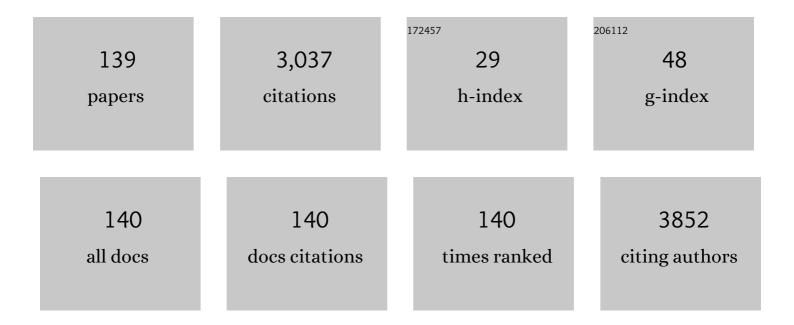
List of Publications by Year in descending order

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MINC-TALLIN

#	Article	IF	CITATIONS
1	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
2	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
3	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
4	Association of Acute Anti-Inflammatory Treatment with Medium-Term Outcomes for Coronary Artery Aneurysms in Kawasaki Disease. , 2022, , .		0
5	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
6	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
7	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
8	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
9	Twin atrioventricular nodes, arrhythmias, and survival in pediatric and adult patients with heterotaxy syndrome. Heart Rhythm, 2021, 18, 605-612.	0.7	9
10	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
11	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
12	Perioperative outcomes of Fontan operation: Impact of heterotaxy syndrome. Journal of the Formosan Medical Association, 2021, 121, 89-89.	1.7	3
13	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
14	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
15	Factors affecting motor development of toddlers who received cardiac corrective procedures during infancy. Early Human Development, 2021, 158, 105392.	1.8	2
16	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
17	Variation in Pharmacologic Management of Patients with Kawasaki Disease with Coronary Artery Aneurysms. Journal of Pediatrics, 2021, , .	1.8	2
18	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

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19	Long-Term Outcomes and Prognosticators of Pediatric Primary Dilated Cardiomyopathy in an Asian Cohort. Frontiers in Pediatrics, 2021, 9, 771283.	1.9	2
20	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
21	Reappraisal of the Subtropical Guidelines on Palivizumab Prophylaxis in Congenital Heart Disease. Frontiers in Pediatrics, 2021, 9, 756787.	1.9	1
22	Preoperative N-terminal pro-brain natriuretic peptide is associated with Fontan outcomes. Journal of Thoracic and Cardiovascular Surgery, 2021, , .	0.8	4
23	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
24	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
25	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
26	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
27	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
28	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
29	Incidence and Postnatal Profile of Fontan Patients by Adolescence from a Nationwide Birth Cohort. Acta Cardiologica Sinica, 2020, 36, 367-374.	0.2	1
30	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
31	Diagnosis of Pulmonary Arterial Hypertension in Children by Using Cardiac Computed Tomography. Korean Journal of Radiology, 2019, 20, 976.	3.4	6
32	Outcomes of Kawasaki Disease Children With Spontaneous Defervescence Within 10 Days. Frontiers in Pediatrics, 2019, 7, 158.	1.9	6
33	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
34	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
35	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
36	Infective endocarditis involving atrial septal occluder. Archives of Disease in Childhood, 2018, 103, 1149-1149.	1.9	0

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37	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
38	The global epidemiology of Kawasaki disease: Review and future perspectives. Global Cardiology Science & Practice, 2018, 2017, e201720.	0.4	85
39	An Unusual Cause of Cyanosis in a Child. Gastroenterology, 2018, 155, e5-e6.	1.3	5
40	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
41	Risk factors and implications of progressive coronary dilatation in children with Kawasaki disease. BMC Pediatrics, 2017, 17, 139.	1.7	29
42	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
43	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
44	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
45	Transcatheter Closure of Perimembranous Ventricular Septal Defects With Amplatzer Duct Occluders. JACC: Cardiovascular Interventions, 2017, 10, 2227-2228.	2.9	9
46	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
47	Special electrophysiological characteristics of pediatric idiopathic ventricular tachycardia. International Journal of Cardiology, 2017, 227, 595-601.	1.7	4
48	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
49	Kawasaki Disease Shock Syndrome. , 2017, , 59-63.		1
50	Tetralogy of Fallot with fifth aortic arch. Acta Cardiologica, 2017, 72, 672-673.	0.9	0
51	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
52	Acute kidney injury in patients with Kawasaki disease. Pediatric Research, 2016, 80, 224-227.	2.3	26
53	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
54	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

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55	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
56	Progressive Coronary Dilatation Predicts Worse Outcome in Kawasaki Disease. Journal of Pediatrics, 2016, 171, 78-82.e1.	1.8	23
57	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
58	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
59	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
60	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
61	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
62	Neonatal tricuspid stenosis caused by device closure of a large coronary fistula. EuroIntervention, 2015, 11, e1-e1.	3.2	0
63	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
64	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
65	Six-Minute Walking Test: Normal Reference Values for Taiwanese Children and Adolescents. Acta Cardiologica Sinica, 2015, 31, 193-201.	0.2	9
66	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
67	Midterm followâ€up results of transcatheter treatment in patients with unroofed coronary sinus. Catheterization and Cardiovascular Interventions, 2014, 83, 243-249.	1.7	9
68	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
69	Changing Spectrum of Infective Endocarditis in Children. Pediatric Infectious Disease Journal, 2014, 33, 467-471.	2.0	16
70	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
71	Viral infections associated with Kawasaki disease. Journal of the Formosan Medical Association, 2014, 113, 148-154.	1.7	190
72	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

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73	Severe Bacterial Infection in Patients with Heterotaxy Syndrome. Journal of Pediatrics, 2014, 164, 99-104.e1.	1.8	33
74	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
75	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
76	Long-Term Outcomes of Pediatric Sinus Bradycardia. Journal of Pediatrics, 2013, 163, 885-889.e1.	1.8	9
77	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
78	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
79	Concomitant Slide Tracheoplasty and Cardiac Operation for Congenital Tracheal Stenosis Associated With VACTERL. Annals of Thoracic Surgery, 2013, 96, 1492-1495.	1.3	6
80	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
81	Lower airway anomalies in children with CATCH 22 syndrome and congenital heart disease. Pediatric Pulmonology, 2013, 48, 587-591.	2.0	13
82	Population-based Study of Kawasaki Disease Shock Syndrome in Taiwan. Pediatric Infectious Disease Journal, 2013, 32, 1384-1386.	2.0	48
83	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
84	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
85	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
86	Increased microvolt T-wave alternans in patients with repaired tetralogy of Fallot. International Journal of Cardiology, 2012, 159, 220-224.	1.7	10
87	Never forget the abdomen in the evaluation of pediatric cardiac computed tomography. International Journal of Cardiology, 2012, 157, e31-e32.	1.7	1
88	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
89	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
90	Natural and unnatural history of tetralogy of Fallot repaired during adolescence and adulthood. Heart and Vessels, 2012, 27, 65-70.	1.2	15

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91	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
92	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
93	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
94	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
95	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
96	Quality of life in adults with congenital heart disease: biopsychosocial determinants and sex-related differences. Heart, 2011, 97, 38-43.	2.9	52
97	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
98	Right ventricular aneurysm and large coronary arteriovenous fistula in a 3100 g infant. European Heart Journal, 2011, 32, 1688-1688.	2.2	0
99	Transcatheter Closure of Moderate-to-Large Patent Ductus Arteriosus in Infants Using Amplatzer Duct Occluder. Circulation Journal, 2010, 74, 361-364.	1.6	39
100	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
101	Fibrinolytic Therapy in Mechanical Valve Thrombosis in a One-Year-Old Girl. Circulation, 2010, 121, e244-5.	1.6	2
102	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
103	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
104	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
105	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
106	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
107	Radiofrequency Catheter Ablation of Supraventricular Tachycardia in Infants and Toddlers. Circulation Journal, 2009, 73, 1717-1721.	1.6	42
108	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

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109	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
110	Transcatheter closure of atrial septal defect without balloon sizing. Catheterization and Cardiovascular Interventions, 2008, 71, 214-221.	1.7	52
111	Idioventricular rhythm induced by therapeutic hypothermia. Resuscitation, 2008, 76, 471-473.	3.0	6
112	Cardiac Conduction Disturbance Detected in a Pediatric Population. Journal of Pediatrics, 2008, 152, 85-89.	1.8	44
113	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
114	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
115	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
116	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
117	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
118	Abnormal Matrix Remodeling in Adolescents and Young Adults with Kawasaki Disease Late after Onset. Clinical Chemistry, 2008, 54, 1815-1822.	3.2	14
119	Progression of aortic regurgitation after surgical repair of outlet-type ventricular septal defects. American Heart Journal, 2007, 153, 336-342.	2.7	33
120	Left Pulmonary Artery Sling Complex: Computed Tomography and Hypothesis of Embryogenesis. Annals of Thoracic Surgery, 2007, 84, 1645-1650.	1.3	46
121	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
122	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
123	Coronary artery anatomy in children with congenital heart disease by computed tomography. International Journal of Cardiology, 2007, 120, 363-370.	1.7	28
124	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
125	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
126	Surgical Outcome of Aortopulmonary Window Repair in Early Infancy. Journal of the Formosan Medical Association, 2006, 105, 813-820.	1.7	13

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127	Clinical Implications of Major Aortopulmonary Collateral Arteries in Patients With Right Isomerism. Annals of Thoracic Surgery, 2006, 82, 153-157.	1.3	14
128	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
129	Noninvasive diagnosis of aortic coarctation in neonates with patent ductus arteriosus. Journal of Pediatrics, 2006, 148, 217-221.	1.8	24
130	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
131	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
132	Association of the C677T methylenetetrahydrofolate reductase mutation with congenital heart diseases. Acta Obstetricia Et Gynecologica Scandinavica, 2005, 84, 1134-1140.	2.8	22
133	Acquired coronary artery fistula after open heart surgery for congenital heart disease. International Journal of Cardiology, 2005, 103, 187-192.	1.7	32
134	Aortic Valve Prolapse Associated With Outlet-Type Ventricular Septal Defect. Annals of Thoracic Surgery, 2005, 79, 1366-1371.	1.3	35
135	Association of the C677T methylenetetrahydrofolate reductase mutation with congenital heart diseases. Acta Obstetricia Et Gynecologica Scandinavica, 2005, 84, 1134-1140.	2.8	4
136	Acute gastric volvulus in a child with asplenia syndrome. Pediatrics International, 2004, 46, 471-473.	0.5	6
137	Postnatal outcome of fetal bradycardia without significant cardiac abnormalities. American Heart Journal, 2004, 147, 540-544.	2.7	33
138	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
139	Long QT Syndrome Manifested as Fetal Ventricular Tachycardia and Intermittent AV Block. American Journal of Perinatology, 1998, 15, 145-147.	1.4	6