Robin Condliffe

List of Publications by Year
in descending order

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Version: 2024-02-01

97	7,048	109321	60623
papers	citations	35 h-index	81 g-index

98	98	98	7029
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Computed tomography lung parenchymal descriptions in routine radiological reporting have diagnostic and prognostic utility in patients with idiopathic pulmonary arterial hypertension and pulmonary hypertension associated with lung disease. ERJ Open Research, 2022, 8, 00549-2021.	2.6	7
2	Comment on â€œExternal validation of the OPALS prediction model for in-hospital mortality in patients with acute decompensated pulmonary hypertensionâ€• ERJ Open Research, 2022, 8, 00066-2022.	2.6	0
3	Elective lower limb orthopedic arthroplasty surgery in patients with pulmonary hypertension. Pulmonary Circulation, 2022, 12, e12019.	1.7	2
4	CMR Measures of Left Atrial Volume Index and Right Ventricular Function Have Prognostic Value in Chronic Thromboembolic Pulmonary Hypertension. Frontiers in Medicine, 2022, 9, 840196.	2.6	2
5	Imaging and Risk Stratification in Pulmonary Arterial Hypertension: Time to Include Right Ventricular Assessment. Frontiers in Cardiovascular Medicine, 2022, 9, 797561.	2.4	7
6	Training and clinical testing of artificial intelligence derived right atrial cardiovascular magnetic resonance measurements. Journal of Cardiovascular Magnetic Resonance, 2022, 24, 25.	3.3	8
7	Severe pulmonary hypertension associated with lung disease is characterised by a loss of small pulmonary vessels on quantitative computed tomography. ERJ Open Research, 2022, 8, 00503-2021.	2.6	10
8	Examining the impact of pulmonary hypertension on nonprofessional caregivers: A mixedâ€methods systematic review. Pulmonary Circulation, 2022, 12, e12077.	1.7	3
9	Machine learning cardiac-MRI features predict mortality in newly diagnosed pulmonary arterial hypertension. European Heart Journal Digital Health, 2022, 3, 265-275.	1.7	11
10	Phenotyping of idiopathic pulmonary arterial hypertension: a registry analysis. Lancet Respiratory Medicine,the, 2022, 10, 937-948.	10.7	57
11	Bayesian Inference Associates Rare KDR Variants With Specific Phenotypes in Pulmonary Arterial Hypertension. Circulation Genomic and Precision Medicine, 2021, 14, .	3.6	29
12	EmPHasis-10 health-related quality of life score predicts outcomes in patients with idiopathic and connective tissue disease-associated pulmonary arterial hypertension: results from a UK multicentre study. European Respiratory Journal, 2021, 57, 2000124.	6.7	29
13	Cardiac-MRI Predicts Clinical Worsening and Mortality in Pulmonary Arterial Hypertension. JACC: Cardiovascular Imaging, 2021, 14, 931-942.	5.3	73
14	Right Ventricular Adaptation Assessed Using Cardiac Magnetic Resonance Predicts Survival in Pulmonary Arterial Hypertension. JACC: Cardiovascular Imaging, 2021, 14, 1271-1272.	5.3	11
15	Maximal Exercise Testing Using the Incremental Shuttle Walking Test Can Be Used to Risk-Stratify Patients with Pulmonary Arterial Hypertension. Annals of the American Thoracic Society, 2021, 18, 34-43.	3.2	13
16	Critical care outcomes in patients with pre-existing pulmonary hypertension: insights from the ASPIRE registry. ERJ Open Research, 2021, 7, 00046-2021.	2.6	15
17	Repeatability and sensitivity to change of non-invasive end points in PAH: the RESPIRE study. Thorax, 2021, 76, 1032-1035.	5.6	13
18	Palliative care in pulmonary hypertension associated with congenital heart disease: systematic review and expert opinion. ESC Heart Failure, 2021, 8, 1901-1914.	3.1	9

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19	Homozygous GDF2 nonsense mutations result in a loss of circulating BMP9 and BMP10 and are associated with either PAH or an â€œHHTâ€likeâ€•syndrome in children. Molecular Genetics & Genomic Medicine, 2021, 9, e1685.	1.2	19
20	Perioperative management of patients with pulmonary hypertension undergoing non-cardiothoracic, non-obstetric surgery: a systematic review and expert consensus statement. British Journal of Anaesthesia, 2021, 126, 774-790.	3.4	45
21	Pulmonary Hypertension in Association with Lung Disease: Quantitative CT and Artificial Intelligence to the Rescue? State-of-the-Art Review. Diagnostics, 2021, 11, 679.	2.6	15
22	Supplementation with Iron in Pulmonary Arterial Hypertension. Two Randomized Crossover Trials. Annals of the American Thoracic Society, 2021, 18, 981-988.	3.2	28
23	Establishing expert consensus for the optimal approach to holistic risk-management in pulmonary arterial hypertension: a Delphi process and narrative review. Expert Review of Respiratory Medicine, 2021, 15, 1493-1503.	2.5	0
24	A diagnostic miRNA signature for pulmonary arterial hypertension using a consensus machine learning approach. EBioMedicine, 2021, 69, 103444.	6.1	30
25	Pulmonary hypertension phenotypes in patients with systemic sclerosis. European Respiratory Review, 2021, 30, 210053.	7.1	27
26	Assessing pulmonary hypertension severity in lung disease is a key step to improving outcomes: embrace resistance and don't be pressurised to go with the flow. European Respiratory Journal, 2021, 58, 2102008.	6.7	6
27	Pulmonary arterial hypertension in adults with congenital heart disease: markers of disease severity, management of advanced heart failure and transplantation. Expert Review of Cardiovascular Therapy, 2021, 19, 837-855.	1.5	2
28	Characterization of GDF2 Mutations and Levels of BMP9 and BMP10 in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 575-585.	5.6	80
29	Identification of Cardiac Magnetic Resonance Imaging Thresholds for Risk Stratification in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 458-468.	5.6	99
30	Mild parenchymal lung disease is still lung disease. European Respiratory Journal, 2020, 56, 2003727.	6.7	3
31	Respiratory follow-up of patients with COVID-19 pneumonia. Thorax, 2020, 75, 1009-1016.	5.6	266
32	Mild parenchymal lung disease and/or low diffusion capacity impacts survival and treatment response in patients diagnosed with idiopathic pulmonary arterial hypertension. European Respiratory Journal, 2020, 55, 2000041.	6.7	48
33	BNP/NT-proBNP in pulmonary arterial hypertension: time for point-of-care testing?. European Respiratory Review, 2020, 29, 200009.	7.1	51
34	Diagnostic accuracy of CT pulmonary angiography in suspected pulmonary hypertension. European Radiology, 2020, 30, 4918-4929.	4.5	29
35	Partial anomalous pulmonary venous drainage in patients presenting with suspected pulmonary hypertension: A series of 90 patients from the ASPIRE registry. Respirology, 2020, 25, 1066-1072.	2.3	10
36	Idiopathic pulmonary arterial hypertension and coâ€existing lung disease: is this a new phenotype?. Pulmonary Circulation, 2020, 10, 1-8.	1.7	16

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37	Adultsâ€™ experiences of living with pulmonary hypertension: a thematic synthesis of qualitative studies. BMJ Open, 2020, 10, e041428.	1.9	10
38	Mildly increased pulmonary arterial pressure: a new disease entity or just a marker of poor prognosis?. European Journal of Heart Failure, 2019, 21, 1057-1061.	7.1	11
39	Discovery of Distinct Immune Phenotypes Using Machine Learning in Pulmonary Arterial Hypertension. Circulation Research, 2019, 124, 904-919.	4.5	141
40	The incremental shuttle walk test predicts mortality in nonâ€group 1 pulmonary hypertension: results from the ASPIRE Registry. Pulmonary Circulation, 2019, 9, 1-9.	1.7	7
41	Traffic exposures, air pollution and outcomes in pulmonary arterial hypertension: a UK cohort study analysis. European Respiratory Journal, 2019, 53, 1801429.	6.7	31
42	Congenital heart disease, pulmonary arterial hypertension and the UKâ€™s Drivers and Vehicle Licensing Agency: controversial new guidance. Pulmonary Circulation, 2019, 9, 1-2.	1.7	0
43	Genetic determinants of risk in pulmonary arterial hypertension: international genome-wide association studies and meta-analysis. Lancet Respiratory Medicine,the, 2019, 7, 227-238.	10.7	122
44	Decision-making in pulmonary endarterectomy surgery. European Respiratory Journal, 2019, 53, 1801973.	6.7	3
45	Diagnosis of Pulmonary Hypertension with Cardiac MRI: Derivation and Validation of Regression Models. Radiology, 2019, 290, 61-68.	7.3	43
46	Diagnostic and prognostic value of a diagnostic CT regression model in suspected pulmonary hypertension. , 2019, , .		1
47	Identification of rare sequence variation underlying heritable pulmonary arterial hypertension. Nature Communications, 2018, 9, 1416.	12.8	279
48	Management of Adults With Congenital Heart Disease and Pulmonary Arterial Hypertension in the UK: Survey of Current Practice, Unmet Needs and Expert Commentary. Heart Lung and Circulation, 2018, 27, 1018-1027.	0.4	7
49	Echocardiographic Screening for Pulmonary Hypertension in CongenitalÂHeart Disease. Journal of the American College of Cardiology, 2018, 72, 2778-2788.	2.8	38
50	Identifying early pulmonary arterial hypertension in patients with systemic sclerosis. European Respiratory Journal, 2018, 51, 1800495.	6.7	6
51	Identifying At-Risk Patients with Combined Pre- and Postcapillary Pulmonary Hypertension Using Interventricular Septal Angle at Cardiac MRI. Radiology, 2018, 289, 61-68.	7.3	27
52	Pulmonary Artery Size in Interstitial Lung Disease and Pulmonary Hypertension: Association with Interstitial Lung Disease Severity and Diagnostic Utility. Frontiers in Cardiovascular Medicine, 2018, 5, 53.	2.4	29
53	Incremental Shuttle Walking Test Distance Is Reduced in Patients With Pulmonary Hypertension in World Health Organisation Functional Class I. Frontiers in Medicine, 2018, 5, 172.	2.6	4
54	Pathophysiology and Diagnosis of Pulmonary Hypertension Due to Left Heart Disease. Frontiers in Medicine, 2018, 5, 174.	2.6	20

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55	Circulating Protein Biomarkers in Systemic Sclerosis Related Pulmonary Arterial Hypertension: A Review of Published Data. Frontiers in Medicine, 2018, 5, 175.	2.6	19
56	The impact of patient choice on survival in chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2018, 52, 1800589.	6.7	87
57	British Thoracic Society Guideline for the initial outpatient management of pulmonary embolism (PE). Thorax, 2018, 73, ii1-ii29.	5.6	58
58	Idiopathic and Systemic Sclerosis-Associated Pulmonary Arterial Hypertension. Chest, 2017, 152, 92-102.	0.8	53
59	Survival in portopulmonary hypertension: Outcomes of the United Kingdom National Pulmonary Arterial Hypertension Registry. Journal of Heart and Lung Transplantation, 2017, 36, 770-779.	0.6	73
60	Incremental shuttle walk test distance and autonomic dysfunction predict survival in pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2017, 36, 871-879.	0.6	16
61	Plasma proteome analysis in patients with pulmonary arterial hypertension: an observational cohort study. Lancet Respiratory Medicine,the, 2017, 5, 717-726.	10.7	99
62	Magnetic Resonance Imaging in the Prognostic Evaluation of Patients with Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2017, 196, 228-239.	5.6	122
63	Phenotypic Characterization of EIF2AK4 Mutation Carriers in a Large Cohort of Patients Diagnosed Clinically With Pulmonary Arterial Hypertension. Circulation, 2017, 136, 2022-2033.	1.6	111
64	The CRASH report: emergency management dilemmas facing acute physicians in patients with pulmonary arterial hypertension. Thorax, 2017, 72, 1035-1045.	5.6	30
65	Unmasking hidden disease: exercise pulmonary haemodynamics in systemic sclerosis. European Respiratory Journal, 2017, 50, 1700885.	6.7	6
66	British Thoracic Society Clinical Statement on Pulmonary Arteriovenous Malformations. Thorax, 2017, 72, 1154-1163.	5.6	94
67	Pulmonary arteriovenous malformations emerge from the shadows. Thorax, 2017, 72, 1071-1073.	5.6	7
68	The use of Macitentan in Fontan circulation: a case report. BMC Cardiovascular Disorders, 2017, 17, 131.	1.7	6
69	Longâ€term outcomes of domiciliary intravenous iloprost in idiopathic and connective tissue diseaseâ€associated pulmonary arterial hypertension. Respirology, 2017, 22, 372-377.	2.3	12
70	An official European Respiratory Society statement: pulmonary haemodynamics during exercise. European Respiratory Journal, 2017, 50, 1700578.	6.7	222
71	Combining creative writing and narrative analysis to deliver new insights into the impact of pulmonary hypertension. BMJ Open Respiratory Research, 2017, 4, e000184.	3.0	4
72	Dynamic Risk Stratification of Patient Long-Term Outcome After Pulmonary Endarterectomy. Circulation, 2016, 133, 1761-1771.	1.6	307

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73	Pulmonary Hypertension in Patients with Heart Failure and Preserved Ejection Fraction: Differential Diagnosis and Management. Pulmonary Circulation, 2016, 6, 3-14.	1.7	20
74	Effect of dual pulmonary vasodilator therapy in pulmonary arterial hypertension associated with congenital heart disease: a retrospective analysis. Open Heart, 2016, 3, e000399.	2.3	7
75	Pulmonary arterial hypertension associated with congenital heart disease: Comparison of clinical and anatomicâ€“pathophysiologic classification. Journal of Heart and Lung Transplantation, 2016, 35, 610-618.	0.6	21
76	Connective tissue disease-associated pulmonary arterial hypertension. F1000prime Reports, 2015, 7, 06.	5.9	41
77	Elevated Plasma CXCL12Î± Is Associated with a Poorer Prognosis in Pulmonary Arterial Hypertension. PLoS ONE, 2015, 10, e0123709.	2.5	27
78	Pulmonary Artery Denervation Reduces Pulmonary Artery Pressure and Induces Histological Changes in an Acute Porcine Model of Pulmonary Hypertension. Circulation: Cardiovascular Interventions, 2015, 8, e002569.	3.9	66
79	Management of acute pulmonary embolism. British Journal of Hospital Medicine (London, England:) Tj ETQq1 1 0.784314 rgBT /Overl	0.5	2
80	Experimental validation of the hyperpolarized ¹²⁹Xe chemical shift saturation recovery technique in healthy volunteers and subjects with interstitial lung disease. Magnetic Resonance in Medicine, 2015, 74, 196-207.	3.0	76
81	Right Ventricular Sex Differences in Patients with Idiopathic Pulmonary Arterial Hypertension Characterised by Magnetic Resonance Imaging: Pair-Matched Case Controlled Study. PLoS ONE, 2015, 10, e0127415.	2.5	33
82	Ambrisentan therapy in pulmonary hypertension: clinical use and tolerability in a referral centre. Therapeutic Advances in Respiratory Disease, 2014, 8, 71-77.	2.6	13
83	LGE Patterns in PulmonaryÂHypertension Do Not ImpactÂOverall Mortality. JACC: Cardiovascular Imaging, 2014, 7, 1209-1217.	5.3	82
84	Management dilemmas in acute pulmonary embolism. Thorax, 2014, 69, 174-180.	5.6	60
85	Definitions and Diagnosis of Pulmonary Hypertension. Journal of the American College of Cardiology, 2013, 62, D42-D50.	2.8	1,467
86	Pregnancy and pulmonary hypertension: a practical approach to management. Obstetric Medicine, 2013, 6, 144-154.	1.1	36
87	Noninvasive Estimation of PA Pressure, Flow, andÂResistance With CMR Imaging. JACC: Cardiovascular Imaging, 2013, 6, 1036-1047.	5.3	129
88	Pulmonary hypertension: diagnosis and management. BMJ, The, 2013, 346, f2028-f2028.	6.0	119
89	Pulmonary hypertension in COPD: results from the ASPIRE registry. European Respiratory Journal, 2013, 41, 1292-1301.	6.7	173
90	Serum Osteoprotegerin is Increased and Predicts Survival in Idiopathic Pulmonary Arterial Hypertension. Pulmonary Circulation, 2012, 2, 21-27.	1.7	24

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91	Changing Demographics, Epidemiology, and Survival of Incident Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2012, 186, 790-796.	5.6	483
92	Survival in Pulmonary Hypertension Registries. Chest, 2011, 139, 1547-1548.	0.8	1
93	CT pulmonary angiography combined with echocardiography in suspected systemic sclerosis-associated pulmonary arterial hypertension. Rheumatology, 2011, 50, 1480-1486.	1.9	32
94	Connective Tissue Diseaseâ€“associated Pulmonary Arterial Hypertension in the Modern Treatment Era. American Journal of Respiratory and Critical Care Medicine, 2009, 179, 151-157.	5.6	576
95	Improved Outcomes in Medically and Surgically Treated Chronic Thromboembolic Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2008, 177, 1122-1127.	5.6	379
96	Management of Suspected Chronic Thromboembolic Pulmonary Hypertension. , 0, , 405-420.		1
97	Unenhanced computed tomography as a diagnostic tool in suspected pulmonary hypertension: a retrospective cross-sectional pilot study. Wellcome Open Research, 0, 6, 249.	1.8	2