Robin Condliffe

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

88 4,659 28 68 g-index

98 5,986 7.6 5.07 ext. papers ext. citations avg, IF L-index

#	Paper	IF	Citations
88	Definitions and diagnosis of pulmonary hypertension. <i>Journal of the American College of Cardiology</i> , 2013 , 62, D42-50	15.1	1163
87	Connective tissue disease-associated pulmonary arterial hypertension in the modern treatment era. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2009 , 179, 151-7	10.2	461
86	Changing demographics, epidemiology, and survival of incident pulmonary arterial hypertension: results from the pulmonary hypertension registry of the United Kingdom and Ireland. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2012 , 186, 790-6	10.2	370
85	Improved outcomes in medically and surgically treated chronic thromboembolic pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2008 , 177, 1122-7	10.2	304
84	Dynamic Risk Stratification of Patient Long-Term Outcome After Pulmonary Endarterectomy: Results From the United Kingdom National Cohort. <i>Circulation</i> , 2016 , 133, 1761-71	16.7	203
83	Identification of rare sequence variation underlying heritable pulmonary arterial hypertension. <i>Nature Communications</i> , 2018 , 9, 1416	17.4	182
82	Respiratory follow-up of patients with COVID-19 pneumonia. <i>Thorax</i> , 2020 , 75, 1009-1016	7.3	139
81	An official European Respiratory Society statement: pulmonary haemodynamics during exercise. <i>European Respiratory Journal</i> , 2017 , 50,	13.6	124
80	Pulmonary hypertension in COPD: results from the ASPIRE registry. <i>European Respiratory Journal</i> , 2013 , 41, 1292-301	13.6	117
79	Noninvasive estimation of PA pressure, flow, and resistance with CMR imaging: derivation and prospective validation study from the ASPIRE registry. <i>JACC: Cardiovascular Imaging</i> , 2013 , 6, 1036-104	7 ^{8.4}	104
78	Pulmonary hypertension: diagnosis and management. <i>BMJ, The</i> , 2013 , 346, f2028	5.9	92
77	Discovery of Distinct Immune Phenotypes Using Machine Learning in Pulmonary Arterial Hypertension. <i>Circulation Research</i> , 2019 , 124, 904-919	15.7	81
76	Magnetic Resonance Imaging in the Prognostic Evaluation of Patients with Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2017 , 196, 228-239	10.2	79
75	Phenotypic Characterization of Mutation Carriers in a Large Cohort of Patients Diagnosed Clinically With Pulmonary Arterial Hypertension. <i>Circulation</i> , 2017 , 136, 2022-2033	16.7	75
74	Plasma proteome analysis in patients with pulmonary arterial hypertension: an observational cohort study. <i>Lancet Respiratory Medicine,the</i> , 2017 , 5, 717-726	35.1	62
73	LGE patterns in pulmonary hypertension do not impact overall mortality. <i>JACC: Cardiovascular Imaging</i> , 2014 , 7, 1209-17	8.4	62
72	British Thoracic Society Clinical Statement on Pulmonary Arteriovenous Malformations. <i>Thorax</i> , 2017 , 72, 1154-1163	7-3	61

(2021-2015)

healthy volunteers and subjects with interstitial lung disease. <i>Magnetic Resonance in Medicine</i> , 2015 , 74, 196-207	4.4	57
Genetic determinants of risk in pulmonary arterial hypertension: international genome-wide association studies and meta-analysis. <i>Lancet Respiratory Medicine,the</i> , 2019 , 7, 227-238	35.1	55
Survival in portopulmonary hypertension: Outcomes of the United Kingdom National Pulmonary Arterial Hypertension Registry. <i>Journal of Heart and Lung Transplantation</i> , 2017 , 36, 770-779	5.8	47
Management dilemmas in acute pulmonary embolism. <i>Thorax</i> , 2014 , 69, 174-80	7.3	47
Pulmonary artery denervation reduces pulmonary artery pressure and induces histological changes in an acute porcine model of pulmonary hypertension. <i>Circulation: Cardiovascular Interventions</i> , 2015 , 8, e002569	6	46
Characterization of Mutations and Levels of BMP9 and BMP10 in Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020 , 201, 575-585	10.2	46
British Thoracic Society Guideline for the initial outpatient management of pulmonary embolism (PE). <i>Thorax</i> , 2018 , 73, ii1-ii29	7.3	43
The impact of patient choice on survival in chronic thromboembolic pulmonary hypertension. <i>European Respiratory Journal</i> , 2018 , 52,	13.6	41
Identification of Cardiac Magnetic Resonance Imaging Thresholds for Risk Stratification in Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020 , 201, 458-468	10.2	37
Connective tissue disease-associated pulmonary arterial hypertension. <i>F1000prime Reports</i> , 2015 , 7, 06		32
Idiopathic and Systemic Sclerosis-Associated Pulmonary Arterial Hypertension: A Comparison of Demographic, Hemodynamic, and MRI Characteristics and Outcomes. <i>Chest</i> , 2017 , 152, 92-102	5.3	28
CT pulmonary angiography combined with echocardiography in suspected systemic sclerosis-associated pulmonary arterial hypertension. <i>Rheumatology</i> , 2011 , 50, 1480-6	3.9	27
Diagnosis of Pulmonary Hypertension with Cardiac MRI: Derivation and Validation of Regression Models. <i>Radiology</i> , 2019 , 290, 61-68	20.5	26
Right ventricular sex differences in patients with idiopathic pulmonary arterial hypertension characterised by magnetic resonance imaging: pair-matched case controlled study. <i>PLoS ONE</i> , 2015 , 10, e0127415	3.7	24
Echocardiographic Screening for Pulmonary Hypertension in Congenital[Heart Disease: JACC Review Topic of the Week. <i>Journal of the American College of Cardiology</i> , 2018 , 72, 2778-2788	15.1	22
The CRASH report: emergency management dilemmas facing acute physicians in patients with pulmonary arterial hypertension. <i>Thorax</i> , 2017 , 72, 1035-1045	7.3	20
Serum osteoprotegerin is increased and predicts survival in idiopathic pulmonary arterial hypertension. <i>Pulmonary Circulation</i> , 2012 , 2, 21-7	2.7	20
Cardiac-MRI Predicts Clinical Worsening and Mortality in Pulmonary Arterial Hypertension: A Systematic Review and Meta-Analysis. <i>JACC: Cardiovascular Imaging</i> , 2021 , 14, 931-942	8.4	20
	Genetic determinants of risk in pulmonary arterial hypertension: international genome-wide association studies and meta-analysis. <i>Lancet Respiratory Medicine, the,</i> 2019, 7, 227-238 Survival in portopulmonary hypertension: Outcomes of the United Kingdom National Pulmonary Arterial Hypertension Registry. <i>Journal of Heart and Lung Transplantation,</i> 2017, 36, 770-779 Management dilemmas in acute pulmonary embolism. <i>Thorax,</i> 2014, 69, 174-80 Pulmonary artery denervation reduces pulmonary artery pressure and induces histological changes in an acute porcine model of pulmonary hypertension. <i>Circulation: Cardiovascular Interventions,</i> 2015, 8, e002569 Characterization of Mutations and Levels of BMP9 and BMP10 in Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine,</i> 2020, 201, 575-585 British Thoracic Society Guideline for the initial outpatient management of pulmonary embolism (PE). <i>Thorax,</i> 2018, 73, iii-iii29 The impact of patient choice on survival in chronic thromboembolic pulmonary hypertension. <i>European Respiratory Journal,</i> 2018, 52, ldentification of Cardiac Magnetic Resonance Imaging Thresholds for Risk Stratification in Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine,</i> 2020, 201, 458-468 Connective tissue disease-associated pulmonary arterial hypertension: A Comparison of Demographic, Hemodynamic, and MRI Characteristics and Outcomes. <i>Chest,</i> 2017, 152, 92-102 CT pulmonary angiography combined with echocardiography in suspected systemic sclerosis-associated pulmonary arterial hypertension: A Comparison of Demographic, Hemodynamic, and MRI Characteristics and Outcomes. <i>Chest,</i> 2017, 152, 92-102 CT pulmonary angiography combined with echocardiography in suspected systemic sclerosis-associated pulmonary arterial hypertension in Congenitall Heart Disease: JACC Review Topic of the Week. <i>Journal of the American College of Cardiology,</i> 2011, 50, 1480-6 Diagnosis of Pulmonary Hypertension with Cardiac MRI: Derivat	healthy volunteers and subjects with interstitial lung disease. Magnetic Resonance in Medicine, 2015, 74, 196-207 Genetic determinants of risk in pulmonary arterial hypertension: international genome-wide association studies and meta-analysis. Lancet Respiratory Medicine, the, 2019, 7, 227-238 Survival in portopulmonary hypertension: Outcomes of the United Kingdom National Pulmonary Arterial Hypertension Registry. Journal of Heart and Lung Transplantation, 2017, 36, 770-779 Management dilemmas in acute pulmonary embolism. Thorax, 2014, 69, 174-80 Pulmonary artery denervation reduces pulmonary artery pressure and induces histological changes in an acute portine model of pulmonary hypertension. Circulation: Cardiovascular Interventions, 2015, 8, e002569 Characterization of Mutations and Levels of BMP9 and BMP10 in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 575-585 British Thoracic Society Guideline for the initial outpatient management of pulmonary embolism (PE). Thorax, 2018, 73, ii1-ii29 The impact of patient choice on survival in chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2018, 52, 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 Connective tissue disease-associated pulmonary arterial hypertension. F1000prime Reports, 2015, 7, 06 Connective tissue disease-associated pulmonary arterial hypertension. A Comparison of Demographic, Hemodynamic, and MRI Characteristics and Outcomes. Chest, 2017, 152, 92-102 The pulmonary angiography combined with echocardiography in suspected systemic sclerosis-associated pulmonary arterial hypertension. Rheumatology, 2011, 50, 1480-6 Diagnosis of Pulmonary Hypertension with Cardiac MRI: Derivation and Validation of Regression Models. Radiology, 2019, 290, 61-68 Right ventricular set differences in patients with idiopathic pu

53	Elevated plasma CXCL12lis associated with a poorer prognosis in pulmonary arterial hypertension. <i>PLoS ONE</i> , 2015 , 10, e0123709	3.7	19
52	Pulmonary Artery Size in Interstitial Lung Disease and Pulmonary Hypertension: Association with Interstitial Lung Disease Severity and Diagnostic Utility. <i>Frontiers in Cardiovascular Medicine</i> , 2018 , 5, 53	5.4	18
51	Traffic exposures, air pollution and outcomes in pulmonary arterial hypertension: a UK cohort study analysis. <i>European Respiratory Journal</i> , 2019 , 53,	13.6	17
50	Mild parenchymal lung disease and/or low diffusion capacity impacts survival and treatment response in patients diagnosed with idiopathic pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2020 , 55,	13.6	17
49	Pregnancy and pulmonary hypertension: a practical approach to management. <i>Obstetric Medicine</i> , 2013 , 6, 144-54	1.5	17
48	Pulmonary arterial hypertension associated with congenital heart disease: Comparison of clinical and anatomic-pathophysiologic classification. <i>Journal of Heart and Lung Transplantation</i> , 2016 , 35, 610-6	8 ^{5.8}	16
47	BNP/NT-proBNP in pulmonary arterial hypertension: time for point-of-care testing?. <i>European Respiratory Review</i> , 2020 , 29,	9.8	15
46	Pulmonary hypertension in patients with heart failure and preserved ejection fraction: differential diagnosis and management. <i>Pulmonary Circulation</i> , 2016 , 6, 3-14	2.7	15
45	Identifying At-Risk Patients with Combined Pre- and Postcapillary Pulmonary Hypertension Using Interventricular Septal Angle at Cardiac MRI. <i>Radiology</i> , 2018 , 289, 61-68	20.5	14
44	Incremental shuttle walk test distance and autonomic dysfunction predict survival in pulmonary arterial hypertension. <i>Journal of Heart and Lung Transplantation</i> , 2017 , 36, 871-879	5.8	13
43	Pathophysiology and Diagnosis of Pulmonary Hypertension Due to Left Heart Disease. <i>Frontiers in Medicine</i> , 2018 , 5, 174	4.9	13
42	Long-term outcomes of domiciliary intravenous iloprost in idiopathic and connective tissue disease-associated pulmonary arterial hypertension. <i>Respirology</i> , 2017 , 22, 372-377	3.6	12
41	Ambrisentan therapy in pulmonary hypertension: clinical use and tolerability in a referral centre. <i>Therapeutic Advances in Respiratory Disease</i> , 2014 , 8, 71-77	4.9	11
40	Circulating Protein Biomarkers in Systemic Sclerosis Related Pulmonary Arterial Hypertension: A Review of Published Data. <i>Frontiers in Medicine</i> , 2018 , 5, 175	4.9	9
39	Bayesian Inference Associates Rare Variants with Specific Phenotypes in Pulmonary Arterial Hypertension. <i>Circulation Genomic and Precision Medicine</i> , 2020 ,	5.2	9
38	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,	13.6	9
37	Idiopathic pulmonary arterial hypertension and co-existing lung disease: is this a new phenotype?. <i>Pulmonary Circulation</i> , 2020 , 10, 2045894020914851	2.7	8
36	Homozygous GDF2 nonsense mutations result in a loss of circulating BMP9 and BMP10 and are associated with either PAH or an "HHT-like" syndrome in children. <i>Molecular Genetics & amp; Genomic Medicine</i> , 2021 , e1685	2.3	7

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35	Mildly increased pulmonary arterial pressure: a new disease entity or just a marker of poor prognosis?. <i>European Journal of Heart Failure</i> , 2019 , 21, 1057-1061	12.3	6
34	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	5.4	6
33	Management of Adults With Congenital Heart Disease and Pulmonary Arterial Hypertension in the UK: Survey of Current Practice, Unmet Needs and Expert Commentary. <i>Heart Lung and Circulation</i> , 2018 , 27, 1018-1027	1.8	6
32	Effect of dual pulmonary vasodilator therapy in pulmonary arterial hypertension associated with congenital heart disease: a retrospective analysis. <i>Open Heart</i> , 2016 , 3, e000399	3	5
31	AdultsRexperiences of living with pulmonary hypertension: a thematic synthesis of qualitative studies. <i>BMJ Open</i> , 2020 , 10, e041428	3	5
30	A diagnostic miRNA signature for pulmonary arterial hypertension using a consensus machine learning approach. <i>EBioMedicine</i> , 2021 , 69, 103444	8.8	5
29	Pulmonary hypertension phenotypes in patients with systemic sclerosis. <i>European Respiratory Review</i> , 2021 , 30,	9.8	5
28	The incremental shuttle walk test predicts mortality in non-group 1 pulmonary hypertension: results from the ASPIRE Registry. <i>Pulmonary Circulation</i> , 2019 , 9, 2045894019848649	2.7	4
27	The use of Macitentan in Fontan circulation: a case report. <i>BMC Cardiovascular Disorders</i> , 2017 , 17, 131	2.3	4
26	Critical care outcomes in patients with pre-existing pulmonary hypertension: insights from the ASPIRE registry. <i>ERJ Open Research</i> , 2021 , 7,	3.5	4
25	Partial anomalous pulmonary venous drainage in patients presenting with suspected pulmonary hypertension: A series of 90 patients from the ASPIRE registry. <i>Respirology</i> , 2020 , 25, 1066-1072	3.6	3
24	Incremental Shuttle Walking Test Distance Is Reduced in Patients With Pulmonary Hypertension in World Health Organisation Functional Class I. <i>Frontiers in Medicine</i> , 2018 , 5, 172	4.9	3
23	Combining creative writing and narrative analysis to deliver new insights into the impact of pulmonary hypertension. <i>BMJ Open Respiratory Research</i> , 2017 , 4, e000184	5.6	3
22	Palliative care in pulmonary hypertension associated with congenital heart disease: systematic review and expert opinion. <i>ESC Heart Failure</i> , 2021 , 8, 1901-1914	3.7	3
21	Supplementation with Iron in Pulmonary Arterial Hypertension. Two Randomized Crossover Trials. <i>Annals of the American Thoracic Society</i> , 2021 , 18, 981-988	4.7	3
20	Decision-making in pulmonary endarterectomy surgery. European Respiratory Journal, 2019, 53,	13.6	3
19	Right Ventricular Adaptation Assessed Using Cardiac Magnetic Resonance Predicts Survival in Pulmonary Arterial Hypertension. <i>JACC: Cardiovascular Imaging</i> , 2021 , 14, 1271-1272	8.4	3
18	Management of acute pulmonary embolism. <i>British Journal of Hospital Medicine (London, England: 2005)</i> , 2015 , 76, C150-5	0.8	2

17	Diagnostic accuracy of CT pulmonary angiography in suspected pulmonary hypertension. <i>European Radiology</i> , 2020 , 30, 4918-4929	8	2
16	Pulmonary Hypertension in Association with Lung Disease: Quantitative CT and Artificial Intelligence to the Rescue? State-of-the-Art Review. <i>Diagnostics</i> , 2021 , 11,	3.8	2
15	Repeatability and sensitivity to change of non-invasive end points in PAH: the RESPIRE study. <i>Thorax</i> , 2021 , 76, 1032-1035	7.3	2
14	Survival in pulmonary hypertension registries: the importance of incident cases. <i>Chest</i> , 2011 , 139, 1547-	1 5,4 8	1
13	Management of Suspected Chronic Thromboembolic Pulmonary Hypertension405-420		1
12	Mild parenchymal lung disease is still lung disease. <i>European Respiratory Journal</i> , 2020 , 56,	13.6	1
11	Maximal Exercise Testing Using the Incremental Shuttle Walking Test Can Be Used to Risk-Stratify Patients with Pulmonary Arterial Hypertension. <i>Annals of the American Thoracic Society</i> , 2021 , 18, 34-43	4.7	1
10	Pulmonary arterial hypertension in adults with congenital heart disease: markers of disease severity, management of advanced heart failure and transplantation. <i>Expert Review of Cardiovascular Therapy</i> , 2021 , 19, 837-855	2.5	1
9	Imaging and Risk Stratification in Pulmonary Arterial Hypertension: Time to Include Right Ventricular Assessment <i>Frontiers in Cardiovascular Medicine</i> , 2022 , 9, 797561	5.4	1
8	Training and clinical testing of artificial intelligence derived right atrial cardiovascular magnetic resonance measurements <i>Journal of Cardiovascular Magnetic Resonance</i> , 2022 , 24, 25	6.9	1
7	Examining the impact of pulmonary hypertension on nonprofessional caregivers: A mixed-methods systematic review <i>Pulmonary Circulation</i> , 2022 , 12, e12077	2.7	1
6	Unenhanced computed tomography as a diagnostic tool in suspected pulmonary hypertension: a retrospective cross-sectional pilot study. <i>Wellcome Open Research</i> ,6, 249	4.8	0
5	CMR Measures of Left Atrial Volume Index and Right Ventricular Function Have Prognostic Value in Chronic Thromboembolic Pulmonary Hypertension <i>Frontiers in Medicine</i> , 2022 , 9, 840196	4.9	0
4	Reply: External validation of the OPALS prediction model for in-hospital mortality in patients with acute decompensated pulmonary hypertension <i>ERJ Open Research</i> , 2022 , 8,	3.5	
3	Establishing expert consensus for the optimal approach to holistic risk-management in pulmonary arterial hypertension: a Delphi process and narrative review. <i>Expert Review of Respiratory Medicine</i> , 2021 , 15, 1493-1503	3.8	
2	Congenital heart disease, pulmonary arterial hypertension and the UKB Drivers and Vehicle Licensing Agency: controversial new guidance. <i>Pulmonary Circulation</i> , 2019 , 9, 2045894019882627	2.7	
1	Elective lower limb orthopedic arthroplasty surgery in patients with pulmonary hypertension <i>Pulmonary Circulation</i> , 2022 , 12, e12019	2.7	