

Mariam Arabi

List of Publications by Year in descending order

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Version: 2024-02-01

49
papers

309
citations

1163117

8
h-index

996975

15
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51
all docs

51
docs citations

51
times ranked

452
citing authors

#	ARTICLE	IF	CITATIONS
1	QTc interval on 24-hour holter monitor: To trust or not to trust?. <i>Annals of Noninvasive Electrocardiology</i> , 2022, 27, e12899.	1.1	6
2	Endovascular Stent Repair of Aortic Coarctation in a Developing Country: A Single-Center Experience. <i>Cardiovascular Revascularization Medicine</i> , 2022, 39, 66-72.	0.8	3
3	Aspirin in COVID-19: Pros and Cons. <i>Frontiers in Pharmacology</i> , 2022, 13, 849628.	3.5	10
4	The use of steroids in treating chylothorax following cardiac surgery in children: a unique perspective. <i>Cardiology in the Young</i> , 2022, , 1-6.	0.8	2
5	Congenital Heart Disease in Syrian Refugee Children: The Experience at a Tertiary Care Center in a Developing Country. <i>Pediatric Cardiology</i> , 2021, 42, 1010-1017.	1.3	2
6	COVID-19 in the MENA Region: Facts and Findings. <i>Journal of Infection in Developing Countries</i> , 2021, 15, 342-349.	1.2	8
7	Pre-operative assessment of pediatric congenital heart disease patients in the COVID-19 era: lessons learned. <i>Cardiology in the Young</i> , 2021, , 1-5.	0.8	0
8	Management of post-operative Junctional Ectopic Tachycardia in symptomatic neonates and infants at a tertiary care center in a developing country: Lessons learned!. <i>Southwest Respiratory and Critical Care Chronicles</i> , 2021, 9, 14-19.	0.2	0
9	Cardiac Manifestations in COVID-19 Patients: A Focus on the Pediatric Population. <i>Canadian Journal of Infectious Diseases and Medical Microbiology</i> , 2021, 2021, 1-12.	1.9	9
10	Acute Kidney Injury Post-cardiac Surgery in Infants and Children: A Single-Center Experience in a Developing Country. <i>Frontiers in Pediatrics</i> , 2021, 9, 637463.	1.9	3
11	COVID-19: potential therapeutics for pediatric patients. <i>Pharmacological Reports</i> , 2021, 73, 1520-1538.	3.3	12
12	Ivabradine: A Potential Therapeutic for Children With Refractory SVT. <i>Frontiers in Cardiovascular Medicine</i> , 2021, 8, 660855.	2.4	6
13	Placement of Labcor Pulmonary Conduit Results in a High Incidence of Postoperative Fever. <i>World Journal for Pediatric & Congenital Heart Surgery</i> , 2021, 12, 55-60.	0.8	0
14	Balloon Valvuloplasty for Congenital Aortic Stenosis: Experience at a Tertiary Center in a Developing Country. <i>Journal of Interventional Cardiology</i> , 2021, 2021, 1-7.	1.2	2
15	Sotalol as an effective adjunct therapy in the management of supraventricular tachycardia induced fetal hydrops fetalis. <i>Journal of Neonatal-Perinatal Medicine</i> , 2020, 13, 267-273.	0.8	3
16	Establishing a High-Quality Congenital Cardiac Surgery Program in a Developing Country: Lessons Learned. <i>Frontiers in Pediatrics</i> , 2020, 8, 357.	1.9	13
17	Cardiac Tamponade Caused by <i>Cutibacterium acnes</i> : An Updated and Comprehensive Review of the Literature. <i>Canadian Journal of Infectious Diseases and Medical Microbiology</i> , 2020, 2020, 1-8.	1.9	14
18	COVID-19 in Pediatric Patients: A Focus on CHD Patients. <i>Frontiers in Cardiovascular Medicine</i> , 2020, 7, 612460.	2.4	20

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19	Large congenital left atrial wall aneurysm: An updated and comprehensive review of the literature. <i>Echocardiography</i> , 2020, 37, 965-970.	0.9	4
20	Port-a-Cath fracture and migration in paediatric cancer patients: incidence and management at a tertiary care centre – a 15-year experience. <i>Cardiology in the Young</i> , 2020, 30, 986-990.	0.8	2
21	Hydroxychloroquine in COVID-19 Patients: Pros and Cons. <i>Frontiers in Pharmacology</i> , 2020, 11, 597985.	3.5	25
22	Novel EIF2AK4 mutations in histologically proven pulmonary capillary hemangiomatosis and hereditary pulmonary arterial hypertension. <i>BMC Medical Genetics</i> , 2019, 20, 176.	2.1	8
23	Aortopulmonary window in adults: A rare entity leading to Eisenmenger syndrome. <i>Echocardiography</i> , 2019, 36, 1173-1178.	0.9	10
24	The first Fetal Echocardiography experience for Prenatal diagnosis of Congenital Heart Disease in Lebanon: Successes and challenges. <i>Journal of the Saudi Heart Association</i> , 2019, 31, 125-129.	0.4	5
25	Fetal Intra-pericardial Morgagni Hernia with effusion affecting one member of a twin gestation. <i>Echocardiography</i> , 2019, 36, 1014-1016.	0.9	2
26	A Novel Somatic Variant in HEY2 Unveils an Alternative Splicing Isoform Linked to Ventricular Septal Defect. <i>Pediatric Cardiology</i> , 2019, 40, 1084-1091.	1.3	4
27	Non-familial cardiomyopathies in Lebanon: exome sequencing results for five idiopathic cases. <i>BMC Medical Genomics</i> , 2019, 12, 33.	1.5	4
28	Steroids as a possible effective therapy in the management of large isolated chylopericardium following open heart surgery. <i>Cardiology in the Young</i> , 2019, 29, 1426-1431.	0.8	2
29	Transcatheter Closure of Atrial Septal Defects: Comparable Experience and Outcomes Between Developing and Developed Countries. <i>Pediatric Cardiology</i> , 2019, 40, 610-615.	1.3	9
30	Lower extremity thrombosis and myocarditis due to Human PVB19 infection. <i>Journal of Pediatric Surgery Case Reports</i> , 2018, 32, 30-31.	0.2	1
31	ECMO is in the air: Long distance air/ground transport of a child on extra corporeal membrane oxygenation. <i>Egyptian Journal of Critical Care Medicine</i> , 2018, 6, 151-153.	0.4	1
32	Degenerated hair follicle cells and partial loss of sebaceous and eccrine glands in a familial case of axenfeld-rieger syndrome: An emerging role for the FOXC1/NFATC1 genetic axis. <i>Journal of Dermatological Science</i> , 2018, 92, 237-244.	1.9	5
33	TARGETED AND EXOME SEQUENCING OF 27 LEBANESE PATIENTS WITH CARDIOMYOPATHIES: NOVEL VARIANTS IN KNOWN GENES, AND POTENTIAL NOVEL GENES. <i>Journal of the American College of Cardiology</i> , 2017, 69, 719.	2.8	0
34	A Novel Mutation in FOXC1 in a Lebanese Family with Congenital Heart Disease and Anterior Segment Dysgenesis: Potential Roles for NFATC1 and DPT in the Phenotypic Variations. <i>Frontiers in Cardiovascular Medicine</i> , 2017, 4, 58.	2.4	15
35	A Novel Role for CSR1P1 in a Lebanese Family with Congenital Cardiac Defects. <i>Frontiers in Genetics</i> , 2017, 8, 217.	2.3	8
36	Incessant Long R-P Tachycardia. <i>Cardiac Electrophysiology Clinics</i> , 2016, 8, 71-74.	1.7	0

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37	The Muscle-Bound Heart. Cardiac Electrophysiology Clinics, 2016, 8, 223-231.	1.7	6
38	A novel nonsense mutation in NPHS1: is aortic stenosis associated with congenital nephropathy?. Journal of Genetics, 2015, 94, 309-312.	0.7	0
39	Surgical repair of partial atrioventricular defect. Multimedia Manual of Cardiothoracic Surgery: MMCTS / European Association for Cardio-Thoracic Surgery, 2015, 2015, mmv037.	0.1	0
40	Surgical repair of complete atrioventricular defect (Nunn technique). Multimedia Manual of Cardiothoracic Surgery: MMCTS / European Association for Cardio-Thoracic Surgery, 2015, 2015, mmv023.	0.1	3
41	Percutaneous closure of patent ductus arteriosus in children using amplatzer duct occluder II: Relationship between <scp>PDA</scp> type and risk of device protrusion into the descending aorta. Catheterization and Cardiovascular Interventions, 2015, 86, E66-72.	1.7	14
42	NKX2-5 Mutations in an Inbred Consanguineous Population: Genetic and Phenotypic Diversity. Scientific Reports, 2015, 5, 8848.	3.3	41
43	Noninvasive Nitric Oxide Therapy in Right Ventricular Systolic Dysfunction Following Arterial Switch Procedure. World Journal for Pediatric & Congenital Heart Surgery, 2014, 5, 460-462.	0.8	0
44	Lack of Cardiac Iron in SCD Patients Despite Severe Iron Overload. Blood, 2014, 124, 4943-4943.	1.4	0
45	Research in Congenital Heart Disease: A Comparative Bibliometric Analysis Between Developing and Developed Countries. Pediatric Cardiology, 2013, 34, 375-382.	1.3	19
46	Elevated Tricuspid Regurgitant Jet Velocity In Lebanese Patients With Sickle Cell Disease Is Associated With Severe Disease and Is Clustered In Families. Blood, 2013, 122, 4684-4684.	1.4	0
47	Cardiac Involvement in Nonketotic Hyperglycinemia. Journal of Child Neurology, 2011, 26, 970-973.	1.4	6
48	The status of pediatric cardiology at a tertiary center in Lebanon. Journal Medical Libanais, 2011, 59, 136-42.	0.0	0
49	Absence of GJA1 gene mutations in four patients with anomalous left coronary artery from the pulmonary artery (ALCAPA). Journal Medical Libanais, 2011, 59, 149-53.	0.0	2