Ardeschir Ghofrani

List of Publications by Citations

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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.

460 papers

40,487 citations

87 h-index

191 g-index

580 ext. papers

48,601 ext. citations

8.4 avg, IF

6.91 L-index

#	Paper	IF	Citations
460	2015 ESC/ERS Guidelines for the diagnosis and treatment of pulmonary hypertension: The Joint Task Force for the Diagnosis and Treatment of Pulmonary Hypertension of the European Society of Cardiology (ESC) and the European Respiratory Society (ERS): Endorsed by: Association for	9.5	3455
459	Updated clinical classification of pulmonary hypertension. <i>Journal of the American College of Cardiology</i> , 2013 , 62, D34-41	15.1	1937
458	Sildenafil citrate therapy for pulmonary arterial hypertension. <i>New England Journal of Medicine</i> , 2005 , 353, 2148-57	59.2	1843
457	2015 ESC/ERS Guidelines for the diagnosis and treatment of pulmonary hypertension: The Joint Task Force for the Diagnosis and Treatment of Pulmonary Hypertension of the European Society of Cardiology (ESC) and the European Respiratory Society (ERS): Endorsed by: Association for European Paediatric and Congenital Cardiology (AEPC), International Society for Heart and Lung	13.6	1672
456	Transplantation (ISHLT). Furnean Respiratory, Journal 2015, 46, 903-75. Inhaled iloprost for severe pulmonary hypertension. New England Journal of Medicine, 2002, 347, 322-9.	59.2	1308
455	Macitentan and morbidity and mortality in pulmonary arterial hypertension. <i>New England Journal of Medicine</i> , 2013 , 369, 809-18	59.2	878
454	Riociguat for the treatment of pulmonary arterial hypertension. <i>New England Journal of Medicine</i> , 2013 , 369, 330-40	59.2	861
453	Riociguat for the treatment of chronic thromboembolic pulmonary hypertension. <i>New England Journal of Medicine</i> , 2013 , 369, 319-29	59.2	852
452	Tadalafil therapy for pulmonary arterial hypertension. <i>Circulation</i> , 2009 , 119, 2894-903	16.7	769
451	Ambrisentan for the treatment of pulmonary arterial hypertension: results of the ambrisentan in pulmonary arterial hypertension, randomized, double-blind, placebo-controlled, multicenter, efficacy (ARIES) study 1 and 2. <i>Circulation</i> , 2008 , 117, 3010-9	16.7	769
450	Reversal of experimental pulmonary hypertension by PDGF inhibition. <i>Journal of Clinical Investigation</i> , 2005 , 115, 2811-21	15.9	764
449	Initial Use of Ambrisentan plus Tadalafil in Pulmonary Arterial Hypertension. <i>New England Journal of Medicine</i> , 2015 , 373, 834-44	59.2	618
448	Sildenafil for treatment of lung fibrosis and pulmonary hypertension: a randomised controlled trial. <i>Lancet, The</i> , 2002 , 360, 895-900	40	590
447	Selexipag for the Treatment of Pulmonary Arterial Hypertension. <i>New England Journal of Medicine</i> , 2015 , 373, 2522-33	59.2	521
446	Mechanisms of disease: pulmonary arterial hypertension. <i>Nature Reviews Cardiology</i> , 2011 , 8, 443-55	14.8	472
445	Bosentan for treatment of inoperable chronic thromboembolic pulmonary hypertension: BENEFiT (Bosentan Effects in iNopErable Forms of chronic Thromboembolic pulmonary hypertension), a randomized, placebo-controlled trial. <i>Journal of the American College of Cardiology</i> , 2008 , 52, 2127-34	15.1	409
444	Complications of right heart catheterization procedures in patients with pulmonary hypertension in experienced centers. <i>Journal of the American College of Cardiology</i> , 2006 , 48, 2546-52	15.1	393

(2004-2009)

443	Updated evidence-based treatment algorithm in pulmonary arterial hypertension. <i>Journal of the American College of Cardiology</i> , 2009 , 54, S78-S84	15.1	379	
442	Imatinib for the treatment of pulmonary arterial hypertension. <i>New England Journal of Medicine</i> , 2005 , 353, 1412-3	59.2	377	
441	Imatinib mesylate as add-on therapy for pulmonary arterial hypertension: results of the randomized IMPRES study. <i>Circulation</i> , 2013 , 127, 1128-38	16.7	368	
440	Sildenafil: from angina to erectile dysfunction to pulmonary hypertension and beyond. <i>Nature Reviews Drug Discovery</i> , 2006 , 5, 689-702	64.1	366	
439	Combination therapy with oral sildenafil and inhaled iloprost for severe pulmonary hypertension. <i>Annals of Internal Medicine</i> , 2002 , 136, 515-22	8	346	
438	Inhaled prostacyclin and iloprost in severe pulmonary hypertension secondary to lung fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 1999 , 160, 600-7	10.2	300	
437	Oral sildenafil as long-term adjunct therapy to inhaled iloprost in severe pulmonary arterial hypertension. <i>Journal of the American College of Cardiology</i> , 2003 , 42, 158-64	15.1	290	
436	Mortality in pulmonary arterial hypertension: prediction by the 2015 European pulmonary hypertension guidelines risk stratification model. <i>European Respiratory Journal</i> , 2017 , 50,	13.6	288	
435	Hypoxia-dependent regulation of nonphagocytic NADPH oxidase subunit NOX4 in the pulmonary vasculature. <i>Circulation Research</i> , 2007 , 101, 258-67	15.7	279	
434	Imatinib in pulmonary arterial hypertension patients with inadequate response to established therapy. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2010 , 182, 1171-7	10.2	276	
433	Aerosolized prostacyclin and iloprost in severe pulmonary hypertension. <i>Annals of Internal Medicine</i> , 1996 , 124, 820-4	8	273	
432	Chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2019, 53,	13.6	263	
431	Classical transient receptor potential channel 6 (TRPC6) is essential for hypoxic pulmonary vasoconstriction and alveolar gas exchange. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006 , 103, 19093-8	11.5	247	
430	A comparison of the acute hemodynamic effects of inhaled nitric oxide and aerosolized iloprost in primary pulmonary hypertension. German PPH study group. <i>Journal of the American College of Cardiology</i> , 2000 , 35, 176-82	15.1	246	
429	Anticoagulation and survival in pulmonary arterial hypertension: results from the Comparative, Prospective Registry of Newly Initiated Therapies for Pulmonary Hypertension (COMPERA). <i>Circulation</i> , 2014 , 129, 57-65	16.7	235	
428	Elderly patients diagnosed with idiopathic pulmonary arterial hypertension: results from the COMPERA registry. <i>International Journal of Cardiology</i> , 2013 , 168, 871-80	3.2	231	
427	Inducible NOS inhibition reverses tobacco-smoke-induced emphysema and pulmonary hypertension in mice. <i>Cell</i> , 2011 , 147, 293-305	56.2	226	
426	Sildenafil increased exercise capacity during hypoxia at low altitudes and at Mount Everest base camp: a randomized, double-blind, placebo-controlled crossover trial. <i>Annals of Internal Medicine</i> , 2004 , 141, 169-77	8	220	

425	Immune and inflammatory cell involvement in the pathology of idiopathic pulmonary arterial hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2012 , 186, 897-908	10.2	219
424	Riociguat for patients with pulmonary hypertension caused by systolic left ventricular dysfunction: a phase IIb double-blind, randomized, placebo-controlled, dose-ranging hemodynamic study. <i>Circulation</i> , 2013 , 128, 502-11	16.7	215
423	Sildenafil for long-term treatment of nonoperable chronic thromboembolic pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2003 , 167, 1139-41	10.2	211
422	Chronic sildenafil treatment inhibits monocrotaline-induced pulmonary hypertension in rats. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2004 , 169, 39-45	10.2	207
421	Long-term ambrisentan therapy for the treatment of pulmonary arterial hypertension. <i>Journal of the American College of Cardiology</i> , 2009 , 54, 1971-81	15.1	191
420	Long-term treatment with sildenafil citrate in pulmonary arterial hypertension: the SUPER-2 study. <i>Chest</i> , 2011 , 140, 1274-1283	5.3	187
419	Riociguat for chronic thromboembolic pulmonary hypertension and pulmonary arterial hypertension: a phase II study. <i>European Respiratory Journal</i> , 2010 , 36, 792-9	13.6	186
418	First acute haemodynamic study of soluble guanylate cyclase stimulator riociguat in pulmonary hypertension. <i>European Respiratory Journal</i> , 2009 , 33, 785-92	13.6	186
417	Expression and function of soluble guanylate cyclase in pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2008 , 32, 881-91	13.6	186
416	Activation of soluble guanylate cyclase reverses experimental pulmonary hypertension and vascular remodeling. <i>Circulation</i> , 2006 , 113, 286-95	16.7	183
415	Identification of rare sequence variation underlying heritable pulmonary arterial hypertension. <i>Nature Communications</i> , 2018 , 9, 1416	17.4	182
414	Riociguat for the treatment of chronic thromboembolic pulmonary hypertension: a long-term extension study (CHEST-2). <i>European Respiratory Journal</i> , 2015 , 45, 1293-302	13.6	175
413	Inhibition of microRNA-17 improves lung and heart function in experimental pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2012 , 185, 409-19	10.2	171
412	Sildenafil treatment for portopulmonary hypertension. <i>European Respiratory Journal</i> , 2006 , 28, 563-7	13.6	168
411	Stress Doppler echocardiography in relatives of patients with idiopathic and familial pulmonary arterial hypertension: results of a multicenter European analysis of pulmonary artery pressure response to exercise and hypoxia. <i>Circulation</i> , 2009 , 119, 1747-57	16.7	164
410	Safety and efficacy of exercise training in various forms of pulmonary hypertension. <i>European Respiratory Journal</i> , 2012 , 40, 84-92	13.6	158
409	Upregulation of NAD(P)H oxidase 1 in hypoxia activates hypoxia-inducible factor 1 via increase in reactive oxygen species. <i>Free Radical Biology and Medicine</i> , 2004 , 36, 1279-88	7.8	156
408	Regulation of hypoxic pulmonary vasoconstriction: basic mechanisms. <i>European Respiratory Journal</i> , 2008 , 32, 1639-51	13.6	152

(2006-2016)

407	2015 ESC/ERS Guidelines for the Diagnosis and Treatment of Pulmonary Hypertension. <i>Revista Espanola De Cardiologia (English Ed)</i> , 2016 , 69, 177	0.7	148
406	Bronchoscopic surfactant administration in patients with severe adult respiratory distress syndrome and sepsis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 1996 , 154, 57-62	10.2	147
405	A functional single-nucleotide polymorphism in the TRPC6 gene promoter associated with idiopathic pulmonary arterial hypertension. <i>Circulation</i> , 2009 , 119, 2313-22	16.7	146
404	Increased levels and reduced catabolism of asymmetric and symmetric dimethylarginines in pulmonary hypertension. <i>FASEB Journal</i> , 2005 , 19, 1175-7	0.9	144
403	The Giessen Pulmonary Hypertension Registry: Survival in pulmonary hypertension subgroups. Journal of Heart and Lung Transplantation, 2017 , 36, 957-967	5.8	138
402	Gull ESC/ERS 2015 sobre diagnilico y tratamiento de la hipertensili pulmonar. <i>Revista Espanola De Cardiologia</i> , 2016 , 69, 177.e1-177.e62	1.5	137
401	Activation of TRPC6 channels is essential for lung ischaemia-reperfusion induced oedema in mice. <i>Nature Communications</i> , 2012 , 3, 649	17.4	137
400	Bosentan added to sildenafil therapy in patients with pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2015 , 46, 405-13	13.6	136
399	Macitentan for the treatment of inoperable chronic thromboembolic pulmonary hypertension (MERIT-1): results from the multicentre, phase 2, randomised, double-blind, placebo-controlled study. <i>Lancet Respiratory Medicine,the</i> , 2017 , 5, 785-794	35.1	133
398	Riociguat for the treatment of pulmonary arterial hypertension: a long-term extension study (PATENT-2). European Respiratory Journal, 2015 , 45, 1303-13	13.6	131
397	Balloon pulmonary angioplasty in chronic thromboembolic pulmonary hypertension. <i>European Respiratory Review</i> , 2017 , 26,	9.8	130
396	Vascular receptor autoantibodies in pulmonary arterial hypertension associated with systemic sclerosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014 , 190, 808-17	10.2	129
395	Long-term treatment with sildenafil in chronic thromboembolic pulmonary hypertension. <i>European Respiratory Journal</i> , 2007 , 30, 922-7	13.6	127
394	Reduced microRNA-150 is associated with poor survival in pulmonary arterial hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2013 , 187, 294-302	10.2	126
393	Effect of exercise and respiratory training on clinical progression and survival in patients with severe chronic pulmonary hypertension. <i>Respiration</i> , 2011 , 81, 394-401	3.7	125
392	Combined tyrosine and serine/threonine kinase inhibition by sorafenib prevents progression of experimental pulmonary hypertension and myocardial remodeling. <i>Circulation</i> , 2008 , 118, 2081-90	16.7	121
391	Phosphodiesterase 1 upregulation in pulmonary arterial hypertension: target for reverse-remodeling therapy. <i>Circulation</i> , 2007 , 115, 2331-9	16.7	118
390	Favorable effects of inhaled treprostinil in severe pulmonary hypertension: results from randomized controlled pilot studies. <i>Journal of the American College of Cardiology</i> , 2006 , 48, 1672-81	15.1	115

389	Balloon pulmonary angioplasty for inoperable patients with chronic thromboembolic pulmonary hypertension: the initial German experience. <i>European Respiratory Journal</i> , 2017 , 49,	13.6	111
388	Nitric oxide pathway and phosphodiesterase inhibitors in pulmonary arterial hypertension. <i>Journal of the American College of Cardiology</i> , 2004 , 43, 68S-72S	15.1	108
387	Targeting cancer with phosphodiesterase inhibitors. <i>Expert Opinion on Investigational Drugs</i> , 2010 , 19, 117-31	5.9	105
386	Targeting non-malignant disorders with tyrosine kinase inhibitors. <i>Nature Reviews Drug Discovery</i> , 2010 , 9, 956-70	64.1	102
385	Prostacyclin and its analogues in the treatment of pulmonary hypertension 2004 , 102, 139-53		102
384	Tadalafil for the treatment of pulmonary arterial hypertension: a double-blind 52-week uncontrolled extension study. <i>Journal of the American College of Cardiology</i> , 2012 , 60, 768-74	15.1	101
383	Phosphodiesterase inhibitors for the treatment of pulmonary hypertension. <i>European Respiratory Journal</i> , 2008 , 32, 198-209	13.6	101
382	Pharmacodynamics and pharmacokinetics of inhaled iloprost, aerosolized by three different devices, in severe pulmonary hypertension. <i>Chest</i> , 2003 , 124, 1294-304	5.3	100
381	Role of epidermal growth factor inhibition in experimental pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2010 , 181, 158-67	10.2	99
380	Predictors of long-term outcomes in patients treated with riociguat for chronic thromboembolic pulmonary hypertension: data from the CHEST-2 open-label, randomised, long-term extension trial. <i>Lancet Respiratory Medicine,the</i> , 2016 , 4, 372-80	35.1	98
379	Initial combination therapy with ambrisentan and tadalafil in connective tissue disease-associated pulmonary arterial hypertension (CTD-PAH): subgroup analysis from the AMBITION trial. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 1219-1227	2.4	96
378	Simvastatin as a treatment for pulmonary hypertension trial. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2010 , 181, 1106-13	10.2	93
377	New trial designs and potential therapies for pulmonary artery hypertension. <i>Journal of the American College of Cardiology</i> , 2013 , 62, D82-91	15.1	90
376	Role of Src tyrosine kinases in experimental pulmonary hypertension. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2012 , 32, 1354-65	9.4	90
375	The soluble guanylate cyclase stimulator riociguat ameliorates pulmonary hypertension induced by hypoxia and SU5416 in rats. <i>PLoS ONE</i> , 2012 , 7, e43433	3.7	89
374	The molecular targets of approved treatments for pulmonary arterial hypertension. <i>Thorax</i> , 2016 , 71, 73-83	7.3	87
373	Pulmonary vascular disease in the developing world. <i>Circulation</i> , 2008 , 118, 1758-66	16.7	87
372	Antiremodeling effects of iloprost and the dual-selective phosphodiesterase 3/4 inhibitor tolafentrine in chronic experimental pulmonary hypertension. <i>Circulation Research</i> , 2004 , 94, 1101-8	15.7	86

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371	Differences in hemodynamic and oxygenation responses to three different phosphodiesterase-5 inhibitors in patients with pulmonary arterial hypertension: a randomized prospective study. Journal of the American College of Cardiology, 2004 , 44, 1488-96	15.1	86
370	Impact of mitochondria and NADPH oxidases on acute and sustained hypoxic pulmonary vasoconstriction. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2006 , 34, 505-13	5.7	83
369	Tadalafil monotherapy and as add-on to background bosentan in patients with pulmonary arterial hypertension. <i>Journal of Heart and Lung Transplantation</i> , 2011 , 30, 632-43	5.8	79
368	Ultrasonic versus jet nebulization of iloprost in severe pulmonary hypertension. <i>European Respiratory Journal</i> , 2001 , 17, 14-9	13.6	79
367	Effects of riociguat in severe experimental pulmonary hypertension. <i>BMC Pharmacology</i> , 2011 , 11,		78
366	The soluble guanylate cyclase stimulator riociguat ameliorates pulmonary hypertension induced by hypoxia and SU5416 in rats. <i>BMC Pharmacology</i> , 2011 , 11,		78
365	Uncertainties in the diagnosis and treatment of pulmonary arterial hypertension. <i>Circulation</i> , 2008 , 118, 1195-201	16.7	78
364	Reserve of Right Ventricular-Arterial Coupling in the Setting of Chronic Overload. <i>Circulation: Heart Failure</i> , 2019 , 12, e005512	7.6	78
363	Validation of the Tricuspid Annular Plane Systolic Excursion/Systolic Pulmonary Artery Pressure Ratio for the Assessment of Right Ventricular-Arterial Coupling in Severe Pulmonary Hypertension. <i>Circulation: Cardiovascular Imaging</i> , 2019 , 12, e009047	3.9	77
362	Riociguat for the treatment of pulmonary arterial hypertension associated with connective tissue disease: results from PATENT-1 and PATENT-2. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 422-426	2.4	76
361	Terguride ameliorates monocrotaline-induced pulmonary hypertension in rats. <i>European Respiratory Journal</i> , 2011 , 37, 1104-18	13.6	75
360	Hypoxic vasoconstriction in intact lungs: a role for NADPH oxidase-derived H(2)O(2)?. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2000 , 279, L683-90	5.8	75
359	RESPITE: switching to riociguat in pulmonary arterial hypertension patients with inadequate response to phosphodiesterase-5 inhibitors. <i>European Respiratory Journal</i> , 2017 , 50,	13.6	74
358	Expression and activity of phosphodiesterase isoforms during epithelial mesenchymal transition: the role of phosphodiesterase 4. <i>Molecular Biology of the Cell</i> , 2009 , 20, 4751-65	3.5	73
357	Long-term safety and efficacy of imatinib in pulmonary arterial hypertension. <i>Journal of Heart and Lung Transplantation</i> , 2015 , 34, 1366-75	5.8	72
356	Fhl-1, a new key protein in pulmonary hypertension. <i>Circulation</i> , 2008 , 118, 1183-94	16.7	71
355	Pathophysiology and treatment of high-altitude pulmonary vascular disease. <i>Circulation</i> , 2015 , 131, 582	-96 .7	70
354	ERS statement on chronic thromboembolic pulmonary hypertension. <i>European Respiratory Journal</i> , 2021 , 57,	13.6	70

353	Riociguat for the treatment of pulmonary hypertension. <i>Expert Opinion on Investigational Drugs</i> , 2011 , 20, 567-76	5.9	69
352	Cyclooxygenase isoenzyme localization and mRNA expression in rat lungs. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 1998 , 18, 479-88	5.7	69
351	Predictors of long-term outcomes in patients treated with riociguat for pulmonary arterial hypertension: data from the PATENT-2 open-label, randomised, long-term extension trial. <i>Lancet Respiratory Medicine,the</i> , 2016 , 4, 361-71	35.1	69
350	Oleic acid inhibits alveolar fluid reabsorption: a role in acute respiratory distress syndrome?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2005 , 171, 469-79	10.2	68
349	Low-dose systemic phosphodiesterase inhibitors amplify the pulmonary vasodilatory response to inhaled prostacyclin in experimental pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 1999 , 160, 1500-6	10.2	68
348	Nocturnal periodic breathing in primary pulmonary hypertension. <i>European Respiratory Journal</i> , 2002 , 19, 658-63	13.6	67
347	Long-term outcome with intravenous iloprost in pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2009 , 34, 132-7	13.6	66
346	Role of the prostanoid EP4 receptor in iloprost-mediated vasodilatation in pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2008 , 178, 188-96	10.2	66
345	Relevance of the TAPSE/PASP ratio in pulmonary arterial hypertension. <i>International Journal of Cardiology</i> , 2018 , 266, 229-235	3.2	65
344	Riociguat for pulmonary arterial hypertension associated with congenital heart disease. <i>Heart</i> , 2015 , 101, 1792-9	5.1	64
343	Anxiety and depression disorders in patients with pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension. <i>Respiratory Research</i> , 2013 , 14, 104	7.3	64
342	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
341	Novel and emerging therapies for pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014 , 189, 394-400	10.2	62
340	Amplification of the pulmonary vasodilatory response to inhaled iloprost by subthreshold phosphodiesterase types 3 and 4 inhibition in severe pulmonary hypertension. <i>Critical Care Medicine</i> , 2002 , 30, 2489-92	1.4	62
339	Hypoxia induces Kv channel current inhibition by increased NADPH oxidase-derived reactive oxygen species. <i>Free Radical Biology and Medicine</i> , 2012 , 52, 1033-42	7.8	60
338	Inflammation, immunological reaction and role of infection in pulmonary hypertension. <i>Clinical Microbiology and Infection</i> , 2011 , 17, 7-14	9.5	60
337	Stimulation of soluble guanylate cyclase prevents cigarette smoke-induced pulmonary hypertension and emphysema. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014 , 189, 1359-73	10.2	59
336	Long-term therapy with inhaled iloprost in patients with pulmonary hypertension. <i>Respiratory Medicine</i> , 2010 , 104, 731-40	4.6	59

335	Safety and tolerability of bosentan in idiopathic pulmonary fibrosis: an open label study. <i>European Respiratory Journal</i> , 2007 , 29, 713-9	13.6	59
334	Mitochondrial Complex IV Subunit 4 Isoform 2 Is Essential for Acute Pulmonary Oxygen Sensing. <i>Circulation Research</i> , 2017 , 121, 424-438	15.7	58
333	Sleep apnea in precapillary pulmonary hypertension. Sleep Medicine, 2013, 14, 247-51	4.6	58
332	Selexipag for the treatment of connective tissue disease-associated pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2017 , 50,	13.6	58
331	Classical transient receptor potential channel 1 in hypoxia-induced pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2013 , 188, 1451-9	10.2	58
330	Impairment of respiratory muscle function in pulmonary hypertension. Clinical Science, 2008, 114, 165-7	71 6.5	58
329	Inhaled iloprost is a potent acute pulmonary vasodilator in HIV-related severe pulmonary hypertension. <i>European Respiratory Journal</i> , 2004 , 23, 321-6	13.6	58
328	Riociguat: Mode of Action and Clinical Development in Pulmonary Hypertension. <i>Chest</i> , 2017 , 151, 468-	48.9	57
327	ASK1 Inhibition Halts Disease Progression in Preclinical Models of Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2018 , 197, 373-385	10.2	57
326	Function of NADPH oxidase 1 in pulmonary arterial smooth muscle cells after monocrotaline-induced pulmonary vascular remodeling. <i>Antioxidants and Redox Signaling</i> , 2013 , 19, 221	3- 3 1	57
325	Effects of hypercapnia with and without acidosis on hypoxic pulmonary vasoconstriction. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2009 , 297, L977-83	5.8	57
324	Acute effects of the combination of sildenafil and inhaled treprostinil on haemodynamics and gas exchange in pulmonary hypertension. <i>Pulmonary Pharmacology and Therapeutics</i> , 2008 , 21, 824-32	3.5	57
323	Pulmonary Hypertension. <i>Deutsches A&#x0308;rzteblatt International</i> , 2017 , 114, 73-84	2.5	57
322	Notch1 signalling regulates endothelial proliferation and apoptosis in pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2016 , 48, 1137-1149	13.6	57
321	Thrombin impairs alveolar fluid clearance by promoting endocytosis of Na+,K+-ATPase. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2005 , 33, 343-54	5.7	56
320	The lectin-like domain of tumor necrosis factor-alpha improves alveolar fluid balance in injured isolated rabbit lungs. <i>Critical Care Medicine</i> , 2008 , 36, 1543-50	1.4	55
319	Inhaled iloprost reverses vascular remodeling in chronic experimental pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2005 , 172, 358-63	10.2	55
318	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

317	Involvement of mast cells in monocrotaline-induced pulmonary hypertension in rats. <i>Respiratory Research</i> , 2011 , 12, 60	7.3	53
316	NO and reactive oxygen species are involved in biphasic hypoxic vasoconstriction of isolated rabbit lungs. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2001 , 280, L638-45	5.8	53
315	PAR-2 inhibition reverses experimental pulmonary hypertension. <i>Circulation Research</i> , 2012 , 110, 1179-	91 15.7	52
314	Basic features of hypoxic pulmonary vasoconstriction in mice. <i>Respiratory Physiology and Neurobiology</i> , 2004 , 139, 191-202	2.8	51
313	Mediator generation and signaling events in alveolar epithelial cells attacked by S. aureus alpha-toxin. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2002 , 282, L207-14	5.8	51
312	Pulmonary Arterial Hypertension-Related Morbidity Is Prognostic for Mortality. <i>Journal of the American College of Cardiology</i> , 2018 , 71, 752-763	15.1	50
311	Mitochondrial hyperpolarization in pulmonary vascular remodeling. Mitochondrial uncoupling protein deficiency as disease model. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2013 , 49, 358-67	5.7	50
310	Lung cancer-associated pulmonary hypertension: Role of microenvironmental inflammation based on tumor cell-immune cell cross-talk. <i>Science Translational Medicine</i> , 2017 , 9,	17.5	50
309	The role of dimethylarginine dimethylaminohydrolase in idiopathic pulmonary fibrosis. <i>Science Translational Medicine</i> , 2011 , 3, 87ra53	17.5	50
308	Future perspectives for the treatment of pulmonary arterial hypertension. <i>Journal of the American College of Cardiology</i> , 2009 , 54, S108-S117	15.1	50
307	Effects of mitochondrial inhibitors and uncouplers on hypoxic vasoconstriction in rabbit lungs. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2003 , 29, 721-32	5.7	50
306	Recovery from circulatory shock in severe primary pulmonary hypertension (PPH) with aerosolization of iloprost. <i>Intensive Care Medicine</i> , 1998 , 24, 631-4	14.5	49
305	Oxygen sensors in hypoxic pulmonary vasoconstriction. <i>Cardiovascular Research</i> , 2006 , 71, 620-9	9.9	49
304	Microcirculatory dysfunction in the brain precedes changes in evoked potentials in endotoxin-induced sepsis syndrome in rats. <i>Cerebrovascular Diseases</i> , 2007 , 23, 140-7	3.2	49
303	Coaerosolization of phosphodiesterase inhibitors markedly enhances the pulmonary vasodilatory response to inhaled iloprost in experimental pulmonary hypertension. Maintenance of lung selectivity. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2001 , 164, 1694-700	10.2	49
302	Traditional and new composite endpoints in heart failure clinical trials: facilitating comprehensive efficacy assessments and improving trial efficiency. <i>European Journal of Heart Failure</i> , 2016 , 18, 482-9	12.3	49
301	p38 MAPK Inhibition Improves Heart Function in Pressure-Loaded Right Ventricular Hypertrophy. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2017 , 57, 603-614	5.7	48
300	5-HT2B receptor antagonists inhibit fibrosis and protect from RV heart failure. <i>BioMed Research International</i> , 2015 , 2015, 438403	3	48

299	Pulmonary hypertension due to chronic lung disease: updated Recommendations of the Cologne Consensus Conference 2011. <i>International Journal of Cardiology</i> , 2011 , 154 Suppl 1, S45-53	3.2	48
298	Cellular pathophysiology and therapy of pulmonary hypertension. <i>Translational Research</i> , 2001 , 138, 367-77		48
297	Hemodynamic and clinical onset in patients with hereditary pulmonary arterial hypertension and BMPR2 mutations. <i>Respiratory Research</i> , 2011 , 12, 99	7.3	47
296	Identification of right heart-enriched genes in a murine model of chronic outflow tract obstruction. Journal of Molecular and Cellular Cardiology, 2010 , 49, 598-605	5.8	47
295	Atrial natriuretic peptide in severe primary and nonprimary pulmonary hypertension: response to iloprost inhalation. <i>Journal of the American College of Cardiology</i> , 2001 , 38, 1130-6	15.1	47
294	Sildenafil improves dynamic vascular function in the brain: studies in patients with pulmonary hypertension. <i>Cerebrovascular Diseases</i> , 2006 , 21, 194-200	3.2	46
293	More on Single-Beat Estimation of Right Ventriculoarterial Coupling in Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018 , 198, 816-818	10.2	45
292	SERAPHIN haemodynamic substudy: the effect of the dual endothelin receptor antagonist macitentan on haemodynamic parameters and NT-proBNP levels and their association with disease progression in patients with pulmonary arterial hypertension. <i>European Heart Journal</i> , 2017 , 38, 1147-1	9.5 155	44
291	Diacylglycerol regulates acute hypoxic pulmonary vasoconstriction via TRPC6. <i>Respiratory Research</i> , 2011 , 12, 20	7.3	44
290	Amplified canonical transforming growth factor-Bignalling heat shock protein 90 in pulmonary fibrosis. <i>European Respiratory Journal</i> , 2017 , 49,	13.6	43
289	Therapeutic efficacy of azaindole-1 in experimental pulmonary hypertension. <i>European Respiratory Journal</i> , 2010 , 36, 808-18	13.6	43
288	Effect of macitentan on hospitalizations: results from the SERAPHIN trial. <i>JACC: Heart Failure</i> , 2015 , 3, 1-8	7.9	42
287	miR-223-IGF-IR signalling in hypoxia- and load-induced right-ventricular failure: a novel therapeutic approach. <i>Cardiovascular Research</i> , 2016 , 111, 184-93	9.9	42
286	Current and future treatments of pulmonary arterial hypertension. <i>British Journal of Pharmacology</i> , 2021 , 178, 6-30	8.6	42
285	Medical management of chronic thromboembolic pulmonary hypertension. <i>European Respiratory Review</i> , 2017 , 26,	9.8	41
284	Arterial hypertension in a murine model of sleep apnea: role of NADPH oxidase 2. <i>Journal of Hypertension</i> , 2014 , 32, 300-5	1.9	41
283	Diagnosis of CTEPH versus IPAH using capillary to end-tidal carbon dioxide gradients. <i>European Respiratory Journal</i> , 2012 , 39, 119-24	13.6	41
282	Prevalence of Pulmonary Hypertension in the General Population: The Rotterdam Study. <i>PLoS ONE</i> , 2015 , 10, e0130072	3.7	41

281	Targeting cyclin-dependent kinases for the treatment of pulmonary arterial hypertension. <i>Nature Communications</i> , 2019 , 10, 2204	17.4	39
280	Targeting the Prostacyclin Pathway with Selexipag in Patients with Pulmonary Arterial Hypertension Receiving Double Combination Therapy: Insights from the Randomized Controlled GRIPHON Study. <i>American Journal of Cardiovascular Drugs</i> , 2018 , 18, 37-47	4	39
279	Balloon pulmonary angioplasty for inoperable patients with chronic thromboembolic disease. <i>Pulmonary Circulation</i> , 2018 , 8, 2045893217753122	2.7	39
278	Inhibition of phosphodiesterase 4 enhances lung alveolarisation in neonatal mice exposed to hyperoxia. <i>European Respiratory Journal</i> , 2009 , 33, 861-70	13.6	39
277	Treatment of pulmonary arterial hypertension (PAH): updated Recommendations of the Cologne Consensus Conference 2011. <i>International Journal of Cardiology</i> , 2011 , 154 Suppl 1, S20-33	3.2	38
276	The science of endothelin-1 and endothelin receptor antagonists in the management of pulmonary arterial hypertension: current understanding and future studies. <i>European Journal of Clinical Investigation</i> , 2009 , 39 Suppl 2, 38-49	4.6	38
275	Hypoxic pulmonary hypertension in mice with constitutively active platelet-derived growth factor receptor- [Pulmonary Circulation, 2011, 1, 259-68]	2.7	38
274	Soluble guanylate cyclase stimulation: an emerging option in pulmonary hypertension therapy. <i>European Respiratory Review</i> , 2009 , 18, 35-41	9.8	38
273	Initial combination therapy with ambrisentan and tadalafil and mortality in patients with pulmonary arterial hypertension: a secondary analysis of the results from the randomised, controlled AMBITION study. <i>Lancet Respiratory Medicine,the</i> , 2016 , 4, 894-901	35.1	37
272	Cardiac Magnetic Resonance Imaging-Based Right Ventricular Strain Analysis for Assessment of Coupling and Diastolic Function in Pulmonary Hypertension. <i>JACC: Cardiovascular Imaging</i> , 2019 , 12, 21	58 :2 16	54 ³⁶
271	Macitentan Improves Health-Related Quality of Life for Patients With Pulmonary Arterial Hypertension: Results From the Randomized Controlled SERAPHIN Trial. <i>Chest</i> , 2017 , 151, 106-118	5.3	36
270	Metered dose inhaler delivery of treprostinil for the treatment of pulmonary hypertension. <i>Pulmonary Pharmacology and Therapeutics</i> , 2009 , 22, 50-6	3.5	36
269	Hypoxia- and non-hypoxia-related pulmonary hypertension - established and new therapies. <i>Cardiovascular Research</i> , 2006 , 72, 30-40	9.9	36
268	Increased neutrophil mediator release in patients with pulmonary hypertensionsuppression by inhaled iloprost. <i>Thrombosis and Haemostasis</i> , 2003 , 90, 1141-9	7	36
267	The role of combination therapy in managing pulmonary arterial hypertension. <i>European Respiratory Review</i> , 2014 , 23, 469-75	9.8	35
266	Psoas muscle architectural design, in vivo sarcomere length range, and passive tensile properties support its role as a lumbar spine stabilizer. <i>Spine</i> , 2011 , 36, E1666-74	3.3	35
265	Burden of pulmonary arterial hypertension in Germany. Respiratory Medicine, 2010, 104, 902-10	4.6	34
264	Heme oxygenase-2 and large-conductance Ca2+-activated K+ channels: lung vascular effects of hypoxia. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2009 , 180, 353-64	10.2	34

263	cAMP phosphodiesterase inhibitors increases nitric oxide production by modulating dimethylarginine dimethylaminohydrolases. <i>Circulation</i> , 2011 , 123, 1194-204	16.7	34
262	Pulmonary hypertension: updated classification and management of pulmonary hypertension. <i>Heart</i> , 2010 , 96, 552-9	5.1	34
261	Prostacyclin enhances stretch-induced surfactant secretion in alveolar epithelial type II cells. <i>American Journal of Respiratory and Critical Care Medicine</i> , 1999 , 160, 846-51	10.2	34
260	Quality of life in patients with chronic thromboembolic pulmonary hypertension. <i>European Respiratory Journal</i> , 2016 , 48, 526-37	13.6	34
259	Telomerecat: A ploidy-agnostic method for estimating telomere length from whole genome sequencing data. <i>Scientific Reports</i> , 2018 , 8, 1300	4.9	33
258	Evidence for a role of protein kinase C in hypoxic pulmonary vasoconstriction. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 1999 , 276, L90-5	5.8	33
257	Glycogen synthase kinase 3beta contributes to proliferation of arterial smooth muscle cells in pulmonary hypertension. <i>PLoS ONE</i> , 2011 , 6, e18883	3.7	33
256	Targeted therapy of pulmonary arterial hypertension: Updated recommendations from the Cologne Consensus Conference 2018. <i>International Journal of Cardiology</i> , 2018 , 272S, 37-45	3.2	33
255	Association of N-Terminal Pro Brain Natriuretic Peptide and Long-Term Outcome in Patients With Pulmonary Arterial Hypertension. <i>Circulation</i> , 2019 , 139, 2440-2450	16.7	32
254	Risk assessment in pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension. <i>European Respiratory Journal</i> , 2019 , 53,	13.6	32
253	The soluble guanylate cyclase activator HMR1766 reverses hypoxia-induced experimental pulmonary hypertension in mice. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2009 , 297, L658-65	5.8	32
252	Eplerenone attenuates pathological pulmonary vascular rather than right ventricular remodeling in pulmonary arterial hypertension. <i>BMC Pulmonary Medicine</i> , 2018 , 18, 41	3.5	31
251	Incident and prevalent cohorts with pulmonary arterial hypertension: insight from SERAPHIN. <i>European Respiratory Journal</i> , 2015 , 46, 1711-20	13.6	31
250	Iloprost-induced desensitization of the prostacyclin receptor in isolated rabbit lungs. <i>Respiratory Research</i> , 2007 , 8, 4	7.3	31
249	Characterization of a murine model of monocrotaline pyrrole-induced acute lung injury. <i>BMC Pulmonary Medicine</i> , 2008 , 8, 25	3.5	31
248	Sequential treatment with riociguat and balloon pulmonary angioplasty for patients with inoperable chronic thromboembolic pulmonary hypertension. <i>Pulmonary Circulation</i> , 2018 , 8, 20458940	1 8783	996
247	Pressure overload leads to an increased accumulation and activity of mast cells in the right ventricle. <i>Physiological Reports</i> , 2017 , 5, e13146	2.6	30
246	Impact of the mitochondria-targeted antioxidant MitoQ on hypoxia-induced pulmonary hypertension. European Respiratory Journal, 2018,	13.6	30

245	Phosphodiesterase 10A upregulation contributes to pulmonary vascular remodeling. <i>PLoS ONE</i> , 2011 , 6, e18136	3.7	30
244	Combination of nonspecific PDE inhibitors with inhaled prostacyclin in experimental pulmonary hypertension. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2001 , 281, L1361-	- § ^{.8}	30
243	Evaluation and Prognostic Relevance of Right Ventricular-Arterial Coupling in Pulmonary Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020 , 201, 116-119	10.2	30
242	Chymase: a multifunctional player in pulmonary hypertension associated with lung fibrosis. <i>European Respiratory Journal</i> , 2015 , 46, 1084-94	13.6	29
241	Late outcomes after acute pulmonary embolism: rationale and design of FOCUS, a prospective observational multicenter cohort study. <i>Journal of Thrombosis and Thrombolysis</i> , 2016 , 42, 600-9	5.1	29
240	Effects of multikinase inhibitors on pressure overload-induced right ventricular remodeling. <i>International Journal of Cardiology</i> , 2013 , 167, 2630-7	3.2	29
239	The Role of Transient Receptor Potential Channel 6 Channels in the Pulmonary Vasculature. <i>Frontiers in Immunology</i> , 2017 , 8, 707	8.4	29
238	New horizons in pulmonary arterial hypertension therapies. <i>European Respiratory Review</i> , 2013 , 22, 503-	- 1 ,48	29
237	Redox signaling and reactive oxygen species in hypoxic pulmonary vasoconstriction. <i>Respiratory Physiology and Neurobiology</i> , 2010 , 174, 282-91	2.8	29
236	Effects of phosphodiesterase 4 inhibition on bleomycin-induced pulmonary fibrosis in mice. <i>BMC Pulmonary Medicine</i> , 2010 , 10, 26	3.5	29
235	Inhaled tolafentrine reverses pulmonary vascular remodeling via inhibition of smooth muscle cell migration. <i>Respiratory Research</i> , 2005 , 6, 128	7.3	29
234	Downregulation of hypoxic vasoconstriction by chronic hypoxia in rabbits: effects of nitric oxide. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2003 , 284, H931-8	5.2	29
233	Congenital erythropoietin over-expression causes "anti-pulmonary hypertensive" structural and functional changes in mice, both in normoxia and hypoxia. <i>Thrombosis and Haemostasis</i> , 2005 , 94, 630-8	7	29
232	Exercise right heart catheterisation before and after pulmonary endarterectomy in patients with chronic thromboembolic disease. <i>European Respiratory Journal</i> , 2018 , 52,	13.6	29
231	Mitochondrial cytochrome redox states and respiration in acute pulmonary oxygen sensing. <i>European Respiratory Journal</i> , 2010 , 36, 1056-66	13.6	28
230	Noninvasive detection of early pulmonary vascular dysfunction in scleroderma. <i>Respiratory Medicine</i> , 2009 , 103, 1713-8	4.6	28
229	Effect of sildenafil on hypoxia-induced changes in pulmonary circulation and right ventricular function. <i>Respiratory Physiology and Neurobiology</i> , 2007 , 159, 196-201	2.8	28
228	HbA1c in pulmonary arterial hypertension: a marker of prognostic relevance?. <i>Journal of Heart and Lung Transplantation</i> , 2012 , 31, 1109-14	5.8	27

(2007-2005)

227	Detection of reactive oxygen species in isolated, perfused lungs by electron spin resonance spectroscopy. <i>Respiratory Research</i> , 2005 , 6, 86	7.3	27
226	Inhaled treprostinil [corrected] for treatment of chronic pulmonary arterial hypertension. <i>Annals of Internal Medicine</i> , 2006 , 144, 149-50	8	27
225	Differential impact of ultrasonically nebulized versus tracheal-instilled surfactant on ventilation-perfusion (VA/Q) mismatch in a model of acute lung injury. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2000 , 161, 152-9	10.2	27
224	Acute effects of riociguat in borderline or manifest pulmonary hypertension associated with chronic obstructive pulmonary disease. <i>Pulmonary Circulation</i> , 2015 , 5, 296-304	2.7	26
223	Inflammatory Mediators Drive Adverse Right Ventricular Remodeling and Dysfunction and Serve as Potential Biomarkers. <i>Frontiers in Physiology</i> , 2018 , 9, 609	4.6	26
222	Histological characterization of mast cell chymase in patients with pulmonary hypertension and chronic obstructive pulmonary disease. <i>Pulmonary Circulation</i> , 2014 , 4, 128-36	2.7	26
221	Partial reversal of experimental pulmonary hypertension by phosphodiesterase-3/4 inhibition. European Respiratory Journal, 2008 , 31, 599-610	13.6	26
220	Endotoxin "priming" potentiates lung vascular abnormalities in response to Escherichia coli hemolysin: an example of synergism between endo- and exotoxin. <i>Journal of Experimental Medicine</i> , 1994 , 180, 1437-43	16.6	26
219	Selexipag treatment for pulmonary arterial hypertension associated with congenital heart disease after defect correction: insights from the randomised controlled GRIPHON study. <i>European Journal of Heart Failure</i> , 2019 , 21, 352-359	12.3	26
218	Riociguat for pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension: Results from a phase II long-term extension study. <i>Respiratory Medicine</i> , 2017 , 128, 50-56	4.6	25
217	Structural and functional prevention of hypoxia-induced pulmonary hypertension by individualized exercise training in mice. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2014 , 306, L986-95	5.8	25
216	Dynamic hyperinflation during exercise in patients with precapillary pulmonary hypertension. <i>Respiratory Medicine</i> , 2012 , 106, 308-13	4.6	25
215	Lung vasodilatory response to inhaled iloprost in experimental pulmonary hypertension: amplification by different type phosphodiesterase inhibitors. <i>Respiratory Research</i> , 2005 , 6, 76	7.3	25
214	Evidence for the Fucoidan/P-Selectin Axis as a Therapeutic Target in Hypoxia-induced Pulmonary Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019 , 199, 1407-1420	10.2	25
213	Riociguat for pulmonary hypertension. <i>Future Cardiology</i> , 2010 , 6, 155-66	1.3	24
212	Hypoxia-induced pulmonary hypertension: different impact of iloprost, sildenafil, and nitric oxide. <i>Respiratory Medicine</i> , 2007 , 101, 2125-32	4.6	24
211	Cigarette Smoke-Induced Emphysema and Pulmonary Hypertension Can Be Prevented by Phosphodiesterase 4 and 5 Inhibition in Mice. <i>PLoS ONE</i> , 2015 , 10, e0129327	3.7	24
2 10	A combination hybrid-based vaccination/adoptive cellular therapy to prevent tumor growth by involvement of T cells. <i>Cancer Research</i> , 2007 , 67, 5443-53	10.1	23

Treatment with low-dose tacrolimus inhibits bleeding complications in a patient with hereditary hemorrhagic telangiectasia and pulmonary arterial hypertension. *Pulmonary Circulation*, **2019**, 9, 2045894018803406

	nemormagic telanglectasia and pulmonary arterial hypertension. Fullmonary circulation, 2019, 9, 20430	- 1010	
208	New potential diagnostic biomarkers for pulmonary hypertension. <i>European Respiratory Journal</i> , 2015 , 46, 1390-6	13.6	22
207	Left ventricular systolic dysfunction associated with pulmonary hypertension riociguat trial (LEPHT): rationale and design. <i>European Journal of Heart Failure</i> , 2012 , 14, 946-53	12.3	22
206	Nitric oxide (NO)-dependent but not NO-independent guanylate cyclase activation attenuates hypoxic vasoconstriction in rabbit lungs. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2000 , 23, 222-7	5.7	22
205	Switching to riociguat versus maintenance therapy with phosphodiesterase-5 inhibitors in patients with pulmonary arterial hypertension (REPLACE): a multicentre, open-label, randomised controlled trial. <i>Lancet Respiratory Medicine,the</i> , 2021 , 9, 573-584	35.1	22
204	REVEAL risk scores applied to riociguat-treated patients in PATENT-2: Impact of changes in risk score on survival. <i>Journal of Heart and Lung Transplantation</i> , 2018 , 37, 513-519	5.8	22
203	Effects of exercise training on pulmonary hemodynamics, functional capacity and inflammation in pulmonary hypertension. <i>Pulmonary Circulation</i> , 2017 , 7, 20-37	2.7	21
202	Right ventricular size and function under riociguat in pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension (the RIVER study). <i>Respiratory Research</i> , 2018 , 19, 258	7.3	21
201	Pulmonary artery to aorta ratio and risk of all-cause mortality in the general population: the Rotterdam Study. <i>European Respiratory Journal</i> , 2017 , 49,	13.6	20
200	Soluble guanylate cyclase stimulator riociguat and phosphodiesterase 5 inhibitor sildenafil ameliorate pulmonary hypertension due to left heart disease in mice. <i>International Journal of Cardiology</i> , 2016 , 216, 85-91	3.2	20
199	An evaluation of fracture stabilization comparing kyphoplasty and titanium mesh repair techniques for vertebral compression fractures: is bone cement necessary?. <i>Spine</i> , 2010 , 35, E768-73	3.3	20
198	Urodilatin, a natriuretic peptide stimulating particulate guanylate cyclase, and the phosphodiesterase 5 inhibitor dipyridamole attenuate experimental pulmonary hypertension: synergism upon coapplication. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2001 , 25, 219	5.7 -25	20
197	Lung cGMP release subsequent to NO inhalation in pulmonary hypertension: responders versus nonresponders. <i>European Respiratory Journal</i> , 2002 , 19, 664-71	13.6	20
196	Pharmacokinetics and metabolism of infused versus inhaled iloprost in isolated rabbit lungs. Journal of Pharmacology and Experimental Therapeutics, 2002, 303, 741-5	4.7	20
195	Effect of nitric oxide synthase (NOS) inhibition on macro- and microcirculation in a model of rat endotoxic shock. <i>Thrombosis and Haemostasis</i> , 2006 , 95, 720-727	7	20
194	Haemodynamic effects of riociguat in inoperable/recurrent chronic thromboembolic pulmonary hypertension. <i>Heart</i> , 2017 , 103, 599-606	5.1	19
193	Sex Differences in Right Ventricular-Pulmonary Arterial Coupling in Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020 , 202, 1042-1046	10.2	19
192	Thin Air Resulting in High Pressure: Mountain Sickness and Hypoxia-Induced Pulmonary Hypertension. <i>Canadian Respiratory Journal</i> , 2017 , 2017, 8381653	2.1	19

191	Updating clinical endpoint definitions. <i>Pulmonary Circulation</i> , 2013 , 3, 206-16	2.7	19	
190	Alveolar epithelial barrier functions in ventilated perfused rabbit lungs. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2001 , 280, L896-904	5.8	19	
189	Pulmonary Hemodynamic Response to Exercise in Chronic Thromboembolic Pulmonary Hypertension before and after Pulmonary Endarterectomy. <i>Respiration</i> , 2015 , 90, 63-73	3.7	18	
188	Biomarkers of tissue remodeling predict survival in patients with pulmonary hypertension. <i>International Journal of Cardiology</i> , 2016 , 223, 821-826	3.2	18	
187	Short-term improvement in pulmonary hemodynamics is strongly predictive of long-term survival in patients with pulmonary arterial hypertension. <i>Pulmonary Circulation</i> , 2013 , 3, 523-32	2.7	18	
186	Zardaverine and aerosolised iloprost in a model of acute respiratory failure. <i>European Respiratory Journal</i> , 2003 , 22, 342-7	13.6	18	
185	Evaluation of the prognostic value of electrocardiography parameters and heart rhythm in patients with pulmonary hypertension. <i>Cardiology Journal</i> , 2016 , 23, 465-72	1.4	18	
184	Right ventricular function correlates of right atrial strain in pulmonary hypertension: a combined cardiac magnetic resonance and conductance catheter study. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2020 , 318, H156-H164	5.2	18	
183	Bypassing mitochondrial complex III using alternative oxidase inhibits acute pulmonary oxygen sensing. <i>Science Advances</i> , 2020 , 6, eaba0694	14.3	18	
182	De Novo Truncating Mutations in WASF1 Cause Intellectual Disability with Seizures. <i>American Journal of Human Genetics</i> , 2018 , 103, 144-153	11	18	
181	Intravenous treprostinil infusion via a fully implantable pump for pulmonary arterial hypertension. <i>Clinical Research in Cardiology</i> , 2017 , 106, 776-783	6.1	17	
180	Short-term venoarterial extracorporeal membrane oxygenation for massive endobronchial hemorrhage after pulmonary endarterectomy. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2018 , 155, 643-649	1.5	17	
179	Impact of S-adenosylmethionine decarboxylase 1 on pulmonary vascular remodeling. <i>Circulation</i> , 2014 , 129, 1510-23	16.7	17	
178	Effects of hypercapnia and NO synthase inhibition in sustained hypoxic pulmonary vasoconstriction. <i>Respiratory Research</i> , 2012 , 13, 7	7.3	17	
177	Phosphodiesterase 6 subunits are expressed and altered in idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2010 , 11, 146	7.3	17	
176	Novel soluble guanylyl cyclase stimulator BAY 41-2272 attenuates ischemia-reperfusion-induced lung injury. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2009 , 296, L462-9	5.8	17	
175	Air travel can be safe and well tolerated in patients with clinically stable pulmonary hypertension. <i>Pulmonary Circulation</i> , 2011 , 1, 239-43	2.7	17	
174	Association between six-minute walk distance and long-term outcomes in patients with pulmonary arterial hypertension: Data from the randomized SERAPHIN trial. <i>PLoS ONE</i> , 2018 , 13, e0193226	3.7	17	

173	Right heart failure in pulmonary hypertension: Diagnosis and new perspectives on vascular and direct right ventricular treatment. <i>British Journal of Pharmacology</i> , 2021 , 178, 90-107	8.6	17
172	Effect of Riociguat and Sildenafil on Right Heart Remodeling and Function in Pressure Overload Induced Model of Pulmonary Arterial Banding. <i>BioMed Research International</i> , 2018 , 2018, 3293584	3	17
171	Comparison of hemodynamic parameters in treatment-nalle and pre-treated patients with pulmonary arterial hypertension in the randomized phase III PATENT-1 study. <i>Journal of Heart and Lung Transplantation</i> , 2017 , 36, 509-519	5.8	16
170	Therapeutic efficacy of TBC3711 in monocrotaline-induced pulmonary hypertension. <i>Respiratory Research</i> , 2011 , 12, 87	7-3	16
169	A case series of patients with severe pulmonary hypertension receiving an implantable pump for intravenous prostanoid therapy. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2012 , 186, 1196-8	10.2	16
168	Apical, but not basolateral, endotoxin preincubation protects alveolar epithelial cells against hydrogen peroxide-induced loss of barrier function: the role of nitric oxide synthesis. <i>Journal of Immunology</i> , 2002 , 169, 1474-81	5-3	16
167	Long-term safety and outcome of intravenous treprostinil via an implanted pump in pulmonary hypertension. <i>Journal of Heart and Lung Transplantation</i> , 2018 , 37, 1235-1244	5.8	16
166	Riociguat for treatment of pulmonary hypertension in COPD: a translational study. <i>European Respiratory Journal</i> , 2019 , 53,	13.6	15
165	Effects of dimethylarginine dimethylaminohydrolase-1 overexpression on the response of the pulmonary vasculature to hypoxia. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2013 , 49, 491-500	5.7	15
164	Cardiovocal Syndrome (Ortnerß Syndrome) Associated with Chronic Thromboembolic Pulmonary Hypertension and Giant Pulmonary Artery Aneurysm: Case Report and Review of the Literature. <i>Case Reports in Medicine</i> , 2012 , 2012, 230736	0.7	15
163	Conebulization of surfactant and urokinase restores gas exchange in perfused lungs with alveolar fibrin formation. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2001 , 280, L79	2 ⁵ 800	15
162	The safety and pharmacokinetics of rapid iloprost aerosol delivery via the BREELIB nebulizer in pulmonary arterial hypertension. <i>Pulmonary Circulation</i> , 2017 , 7, 505-513	2.7	14
161	Maintained right ventricular pressure overload induces ventricular-arterial decoupling in mice. <i>Experimental Physiology</i> , 2017 , 102, 180-189	2.4	14
160	A comprehensive echocardiographic method for risk stratification in pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2020 , 56,	13.6	14
159	FHL-1 is not involved in pressure overload-induced maladaptive right ventricular remodeling and dysfunction. <i>Basic Research in Cardiology</i> , 2020 , 115, 17	11.8	14
158	REVEAL risk score in patients with chronic thromboembolic pulmonary hypertension receiving riociguat. <i>Journal of Heart and Lung Transplantation</i> , 2018 , 37, 836-843	5.8	14
157	Riociguat treatment for portopulmonary hypertension: a subgroup analysis from the PATENT-1/-2 studies. <i>Pulmonary Circulation</i> , 2018 , 8, 2045894018769305	2.7	14
156	The prognostic impact of thyroid function in pulmonary hypertension. <i>Journal of Heart and Lung Transplantation</i> , 2016 , 35, 1427-1434	5.8	14

155	Cofilin, a hypoxia-regulated protein in murine lungs identified by 2DE: role of the cytoskeletal protein cofilin in pulmonary hypertension. <i>Proteomics</i> , 2013 , 13, 75-88	4.8	14	
154	Procedural safety of a fully implantable intravenous prostanoid pump for pulmonary hypertension. <i>Clinical Research in Cardiology</i> , 2017 , 106, 174-182	6.1	14	
153	Nitric Oxide Synthase 2 Induction Promotes Right Ventricular Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2019 , 60, 346-356	5.7	14	
152	Impaired right ventricular lusitropy is associated with ventilatory inefficiency in pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2019 , 54,	13.6	13	
151	Intravenous treprostinil as an add-on therapy in patients with pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2019 , 38, 748-756	5.8	13	
150	A simple echocardiographic estimate of right ventricular-arterial coupling to assess severity and outcome in pulmonary hypertension on chronic lung disease. <i>European Respiratory Journal</i> , 2019 , 54,	13.6	13	
149	Severe Emphysema in the SU5416/Hypoxia Rat Model of Pulmonary Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019 , 200, 515-518	10.2	13	
148	Sildenafil versus nitric oxide for acute vasodilator testing in pulmonary arterial hypertension. <i>Pulmonary Circulation</i> , 2015 , 5, 305-12	2.7	13	
147	Prolonged vasodilatory response to nanoencapsulated sildenafil in pulmonary hypertension. <i>Nanomedicine: Nanotechnology, Biology, and Medicine</i> , 2016 , 12, 63-8	6	13	
146	Riociguat for pulmonary hypertension. New England Journal of Medicine, 2013, 369, 2268	59.2	13	
145	Long-term effects of intravenous iloprost in patients with idiopathic pulmonary arterial hypertension deteriorating on non-parenteral therapy. <i>BMC Pulmonary Medicine</i> , 2011 , 11, 56	3.5	13	
144	Acute hemodynamic response to single oral doses of BAY 60-4552, a soluble guanylate cyclase stimulator, in patients with biventricular heart failure. <i>BMC Pharmacology</i> , 2009 , 9,		13	
143	Comparison of pharmacokinetics and vasodilatory effect of nebulized and infused iloprost in experimental pulmonary hypertension: rapid tolerance development. <i>Journal of Aerosol Medicine and Pulmonary Drug Delivery</i> , 2006 , 19, 353-63		13	
142	AMBRISENTAN HAS NO CLINICALLY RELEVANT EFFECT ON THE PHARMACOKINETICS OR PHARMACODYNAMICS OF WARFARIN. <i>Chest</i> , 2006 , 130, 256S	5.3	13	
141	Increase in alveolar antioxidant levels in hyperoxic and anoxic ventilated rabbit lungs during ischemia. <i>Free Radical Biology and Medicine</i> , 2004 , 36, 78-89	7.8	13	
140	Mast cell chymase: an indispensable instrument in the pathological symphony of idiopathic pulmonary fibrosis?. <i>Histology and Histopathology</i> , 2013 , 28, 691-9	1.4	13	
139	Prevalence of Mental Disorders and Impact on Quality of Life in Patients With Pulmonary Arterial Hypertension. <i>Frontiers in Psychiatry</i> , 2021 , 12, 667602	5	13	
138	Protection against pressure overload-induced right heart failure by uncoupling protein 2 silencing. <i>Cardiovascular Research</i> , 2019 , 115, 1217-1227	9.9	12	

137	Temporary treatment interruptions with oral selexipag in pulmonary arterial hypertension: Insights from the Prostacyclin (PGI) Receptor Agonist in Pulmonary Arterial Hypertension (GRIPHON) study. Journal of Heart and Lung Transplantation, 2018, 37, 401-408	5.8	12
136	The prognostic significance of inspiratory capacity in pulmonary arterial hypertension. <i>Respiration</i> , 2014 , 88, 24-30	3.7	12
135	Update on pulmonary hypertension 2009. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2010 , 181, 1020-6	10.2	12
134	Acute hemodynamic effects of nebulized iloprost via the I-neb Adaptive Aerosol Delivery system in pulmonary hypertension. <i>Pulmonary Circulation</i> , 2015 , 5, 162-70	2.7	11
133	Use of responder threshold criteria to evaluate the response to treatment in the phase III CHEST-1 study. <i>Journal of Heart and Lung Transplantation</i> , 2015 , 34, 348-55	5.8	11
132	Nebulization of the acidified sodium nitrite formulation attenuates acute hypoxic pulmonary vasoconstriction. <i>Respiratory Research</i> , 2010 , 11, 81	7.3	11
131	Congestive nephropathy: a neglected entity? Proposal for diagnostic criteria and future perspectives. <i>ESC Heart Failure</i> , 2021 , 8, 183-203	3.7	11
130	Nintedanib in Severe Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018 , 198, 808-810	10.2	11
129	Hemodynamic phenotyping based on exercise catheterization predicts outcome in patients with heart failure and reduced ejection fraction. <i>Journal of Heart and Lung Transplantation</i> , 2017 , 36, 880-88	9 ^{5.8}	10
128	Plasma MMP2/TIMP4 Ratio at Follow-up Assessment Predicts Disease Progression of Idiopathic Pulmonary Arterial Hypertension. <i>Lung</i> , 2017 , 195, 489-496	2.9	10
127	A retrospective review of long anterior fusions to the sacrum. Spine Journal, 2011, 11, 290-4	4	10
126	Stem cell-mediated natural tissue engineering. <i>Journal of Cellular and Molecular Medicine</i> , 2011 , 15, 52-	63 .6	10
125	2015 ESC/ERS GUIDELINES FOR THE DIAGNOSIS AND TREATMENT OF PULMONARY HYPERTENSION. <i>Russian Journal of Cardiology</i> , 2016 , 5-64	1.3	10
124	Individual dose adjustment of riociguat in patients with pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension. <i>Respiratory Medicine</i> , 2017 , 129, 124-129	4.6	9
123	Use of clinically relevant responder threshold criteria to evaluate the response to treatment in the phase III PATENT-1 study. <i>Journal of Heart and Lung Transplantation</i> , 2015 , 34, 338-47	5.8	9
122	Ventricular tachycardias in patients with pulmonary hypertension: an underestimated prevalence? A prospective clinical study. <i>Herzschrittmachertherapie Und Elektrophysiologie</i> , 2015 , 26, 155-62	0.8	9
121	Heart Rate Variability is Related to Disease Severity in Children and Young Adults with Pulmonary Hypertension. <i>Frontiers in Pediatrics</i> , 2015 , 3, 63	3.4	9
120	Electrophysiological studies in patients with pulmonary hypertension: a retrospective investigation. <i>BioMed Research International</i> , 2014 , 2014, 617565	3	9

(2017-2020)

119	Impact of SARS-CoV-2 pandemic on pulmonary hypertension out-patient clinics in Germany: a multi-centre study. <i>Pulmonary Circulation</i> , 2020 , 10, 2045894020941682	2.7	9
118	Initial combination therapy with ambrisentan + tadalafil on pulmonary arterial hypertension-related hospitalization in the AMBITION trial. <i>Journal of Heart and Lung Transplantation</i> , 2019 , 38, 194-202	5.8	9
117	Clinical outcomes stratified by baseline functional class after initial combination therapy for pulmonary arterial hypertension. <i>Respiratory Research</i> , 2019 , 20, 208	7.3	8
116	Right ventricular function in pulmonary (arterial) hypertension. <i>Herz</i> , 2019 , 44, 509-516	2.6	8
115	Pulmonary Hypertension in Adults with Congenital Heart Disease: Real-World Data from the International COMPERA-CHD Registry. <i>Journal of Clinical Medicine</i> , 2020 , 9,	5.1	8
114	Survival with sildenafil and inhaled iloprost in a cohort with pulmonary hypertension: an observational study. <i>BMC Pulmonary Medicine</i> , 2016 , 16, 5	3.5	8
113	Animal models of pulmonary hypertension: role in translational research. <i>Drug Discovery Today: Disease Models</i> , 2010 , 7, 89-97	1.3	8
112	Synergism of alveolar endotoxin "priming" and intravascular exotoxin challenge in lung injury. <i>American Journal of Respiratory and Critical Care Medicine</i> , 1996 , 154, 460-8	10.2	8
111	Prostanoids and phosphodiesterase inhibitors in experimental pulmonary hypertension. <i>Current Topics in Developmental Biology</i> , 2005 , 67, 251-84	5.3	8
110	The Clinical Significance of HbA1c in Operable Chronic Thromboembolic Pulmonary Hypertension. <i>PLoS ONE</i> , 2016 , 11, e0152580	3.7	8
109	Modulating cGMP to treat lung diseases. Handbook of Experimental Pharmacology, 2009, 469-83	3.2	8
108	Multibeat Right Ventricular-Arterial Coupling during a Positive Acute Vasoreactivity Test. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019 , 199, e41-e42	10.2	8
107	Hypoxic Pulmonary Vasoconstriction II riggered by an Increase in Reactive Oxygen Species?. <i>Novartis Foundation Symposium</i> , 196-213		8
106	Resistant hypertension: medical management and alternative therapies. <i>Cardiology Clinics</i> , 2015 , 33, 75-87	2.5	7
105	Effects of carbon monoxide-releasing molecules on pulmonary vasoreactivity in isolated perfused lungs. <i>Journal of Applied Physiology</i> , 2016 , 120, 271-81	3.7	7
104	Enhanced circulating levels of CD3 cells-derived extracellular vesicles in different forms of pulmonary hypertension. <i>Pulmonary Circulation</i> , 2019 , 9, 2045894019864357	2.7	7
103	Measurement of exhaled hydrogen peroxide from rabbit lungs. <i>Biological Chemistry</i> , 2004 , 385, 259-64	4.5	7
102	Macitentan for inoperable chronic thromboembolic pulmonary hypertension (CTEPH): results from the randomised controlled MERIT study 2017 ,		7

101	COMPERA 2.0: A refined 4-strata risk assessment model for pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2021 ,	13.6	7
100	Association of right atrial conduit phase with right ventricular lusitropic function in pulmonary hypertension. <i>International Journal of Cardiovascular Imaging</i> , 2020 , 36, 633-642	2.5	7
99	Response by Tello et al to Letter Regarding Article, "Validation of the Tricuspid Annular Plane Systolic Excursion/Systolic Pulmonary Artery Pressure Ratio for the Assessment of Right Ventricular-Arterial Coupling in Severe Pulmonary Hypertension". <i>Circulation: Cardiovascular</i>	3.9	7
98	Imaging, 2019 , 12, e010059 Pulmonary Hypertension in Acute and Chronic High Altitude Maladaptation Disorders. <i>International Journal of Environmental Research and Public Health</i> , 2021 , 18,	4.6	7
97	Hypoxic pulmonary vasoconstriction in isolated mouse pulmonary arterial vessels. <i>Experimental Physiology</i> , 2018 , 103, 1185-1191	2.4	7
96	Hypoxic pulmonary vasoconstrictiontriggered by an increase in reactive oxygen species?. <i>Novartis Foundation Symposium</i> , 2006 , 272, 196-208; discussion 208-17		7
95	Interaction of ambrisentan and phenprocoumon in patients with pulmonary hypertension. <i>Pulmonary Pharmacology and Therapeutics</i> , 2014 , 28, 87-89	3.5	6
94	Subthreshold doses of nebulized prostacyclin and rolipram synergistaically protect against lung ischemia-reperfusion. <i>Transplantation</i> , 2003 , 75, 814-21	1.8	6
93	Chronic intratracheal application of the soluble guanylyl cyclase stimulator BAY 41-8543 ameliorates experimental pulmonary hypertension. <i>Oncotarget</i> , 2017 , 8, 29613-29624	3.3	6
92	Temporal trends in pulmonary arterial hypertension: Results from the COMPERA registry. <i>European Respiratory Journal</i> , 2021 ,	13.6	6
91	Risk assessment in severe pulmonary hypertension due to interstitial lung disease. <i>Journal of Heart and Lung Transplantation</i> , 2020 , 39, 1118-1125	5.8	6
90	Prognostic Relevance of Nonsustained Ventricular Tachycardia in Patients with Pulmonary Hypertension. <i>BioMed Research International</i> , 2016 , 2016, 1327265	3	6
89	Impairment of hypoxic pulmonary vasoconstriction in acute respiratory distress syndrome. <i>European Respiratory Review</i> , 2021 , 30,	9.8	6
88	Effect of nitric oxide synthase (NOS) inhibition on macro- and microcirculation in a model of rat endotoxic shock. <i>Thrombosis and Haemostasis</i> , 2006 , 95, 720-7	7	6
87	The prognostic relevance of oxygen uptake in inoperable chronic thromboembolic pulmonary hypertension. <i>Clinical Respiratory Journal</i> , 2017 , 11, 682-690	1.7	5
86	Integrating Data From Randomized Controlled Trials and Observational Studies to Assess Survival in Rare Diseases. <i>Circulation: Cardiovascular Quality and Outcomes</i> , 2019 , 12, e005095	5.8	5
85	Altered proteasome function in right ventricular hypertrophy. Cardiovascular Research, 2020, 116, 406-4	1959	5
84	Flow rate variance of a fully implantable pump for the delivery of intravenous treprostinil in pulmonary arterial hypertension. <i>Pulmonary Circulation</i> , 2020 , 10, 2045894020910136	2.7	5

(2016-2016)

83	Circulating Angiopoietin-1 Is Not a Biomarker of Disease Severity or Prognosis in Pulmonary Hypertension. <i>PLoS ONE</i> , 2016 , 11, e0165982	3.7	5
82	Advanced risk stratification of intermediate risk group in pulmonary arterial hypertension. <i>Pulmonary Circulation</i> , 2020 , 10, 2045894020961739	2.7	5
81	Relevance of angiopoietin-2 and soluble P-selectin levels in patients with pulmonary arterial hypertension receiving combination therapy with oral treprostinil: a FREEDOM-C2 biomarker substudy. <i>Pulmonary Circulation</i> , 2016 , 6, 516-523	2.7	5
80	Evaluating Systolic and Diastolic Cardiac Function in Rodents Using Microscopic Computed Tomography. <i>Circulation: Cardiovascular Imaging</i> , 2018 , 11, e007653	3.9	5
79	Poor sleep quality is associated with exercise limitation in precapillary pulmonary hypertension. BMC Pulmonary Medicine, 2015 , 15, 11	3.5	4
78	Physical Activity and Mental Health of Patients with Pulmonary Hypertension during the COVID-19 Pandemic. <i>Journal of Clinical Medicine</i> , 2020 , 9,	5.1	4
77	Evaluation of pulmonary hypertension by right heart catheterisation: does timing matter?. European Respiratory Journal, 2020 , 56,	13.6	4
76	Influence of gender in monocrotaline and chronic hypoxia induced pulmonary hypertension in obese rats and mice. <i>Respiratory Research</i> , 2020 , 21, 136	7.3	4
75	NADPH oxidase subunit NOXO1 is a target for emphysema treatment in COPD. <i>Nature Metabolism</i> , 2020 , 2, 532-546	14.6	4
74	Cytochrome P450 epoxygenase-derived 5,6-epoxyeicosatrienoic acid relaxes pulmonary arteries in normoxia but promotes sustained pulmonary vasoconstriction in hypoxia. <i>Acta Physiologica</i> , 2020 , 230, e13521	5.6	4
73	Switching inhaled iloprost formulations in patients with pulmonary arterial hypertension: the VENTASWITCH Trial. <i>Pulmonary Circulation</i> , 2018 , 8, 2045894018798921	2.7	4
72	Reply to Bogaard: Emphysema Is-at the Most-Only a Mild Phenotype in the Sugen/Hypoxia Rat Model of Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019 , 200, 1450-1452	10.2	4
71	Long-term Riociguat Treatment in Inoperable and Persistent/Recurrent CTEPH Patients in WHO Functional Class (FC) I/II Versus FC III/IV at Baseline: Results From the 16-Week Phase III CHEST-1 Study and CHEST-2 Open-Label Extension. <i>Chest</i> , 2014 , 145, 535B	5.3	4
70	Response to letters regarding article, "Anticoagulation and survival in pulmonary arterial hypertension: results from the Comparative, Prospective Registry of Newly Initiated Therapies for Pulmonary Hypertension (COMPERA)". <i>Circulation</i> , 2014 , 130, e110-2	16.7	4
69	Cologne Consensus Conference on pulmonary hypertension. <i>International Journal of Cardiology</i> , 2011 , 154 Suppl 1, S1-2	3.2	4
68	Right ventricular pressure-volume loop shape and systolic pressure change in pulmonary hypertension. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2021 , 320, L715-l	<i>.7</i> 285	4
67	Impact of SARS-CoV-2-Pandemic on Mental Disorders and Quality of Life in Patients With Pulmonary Arterial Hypertension. <i>Frontiers in Psychiatry</i> , 2021 , 12, 668647	5	4
66	Heart rate response during 6-minute walking testing predicts outcome in operable chronic thromboembolic pulmonary hypertension. <i>BMC Pulmonary Medicine</i> , 2016 , 16, 96	3.5	4

65	COPD-associated pulmonary hypertension: clinical implications and current methods for treatment. <i>Expert Review of Respiratory Medicine</i> , 2016 , 10, 755-66	3.8	4
64	Comparison of MRI and VQ-SPECT as a Screening Test for Patients With Suspected CTEPH: CHANGE-MRI Study Design and Rationale. <i>Frontiers in Cardiovascular Medicine</i> , 2020 , 7, 51	5.4	4
63	Pulmonary function and diffusion capacity are associated with pulmonary arterial systolic pressure in the general population: The Rotterdam Study. <i>Respiratory Medicine</i> , 2017 , 132, 50-55	4.6	3
62	Circulating Apoptotic Signals During Acute and Chronic Exposure to High Altitude in Kyrgyz Population. <i>Frontiers in Physiology</i> , 2019 , 10, 54	4.6	3
61	Right ventricular dyssynchrony: from load-independent right ventricular function to wall stress in severe pulmonary arterial hypertension. <i>Pulmonary Circulation</i> , 2020 , 10, 2045894020925759	2.7	3
60	Measures of subclinical cardiac dysfunction and increased filling pressures associate with pulmonary arterial pressure in the general population: results from the population-based Rotterdam Study. <i>European Journal of Epidemiology</i> , 2018 , 33, 403-413	12.1	3
59	Acute response to rapid iloprost inhalation using the BreelibIhebulizer in pulmonary arterial hypertension: the BreelibIacute study. <i>Pulmonary Circulation</i> , 2019 , 9, 2045894019875342	2.7	3
58	Long-term Riociguat Treatment in PAH Patients in WHO Functional Class (FC) I/II Versus FC III/IV at Baseline: Results From the 12-Week Phase III PATENT-1 Study and PATENT-2 Open-Label Extension. <i>Chest</i> , 2014 , 145, 513A	5.3	3
57	Individualized Dosing of Selexipag Based on Tolerability in the GRIPHON Study Shows Consistent Efficacy and Safety in Patients With Pulmonary Arterial Hypertension (PAH). <i>Chest</i> , 2015 , 148, 961A	5.3	3
56	Riociguat for the Treatment of Pulmonary Arterial Hypertension (PAH): 1-Year Results from the PATENT-2 Long-term Extension Study. <i>Chest</i> , 2013 , 144, 1024A	5.3	3
55	Riociguat for the Treatment of Chronic Thromboembolic Pulmonary Hypertension (CTEPH): 1-Year Results from the CHEST-2 Long-term Extension Study. <i>Chest</i> , 2013 , 144, 1023A	5.3	3
54	Diagnostik und Therapie der pulmonalen Hypertonie. <i>Kardiologe</i> , 2010 , 4, 189-207	0.6	3
53	TORREY, a Phase 2 study to evaluate the efficacy and safety of inhaled seralutinib for the treatment of pulmonary arterial hypertension. <i>Pulmonary Circulation</i> , 2021 , 11, 20458940211057071	2.7	3
52	Validity of echocardiographic tricuspid regurgitation gradient to screen for new definition of pulmonary hypertension. <i>EClinicalMedicine</i> , 2021 , 34, 100822	11.3	3
51	Protein expression profiling suggests relevance of noncanonical pathways in isolated pulmonary embolism. <i>Blood</i> , 2021 , 137, 2681-2693	2.2	3
50	CILP1 as a biomarker for right ventricular maladaptation in pulmonary hypertension. <i>European Respiratory Journal</i> , 2021 , 57,	13.6	3
49	A novel non-invasive and echocardiography-derived method for quantification of right ventricular pressure-volume loops. <i>European Heart Journal Cardiovascular Imaging</i> , 2021 ,	4.1	3
48	Inspiratory capacity is not altered in operable chronic thromboembolic pulmonary hypertension. <i>Pulmonary Circulation</i> , 2017 , 7, 543-546	2.7	2

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47	Is PKM2 Phosphorylation a Prerequisite for Oligomer Disassembly in Pulmonary Arterial Hypertension?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019 , 200, 1550-1554	10.2	2
46	Pulmonary Vascular Pressure Response to Acute Cold Exposure in Kyrgyz Highlanders. <i>High Altitude Medicine and Biology</i> , 2019 , 20, 375-382	1.9	2
45	Effect of Selexipag on Morbidity/Mortality in Pulmonary Arterial Hypertension: Results of the GRIPHON Study. <i>Journal of Heart and Lung Transplantation</i> , 2015 , 34, S163	5.8	2
44	Genetic Deletion of p66shc and/or Cyclophilin D Results in Decreased Pulmonary Vascular Tone. <i>Cardiovascular Research</i> , 2020 ,	9.9	2
43	Beyond interleukin-6 in right ventricular function: Evidence for another biomarker. <i>Journal of Heart and Lung Transplantation</i> , 2018 , 37, 674-675	5.8	2
42	Riociguat in Combination With Prostacyclin Analogs for the Treatment of Pulmonary Arterial Hypertension (PAH): A Subgroup Analysis of the PATENT Studies. <i>Chest</i> , 2015 , 148, 922A	5.3	2
41	Riociguat dose titration in patients with chronic thromboembolic pulmonary hypertension (CTEPH) or pulmonary arterial hypertension (PAH). <i>BMC Pharmacology</i> , 2009 , 9,		2
40	Effects Of Inhaled Aviptadil (Vasoactive Intestinal Peptide) In Patients With Pulmonary Arterial Hypertension (PAH) 2010 ,		2
39	Initial combination therapy with ambrisentan (AMB) and tadalafil (TAD) in treatment naWe patients with pulmonary arterial hypertension (PAH): Efficacy and safety in the AMBITION study intent to treat (ITT) population 2015 ,		2
38	IRAG1 Deficient Mice Develop PKG1 Dependent Pulmonary Hypertension. <i>Cells</i> , 2020 , 9,	7.9	2
37	Effects of macitentan and tadalafil monotherapy or their combination on the right ventricle and plasma metabolites in pulmonary hypertensive rats. <i>Pulmonary Circulation</i> , 2020 , 10, 204589402094728	3 3 .7	2
36	Genetic Deficiency and Pharmacological Stabilization of Mast Cells Ameliorate Pressure Overload-Induced Maladaptive Right Ventricular Remodeling in Mice. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	2
35	Selexipag for the treatment of pulmonary arterial hypertension. <i>Expert Opinion on Pharmacotherapy</i> , 2016 , 17, 1825-34	4	2
34	Amelioration of elastase-induced lung emphysema and reversal of pulmonary hypertension by pharmacological iNOS inhibition in mice. <i>British Journal of Pharmacology</i> , 2021 , 178, 152-171	8.6	2
33	Prevalence of Mental Disorders in Patients With Chronic Thromboembolic Pulmonary Hypertension <i>Frontiers in Psychiatry</i> , 2022 , 13, 821466	5	2
32	Cardiopulmonary haemodynamics in portopulmonary hypertension. <i>Lancet Respiratory Medicine,the</i> , 2019 , 7, 556-558	35.1	1
31	Reply to "a pediatric perspective on the TAPSE/PASP ratio in pulmonary arterial hypertension". <i>International Journal of Cardiology</i> , 2019 , 278, 240-241	3.2	1
30	is a Promising Therapeutic Option for Treatment of Pulmonary Hypertension due to the Potent Anti-Proliferative and Vasorelaxant Properties. <i>Medicina (Lithuania)</i> , 2020 , 56,	3.1	1

29	SPARCL1 as a biomarker of maladaptive right ventricular remodelling in pulmonary hypertension. <i>Biomarkers</i> , 2020 , 25, 290-295	2.6	1
28	Impact of Pulmonary Arterial Hypertension on Employment, Work Productivity, and Quality of Life - Results of a Cross-Sectional Multi-Center Study <i>Frontiers in Psychiatry</i> , 2021 , 12, 781532	5	1
27	Clinical Relevance of Right Atrial Functional Response to Treatment in Pulmonary Arterial Hypertension <i>Frontiers in Cardiovascular Medicine</i> , 2021 , 8, 775039	5.4	1
26	Subcutaneous treprostinil: a new treatment for chronic thromboembolic pulmonary hypertension?. <i>Lancet Respiratory Medicine,the</i> , 2019 , 7, 191-193	35.1	1
25	EXPRESS: Switching to riociguat: A potential treatment strategy for the management of CTEPH and PAH. <i>Pulmonary Circulation</i> , 2019 , 2045894019837849	2.7	1
24	PINK1-mediated Mitophagy Contributes to Pulmonary Vascular Remodeling in Pulmonary Hypertension. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2021 , 65, 226-228	5.7	1
23	Osteopontin and galectin-3 as biomarkers of maladaptive right ventricular remodeling în pulmonary hypertension. <i>Biomarkers in Medicine</i> , 2021 , 15, 1021-1034	2.3	1
22	Exercise Hemodynamic Profiling Is Associated With Outcome in Patients Undergoing Percutaneous Mitral Valve Repair. <i>Circulation: Cardiovascular Interventions</i> , 2021 , 14, e010453	6	1
21	Phosphodiesterase-5 Inhibitors in Pulmonary Arterial Hypertension105-125		1
20	Childhood Maltreatment, Mental Well-Being, and Healthy Lifestyle in Patients With Chronic Thromboembolic Pulmonary Hypertension <i>Frontiers in Psychiatry</i> , 2022 , 13, 821468	5	1
19	Follow-Up (Measurement) of Corrected QT Interval in Adult Patients before and after Lung Transplantation. <i>BioMed Research International</i> , 2017 , 2017, 4519796	3	О
18	Diagnostik und Therapie der pulmonalen Hypertonie: Europßche Leitlinien 2009. <i>Deutsche Medizinische Wochenschrift</i> , 2010 , 135, e2-e15	O	О
17	Exercise hemodynamics in heart failure patients with preserved and mid-range ejection fraction: key role of the right heart. <i>Clinical Research in Cardiology</i> , 2021 , 1	6.1	0
16	Medical treatment of pulmonary hypertension in adults with congenital heart disease: updated and extended results from the International COMPERA-CHD Registry <i>Cardiovascular Diagnosis and Therapy</i> , 2021 , 11, 1255-1268	2.6	O
15	Relevance of Cor Pulmonale in COPD With and Without Pulmonary Hypertension: A Retrospective Cohort Study <i>Frontiers in Cardiovascular Medicine</i> , 2022 , 9, 826369	5.4	0
14	Tidal Carbon Dioxide as a Prognostic Feature in Inoperable Chronic Thromboembolic Pulmonary Hypertension. <i>Annals of the American Thoracic Society</i> , 2017 , 14, 1603-1604	4.7	
13	Therapie der pulmonal arteriellen Hypertonie. Klinikarzt, 2017 , 46, 374-381	Ο	
12	Aktuelle Therapie der pulmonal-arteriellen Hypertonie. <i>Pneumologe</i> , 2010 , 7, 192-199	0.1	

LIST OF PUBLICATIONS

11	Prevention of pulmonary vascular and myocardial remodeling by the combined tyrosine and serine-/threonine kinase inhibitor, sorafenib, in pulmonary hypertension and right heart failure. <i>European Respiratory Review</i> , 2008 , 17, 72-73	9.8
10	Sildenafil for lung fibrosis and pulmonary hypertension. <i>Lancet, The</i> , 2003 , 361, 263	40
9	Schwere pulmonale Hypertonie - Vasodilative Therapie in der Lungenstrombahn. <i>Pneumologie</i> , 2000 , 54, 160-169	0.5
8	Childhood Trauma in Patients With PAH-Prevalence, Impact on QoL, and Mental Health-A Preliminary Report <i>Frontiers in Psychiatry</i> , 2022 , 13, 812862	5
7	NO pathway and phosphodiesterase inhibitors in pulmonary arterial hypertension 2004 , 163-168	
6	Classical transient receptor potential channel 6 (TRPC6) is essential for ischemia-reperfusion injury of the lung. <i>FASEB Journal</i> , 2010 , 24, 591.2	0.9
5	Pulmonalarterielle Hypertonie bei angeborenen Herzfehlern: Problemstellung und Versorgungslage. <i>Deutsche Medizinische Wochenschrift</i> , 2013 , 138, e27-e27	0
4	Reply to "Risk stratification in PH associated with interstitial lung disease: The Holy Grail?". <i>Journal of Heart and Lung Transplantation</i> , 2021 , 40, 317	5.8
3	Response to: Comment on "Effect of Riociguat and Sildenafil on Right Heart Remodeling and Function in Pressure Overload Induced Model of Pulmonary Arterial Banding". <i>BioMed Research International</i> , 2018 , 2018, 7491284	3
2	Deficiency of Axl aggravates pulmonary arterial hypertension via BMPR2. <i>Communications Biology</i> , 2021 , 4, 1002	6.7
1	Metacognitions in Patients With Frequent Mental Disorders After Diagnosis of Pulmonary Arterial Hypertension <i>Frontiers in Psychiatry</i> , 2022 , 13, 812812	5