Joseph Atallah

List of Publications by Year
in descending order

32 papers	1,026 citations	759233 12 h-index	477307 29 g-index
34 all docs	34 docs citations	34 times ranked	1473 citing authors

#	Article	IF	Citations
1	Neurodevelopmental Outcomes After Cardiac Surgery in Infancy. Pediatrics, 2015, 135, 816-825.	2.1	392
2	Multi-Institutional Study of Implantable Defibrillator Lead Performance in Children and Young Adults. Circulation, 2013, 127, 2393-2402.	1.6	158
3	Two-Year Survival and Mental and Psychomotor Outcomes After the Norwood Procedure. Circulation, 2008, 118, 1410-1418.	1.6	92
4	Implantable cardioverter-defibrillator use in catecholaminergic polymorphic ventricular tachycardia: A systematic review. Heart Rhythm, 2018, 15, 1791-1799.	0.7	77
5	Two-year general and neurodevelopmental outcome after neonatal complex cardiac surgery in patients with deletion 22q11.2: A comparative study. Journal of Thoracic and Cardiovascular Surgery, 2007, 134, 772-779.	0.8	52
6	Strain Rate in Children and Young Piglets Mirrors Changes in Contractility and Demonstrates a Force-Frequency Relationship. Journal of the American Society of Echocardiography, 2017, 30, 797-806.	2.8	31
7	Ventricular Arrhythmia and Life-Threatening Events in Patients With Repaired Tetralogy of Fallot. American Journal of Cardiology, 2020, 132, 126-132.	1.6	29
8	An International Multicenter Cohort Study on Î²-Blockers for the Treatment of Symptomatic Children With Catecholaminergic Polymorphic Ventricular Tachycardia. Circulation, 2022, 145, 333-344.	1.6	28
9	Initially unexplained cardiac arrest in children and adolescents: A national experience from the Canadian Pediatric Heart Rhythm Network. Heart Rhythm, 2020, 17, 975-981.	0.7	21
10	The Hearts in Rhythm Organization: A Canadian National Cardiogenetics Network. CJC Open, 2020, 2, 652-662.	1.5	14
11	Uptake of Predictive Genetic Testing and Cardiac Evaluation for Children at Risk for an Inherited Arrhythmia or Cardiomyopathy. Journal of Genetic Counseling, 2018, 27, 124-130.	1.6	13
12	The Bayley-III scale may underestimate neurodevelopmental disability after cardiac surgery in infants. European Journal of Cardio-thoracic Surgery, 2020, 57, 63-71.	1.4	13
13	Biophysical Characterization of a Novel SCN5A Mutation Associated With an Atypical Phenotype of Atrial and Ventricular Arrhythmias and Sudden Death. Frontiers in Physiology, 2020, 11, 610436.	2.8	12
14	Fetal Umbilical Arterial Pulsatility Correlates With 2-Year Growth and Neurodevelopmental Outcomes in Congenital Heart Disease. Canadian Journal of Cardiology, 2021, 37, 425-432.	1.7	10
15	Clinical and Functional Developmental Outcomes in Neonates Undergoing Truncus Arteriosus Repair: A Cohort Study. Annals of Thoracic Surgery, 2016, 101, 1827-1833.	1.3	9
16	Insights on Atrial Fibrillation in Congenital Heart Disease. Canadian Journal of Cardiology, 2018, 34, 1531-1533.	1.7	9
17	Physical activity restriction for children and adolescents diagnosed with an inherited arrhythmia or cardiomyopathy and its impact on body mass index. Journal of Cardiovascular Electrophysiology, 2018, 29, 1648-1653.	1.7	8
18	Antenatal Diagnosis and Successful Surgical Removal of a Large Right Ventricular Fibroma. Pediatric Cardiology, 2006, 27, 493-496.	1.3	7

#	Article	IF	Citations
19	Outcomes of Pediatric Patients With Defibrillators Following Initial Presentation With Sudden Cardiac Arrest. Circulation: Arrhythmia and Electrophysiology, 2021, 14, e008517.	4.8	7
20	The Survival of a Newborn with an Interrupted Aortic Arch and a Closed Ductus Arteriosus. Congenital Heart Disease, 2008, 3, 144-145.	0.2	6
21	Exercise and Î²-blocker therapy recommendations for inherited arrhythmogenic conditions. Cardiology in the Young, 2016, 26, 1123-1129.	0.8	6
22	Riata lead failure in pediatric and congenital heart disease patients. Journal of Cardiovascular Electrophysiology, 2019, 30, 320-325.	1.7	6
23	When to Offer Predictive Genetic Testing to Children at Risk of an Inherited Arrhythmia or Cardiomyopathy. Circulation Genomic and Precision Medicine, 2018, 11, e002300.	3.6	5
24	The impact of physical activity modification on the well-being of a cohort of children with an inherited arrhythmia or cardiomyopathy. Cardiology in the Young, 2020, 30, 692-697.	0.8	5
25	Predicting Post-Fontan Length of Stay: The Limits of Measured Variables. Pediatric Cardiology, 2019, 40, 1208-1216.	1.3	4
26	Correlation of electrocardiogram parameters and hemodynamic outcomes in patients with isolated secundum atrial septal defects. Annals of Pediatric Cardiology, 2017, 10, 152.	0.5	4
27	Return of Results Policies for Genomic Research: Current Practices and the Hearts in Rhythm Organization (HiRO) Approach. Canadian Journal of Cardiology, 2022, 38, 526-535.	1.7	3
28	Evolution of the Fetal Atrioventricular Interval from 6 to 40 Weeks of Gestation. American Journal of Cardiology, 2019, 123, 1709-1714.	1.6	2
29	Leuprolide Acetate and QTc Interval in Gender-Diverse Youth. Transgender Health, 2023, 8, 84-88.	2.5	2
30	A Novel Surgical Technique for Repair of Congenitally Corrected Transposition of the Great Arteries With Atrioventricular Septal Defect: Avoiding Damage to the Conduction System. Annals of Thoracic Surgery, 2015, 100, 1121-1123.	1.3	1
31	Right arch and bilateral arterial ducts in a patient with discordant atrioventricular connections with pulmonary atresia and non-confluent pulmonary arteries. Cardiology in the Young, 2007, 17, 111.	0.8	0
32	Use of topical lidocaine in eliminating mechanically stimulated ventricular fibrillation in a patient with short QT syndrome. HeartRhythm Case Reports, 2019, 5, 152-154.	0.4	0