

Erwin Oechslin

List of Publications by Year in descending order

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Version: 2024-02-01

94
papers

6,659
citations

218381

26
h-index

62479

80
g-index

119
all docs

119
docs citations

119
times ranked

5994
citing authors

#	ARTICLE	IF	CITATIONS
1	ESC Guidelines for the management of grown-up congenital heart disease (new version 2010): The Task Force on the Management of Grown-up Congenital Heart Disease of the European Society of Cardiology (ESC). <i>European Heart Journal</i> , 2010, 31, 2915-2957.	1.0	2,134
2	2020 ESC Guidelines for the management of adult congenital heart disease. <i>European Heart Journal</i> , 2021, 42, 563-645.	1.0	971
3	Mutations in Sarcomere Protein Genes in Left Ventricular Noncompaction. <i>Circulation</i> , 2008, 117, 2893-2901.	1.6	414
4	Left ventricular non-compaction revisited: a distinct phenotype with genetic heterogeneity?. <i>European Heart Journal</i> , 2011, 32, 1446-1456.	1.0	376
5	The spectrum of adult congenital heart disease in Europe: morbidity and mortality in a 5 year follow-up period. <i>European Heart Journal</i> , 2005, 26, 2325-2333.	1.0	370
6	Depression and anxiety in adult congenital heart disease: Predictors and prevalence. <i>International Journal of Cardiology</i> , 2009, 137, 158-164.	0.8	276
7	Canadian Cardiovascular Society 2009 Consensus Conference on the management of adults with congenital heart disease: Executive summary. <i>Canadian Journal of Cardiology</i> , 2010, 26, 143-150.	0.8	175
8	Sarcomere Gene Mutations in Isolated Left Ventricular Noncompaction Cardiomyopathy Do Not Predict Clinical Phenotype. <i>Circulation: Cardiovascular Genetics</i> , 2011, 4, 367-374.	5.1	167
9	Transition Intervention for Adolescents With Congenital Heart Disease. <i>Journal of the American College of Cardiology</i> , 2018, 71, 1768-1777.	1.2	107
10	Canadian Cardiovascular Society 2009 Consensus Conference on the management of adults with congenital heart disease: Complex congenital cardiac lesions. <i>Canadian Journal of Cardiology</i> , 2010, 26, e98-e117.	0.8	97
11	Delivery of care for adult patients with congenital heart disease in Europe: results from the Euro Heart Survey. <i>European Heart Journal</i> , 2006, 27, 1324-1330.	1.0	92
12	Systemic Endothelial Dysfunction in Adults With Cyanotic Congenital Heart Disease. <i>Circulation</i> , 2005, 112, 1106-1112.	1.6	90
13	Hepatocellular Carcinoma After Fontan Operation. <i>Circulation</i> , 2018, 138, 746-748.	1.6	82
14	Pheochromocytoma and Paraganglioma in Cyanotic Congenital Heart Disease. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, 1325-1334.	1.8	77
15	Hematological management of the cyanotic adult with congenital heart disease. <i>International Journal of Cardiology</i> , 2004, 97, 109-115.	0.8	75
16	Left Ventricular Noncompaction. <i>Journal of the American College of Cardiology</i> , 2018, 71, 723-726.	1.2	61
17	Haploinsufficiency of vascular endothelial growth factor related signaling genes is associated with tetralogy of Fallot. <i>Genetics in Medicine</i> , 2019, 21, 1001-1007.	1.1	58
18	The Adult Patient with Eisenmenger Syndrome: A Medical Update After Dana Point Part I: Epidemiology, Clinical Aspects and Diagnostic Options. <i>Current Cardiology Reviews</i> , 2010, 6, 343-355.	0.6	56

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19	Adults with Congenital Heart Disease: Psychological Needs and Treatment Preferences. <i>Congenital Heart Disease</i> , 2009, 4, 139-146.	0.0	52
20	Consensus recommendations for echocardiography in adults with congenital heart defects from the International Society of Adult Congenital Heart Disease (ISACHD). <i>International Journal of Cardiology</i> , 2018, 272, 77-83.	0.8	49
21	The Adult Patient with Eisenmenger Syndrome: A Medical Update after Dana Point Part III: Specific Management and Surgical Aspects. <i>Current Cardiology Reviews</i> , 2010, 6, 363-372.	0.6	48
22	Hepatocellular carcinoma and the Fontan circulation: Clinical presentation and outcomes. <i>International Journal of Cardiology</i> , 2021, 322, 142-148.	0.8	45
23	Reaching consensus for unified medical language in Fontan care. <i>ESC Heart Failure</i> , 2021, 8, 3894-3905.	1.4	35
24	Heart University: a new online educational forum in paediatric and adult congenital cardiac care. The future of virtual learning in a post-pandemic world?. <i>Cardiology in the Young</i> , 2020, 30, 560-567.	0.4	34
25	Pregnancy-Related Obstetric and Cardiologic Problems in Women After Atrial Switch Operation for Transposition of the Great Arteries. <i>Circulation Journal</i> , 2014, 78, 443-449.	0.7	31
26	Prevention of Sudden Cardiac Death in Adults With Congenital Heart Disease. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2017, 10, .	2.1	29
27	Interaction of Sildenafil With cAMP-Mediated Vasodilation In Vivo. <i>Hypertension</i> , 2002, 40, 763-767.	1.3	28
28	Management of adults with cyanotic congenital heart disease. <i>Heart</i> , 2015, 101, 485-494.	1.2	28
29	Canadian Cardiovascular Society 2022 Guidelines for Cardiovascular Interventions in Adults With Congenital Heart Disease. <i>Canadian Journal of Cardiology</i> , 2022, 38, 862-896.	0.8	28
30	Improving medical care and prevention in adults with congenital heart disease—reflections on a global problem—part I: development of congenital cardiology, epidemiology, clinical aspects, heart failure, cardiac arrhythmia. <i>Cardiovascular Diagnosis and Therapy</i> , 2018, 8, 705-715.	0.7	26
31	Return of genetic and genomic research findings: experience of a pediatric biorepository. <i>BMC Medical Genomics</i> , 2019, 12, 173.	0.7	24
32	Adherence to guidelines in the clinical care for adults with congenital heart disease: The Euro Heart Survey on Adult Congenital Heart Disease. <i>European Heart Journal</i> , 2006, 27, 737-745.	1.0	23
33	A call for adult congenital heart disease patient participation in cardiac rehabilitation. <i>International Journal of Cardiology</i> , 2011, 150, 345-346.	0.8	23
34	Eisenmenger Syndrome: A Multisystem Disorder—Do Not Destabilize the Balanced but Fragile Physiology. <i>Canadian Journal of Cardiology</i> , 2019, 35, 1664-1674.	0.8	23
35	Genes and Pathways Implicated in Tetralogy of Fallot Revealed by Ultra-Rare Variant Burden Analysis in 231 Genome Sequences. <i>Frontiers in Genetics</i> , 2020, 11, 957.	1.1	23
36	Nosology of Noncompaction Cardiomyopathy: The Emperor Still Wears Clothes!. <i>Canadian Journal of Cardiology</i> , 2017, 33, 701-704.	0.8	22

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37	Factors Influencing Participation in a Population-based Biorepository for Childhood Heart Disease. <i>Pediatrics</i> , 2012, 130, e1198-e1205.	1.0	21
38	Self-efficacy as a predictor of patient-reported outcomes in adults with congenital heart disease. <i>European Journal of Cardiovascular Nursing</i> , 2018, 17, 619-626.	0.4	21
39	Risk prediction models for heart failure admissions in adults with congenital heart disease. <i>International Journal of Cardiology</i> , 2021, 322, 149-157.	0.8	21
40	Cardiovascular magnetic resonance based diagnosis of left ventricular non-compaction cardiomyopathy: impact of cine bSSFP strain analysis. <i>Journal of Cardiovascular Magnetic Resonance</i> , 2020, 22, 9.	1.6	21
41	Sense of coherence in adults with congenital heart disease in 15 countries: Patient characteristics, cultural dimensions and quality of life. <i>European Journal of Cardiovascular Nursing</i> , 2021, 20, 48-55.	0.4	20
42	A cluster randomized trial of a transition intervention for adolescents with congenital heart disease: rationale and design of the CHAPTER 2 study. <i>BMC Cardiovascular Disorders</i> , 2016, 16, 127.	0.7	19
43	Transition of Care in Congenital Heart Disease: Ensuring the Proper Handoff. <i>Current Cardiology Reports</i> , 2017, 19, 55.	1.3	17
44	PREVENTION-ACHD: PRospEctIVE study on implaNTable cardioverter-defibrillator therapy and sudden cardiac death in Adults with Congenital Heart Disease; Rationale and Design. <i>Netherlands Heart Journal</i> , 2019, 27, 474-479.	0.3	17
45	Impact of a 22q11.2 Microdeletion on Adult All-Cause Mortality in Tetralogy of Fallot Patients. <i>Canadian Journal of Cardiology</i> , 2020, 36, 1091-1097.	0.8	17
46	Double aortic and pulmonary valves: an artifact generated by ultrasound refraction. <i>Journal of the American Society of Echocardiography</i> , 2004, 17, 786-787.	1.2	16
47	Common Genetic Variants Contribute to Risk of Transposition of the Great Arteries. <i>Circulation Research</i> , 2022, 130, 166-180.	2.0	15
48	Improving medical care and prevention in adults with congenital heart disease—reflections on a global problem—part II: infective endocarditis, pulmonary hypertension, pulmonary arterial hypertension and aortopathy. <i>Cardiovascular Diagnosis and Therapy</i> , 2018, 8, 716-724.	0.7	14
49	Whole genome sequencing delineates regulatory, copy number, and cryptic splice variants in early onset cardiomyopathy. <i>Npj Genomic Medicine</i> , 2022, 7, 18.	1.7	14
50	Prolonged T _{peak} – T _{end} interval is a risk factor for sudden cardiac death in adults with congenital heart disease. <i>Congenital Heart Disease</i> , 2019, 14, 952-957.	0.0	12
51	2020 ESC Guidelines for the management of adult congenital heart disease. <i>Revista Espanola De Cardiologia (English Ed)</i> , 2021, 74, 436.	0.4	12
52	Heart Failure and Patient-Reported Outcomes in Adults With Congenital Heart Disease from 15 Countries. <i>Journal of the American Heart Association</i> , 2022, 11, e024993.	1.6	10
53	Left Ventricular Noncompaction. <i>Journal of the American College of Cardiology</i> , 2019, 73, 1612-1615.	1.2	9
54	Bleeding and thrombotic risk in pregnant women with Fontan physiology. <i>Heart</i> , 2021, 107, 1390-1397.	1.2	9

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55	Left Ventricular Noncompaction Is a Myocardial Phenotype: Cardiomyopathyâ€”Yes or No?. Canadian Journal of Cardiology, 2021, 37, 366-369.	0.8	9
56	Renal dysfunction in adults with congenital heart defects. Progress in Pediatric Cardiology, 2016, 41, 51-57.	0.2	8
57	Reduced Systolic Function and Not Genetic Variants Determine Outcome in Pediatric and Adult Left Ventricular Noncompaction Cardiomyopathy. Frontiers in Pediatrics, 2021, 9, 722926.	0.9	8
58	Advance care planning and palliative care in ACHD: the healthcare providersâ€™ perspective. Cardiology in the Young, 2020, 30, 402-408.	0.4	7
59	The role of 22q11.2 deletion syndrome in the relationship between congenital heart disease and scoliosis. Spine Journal, 2020, 20, 956-963.	0.6	7
60	Augmentation of pulmonary blood flow and cardiac output by non-invasive external ventilation late after Fontan palliation. Heart, 2021, 107, 142-149.	1.2	7
61	Patient-Reported Outcomes in Adults With Congenital Heart Disease Following Hospitalization (from) Tj ETQq1 1 0,784314 rgBT /Overl	0.7	7
62	Rationale and Perspective of Endothelin-1 Antagonism in Acute Heart Failure. Journal of Cardiovascular Pharmacology, 2001, 38, S53-S57.	0.8	5
63	Primary lymphedema and other lymphatic anomalies are associated with 22q11.2 deletion syndrome. European Journal of Medical Genetics, 2018, 61, 411-415.	0.7	5
64	Healthcare system inputs and patient-reported outcomes: a study in adults with congenital heart defect from 15 countries. BMC Health Services Research, 2020, 20, 496.	0.9	5
65	Left Anterior Descending Coronary Artery Pseudoaneurysm Compressing the Main Pulmonary Artery in a Patient with Behçet's Disease. Echocardiography, 2012, 29, E91-3.	0.3	4
66	Anatomical complexity does not predict outcomes after COVID-19 in adults with congenital heart disease. Heart, 2021, 107, 1193-1195.	1.2	4
67	Transbaffle Multielectrode Mapping of Atrial Flutter Postâ€”Double Switch Operation. Journal of Cardiovascular Electrophysiology, 2016, 27, 1240-1241.	0.8	3
68	Cor Triatriatum Sinister with Secundum Atrial Septal Defect. Case, 2017, 1, 141-146.	0.1	3
69	Learning strategies among adult CHD fellows. Cardiology in the Young, 2019, 29, 1356-1360.	0.4	3
70	Lateral tunnel Fontan atrial tachycardia ablation trans-baffle access is not mandatory as the initial strategy. Journal of Interventional Cardiac Electrophysiology, 2020, 58, 299-306.	0.6	3
71	Cardiovascular and abdominal flow alterations in adults with morphologic evidence of liver disease post Fontan palliation. International Journal of Cardiology, 2020, 317, 63-69.	0.8	3
72	Eisenmenger Syndrome. , 2011, , 358-370.		2

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73	Management of Bilateral Axillary Aneurysms, Threatened Limb, and Diffuse Vasculopathy in a Patient With ACTA2 Mutation. Canadian Journal of Cardiology, 2017, 33, 554.e1-554.e3.	0.8	2
74	Congenital Aortic Arch Anomalies: Lessons Learned and to Learn!. Canadian Journal of Cardiology, 2019, 35, 373-375.	0.8	2
75	Misperception of Survival in Adult Congenital Heart Disease and Importance of Both Anatomic and Functional Indices: Educate Your Patients!. Canadian Journal of Cardiology, 2019, 35, 1635-1639.	0.8	2
76	Abnormal spirometry in adults with 22q11.2 microdeletion and congenital heart disease. International Journal of Cardiology Congenital Heart Disease, 2021, 3, 100085.	0.2	2
77	Anomalous origin of a coronary artery from the pulmonary artery presenting in adulthood: Experience from a tertiary center. International Journal of Cardiology Congenital Heart Disease, 2021, 4, 100169.	0.2	2
78	Treatment of adults with Eisenmenger syndrome – state of the art in the 21st century: a short overview. Cardiovascular Diagnosis and Therapy, 2021, 11, 1190-1199.	0.7	2
79	Outcome and right ventricle remodelling after valve replacement for pulmonic stenosis. Heart, 2022, 108, 1290-1295.	1.2	2
80	Bilateral Perirenal Lymphangioma: A Conservative Approach. Urologia Internationalis, 2005, 74, 188-189.	0.6	1
81	Feasibility of Transvenous Coronary Sinus Lead Implantation in Congenitally Corrected Transposition of the Great Arteries. Canadian Journal of Cardiology, 2014, 30, 248.e11.	0.8	1
82	Transmural circumflex infarction in a young Fontan patient: to leave home to find it. European Heart Journal Cardiovascular Imaging, 2018, 19, 953-954.	0.5	1
83	Echocardiography in adults with congenital heart disease: Combining the best of both worlds. International Journal of Cardiology, 2018, 272, 84-85.	0.8	1
84	Perception is not reality when risk stratifying adults with congenital heart disease for COVID-19. Open Heart, 2021, 8, e001660.	0.9	1
85	Pain in adults with congenital heart disease - An international perspective. International Journal of Cardiology Congenital Heart Disease, 2021, 5, 100200.	0.2	1
86	Recomendações da ESC para o tratamento da cardiopatia congênita no adulto (nova versão de 2010). Revista Portuguesa De Cardiologia, 2012, 31, 541.e1-541.e53.	0.2	0
87	Worsening Hypoxia Post Lung Transplant: What has Changed?. Heart Lung and Circulation, 2016, 25, e155-e157.	0.2	0
88	LATE COMPLICATIONS AND CARDIOPULMONARY PERFORMANCE IN ADULTS WITH FONTAN PHYSIOLOGY: RV-DOMINANT VERSUS THOSE WITH LV-DOMINANT OR UNSEPTATED VENTRICLES. Journal of the American College of Cardiology, 2017, 69, 655.	1.2	0
89	Hypertensive Response With Exercise to Reveal Increased Cardiovascular Risk in Adults With Aortic Coarctation Repair: Value and Caution. Canadian Journal of Cardiology, 2018, 34, 536-539.	0.8	0
90	Looking to the Left to Get It Right: Left Ventricular Systolic Dysfunction and Risk Stratification Late After Tetralogy of Fallot Repair. Canadian Journal of Cardiology, 2019, 35, 1623-1625.	0.8	0

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91	Intraoperative Transesophageal Echocardiography in a Rare Case of Dextrocardia, Situs Inversus Totalis, and Double-Outlet Right Ventricle. <i>A&A Practice</i> , 2020, 14, 123-126.	0.2	0
92	Biventricular diverticula in a patient with restrictive cardiomyopathy. <i>British Heart Journal</i> , 2000, 84, 333-334.	2.2	0
93	The "Forgotten Ones": The Natural and Unnatural History of Univentricular Physiology Without Fontan Palliation. <i>Canadian Journal of Cardiology</i> , 2022, 38, 858-861.	0.8	0
94	Standing on the Shoulders of a Giant: Dr Gary Douglas Webb, 1943-2021. <i>Canadian Journal of Cardiology</i> , 2022, 38, 852-854.	0.8	0