Kamel Shibbani

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3661916/publications.pdf

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17	216	1478505 6	1199594 12
papers	citations	h-index	g-index

17	17	17	466
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Feasibility, Safety, and Shortâ€Term Outcomes of Transcatheter Patent Ductus Arteriosus Closure in Premature Infants on Highâ€Frequency Jet Ventilation. Journal of the American Heart Association, 2022, 11, e025343.	3.7	5
2	Anomalous Left Anterior Descending Coronary Artery With Retroaortic LeftÂCircumflex Artery. JACC: Case Reports, 2021, 3, 546-549.	0.6	1
3	The road less travelled: percutaneous closure of a secundum atrial septal defect through the left internal jugular vein. Cardiology in the Young, 2021, 31, 1696-1697.	0.8	1
4	Aspirin Use and Transcatheter Pulmonary Valve Replacement, the Need for Consistency. Pediatric Cardiology, 2021, 42, 1640-1646.	1.3	6
5	Double-Outlet Right Ventricle, Pulmonary Atresia, and Discontinuous Branch Pulmonary Arteries Supplied byÂBilateralÂDucti. JACC: Case Reports, 2021, 3, 1236-1240.	0.6	0
6	This and that: management of Tetralogy of Fallot and pulmonary vein stenosis in an infantâ€”a case report. European Heart Journal - Case Reports, 2021, 5, ytab429.	0.6	0
7	Dissecting transcriptomic signatures of neuronal differentiation and maturation using iPSCs. Nature Communications, 2020, 11, 462.	12.8	96
8	Preclinical comparative assessment of a dedicated pediatric polyâ€Lâ€lacticâ€acidâ€based bioresorbable scaffold with a lowâ€profile bare metal stent. Catheterization and Cardiovascular Interventions, 2020, 96, 878-888.	1.7	7
9	SGLT1 and Sweet Genetic Insights Into Cardiometabolic Risk. Journal of the American College of Cardiology, 2018, 72, 1774-1777.	2.8	0
10	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. Frontiers in Cardiovascular Medicine, 2017, 4, 58.	2.4	15
11	A Novel Role for CSRP1 in a Lebanese Family with Congenital Cardiac Defects. Frontiers in Genetics, 2017, 8, 217.	2.3	8
12	Premature Valvular Heart Disease in Homozygous Familial Hypercholesterolemia. Cholesterol, 2017, 2017, 1-7.	1.6	17
13	Identifying Gaps in Technology for Congenital Interventions: Analysis of a Needs Survey from Congenital Interventional Cardiologists. Pediatric Cardiology, 2016, 37, 925-931.	1.3	13
14	Molecular Pathways and Animal Models of Tricuspid Atresia and Univentricular Heart. , 2016, , 591-605.		0
15	Perventricular and Percutaneous Closure of Traumatic Ventricular Septal Defects Following Blunt Chest Trauma. Journal of Structural Heart Disease, 2016, 2, 98-101.	0.1	0
16	Proteomics studies in inner ear disorders: pathophysiology and biomarkers. Expert Review of Proteomics, 2015, 12, 185-196.	3.0	14
17	Two Heterozygous Mutations in NFATC1 in a Patient with Tricuspid Atresia. PLoS ONE, 2012, 7, e49532.	2.5	33