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	Cardiovascular Function and Exercise Capacity in Childhood Cancer Survivors. Journal of Clinical Medicine, 2022, 11, 628.	2.4	2
2	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
3	Cell cycle defects underlie childhood-onset cardiomyopathy associated with Noonan syndrome. IScience, 2022, 25, 103596.	4.1	9
4	Extensive identification of genes involved in congenital and structural heart disorders and cardiomyopathy. , 2022, 1, 157-173.		22
5	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
6	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
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	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
9	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
10	NADPH oxidases and HIF1 promote cardiac dysfunction and pulmonary hypertension in response to glucocorticoid excess. Redox Biology, 2020, 34, 101536.	9.0	11
11	Reclassification of genetic variants in children with long QT syndrome. Molecular Genetics & Genomic Medicine, 2020, 8, e1300.	1.2	23
12	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
13	Surgical septal myectomy for obstructive hypertrophic cardiomyopathy during infancy. Translational Pediatrics, 2019, 8, 90-91.	1.2	0
14	Long-term outcomes of childhood onset Noonan compared to sarcomere hypertrophic cardiomyopathy. Cardiovascular Diagnosis and Therapy, 2019, 9, S299-S309.	1.7	16
15	Reasons for Failure of Systemic-to-Pulmonary Artery Shunts in Neonates. Thoracic and Cardiovascular Surgeon, 2019, 67, 002-007.	1.0	13
16	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
17	Risk Factors for Failure of Systemic-to-Pulmonary Artery Shunts in Biventricular Circulation. Pediatric Cardiology, 2018, 39, 1323-1329.	1.3	5
18	A case of Uhlâ€™s anomaly presenting with ventricular tachycardia. European Heart Journal Cardiovascular Imaging, 2018, 19, 1312-1312.	1.2	2