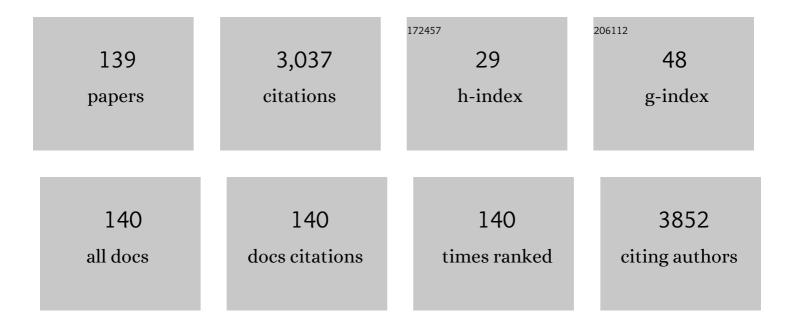
List of Publications by Year in descending order

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MINC-TALLIN

#	Article	IF	CITATIONS
1	Trends in the utilization of computed tomography and cardiac catheterization among children with congenital heart disease. Journal of the Formosan Medical Association, 2015, 114, 1061-1068.	1.7	286
2	Viral infections associated with Kawasaki disease. Journal of the Formosan Medical Association, 2014, 113, 148-154.	1.7	190
3	Losartan Added to β-Blockade Therapy for Aortic Root Dilation in Marfan Syndrome: A Randomized, Open-Label Pilot Study. Mayo Clinic Proceedings, 2013, 88, 271-276.	3.0	153
4	Acute and late coronary outcomes in 1073 patients with Kawasaki disease with and without intravenous Î ³ -immunoglobulin therapy. Archives of Disease in Childhood, 2015, 100, 542-547.	1.9	86
5	The global epidemiology of Kawasaki disease: Review and future perspectives. Global Cardiology Science & Practice, 2018, 2017, e201720.	0.4	85
6	Short- and intermediate-term results of transcatheter closure of atrial septal defect with the Amplatzer Septal Occluder. American Heart Journal, 2004, 148, 511-517.	2.7	75
7	Risk factors and derived formosa score forÂintravenous immunoglobulin unresponsiveness in Taiwanese children with Kawasaki disease. Journal of the Formosan Medical Association, 2016, 115, 350-355.	1.7	71
8	Long-Term Survival and Unnatural Deaths of Patients With Repaired Tetralogy of Fallot in an Asian Cohort. Circulation: Cardiovascular Quality and Outcomes, 2012, 5, 120-125.	2.2	69
9	Clinical Implication of the C Allele of the ITPKC Gene SNP rs28493229 in Kawasaki Disease. Pediatric Infectious Disease Journal, 2011, 30, 148-152.	2.0	66
10	Reversal of Cardiac Dysfunction after Enzyme Replacement in Patients with Infantile-Onset Pompe Disease. Journal of Pediatrics, 2009, 155, 271-275.e2.	1.8	56
11	Transcatheter closure of moderate to large patent ductus arteriosus with the Amplatzer duct occluder. Catheterization and Cardiovascular Interventions, 2007, 69, 572-578.	1.7	55
12	Comparison of 3 Risk Estimation Methods for Predicting Cardiac Outcomes in Pregnant Women With Congenital Heart Disease. Circulation Journal, 2015, 79, 1609-1617.	1.6	53
13	Transcatheter closure of atrial septal defect without balloon sizing. Catheterization and Cardiovascular Interventions, 2008, 71, 214-221.	1.7	52
14	Quality of life in adults with congenital heart disease: biopsychosocial determinants and sex-related differences. Heart, 2011, 97, 38-43.	2.9	52
15	Population-based Study of Kawasaki Disease Shock Syndrome in Taiwan. Pediatric Infectious Disease Journal, 2013, 32, 1384-1386.	2.0	48
16	Acute metabolic decompensation and sudden death in Barth syndrome: report of a family and a literature review. European Journal of Pediatrics, 2008, 167, 941-944.	2.7	47
17	Left Pulmonary Artery Sling Complex: Computed Tomography and Hypothesis of Embryogenesis. Annals of Thoracic Surgery, 2007, 84, 1645-1650.	1.3	46
18	Cardiac Conduction Disturbance Detected in a Pediatric Population. Journal of Pediatrics, 2008, 152, 85-89.	1.8	44

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19	A strategic approach to transcatheter closure of patent ductus: Gianturco coils for small-to-moderate ductus and Amplatzer duct occluder for large ductus. International Journal of Cardiology, 2006, 106, 10-15.	1.7	43
20	Radiofrequency Catheter Ablation of Supraventricular Tachycardia in Infants and Toddlers. Circulation Journal, 2009, 73, 1717-1721.	1.6	42
21	Transcatheter Closure of Moderate-to-Large Patent Ductus Arteriosus in Infants Using Amplatzer Duct Occluder. Circulation Journal, 2010, 74, 361-364.	1.6	39
22	Epidemiological Features of Kawasaki Disease in Taiwan, 1976–2007: Results of Five Nationwide Questionnaire Hospital Surveys. Pediatrics and Neonatology, 2014, 55, 92-96.	0.9	39
23	Aortic Valve Prolapse Associated With Outlet-Type Ventricular Septal Defect. Annals of Thoracic Surgery, 2005, 79, 1366-1371.	1.3	35
24	Postnatal outcome of fetal bradycardia without significant cardiac abnormalities. American Heart Journal, 2004, 147, 540-544.	2.7	33
25	Progression of aortic regurgitation after surgical repair of outlet-type ventricular septal defects. American Heart Journal, 2007, 153, 336-342.	2.7	33
26	Severe Bacterial Infection in Patients with Heterotaxy Syndrome. Journal of Pediatrics, 2014, 164, 99-104.e1.	1.8	33
27	Acquired coronary artery fistula after open heart surgery for congenital heart disease. International Journal of Cardiology, 2005, 103, 187-192.	1.7	32
28	Long-term outcome of twin atrioventricular node and supraventricular tachycardia in patients with right isomerism of the atrial appendage. Heart Rhythm, 2008, 5, 224-229.	0.7	31
29	Circulating biomarkers of collagen type I metabolism mark the right ventricular fibrosis and adverse markers of clinical outcome in adults with repaired tetralogy of Fallot. International Journal of Cardiology, 2013, 167, 2963-2968.	1.7	31
30	Risk factors for infective endocarditis in children with congenital heart diseases - A nationwide population-based case control study. International Journal of Cardiology, 2017, 248, 126-130.	1.7	31
31	Ventricular Septal Defect With Secondary Left Ventricular-to-Right Atrial Shunt Is Associated With a Higher Risk for Infective Endocarditis and a Lower Late Chance of Closure. Pediatrics, 2006, 117, e262-e267.	2.1	29
32	Risk factors and implications of progressive coronary dilatation in children with Kawasaki disease. BMC Pediatrics, 2017, 17, 139.	1.7	29
33	Coronary Diameters in Taiwanese Children Younger than 6 Years Old: Z-Score Regression Equations Derived from Body Surface Area. Acta Cardiologica Sinica, 2014, 30, 266-73.	0.2	29
34	Coronary artery anatomy in children with congenital heart disease by computed tomography. International Journal of Cardiology, 2007, 120, 363-370.	1.7	28
35	Detection of pulmonary arterial morphology in tetralogy of Fallot with pulmonary atresia by computed tomography: 12Âyears of experience. European Journal of Pediatrics, 2012, 171, 579-586.	2.7	28
36	Heart rate variability monitoring in the detection of central nervous system complications in children with enterovirus infection. Journal of Critical Care, 2006, 21, 280-286.	2.2	27

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37	Prognostic Value of Submaximal Exercise Data for Cardiac Morbidity in Fontan Patients. Medicine and Science in Sports and Exercise, 2014, 46, 10-15.	0.4	27
38	Acute kidney injury in patients with Kawasaki disease. Pediatric Research, 2016, 80, 224-227.	2.3	26
39	Left Ventricular Geometry, Global Function, and Dyssynchrony in Infants and Children With Pompe Cardiomyopathy Undergoing Enzyme Replacement Therapy. Journal of Cardiac Failure, 2011, 17, 930-936.	1.7	25
40	Noninvasive diagnosis of aortic coarctation in neonates with patent ductus arteriosus. Journal of Pediatrics, 2006, 148, 217-221.	1.8	24
41	Usefulness of 3D Reconstructed Computed Tomography Imaging for Double Outlet Right Ventricle. Journal of the Formosan Medical Association, 2008, 107, 371-380.	1.7	23
42	Progressive Coronary Dilatation Predicts Worse Outcome in Kawasaki Disease. Journal of Pediatrics, 2016, 171, 78-82.e1.	1.8	23
43	State-of-the-art acute phase management of Kawasaki disease after 2017 scientific statement from the American Heart Association. Pediatrics and Neonatology, 2018, 59, 543-552.	0.9	23
44	Association of the C677T methylenetetrahydrofolate reductase mutation with congenital heart diseases. Acta Obstetricia Et Gynecologica Scandinavica, 2005, 84, 1134-1140.	2.8	22
45	Clinical spectrum and long-term outcome of Ebstein's anomaly based on a 26-year experience in an Asian cohort. European Journal of Pediatrics, 2009, 168, 685-690.	2.7	22
46	In utero onset of long QT syndrome with atrioventricular block and spontaneous or lidocaine-induced ventricular tachycardia: Compound effects of hERG pore region mutation and SCN5A N-terminus variant. Heart Rhythm, 2008, 5, 1567-1574.	0.7	20
47	Coexisting mutations/polymorphisms of the long QT syndrome genes in patients with repaired Tetralogy of Fallot are associated with the risks of life-threatening events. Human Genetics, 2012, 131, 1295-1304.	3.8	19
48	Device deformation and left pulmonary artery obstruction after transcatheter patent ductus arteriosus closure in preterm infants. International Journal of Cardiology, 2020, 312, 50-55.	1.7	19
49	Genetic Syndromes and Outcome After Surgical Repair of Pulmonary Atresia and Ventricular Septal Defect. Annals of Thoracic Surgery, 2012, 94, 1627-1633.	1.3	18
50	Total Anomalous Pulmonary Venous Connection: 15 Years' Experience of a Tertiary Care Center in Taiwan. Pediatrics and Neonatology, 2012, 53, 164-170.	0.9	18
51	Implantable cardioverter defibrillator therapy in repaired tetralogy of Fallot after pulmonary valve replacement: Implications for the mechanism of ventricular arrhythmia. International Journal of Cardiology, 2017, 249, 156-160.	1.7	18
52	Ventricular geometric characteristics and functional benefit of mild right ventricular outflow tract obstruction in patients with significant pulmonary regurgitation after repair of tetralogy of Fallot. American Heart Journal, 2014, 167, 555-561.	2.7	17
53	Changing Spectrum of Infective Endocarditis in Children. Pediatric Infectious Disease Journal, 2014, 33, 467-471.	2.0	16
54	Late Cardiovascular Complications After Surgical or Balloon Angioplasty of Coarctation of Aorta in an Asian Cohort. American Journal of Cardiology, 2009, 104, 1139-1144.	1.6	15

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55	Natural and unnatural history of tetralogy of Fallot repaired during adolescence and adulthood. Heart and Vessels, 2012, 27, 65-70.	1.2	15
56	Low-Molecular-Weight Heparin vs Warfarin for Thromboprophylaxis in Children With Coronary Artery Aneurysms After Kawasaki Disease: A Pragmatic Registry Trial. Canadian Journal of Cardiology, 2020, 36, 1598-1607.	1.7	15
57	Clinical Implications of Major Aortopulmonary Collateral Arteries in Patients With Right Isomerism. Annals of Thoracic Surgery, 2006, 82, 153-157.	1.3	14
58	Abnormal Matrix Remodeling in Adolescents and Young Adults with Kawasaki Disease Late after Onset. Clinical Chemistry, 2008, 54, 1815-1822.	3.2	14
59	Implication of QRS Prolongation and Its Relation to Mechanical Dyssynchrony in Idiopathic Dilated Cardiomyopathy in Childhood. American Journal of Cardiology, 2009, 103, 103-109.	1.6	14
60	Computed tomography predict regression of coronary artery aneurysm in patients with Kawasaki disease. Journal of the Formosan Medical Association, 2017, 116, 806-814.	1.7	14
61	Surgical Outcome of Aortopulmonary Window Repair in Early Infancy. Journal of the Formosan Medical Association, 2006, 105, 813-820.	1.7	13
62	Lower airway anomalies in children with CATCH 22 syndrome and congenital heart disease. Pediatric Pulmonology, 2013, 48, 587-591.	2.0	13
63	The prevention of coronary arterial abnormalities in Kawasaki disease: AÂmeta-analysis of the corticosteroid effectiveness. Journal of Microbiology, Immunology and Infection, 2018, 51, 321-331.	3.1	13
64	A genome-wide association analysis identifies novel susceptibility loci for coronary arterial lesions in patients with Kawasaki disease. Translational Research, 2013, 161, 513-515.	5.0	12
65	Mid-to-long-term follow-up results of transcatheter closure of atrial septal defect in patients older than 40Âyears. Heart and Vessels, 2017, 32, 467-473.	1.2	12
66	Left ventricular reverse remodeling after successful cardiac resynchronization therapy in a 3-year-old girl with idiopathic dilated cardiomyopathy. International Journal of Cardiology, 2007, 117, e7-e9.	1.7	11
67	Coronary Artery Diameters in Infants and Children With Congenital Heart Disease as Determined by Computed Tomography. American Journal of Cardiology, 2007, 100, 1696-1701.	1.6	11
68	Restoration of cardiac function by setting the ventricular pacing at a lower range in an infant with congenital complete atrioventricular block and dilated cardiomyopathy. International Journal of Cardiology, 2008, 131, e38-e40.	1.7	11
69	Alternative approach for selected severe pulmonary hypertension of congenital heart defect without initial correction — Palliative surgical treatment. International Journal of Cardiology, 2011, 151, 313-317.	1.7	11
70	Shock and unresponsiveness to repeated courses of intravenous immunoglobulin in Kawasaki disease: a nationwide database study. Pediatric Research, 2020, 87, 961-966.	2.3	11
71	Safety and efficacy of transcatheter closure of outlet-type ventricular septal defects in children and adults with Amplatzer Duct Occluder II. Journal of the Formosan Medical Association, 2021, 120, 180-188.	1.7	11
72	Increased microvolt T-wave alternans in patients with repaired tetralogy of Fallot. International Journal of Cardiology, 2012, 159, 220-224.	1.7	10

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73	Estimation of the Incidence of Kawasaki Disease in Taiwan. A Comparison of Two Data Sources: Nationwide Hospital Survey and National Health Insurance Claims. Pediatrics and Neonatology, 2014, 55, 97-100.	0.9	10
74	Successful Transcatheter Handmade-Valved Graft Stent for Branch Pulmonary Regurgitation: Novel Approach in a Special Event. Annals of Thoracic Surgery, 2016, 102, e541-e543.	1.3	10
75	Low immunoglobulin M memory B-cell percentage in patients with heterotaxy syndrome correlates with the risk of severe bacterial infection. Pediatric Research, 2016, 79, 271-277.	2.3	10
76	Long-term outcomes of arrhythmia and distinct electrophysiological features in congenitally corrected transposition of the great arteries in an Asian cohort. American Heart Journal, 2021, 231, 73-81.	2.7	10
77	Resolution of pathologic Q wave, left ventricular dysfunction and mitral regurgitation after dual coronary repair of the anomalous origin of the left coronary artery from the pulmonary artery. European Journal of Pediatrics, 2008, 167, 1277-1282.	2.7	9
78	Long-Term Outcomes of Pediatric Sinus Bradycardia. Journal of Pediatrics, 2013, 163, 885-889.e1.	1.8	9
79	Midterm followâ€up results of transcatheter treatment in patients with unroofed coronary sinus. Catheterization and Cardiovascular Interventions, 2014, 83, 243-249.	1.7	9
80	Risk of Respiratory Syncytial Virus Infection in Cyanotic Congenital Heart Disease in a Subtropical Area. Journal of Pediatrics, 2016, 171, 25-30.e1.	1.8	9
81	Transcatheter Closure of Perimembranous Ventricular Septal Defects With Amplatzer Duct Occluders. JACC: Cardiovascular Interventions, 2017, 10, 2227-2228.	2.9	9
82	Atrial flutter/fibrillation in patients receiving transcatheter closure of atrial septal defect. Journal of the Formosan Medical Association, 2017, 116, 522-528.	1.7	9
83	Twin atrioventricular nodes, arrhythmias, and survival in pediatric and adult patients with heterotaxy syndrome. Heart Rhythm, 2021, 18, 605-612.	0.7	9
84	Six-Minute Walking Test: Normal Reference Values for Taiwanese Children and Adolescents. Acta Cardiologica Sinica, 2015, 31, 193-201.	0.2	9
85	Pulmonary Hypertension in Adult Congenital Heart Disease in Asia: A Distinctive Feature of Complex Congenital Heart Disease. Journal of the American Heart Association, 2022, 11, e022596.	3.7	9
86	Long-term outcome of repaired tetralogy of Fallot: Survival, tachyarrhythmia, and impact of pulmonary valve replacement. Heart Rhythm, 2022, 19, 1856-1863.	0.7	9
87	Efficacy of a Novel Palivizumab Prophylaxis Protocol for Respiratory Syncytial Virus Infection in Congenital Heart Disease: A Multicenter Study. Journal of Pediatrics, 2018, 195, 108-114.e1.	1.8	8
88	Comparison of risk scores for predicting intravenous immunoglobulin resistance in Taiwanese patients with Kawasaki disease. Journal of the Formosan Medical Association, 2021, 120, 1884-1889.	1.7	8
89	Left Ventricular Outflow Tract Obstruction in Complete Transposition of the Great Arteries - Echocardiography Criteria for Surgical Strategies Circulation Journal, 2010, 74, 1214-1218.	1.6	7
90	The Role of Mechanical-Electrical Interaction in Ventricular Arrhythmia: Evidence From a Novel Animal Model for Repaired Tetralogy of Fallot. Pediatric Research, 2011, 70, 247-252.	2.3	7

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91	High Sensitivity C Reactive Protein (hs-CRP) in Adolescent and Young Adult Patients with History of Kawasaki Disease. Acta Cardiologica Sinica, 2015, 31, 473-7.	0.2	7
92	Long QT Syndrome Manifested as Fetal Ventricular Tachycardia and Intermittent AV Block. American Journal of Perinatology, 1998, 15, 145-147.	1.4	6
93	Acute gastric volvulus in a child with asplenia syndrome. Pediatrics International, 2004, 46, 471-473.	0.5	6
94	Idioventricular rhythm induced by therapeutic hypothermia. Resuscitation, 2008, 76, 471-473.	3.0	6
95	Dilated Cardiomyopathy After Long-Term Right Ventricular Apical Pacing in Children With Complete Atrioventricular Block: Role of Setting of Ventricular Pacing. Journal of Cardiac Failure, 2009, 15, 681-688.	1.7	6
96	Concomitant Slide Tracheoplasty and Cardiac Operation for Congenital Tracheal Stenosis Associated With VACTERL. Annals of Thoracic Surgery, 2013, 96, 1492-1495.	1.3	6
97	Congenital Right Intermediate Bronchial Stenosis With Carina Trifurcation: Successful Management With Slide Tracheobronchial Plasty. Annals of Thoracic Surgery, 2014, 98, 357-359.	1.3	6
98	Diagnosis of Pulmonary Arterial Hypertension in Children by Using Cardiac Computed Tomography. Korean Journal of Radiology, 2019, 20, 976.	3.4	6
99	Outcomes of Kawasaki Disease Children With Spontaneous Defervescence Within 10 Days. Frontiers in Pediatrics, 2019, 7, 158.	1.9	6
100	Congenital heart disease with pulmonary artery hypertension in an Asian cohort-initial report from TACHYON (TAiwan congenital heart disease associated with pulmonarY arterial hypertension) registry. International Journal of Cardiology, 2020, 317, 49-55.	1.7	6
101	Increased prevalence of inattention-related symptoms in a large cohort of patients with congenital heart disease. European Child and Adolescent Psychiatry, 2021, 30, 647-655.	4.7	6
102	Hypoplastic Left Heart Syndrome With Valvular Pulmonary Stenosis: Successful Management With Norwood Staged Reconstruction. Annals of Thoracic Surgery, 2011, 92, 1115-1116.	1.3	5
103	An Unusual Cause of Cyanosis in a Child. Gastroenterology, 2018, 155, e5-e6.	1.3	5
104	Energy Balance-Related Behaviors and Body Mass Index in Asian School-Aged Children With Congenital Heart Disease. Journal of Cardiovascular Nursing, 2020, 35, 291-299.	1.1	5
105	Transcatheter Closure of Atrial Septal Defect Associated With Pulmonary Artery Hypertension using Fenestrated Devices. American Journal of Cardiology, 2021, 147, 122-128.	1.6	5
106	Zero Fluoroscopy During Ablation of Right-Sided Supraventricular Tachycardia Substrates in a Pediatric Population - Initial Experience in Taiwan. Acta Cardiologica Sinica, 2019, 35, 476-483.	0.2	5
107	Special electrophysiological characteristics of pediatric idiopathic ventricular tachycardia. International Journal of Cardiology, 2017, 227, 595-601.	1.7	4
108	Diagnosis and Treatment for embolic stroke of undetermined source: Consensus statement from the Taiwan stroke society and Taiwan society of cardiology. Journal of the Formosan Medical Association, 2021, 120, 93-106.	1.7	4

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109	Prognostic markers in patients undergoing transcatheter implantation of Venus P-valve: Experience in Taiwan. Journal of the Formosan Medical Association, 2021, 120, 1202-1211.	1.7	4
110	Association of the C677T methylenetetrahydrofolate reductase mutation with congenital heart diseases. Acta Obstetricia Et Gynecologica Scandinavica, 2005, 84, 1134-1140.	2.8	4
111	Long-Term Outcomes of Native Coarctation of the Aorta after Balloon Angioplasty or Surgical Aortoplasty in Newborns and Young Infants Less Than 3 Months of Age. Acta Cardiologica Sinica, 2013, 29, 168-74.	0.2	4
112	Preoperative N-terminal pro-brain natriuretic peptide is associated with Fontan outcomes. Journal of Thoracic and Cardiovascular Surgery, 2021, , .	0.8	4
113	Assessing utility of exercise test in determining exercise prescription in adolescent and adult patients with repaired tetralogy of fallot. Heart and Vessels, 2017, 32, 201-207.	1.2	3
114	Improved Left Ventricular Strain and Dyssynchrony After Pulmonary Artery Banding in an Infant with End-Stage Dilated Cardiomyopathy: Insights from Three-Dimensional Speckle Tracking. Pediatric Cardiology, 2019, 40, 1317-1319.	1.3	3
115	Exercise Capacity and Ventricular Remodeling After Transcatheter Ventricular Septal Defect Closure in Asymptomatic or Minimally Symptomatic Adolescents and Adults. Circulation: Cardiovascular Interventions, 2020, 13, e008813.	3.9	3
116	Perioperative outcomes of Fontan operation: Impact of heterotaxy syndrome. Journal of the Formosan Medical Association, 2021, 121, 89-89.	1.7	3
117	Diffuse midline glioma presenting with central sleep apnea and pulmonary hypertension in a 4-year-old patient: a case report. Journal of Clinical Sleep Medicine, 2021, 17, 325-328.	2.6	3
118	Rapid Two-stage Versus One-stage Surgical Repair of Interrupted Aortic Arch with Ventricular Septal Defect in Neonates. Journal of the Formosan Medical Association, 2008, 107, 876-884.	1.7	2
119	Fibrinolytic Therapy in Mechanical Valve Thrombosis in a One-Year-Old Girl. Circulation, 2010, 121, e244-5.	1.6	2
120	Factors affecting motor development of toddlers who received cardiac corrective procedures during infancy. Early Human Development, 2021, 158, 105392.	1.8	2
121	Variation in Pharmacologic Management of Patients with Kawasaki Disease with Coronary Artery Aneurysms. Journal of Pediatrics, 2021, , .	1.8	2
122	Long-Term Outcomes and Prognosticators of Pediatric Primary Dilated Cardiomyopathy in an Asian Cohort. Frontiers in Pediatrics, 2021, 9, 771283.	1.9	2
123	S12-3 FAVORABLE OUTCOME FOR TOTAL REPAIR OF TETRALOGY OF FALLOT IN YOUNG INFANT STAGE. International Journal of Cardiology, 2007, 122, S36.	1.7	1
124	Never forget the abdomen in the evaluation of pediatric cardiac computed tomography. International Journal of Cardiology, 2012, 157, e31-e32.	1.7	1
125	Rotational Atherectomy and Stent Implantation in an 11-Year-Old Boy with a History of Kawasaki Disease. Pediatrics and Neonatology, 2016, 57, 248-251.	0.9	1
126	Partial anomalous pulmonary venous return to the azygos vein: A case report. Journal of the Formosan Medical Association, 2016, 115, 481-482.	1.7	1

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127	Kawasaki Disease Shock Syndrome. , 2017, , 59-63.		1
128	Assessing Late Cardiopulmonary Function in Patients with Repaired Tetralogy of Fallot Using Exercise Cardiopulmonary Function Test and Cardiac Magnetic Resonance. Acta Cardiologica Sinica, 2015, 31, 478-84.	0.2	1
129	Incidence and Postnatal Profile of Fontan Patients by Adolescence from a Nationwide Birth Cohort. Acta Cardiologica Sinica, 2020, 36, 367-374.	0.2	1
130	Changing Spectrum of Cardiac Diseases in Children: An Extended Longitudinal Observation Study of a Pediatric Cardiac Screening Program. Acta Cardiologica Sinica, 2021, 37, 420-426.	0.2	1
131	Reappraisal of the Subtropical Guidelines on Palivizumab Prophylaxis in Congenital Heart Disease. Frontiers in Pediatrics, 2021, 9, 756787.	1.9	1
132	Zero-fluoroscopy ablation of left-sided arrhythmia substrates in children – Mid-term safety and feasibility study from transaortic approach. Journal of the Formosan Medical Association, 2022, 121, 2035-2043.	1.7	1
133	Right ventricular aneurysm and large coronary arteriovenous fistula in a 3100 g infant. European Heart Journal, 2011, 32, 1688-1688.	2.2	0
134	Tetralogy of Fallot with fifth aortic arch. Acta Cardiologica, 2017, 72, 672-673.	0.9	0
135	Infective endocarditis involving atrial septal occluder. Archives of Disease in Childhood, 2018, 103, 1149-1149.	1.9	0
136	Early diagnosis and treatment of anomalous origin of left coronary artery from pulmonary artery in an asymptomatic 1â€monthâ€old infant. Kaohsiung Journal of Medical Sciences, 2021, 37, 922-923.	1.9	0
137	Neonatal tricuspid stenosis caused by device closure of a large coronary fistula. EuroIntervention, 2015, 11, e1-e1.	3.2	0
138	Lung transplantation with concomitant cardiac repair for congenital hypoplasia of bilateral pulmonary arteries and patent ductus arteriosus. Journal of Cardiothoracic Surgery, 2022, 17, 49.	1.1	0
139	Association of Acute Anti-Inflammatory Treatment with Medium-Term Outcomes for Coronary Artery Aneurysms in Kawasaki Disease. , 2022, , .		0