews</i> , 2006, 11, 109-123. | 1.7 | 29 |
| 36 | Abnormal Vasomotor Function of the Epicardial Coronary Arteries in Children Five to Eight Years After Arterial Switch Operation. <i>Journal of the American College of Cardiology</i> , 2005, 46, 1565-1572. | 1.2 | 60 |

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|----|--|-----|-----------|
| 37 | Harmonic Imaging with Levovist for Transthoracic Echocardiographic Reconstruction of Left Ventricle in Patients with Post-Ischemic Left Ventricular Dysfunction and Suboptimal Acoustic Windows. <i>Journal of the American Society of Echocardiography</i> , 2000, 13, 139-145. | 1.2 | 9 |
| 38 | Cardiomyopathy in Duchenne Muscular Distrophy: Clinical Insights and Therapeutic Implications. , 0, , . | | 1 |