Cordula M Wolf

List of Publications by Year
in descending order

18	150	1478505 6	1372567 10
papers	citations	h-index	g-index

20	20	20	163
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Reclassification of genetic variants in children with long QT syndrome. Molecular Genetics & Genomic Medicine, 2020, 8, e1300.	1.2	23
2	Extensive identification of genes involved in congenital and structural heart disorders and cardiomyopathy. , 2022, 1, 157-173.		22
3	Clinical long-term outcome of septal myectomy for obstructive hypertrophic cardiomyopathy in infants. European Journal of Cardio-thoracic Surgery, 2018, 53, 538-544.	1.4	19
4	Long-term outcomes of childhood onset Noonan compared to sarcomere hypertrophic cardiomyopathy. Cardiovascular Diagnosis and Therapy, 2019, 9, S299-S309.	1.7	16
5	Reasons for Failure of Systemic-to-Pulmonary Artery Shunts in Neonates. Thoracic and Cardiovascular Surgeon, 2019, 67, 002-007.	1.0	13
6	NADPH oxidases and HIF1 promote cardiac dysfunction and pulmonary hypertension in response to glucocorticoid excess. Redox Biology, 2020, 34, 101536.	9.0	11
7	Management of cardiac aspects in children with Noonan syndrome â€“ results from a European clinical practice survey among paediatric cardiologists. European Journal of Medical Genetics, 2021, 65, 104372.	1.3	11
8	Cell cycle defects underlie childhood-onset cardiomyopathy associated with Noonan syndrome. IScience, 2022, 25, 103596.	4.1	9
9	Management of growth failure and other endocrine aspects in patients with Noonan syndrome across Europe: A sub-analysis of a European clinical practice survey. European Journal of Medical Genetics, 2022, 65, 104404.	1.3	6
10	Risk Factors for Failure of Systemic-to-Pulmonary Artery Shunts in Biventricular Circulation. Pediatric Cardiology, 2018, 39, 1323-1329.	1.3	5
11	Compound Mutation in Cardiac Sarcomere Proteins Is Associated with Increased Risk for Major Arrhythmic Events in Pediatric Onset Hypertrophic Cardiomyopathy. Journal of Clinical Medicine, 2021, 10, 5256.	2.4	4
12	European Medical Education Initiative on Noonan syndrome: A clinical practice survey assessing the diagnosis and clinical management of individuals with Noonan syndrome across Europe. European Journal of Medical Genetics, 2021, 65, 104371.	1.3	3
13	A case of Uhlâ€™s anomaly presenting with ventricular tachycardia. European Heart Journal Cardiovascular Imaging, 2018, 19, 1312-1312.	1.2	2
14	Low-molecular-weight heparin administered by subcutaneous catheter is a safe and effective anti-coagulation regimen in selected inpatient infants and children with complex congenital heart disease. Cardiology in the Young, 2021, 31, 1439-1444.	0.8	2
15	Cardiovascular Function and Exercise Capacity in Childhood Cancer Survivors. Journal of Clinical Medicine, 2022, 11, 628.	2.4	2
16	No increased extracellular volume fraction or conduction time after childhood septal myectomy. European Journal of Cardio-thoracic Surgery, 2020, 57, 958-964.	1.4	1
17	Surgical septal myectomy for obstructive hypertrophic cardiomyopathy during infancy. Translational Pediatrics, 2019, 8, 90-91.	1.2	0
18	Peak Oxygen Uptake on Cardiopulmonary Exercise Test Is a Predictor for Severe Arrhythmic Events during Three-Year Follow-Up in Patients with Complex Congenital Heart Disease. Journal of Cardiovascular Development and Disease, 2022, 9, 215.	1.6	0