## Ardeschir Ghofrani

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3367351/publications.pdf

Version: 2024-02-01

510 papers

54,821 citations

98 h-index 219 g-index

580 all docs

580 docs citations

580 times ranked

25311 citing authors

#	Article	IF	CITATIONS
1	2015 ESC/ERS Guidelines for the diagnosis and treatment of pulmonary hypertension. European Heart Journal, 2016, 37, 67-119.	1.0	5,074
2	Updated Clinical Classification of Pulmonary Hypertension. Journal of the American College of Cardiology, 2013, 62, D34-D41.	1.2	2,865
3	2015 ESC/ERS Guidelines for the diagnosis and treatment of pulmonary hypertension. European Respiratory Journal, 2015, 46, 903-975.	3.1	2,415
4	Sildenafil Citrate Therapy for Pulmonary Arterial Hypertension. New England Journal of Medicine, 2005, 353, 2148-2157.	13.9	2,237
5	Inhaled Iloprost for Severe Pulmonary Hypertension. New England Journal of Medicine, 2002, 347, 322-329.	13.9	1,626
6	Macitentan and Morbidity and Mortality in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2013, 369, 809-818.	13.9	1,168
7	Riociguat for the Treatment of Chronic Thromboembolic Pulmonary Hypertension. New England Journal of Medicine, 2013, 369, 319-329.	13.9	1,144
8	Riociguat for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2013, 369, 330-340.	13.9	1,120
9	Ambrisentan for the Treatment of Pulmonary Arterial Hypertension. Circulation, 2008, 117, 3010-3019.	1.6	967
10	Tadalafil Therapy for Pulmonary Arterial Hypertension. Circulation, 2009, 119, 2894-2903.	1.6	956
10	Tadalafil Therapy for Pulmonary Arterial Hypertension. Circulation, 2009, 119, 2894-2903.  Reversal of experimental pulmonary hypertension by PDGF inhibition. Journal of Clinical Investigation, 2005, 115, 2811-2821.	1.6 3.9	956 917
	Reversal of experimental pulmonary hypertension by PDGF inhibition. Journal of Clinical Investigation,		
11	Reversal of experimental pulmonary hypertension by PDGF inhibition. Journal of Clinical Investigation, 2005, 115, 2811-2821.  Initial Use of Ambrisentan plus Tadalafil in Pulmonary Arterial Hypertension. New England Journal of	3.9	917
11 12	Reversal of experimental pulmonary hypertension by PDGF inhibition. Journal of Clinical Investigation, 2005, 115, 2811-2821.  Initial Use of Ambrisentan plus Tadalafil in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 834-844.  Selexipag for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine,	3.9	917 906
11 12 13	Reversal of experimental pulmonary hypertension by PDGF inhibition. Journal of Clinical Investigation, 2005, 115, 2811-2821.  Initial Use of Ambrisentan plus Tadalafil in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 834-844.  Selexipag for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 2522-2533.  Sildenafil for treatment of lung fibrosis and pulmonary hypertension: a randomised controlled trial.	3.9 13.9 13.9	917 906 790
11 12 13	Reversal of experimental pulmonary hypertension by PDGF inhibition. Journal of Clinical Investigation, 2005, 115, 2811-2821.  Initial Use of Ambrisentan plus Tadalafil in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 834-844.  Selexipag for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 2522-2533.  Sildenafil for treatment of lung fibrosis and pulmonary hypertension: a randomised controlled trial. Lancet, The, 2002, 360, 895-900.	3.9 13.9 13.9 6.3	917 906 790 720
11 12 13 14	Reversal of experimental pulmonary hypertension by PDGF inhibition. Journal of Clinical Investigation, 2005, 115, 2811-2821.  Initial Use of Ambrisentan plus Tadalafil in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 834-844.  Selexipag for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 2522-2533.  Sildenafil for treatment of lung fibrosis and pulmonary hypertension: a randomised controlled trial. Lancet, The, 2002, 360, 895-900.  Chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2019, 53, 1801915.	3.9 13.9 13.9 6.3	917 906 790 720

#	Article	IF	CITATIONS
19	Mortality in pulmonary arterial hypertension: prediction by the 2015 European pulmonary hypertension guidelines risk stratification model. European Respiratory Journal, 2017, 50, 1700740.	3.1	489
20	Imatinib Mesylate as Add-on Therapy for Pulmonary Arterial Hypertension. Circulation, 2013, 127, 1128-1138.	1.6	482
21	Sildenafil: from angina to erectile dysfunction to pulmonary hypertension and beyond. Nature Reviews Drug Discovery, 2006, 5, 689-702.	21.5	471
22	Updated Evidence-Based Treatment Algorithm in Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2009, 54, S78-S84.	1.2	463
23	Combination Therapy with Oral Sildenafil and Inhaled Iloprost for Severe Pulmonary Hypertension. Annals of Internal Medicine, 2002, 136, 515.	2.0	446
24	Imatinib for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2005, 353, 1412-1413.	13.9	440
25	Inhaled Prostacyclin and Iloprost in Severe Pulmonary Hypertension Secondary to Lung Fibrosis. American Journal of Respiratory and Critical Care Medicine, 1999, 160, 600-607.	2.5	369
26	Oral sildenafil as long-term adjunct therapy to inhaled iloprost in severe pulmonary arterial hypertension. Journal of the American College of Cardiology, 2003, 42, 158-164.	1.2	359
27	Elderly patients diagnosed with idiopathic pulmonary arterial hypertension: Results from the COMPERA registry. International Journal of Cardiology, 2013, 168, 871-880.	0.8	357
28	Aerosolized Prostacyclin and Iloprost in Severe Pulmonary Hypertension. Annals of Internal Medicine, 1996, 124, 820.	2.0	347
29	Imatinib in Pulmonary Arterial Hypertension Patients with Inadequate Response to Established Therapy. American Journal of Respiratory and Critical Care Medicine, 2010, 182, 1171-1177.	2.5	331
30	Hypoxia-Dependent Regulation of Nonphagocytic NADPH Oxidase Subunit NOX4 in the Pulmonary Vasculature. Circulation Research, 2007, 101, 258-267.	2.0	317
31	Anticoagulation and Survival in Pulmonary Arterial Hypertension. Circulation, 2014, 129, 57-65.	1.6	317
32	A comparison of the acute hemodynamic effects of inhaled nitric oxide and aerosolized iloprost in primary pulmonary hypertension. Journal of the American College of Cardiology, 2000, 35, 176-182.	1.2	296
33	Immune and Inflammatory Cell Involvement in the Pathology of Idiopathic Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2012, 186, 897-908.	2.5	296
34	Inducible NOS Inhibition Reverses Tobacco-Smoke-Induced Emphysema and Pulmonary Hypertension in Mice. Cell, 2011, 147, 293-305.	13.5	293
35	ERS statement on chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2021, 57, 2002828.	3.1	287
36	Riociguat for Patients With Pulmonary Hypertension Caused by Systolic Left Ventricular Dysfunction. Circulation, 2013, 128, 502-511.	1.6	286

#	Article	IF	Citations
37	Identification of rare sequence variation underlying heritable pulmonary arterial hypertension. Nature Communications, 2018, 9, 1416.	5.8	279
38	Classical transient receptor potential channel 6 (TRPC6) is essential for hypoxic pulmonary vasoconstriction and alveolar gas exchange. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 19093-19098.	3.3	273
39	Sildenafil Increased Exercise Capacity during Hypoxia at Low Altitudes and at Mount Everest Base Camp. Annals of Internal Medicine, 2004, 141, 169.	2.0	271
40	Sildenafil for Long-Term Treatment of Nonoperable Chronic Thromboembolic Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2003, 167, 1139-1141.	2.5	265
41	Riociguat for the treatment of chronic thromboembolic pulmonary hypertension: a long-term extension study (CHEST-2). European Respiratory Journal, 2015, 45, 1293-1302.	3.1	247
42	Long-term Treatment With Sildenafil Citrate in Pulmonary Arterial Hypertension. Chest, 2011, 140, 1274-1283.	0.4	237
43	Chronic Sildenafil Treatment Inhibits Monocrotaline-induced Pulmonary Hypertension in Rats. American Journal of Respiratory and Critical Care Medicine, 2004, 169, 39-45.	2.5	230
44	Long-Term Ambrisentan Therapy for the Treatment of Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2009, 54, 1971-1981.	1.2	227
45	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. Circulation: Cardiovascular Imaging, 2019, 12, e009047.	1.3	222
46	The Giessen Pulmonary Hypertension Registry: Survival in pulmonary hypertension subgroups. Journal of Heart and Lung Transplantation, 2017, 36, 957-967.	0.3	221
47	Expression and function of soluble guanylate cyclase in pulmonary arterial hypertension. European Respiratory Journal, 2008, 32, 881-891.	3.1	216
48	First acute haemodynamic study of soluble guanylate cyclase stimulator riociguat in pulmonary hypertension. European Respiratory Journal, 2009, 33, 785-792.	3.1	216
49	Riociguat for chronic thromboembolic pulmonary hypertension and pulmonary arterial hypertension: a phase II study. European Respiratory Journal, 2010, 36, 792-799.	3.1	212
50	2015 ESC/ERS Guidelines for the Diagnosis and Treatment of Pulmonary Hypertension. Revista Espanola De Cardiologia (English Ed ), 2016, 69, 177.	0.4	210
51	Activation of Soluble Guanylate Cyclase Reverses Experimental Pulmonary Hypertension and Vascular Remodeling. Circulation, 2006, 113, 286-295.	1.6	208
52	Inhibition of MicroRNA-17 Improves Lung and Heart Function in Experimental Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2012, 185, 409-419.	2.5	206
53	Stress Doppler Echocardiography in Relatives of Patients With Idiopathic and Familial Pulmonary Arterial Hypertension. Circulation, 2009, 119, 1747-1757.	1.6	205
54	Macitentan for the treatment of inoperable chronic thromboembolic pulmonary hypertension (MERIT-1): results from the multicentre, phase 2, randomised, double-blind, placebo-controlled study. Lancet Respiratory Medicine, the, 2017, 5, 785-794.	5.2	201

#	Article	IF	CITATIONS
55	Sildenafil treatment for portopulmonary hypertension. European Respiratory Journal, 2006, 28, 563-567.	3.1	199
56	Safety and efficacy of exercise training in various forms of pulmonary hypertension. European Respiratory Journal, 2012, 40, 84-92.	3.1	199
57	Regulation of hypoxic pulmonary vasoconstriction: basic mechanisms. European Respiratory Journal, 2008, 32, 1639-1651.	3.1	184
58	Bosentan added to sildenafil therapy in patients with pulmonary arterial hypertension. European Respiratory Journal, 2015, 46, 405-413.	3.1	184
59	Upregulation of NAD(P)H oxidase 1 in hypoxia activates hypoxia-inducible factor 1 via increase in reactive oxygen species. Free Radical Biology and Medicine, 2004, 36, 1279-1288.	1.3	183
60	Balloon pulmonary angioplasty in chronic thromboembolic pulmonary hypertension. European Respiratory Review, 2017, 26, 160119.	3.0	183
61	Balloon pulmonary angioplasty for inoperable patients with chronic thromboembolic pulmonary hypertension: the initial German experience. European Respiratory Journal, 2017, 49, 1602409.	3.1	178
62	Riociguat for the treatment of pulmonary arterial hypertension: a long-term extension study (PATENT-2). European Respiratory Journal, 2015, 45, 1303-1313.	3.1	174
63	A Functional Single-Nucleotide Polymorphism in the <i>TRPC6 &lt; /i&gt; Gene Promoter Associated With Idiopathic Pulmonary Arterial Hypertension. Circulation, 2009, 119, 2313-2322.</i>	1.6	173
64	Bronchoscopic surfactant administration in patients with severe adult respiratory distress syndrome and sepsis American Journal of Respiratory and Critical Care Medicine, 1996, 154, 57-62.	2.5	171
65	Vascular Receptor Autoantibodies in Pulmonary Arterial Hypertension Associated with Systemic Sclerosis. American Journal of Respiratory and Critical Care Medicine, 2014, 190, 808-817.	2.5	170
66	Long-term treatment with sildenafil in chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2007, 30, 922-927.	3.1	163
67	Activation of TRPC6 channels is essential for lung ischaemia–reperfusion induced oedema in mice. Nature Communications, 2012, 3, 649.	5.8	162
68	Increased levels and reduced catabolism of asymmetric and symmetric dimethylarginines in pulmonary hypertension. FASEB Journal, 2005, 19, 1175-1177.	0.2	158
69	Reserve of Right Ventricular-Arterial Coupling in the Setting of Chronic Overload. Circulation: Heart Failure, 2019, 12, e005512.	1.6	158
70	Relevance of the TAPSE/PASP ratio in pulmonary arterial hypertension. International Journal of Cardiology, 2018, 266, 229-235.	0.8	154
71	Reduced MicroRNA-150 Is Associated with Poor Survival in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2013, 187, 294-302.	2.5	153
72	Effect of Exercise and Respiratory Training on Clinical Progression and Survival in Patients with Severe Chronic Pulmonary Hypertension. Respiration, 2011, 81, 394-401.	1.2	151

#	Article	IF	Citations
73	Phosphodiesterase 1 Upregulation in Pulmonary Arterial Hypertension. Circulation, 2007, 115, 2331-2339.	1.6	139
74	Combined Tyrosine and Serine/Threonine Kinase Inhibition by Sorafenib Prevents Progression of Experimental Pulmonary Hypertension and Myocardial Remodeling. Circulation, 2008, 118, 2081-2090.	1.6	139
75	Favorable Effects of Inhaled Treprostinil in Severe Pulmonary Hypertension. Journal of the American College of Cardiology, 2006, 48, 1672-1681.	1.2	135
76	Initial combination therapy with ambrisentan and tadalafil in connective tissue disease-associated pulmonary arterial hypertension (CTD-PAH): subgroup analysis from the AMBITION trial. Annals of the Rheumatic Diseases, 2017, 76, 1219-1227.	0.5	135
77	Differences in hemodynamic and oxygenation responses to three different phosphodiesterase-5 inhibitors in patients with pulmonary arterial hypertension. Journal of the American College of Cardiology, 2004, 44, 1488-1496.	1.2	134
78	Nitric oxide pathway and phosphodiesterase inhibitors in pulmonary arterial hypertension. Journal of the American College of Cardiology, 2004, 43, S68-S72.	1.2	131
79	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. Lancet Respiratory Medicine,the, 2016, 4, 372-380.	5.2	130
80	Phosphodiesterase inhibitors for the treatment of pulmonary hypertension. European Respiratory Journal, 2008, 32, 198-209.	3.1	129
81	The molecular targets of approved treatments for pulmonary arterial hypertension. Thorax, 2016, 71, 73-83.	2.7	126
82	Pharmacodynamics and Pharmacokinetics of Inhaled Iloprost, Aerosolized by Three Different Devices, in Severe Pulmonary Hypertension. Chest, 2003, 124, 1294-1304.	0.4	124
83	Tadalafil for the Treatment of Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2012, 60, 768-774.	1.2	124
84	COMPERA 2.0: a refined four-stratum risk assessment model for pulmonary arterial hypertension. European Respiratory Journal, 2022, 60, 2102311.	3.1	124
85	Targeting cancer with phosphodiesterase inhibitors. Expert Opinion on Investigational Drugs, 2010, 19, 117-131.	1.9	123
86	Genetic determinants of risk in pulmonary arterial hypertension: international genome-wide association studies and meta-analysis. Lancet Respiratory Medicine, the, 2019, 7, 227-238.	5.2	122
87	Prostacyclin and its analogues in the treatment of pulmonary hypertension. , 2004, 102, 139-153.		119
88	Targeting non-malignant disorders with tyrosine kinase inhibitors. Nature Reviews Drug Discovery, 2010, 9, 956-970.	21.5	118
89	Role of Epidermal Growth Factor Inhibition in Experimental Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2010, 181, 158-167.	2.5	118
90	New Trial Designs and Potential Therapies for Pulmonary Artery Hypertension. Journal of the American College of Cardiology, 2013, 62, D82-D91.	1.2	113

#	Article	IF	Citations
91	RESPITE: switching to riociguat in pulmonary arterial hypertension patients with inadequate response to phosphodiesterase-5 inhibitors. European Respiratory Journal, 2017, 50, 1602425.	3.1	113
92	Simvastatin as a Treatment for Pulmonary Hypertension Trial. American Journal of Respiratory and Critical Care Medicine, 2010, 181, 1106-1113.	2.5	112
93	Role of Src Tyrosine Kinases in Experimental Pulmonary Hypertension. Arteriosclerosis, Thrombosis, and Vascular Biology, 2012, 32, 1354-1365.	1.1	108
94	Pathophysiology and Treatment of High-Altitude Pulmonary Vascular Disease. Circulation, 2015, 131, 582-590.	1.6	108
95	Riociguat for the treatment of pulmonary arterial hypertension associated with connective tissue disease: results from PATENT-1 and PATENT-2. Annals of the Rheumatic Diseases, 2017, 76, 422-426.	0.5	108
96	Current and future treatments of pulmonary arterial hypertension. British Journal of Pharmacology, 2021, 178, 6-30.	2.7	104
97	Long-term safety and efficacy of imatinib in pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2015, 34, 1366-1375.	0.3	103
98	Ultrasonic versus jet nebulization of iloprost in severe pulmonary hypertension. European Respiratory Journal, 2001, 17, 14-19.	3.1	100
99	The Soluble Guanylate Cyclase Stimulator Riociguat Ameliorates Pulmonary Hypertension Induced by Hypoxia and SU5416 in Rats. PLoS ONE, 2012, 7, e43433.	1.1	100
100	Plasma proteome analysis in patients with pulmonary arterial hypertension: an observational cohort study. Lancet Respiratory Medicine, the, 2017, 5, 717-726.	5.2	99
101	Antiremodeling Effects of Iloprost and the Dual-Selective Phosphodiesterase 3/4 Inhibitor Tolafentrine in Chronic Experimental Pulmonary Hypertension. Circulation Research, 2004, 94, 1101-1108.	2.0	97
102	Pulmonary Vascular Disease in the Developing World. Circulation, 2008, 118, 1758-1766.	1.6	97
103	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. Lancet Respiratory Medicine, the, 2016, 4, 361-371.	5.2	97
104	Selexipag for the treatment of connective tissue disease-associated pulmonary arterial hypertension. European Respiratory Journal, 2017, 50, 1602493.	3.1	97
105	Tadalafil monotherapy and as add-on to background bosentan in patients with pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2011, 30, 632-643.	0.3	95
106	Terguride ameliorates monocrotaline-induced pulmonary hypertension in rats. European Respiratory Journal, 2011, 37, 1104-1118.	3.1	93
107	Impact of Mitochondria and NADPH Oxidases on Acute and Sustained Hypoxic Pulmonary Vasoconstriction. American Journal of Respiratory Cell and Molecular Biology, 2006, 34, 505-513.	1.4	90
108	Uncertainties in the Diagnosis and Treatment of Pulmonary Arterial Hypertension. Circulation, 2008, 118, 1195-1201.	1.6	90

#	Article	IF	CITATIONS
109	Mitochondrial Complex IV Subunit 4 Isoform 2 Is Essential for Acute Pulmonary Oxygen Sensing. Circulation Research, 2017, 121, 424-438.	2.0	90
110	Notch1 signalling regulates endothelial proliferation and apoptosis in pulmonary arterial hypertension. European Respiratory Journal, 2016, 48, 1137-1149.	3.1	89
111	Hypoxic vasoconstriction in intact lungs: a role for NADPH oxidase-derived H <sub>2</sub> O <sub>2</sub> ?. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2000, 279, L683-L690.	1.3	87
112	Riociguat for pulmonary arterial hypertension associated with congenital heart disease. Heart, 2015, 101, 1792-1799.	1.2	87
113	Pulmonary Hypertension. Deutsches Ärzteblatt International, 2017, 114, 73-84.	0.6	87
114	Switching to riociguat versus maintenance therapy with phosphodiesterase-5 inhibitors in patients with pulmonary arterial hypertension (REPLACE): a multicentre, open-label, randomised controlled trial. Lancet Respiratory Medicine, the, 2021, 9, 573-584.	<b>5.</b> 2	85
115	Expression and Activity of Phosphodiesterase Isoforms during Epithelial Mesenchymal Transition: The Role of Phosphodiesterase 4. Molecular Biology of the Cell, 2009, 20, 4751-4765.	0.9	84
116	Long-term outcome with intravenous iloprost in pulmonary arterial hypertension. European Respiratory Journal, 2009, 34, 132-137.	3.1	83
117	Anxiety and depression disorders in patients with pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension. Respiratory Research, 2013, 14, 104.	1.4	83
118	Role of the Prostanoid EP4 Receptor in Iloprost-mediated Vasodilatation in Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2008, 178, 188-196.	2.5	82
119	Pulmonary Arterial Hypertension-Related Morbidity Is Prognostic for Mortality. Journal of the American College of Cardiology, 2018, 71, 752-763.	1.2	82
120	Congestive nephropathy: a neglected entity? Proposal for diagnostic criteria and future perspectives. ESC Heart Failure, 2021, 8, 183-203.	1.4	82
121	Riociguat for the treatment of pulmonary hypertension. Expert Opinion on Investigational Drugs, 2011, 20, 567-576.	1.9	81
122	Stimulation of Soluble Guanylate Cyclase Prevents Cigarette Smoke–induced Pulmonary Hypertension and Emphysema. American Journal of Respiratory and Critical Care Medicine, 2014, 189, 1359-1373.	2.5	80
123	Nocturnal periodic breathing in primary pulmonary hypertension. European Respiratory Journal, 2002, 19, 658-663.	3.1	79
124	Fhl-1, a New Key Protein in Pulmonary Hypertension. Circulation, 2008, 118, 1183-1194.	1.6	79
125	Riociguat: Mode of Action and Clinical Development in Pulmonary Hypertension. Chest, 2017, 151, 468-480.	0.4	79
126	ASK1 Inhibition Halts Disease Progression in Preclinical Models of Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2018, 197, 373-385.	2.5	78

#	Article	IF	Citations
127	Classical Transient Receptor Potential Channel 1 in Hypoxia-induced Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2013, 188, 1451-1459.	2.5	77
128	Cyclooxygenase Isoenzyme Localization and mRNA Expression in Rat Lungs. American Journal of Respiratory Cell and Molecular Biology, 1998, 18, 479-488.	1.4	76
129	Inflammation, immunological reaction and role of infection in pulmonary hypertension. Clinical Microbiology and Infection, 2011, 17, 7-14.	2.8	<b>7</b> 5
130	Sleep apnea in precapillary pulmonary hypertension. Sleep Medicine, 2013, 14, 247-251.	0.8	75
131	Novel and Emerging Therapies for Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2014, 189, 394-400.	2.5	<b>7</b> 5
132	Cardiac Magnetic Resonance Imaging-Based Right Ventricular Strain Analysis for Assessment of Coupling and Diastolic Function in Pulmonary Hypertension. JACC: Cardiovascular Imaging, 2019, 12, 2155-2164.	2.3	75
133	Safety and tolerability of bosentan in idiopathic pulmonary fibrosis: an open label study. European Respiratory Journal, 2007, 29, 713-719.	3.1	74
134	Traditional and new composite endpoints inÂheart failure clinical trials: facilitating comprehensive efficacy assessments and improving trial efficiency. European Journal of Heart Failure, 2016, 18, 482-489.	2.9	74
135	Low-dose Systemic Phosphodiesterase Inhibitors Amplify the Pulmonary Vasodilatory Response to Inhaled Prostacyclin in Experimental Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 1999, 160, 1500-1506.	2.5	<b>7</b> 3
136	Inhaled iloprost is a potent acute pulmonary vasodilator in HIVâ€related severe pulmonary hypertension. European Respiratory Journal, 2004, 23, 321-326.	3.1	72
137	Oleic Acid Inhibits Alveolar Fluid Reabsorption. American Journal of Respiratory and Critical Care Medicine, 2005, 171, 469-479.	2.5	72
138	Impairment of respiratory muscle function in pulmonary hypertension. Clinical Science, 2008, 114, 165-171.	1.8	72
139	Long-term therapy with inhaled iloprost in patients with pulmonary hypertension. Respiratory Medicine, 2010, 104, 731-740.	1.3	72
140	p38 MAPK Inhibition Improves Heart Function in Pressure-Loaded Right Ventricular Hypertrophy. American Journal of Respiratory Cell and Molecular Biology, 2017, 57, 603-614.	1.4	72
141	Amplification of the pulmonary vasodilatory response to inhaled iloprost by subthreshold phosphodiesterase types 3 and 4 inhibition in severe pulmonary hypertension. Critical Care Medicine, 2002, 30, 2489-2492.	0.4	69
142	Lung cancer–associated pulmonary hypertension: Role of microenvironmental inflammation based on tumor cell–immune cell cross-talk. Science Translational Medicine, 2017, 9, .	5.8	69
143	Targeting the Prostacyclin Pathway with Selexipag in Patients with Pulmonary Arterial Hypertension Receiving Double Combination Therapy: Insights from the Randomized Controlled GRIPHON Study. American Journal of Cardiovascular Drugs, 2018, 18, 37-47.	1.0	69
144	Targeting cyclin-dependent kinases for the treatment of pulmonary arterial hypertension. Nature Communications, 2019, 10, 2204.	5.8	69

#	Article	IF	CITATIONS
145	Hypoxia induces Kv channel current inhibition by increased NADPH oxidase-derived reactive oxygen species. Free Radical Biology and Medicine, 2012, 52, 1033-1042.	1.3	68
146	Risk assessment in pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2019, 53, 1802004.	3.1	68
147	Evaluation and Prognostic Relevance of Right Ventricular–Arterial Coupling in Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 116-119.	2.5	68
148	Association of N-Terminal Pro Brain Natriuretic Peptide and Long-Term Outcome in Patients With Pulmonary Arterial Hypertension. Circulation, 2019, 139, 2440-2450.	1.6	67
149	Involvement of mast cells in monocrotaline-induced pulmonary hypertension in rats. Respiratory Research, 2011, 12, 60.	1.4	66
150	Mitochondrial Hyperpolarization in Pulmonary Vascular Remodeling. Mitochondrial Uncoupling Protein Deficiency as Disease Model. American Journal of Respiratory Cell and Molecular Biology, 2013, 49, 358-367.	1.4	66
151	Amplified canonical transforming growth factor-β signalling <i>via</i> heat shock protein 90 in pulmonary fibrosis. European Respiratory Journal, 2017, 49, 1501941.	3.1	66
152	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. European Heart Journal, 2017, 38, 1147-1155.	1.0	65
153	Thrombin Impairs Alveolar Fluid Clearance by Promoting Endocytosis of Na+,K+-ATPase. American Journal of Respiratory Cell and Molecular Biology, 2005, 33, 343-354.	1.4	64
154	Acute effects of the combination of sildenafil and inhaled treprostinil on haemodynamics and gas exchange in pulmonary hypertension. Pulmonary Pharmacology and Therapeutics, 2008, 21, 824-832.	1.1	64
155	Impact of the mitochondria-targeted antioxidant MitoQ on hypoxia-induced pulmonary hypertension. European Respiratory Journal, 2018, 51, 1701024.	3.1	64
156	More on Single-Beat Estimation of Right Ventriculoarterial Coupling in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2018, 198, 816-818.	2.5	63
157	Inhaled Iloprost Reverses Vascular Remodeling in Chronic Experimental Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2005, 172, 358-363.	2.5	62
158	Future Perspectives for the Treatment of Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2009, 54, S108-S117.	1.2	62
159	Function of NADPH Oxidase 1 in Pulmonary Arterial Smooth Muscle Cells After Monocrotaline-Induced Pulmonary Vascular Remodeling. Antioxidants and Redox Signaling, 2013, 19, 2213-2231.	2.5	62
160	5-HT2B Receptor Antagonists Inhibit Fibrosis and Protect from RV Heart Failure. BioMed Research International, 2015, 2015, 1-8.	0.9	62
161	Oxygen sensors in hypoxic pulmonary vasoconstriction. Cardiovascular Research, 2006, 71, 620-629.	1.8	61
162	The lectin-like domain of tumor necrosis factor-α improves alveolar fluid balance in injured isolated rabbit lungs*. Critical Care Medicine, 2008, 36, 1543-1550.	0.4	61

#	Article	IF	Citations
163	Pulmonary hypertension due to chronic lung disease: Updated Recommendations of the Cologne Consensus Conference 2011. International Journal of Cardiology, 2011, 154, S45-S53.	0.8	61
164	PAR-2 Inhibition Reverses Experimental Pulmonary Hypertension. Circulation Research, 2012, 110, 1179-1191.	2.0	61
165	Effects of hypercapnia with and without acidosis on hypoxic pulmonary vasoconstriction. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2009, 297, L977-L983.	1.3	60
166	Diagnosis of CTEPH versus IPAH using capillary to end-tidal carbon dioxide gradients. European Respiratory Journal, 2012, 39, 119-124.	3.1	60
167	Recovery from circulatory shock in severe primary pulmonary hypertension (PPH) with aerosolization of iloprost. Intensive Care Medicine, 1998, 24, 631-634.	3.9	59
168	NO and reactive oxygen species are involved in biphasic hypoxic vasoconstriction of isolated rabbit lungs. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2001, 280, L638-L645.	1.3	59
169	Microcirculatory Dysfunction in the Brain Precedes Changes in Evoked Potentials in Endotoxin-Induced Sepsis Syndrome in Rats. Cerebrovascular Diseases, 2007, 23, 140-147.	0.8	59
170	Hemodynamic and clinical onset in patients with hereditary pulmonary arterial hypertension and BMPR2 mutations. Respiratory Research, 2011, 12, 99.	1.4	59
171	The Role of Dimethylarginine Dimethylaminohydrolase in Idiopathic Pulmonary Fibrosis. Science Translational Medicine, 2011, 03, 87ra53.	5.8	59
172	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. Lancet Respiratory Medicine,the, 2016, 4, 894-901.	5.2	59
173	Basic features of hypoxic pulmonary vasoconstriction in mice. Respiratory Physiology and Neurobiology, 2004, 139, 191-202.	0.7	58
174	Cellular pathophysiology and therapy of pulmonary hypertension. Translational Research, 2001, 138, 367-377.	2.4	57
175	Mediator generation and signaling events in alveolar epithelial cells attacked by <i>S. aureus</i> ݱ-toxin. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2002, 282, L207-L214.	1.3	57
176	Exercise right heart catheterisation before and after pulmonary endarterectomy in patients with chronic thromboembolic disease. European Respiratory Journal, 2018, 52, 1800458.	3.1	57
177	Prevalence of Pulmonary Hypertension in the General Population: The Rotterdam Study. PLoS ONE, 2015, 10, e0130072.	1.1	57
178	Temporal trends in pulmonary arterial hypertension: results from the COMPERA registry. European Respiratory Journal, 2022, 59, 2102024.	3.1	57
179	Phenotyping of idiopathic pulmonary arterial hypertension: a registry analysis. Lancet Respiratory Medicine, the, 2022, 10, 937-948.	5.2	57
180	Atrial natriuretic peptide in severe primary and nonprimary pulmonary hypertension. Journal of the American College of Cardiology, 2001, 38, 1130-1136.	1.2	56

#	Article	IF	CITATIONS
181	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.	0.9	56
182	Targeted therapy of pulmonary arterial hypertension: Updated recommendations from the Cologne Consensus Conference 2018. International Journal of Cardiology, 2018, 272, 37-45.	0.8	56
183	Soluble guanylate cyclase stimulation: an emerging option in pulmonary hypertension therapy. European Respiratory Review, 2009, 18, 35-41.	3.0	55
184	Coaerosolization of Phosphodiesterase Inhibitors Markedly Enhances the Pulmonary Vasodilatory Response to Inhaled Iloprost in Experimental Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2001, 164, 1694-1700.	2.5	54
185	Sildenafil Improves Dynamic Vascular Function in the Brain: Studies in Patients with Pulmonary Hypertension. Cerebrovascular Diseases, 2006, 21, 194-200.	0.8	54
186	miR-223–IGF-IR signalling in hypoxia- and load-induced right-ventricular failure: a novel therapeutic approach. Cardiovascular Research, 2016, 111, 184-193.	1.8	54
187	Balloon pulmonary angioplasty for inoperable patients with chronic thromboembolic disease. Pulmonary Circulation, 2018, 8, 1-6.	0.8	54
188	Quality of life in patients with chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2016, 48, 526-537.	3.1	52
189	Medical management of chronic thromboembolic pulmonary hypertension. European Respiratory Review, 2017, 26, 160107.	3.0	52
190	Effects of Mitochondrial Inhibitors and Uncouplers on Hypoxic Vasoconstriction in Rabbit Lungs. American Journal of Respiratory Cell and Molecular Biology, 2003, 29, 721-732.	1.4	51
191	Diacylglycerol regulates acute hypoxic pulmonary vasoconstriction via TRPC6. Respiratory Research, 2011, 12, 20.	1.4	51
192	Effect of Macitentan on Hospitalizations. JACC: Heart Failure, 2015, 3, 1-8.	1.9	51
193	Late outcomes after acute pulmonary embolism: rationale and design of FOCUS, a prospective observational multicenter cohort study. Journal of Thrombosis and Thrombolysis, 2016, 42, 600-609.	1.0	50
194	Hypoxia- and non-hypoxia-related pulmonary hypertension â€" Established and new therapies. Cardiovascular Research, 2006, 72, 30-40.	1.8	49
195	Therapeutic efficacy of azaindole-1 in experimental pulmonary hypertension. European Respiratory Journal, 2010, 36, 808-818.	3.1	48
196	Psoas Muscle Architectural Design, In Vivo Sarcomere Length Range, and Passive Tensile Properties Support Its Role as a Lumbar Spine Stabilizer. Spine, 2011, 36, E1666-E1674.	1.0	48
197	Telomerecat: A ploidy-agnostic method for estimating telomere length from whole genome sequencing data. Scientific Reports, 2018, 8, 1300.	1.6	48
198	Sex Differences in Right Ventricular–Pulmonary Arterial Coupling in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 1042-1046.	2.5	48

#	Article	IF	Citations
199	Increased neutrophil mediator release in patients with pulmonary hypertension $\hat{a}\in$ suppression by inhaled iloprost. Thrombosis and Haemostasis, 2003, 90, 1141-1149.	1.8	47
200	Arterial hypertension in a murine model of sleep apnea. Journal of Hypertension, 2014, 32, 300-305.	0.3	47
201	Treatment of pulmonary arterial hypertension (PAH): Updated Recommendations of the Cologne Consensus Conference 2011. International Journal of Cardiology, 2011, 154, S20-S33.	0.8	46
202	Macitentan Improves Health-Related QualityÂof Life for Patients With Pulmonary Arterial Hypertension. Chest, 2017, 151, 106-118.	0.4	46
203	Eplerenone attenuates pathological pulmonary vascular rather than right ventricular remodeling in pulmonary arterial hypertension. BMC Pulmonary Medicine, 2018, 18, 41.	0.8	46
204	Chymase: a multifunctional player in pulmonary hypertension associated with lung fibrosis. European Respiratory Journal, 2015, 46, 1084-1094.	3.1	45
205	The science of endothelinâ€1 and endothelin receptor antagonists in the management of pulmonary arterial hypertension: current understanding and future studies. European Journal of Clinical Investigation, 2009, 39, 38-49.	1.7	44
206	Updated classification and management of pulmonary hypertension. Heart, 2010, 96, 552-559.	1.2	44
207	Hypoxic Pulmonary Hypertension in Mice with Constitutively Active Plateletâ€Derived Growth Factor Receptorâ€Î². Pulmonary Circulation, 2011, 1, 259-268.	0.8	44
208	The role of combination therapy in managing pulmonary arterial hypertension. European Respiratory Review, 2014, 23, 469-475.	3.0	44
209	Sequential treatment with riociguat and balloon pulmonary angioplasty for patients with inoperable chronic thromboembolic pulmonary hypertension. Pulmonary Circulation, 2018, 8, 1-7.	0.8	44
210	Evidence for a role of protein kinase C in hypoxic pulmonary vasoconstriction. American Journal of Physiology - Lung Cellular and Molecular Physiology, 1999, 276, L90-L95.	1.3	43
211	Metered dose inhaler delivery of treprostinil for the treatment of pulmonary hypertension. Pulmonary Pharmacology and Therapeutics, 2009, 22, 50-56.	1.1	43
212	Burden of pulmonary arterial hypertension in Germany. Respiratory Medicine, 2010, 104, 902-910.	1.3	43
213	Pulmonary Hypertension in Acute and Chronic High Altitude Maladaptation Disorders. International Journal of Environmental Research and Public Health, 2021, 18, 1692.	1.2	43
214	Inhibition of phosphodiesterase 4 enhances lung alveolarisation in neonatal mice exposed to hyperoxia. European Respiratory Journal, 2009, 33, 861-870.	3.1	42
215	cAMP Phosphodiesterase Inhibitors Increases Nitric Oxide Production by Modulating Dimethylarginine Dimethylaminohydrolases. Circulation, 2011, 123, 1194-1204.	1.6	42
216	Inflammatory Mediators Drive Adverse Right Ventricular Remodeling and Dysfunction and Serve as Potential Biomarkers. Frontiers in Physiology, 2018, 9, 609.	1.3	42

#	Article	IF	CITATIONS
217	Right ventricular function correlates of right atrial strain in pulmonary hypertension: a combined cardiac magnetic resonance and conductance catheter study. American Journal of Physiology - Heart and Circulatory Physiology, 2020, 318, H156-H164.	1.5	42
218	A comprehensive echocardiographic method for risk stratification in pulmonary arterial hypertension. European Respiratory Journal, 2020, 56, 2000513.	3.1	42
219	Selexipag treatment for pulmonary arterial hypertension associated with congenital heart disease after defect correction: insights from the randomised controlled GRIPHON study. European Journal of Heart Failure, 2019, 21, 352-359.	2.9	40
220	Right heart failure in pulmonary hypertension: Diagnosis and new perspectives on vascular and direct right ventricular treatment. British Journal of Pharmacology, 2021, 178, 90-107.	2.7	40
221	Incident and prevalent cohorts with pulmonary arterial hypertension: insight from SERAPHIN. European Respiratory Journal, 2015, 46, 1711-1720.	3.1	39
222	The Role of Transient Receptor Potential Channel 6 Channels in the Pulmonary Vasculature. Frontiers in Immunology, 2017, 8, 707.	2.2	39
223	Right ventricular size and function under riociguat in pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension (the RIVER study). Respiratory Research, 2018, 19, 258.	1.4	39
224	Evidence for the Fucoidan/P-Selectin Axis as a Therapeutic Target in Hypoxia-induced Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2019, 199, 1407-1420.	2.5	39
225	Bypassing mitochondrial complex III using alternative oxidase inhibits acute pulmonary oxygen sensing. Science Advances, 2020, 6, eaba0694.	4.7	39
226	Effects of phosphodiesterase 4 inhibition on bleomycin-induced pulmonary fibrosis in mice. BMC Pulmonary Medicine, 2010, 10, 26.	0.8	38
227	Combination of nonspecific PDE inhibitors with inhaled prostacyclin in experimental pulmonary hypertension. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2001, 281, L1361-L1368.	1.3	37
228	Heme Oxygenase-2 and Large-Conductance Ca2+-activated K+Channels. American Journal of Respiratory and Critical Care Medicine, 2009, 180, 353-364.	2.5	37
229	Prostacyclin Enhances Stretch-induced Surfactant Secretion in Alveolar Epithelial Type II Cells. American Journal of Respiratory and Critical Care Medicine, 1999, 160, 846-851.	2.5	36
230	Characterization of a murine model of monocrotaline pyrrole-induced acute lung injury. BMC Pulmonary Medicine, 2008, 8, 25.	0.8	36
231	Phosphodiesterase 10A Upregulation Contributes to Pulmonary Vascular Remodeling. PLoS ONE, 2011, 6, e18136.	1.1	36
232	New horizons in pulmonary arterial hypertension therapies. European Respiratory Review, 2013, 22, 503-514.	3.0	36
233	Histological Characterization of Mast Cell Chymase in Patients with Pulmonary Hypertension and Chronic Obstructive Pulmonary Disease. Pulmonary Circulation, 2014, 4, 128-136.	0.8	36
234	Pressure overload leads to an increased accumulation and activity of mast cells in the right ventricle. Physiological Reports, 2017, 5, e13146.	0.7	36

#	Article	IF	CITATIONS
235	De Novo Truncating Mutations in WASF1 Cause Intellectual Disability with Seizures. American Journal of Human Genetics, 2018, 103, 144-153.	2.6	36
236	Glycogen Synthase Kinase 3beta Contributes to Proliferation of Arterial Smooth Muscle Cells in Pulmonary Hypertension. PLoS ONE, 2011, 6, e18883.	1.1	36
237	The soluble guanylate cyclase activator HMR1766 reverses hypoxia-induced experimental pulmonary hypertension in mice. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2009, 297, L658-L665.	1.3	35
238	Redox signaling and reactive oxygen species in hypoxic pulmonary vasoconstriction. Respiratory Physiology and Neurobiology, 2010, 174, 282-291.	0.7	35
239	Effects of multikinase inhibitors on pressure overload-induced right ventricular remodeling. International Journal of Cardiology, 2013, 167, 2630-2637.	0.8	35
240	Downregulation of hypoxic vasoconstriction by chronic hypoxia in rabbits: effects of nitric oxide. American Journal of Physiology - Heart and Circulatory Physiology, 2003, 284, H931-H938.	1.5	34
241	Inhaled tolafentrine reverses pulmonary vascular remodeling via inhibition of smooth muscle cell migration. Respiratory Research, 2005, 6, 128.	1.4	34
242	lloprost-induced desensitization of the prostacyclin receptor in isolated rabbit lungs. Respiratory Research, 2007, 8, 4.	1.4	34
243	Mitochondrial cytochrome redox states and respiration in acute pulmonary oxygen sensing. European Respiratory Journal, 2010, 36, 1056-1066.	3.1	34
244	Haemodynamic effects of riociguat in inoperable/recurrent chronic thromboembolic pulmonary hypertension. Heart, 2017, 103, 599-606.	1.2	34
245	Treatment with lowâ€dose tacrolimus inhibits bleeding complications in a patient with hereditary hemorrhagic telangiectasia and pulmonary arterial hypertension. Pulmonary Circulation, 2019, 9, 1-3.	0.8	34
246	Inhaled Treprostinil for Treatment of Chronic Pulmonary Arterial Hypertension. Annals of Internal Medicine, 2006, 144, 149.	2.0	33
247	HbA1c in pulmonary arterial hypertension: A marker of prognostic relevance?. Journal of Heart and Lung Transplantation, 2012, 31, 1109-1114.	0.3	33
248	Effects of exercise training on pulmonary hemodynamics, functional capacity and inflammation in pulmonary hypertension. Pulmonary Circulation, 2017, 7, 20-37.	0.8	33
249	Short-term venoarterial extracorporeal membrane oxygenation for massive endobronchial hemorrhage after pulmonary endarterectomy. Journal of Thoracic and Cardiovascular Surgery, 2018, 155, 643-649.	0.4	33
250	Association between six-minute walk distance and long-term outcomes in patients with pulmonary arterial hypertension: Data from the randomized SERAPHIN trial. PLoS ONE, 2018, 13, e0193226.	1.1	33
251	Detection of reactive oxygen species in isolated, perfused lungs by electron spin resonance spectroscopy. Respiratory Research, 2005, 6, 86.	1.4	32
252	Effect of sildenafil on hypoxia-induced changes in pulmonary circulation and right ventricular function. Respiratory Physiology and Neurobiology, 2007, 159, 196-201.	0.7	32

#	Article	IF	CITATIONS
253	New potential diagnostic biomarkers for pulmonary hypertension. European Respiratory Journal, 2015, 46, 1390-1396.	3.1	32
254	Thin Air Resulting in High Pressure: Mountain Sickness and Hypoxia-Induced Pulmonary Hypertension. Canadian Respiratory Journal, 2017, 2017, 1-17.	0.8	32
255	Congenital erythropoietin over-expression causes "anti-pulmonary hypertensive―structural and functional changes in mice, both in normoxia and hypoxia. Thrombosis and Haemostasis, 2005, 94, 630-638.	1.8	31
256	Lung vasodilatory response to inhaled iloprost in experimental pulmonary hypertension: amplification by different type phosphodiesterase inhibitors. Respiratory Research, 2005, 6, 76.	1.4	31
257	Structural and functional prevention of hypoxia-induced pulmonary hypertension by individualized exercise training in mice. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2014, 306, L986-L995.	1.3	31
258	Acute Effects of Riociguat in Borderline or Manifest Pulmonary Hypertension Associated with Chronic Obstructive Pulmonary Disease. Pulmonary Circulation, 2015, 5, 296-304.	0.8	31
259	Riociguat for pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension: Results from a phase II long-term extension study. Respiratory Medicine, 2017, 128, 50-56.	1.3	31
260	Endotoxin "priming" potentiates lung vascular abnormalities in response to Escherichia coli hemolysin: an example of synergism between endo- and exotoxin Journal of Experimental Medicine, 1994, 180, 1437-1443.	4.2	30
261	Riociguat for pulmonary hypertension. Future Cardiology, 2010, 6, 155-166.	0.5	30
262	A simple echocardiographic estimate of right ventricular-arterial coupling to assess severity and outcome in pulmonary hypertension on chronic lung disease. European Respiratory Journal, 2019, 54, 1802435.	3.1	30
263	Prevalence of Mental Disorders and Impact on Quality of Life in Patients With Pulmonary Arterial Hypertension. Frontiers in Psychiatry, 2021, 12, 667602.	1.3	30
264	Differential Impact of Ultrasonically Nebulized Versus Tracheal-instilled Surfactant on Ventilation-Perfusion (V˙A/Q˙) Mismatch in a Model of Acute Lung Injury. American Journal of Respiratory and Critical Care Medicine, 2000, 161, 152-159.	2.5	29
265	Noninvasive detection of early pulmonary vascular dysfunction in scleroderma. Respiratory Medicine, 2009, 103, 1713-1718.	1.3	29
266	Dynamic hyperinflation during exercise in patients with precapillary pulmonary hypertension. Respiratory Medicine, 2012, 106, 308-313.	1.3	29
267	Pulmonary artery to aorta ratio and risk of all-cause mortality in the general population: the Rotterdam Study. European Respiratory Journal, 2017, 49, 1602168.	3.1	29
268	REVEAL risk score in patients with chronic thromboembolic pulmonary hypertension receiving riociguat. Journal of Heart and Lung Transplantation, 2018, 37, 836-843.	0.3	29
269	REVEAL risk scores applied to riociguat-treated patients in PATENT-2: Impact of changes in risk score on survival. Journal of Heart and Lung Transplantation, 2018, 37, 513-519.	0.3	29
270	Effect of Riociguat and Sildenafil on Right Heart Remodeling and Function in Pressure Overload Induced Model of Pulmonary Arterial Banding. BioMed Research International, 2018, 2018, 1-9.	0.9	29

#	Article	IF	CITATIONS
271	Intravenous treprostinil as an add-on therapy in patients with pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2019, 38, 748-756.	0.3	29
272	Cigarette Smoke-Induced Emphysema and Pulmonary Hypertension Can Be Prevented by Phosphodiesterase 4 and 5 Inhibition in Mice. PLoS ONE, 2015, 10, e0129327.	1.1	29
273	Partial reversal of experimental pulmonary hypertension by phosphodiesterase-3/4 inhibition. European Respiratory Journal, 2008, 31, 599-610.	3.1	28
274	Soluble guanylate cyclase stimulator riociguat and phosphodiesterase 5 inhibitor sildenafil ameliorate pulmonary hypertension due to left heart disease in mice. International Journal of Cardiology, 2016, 216, 85-91.	0.8	28
275	Hypoxia-induced pulmonary hypertension: Different impact of iloprost, sildenafil, and nitric oxide. Respiratory Medicine, 2007, 101, 2125-2132.	1.3	27
276	A Combination Hybrid-Based Vaccination/Adoptive Cellular Therapy to Prevent Tumor Growth by Involvement of T Cells. Cancer Research, 2007, 67, 5443-5453.	0.4	26
277	Riociguat treatment for portopulmonary hypertension: a subgroup analysis from the PATENTâ $\in$ 1/â $\in$ 2 studies. Pulmonary Circulation, 2018, 8, 1-4.	0.8	26
278	Long-term safety and outcome of intravenous treprostinil via an implanted pump in pulmonary hypertension. Journal of Heart and Lung Transplantation, 2018, 37, 1235-1244.	0.3	26
279	Effect of nitric oxide synthase (NOS) inhibition on macro- and microcirculation in a model of rat endotoxic shock. Thrombosis and Haemostasis, 2006, 95, 720-727.	1.8	26
280	Left ventricular systolic dysfunction associated with pulmonary hypertension riociguat trial (LEPHT): rationale and design. European Journal of Heart Failure, 2012, 14, 946-953.	2.9	25
281	The prognostic impact of thyroid function in pulmonary hypertension. Journal of Heart and Lung Transplantation, 2016, 35, 1427-1434.	0.3	25
282	Riociguat for treatment of pulmonary hypertension in COPD: a translational study. European Respiratory Journal, 2019, 53, 1802445.	3.1	25
283	Nitric Oxide (NO)–Dependent but Not NO-Independent Guanylate Cyclase Activation Attenuates Hypoxic Vasoconstriction in Rabbit Lungs. American Journal of Respiratory Cell and Molecular Biology, 2000, 23, 222-227.	1.4	24
284	Lung cGMP release subsequent to NO inhalation in pulmonary hypertension: respondersversusnonresponders. European Respiratory Journal, 2002, 19, 664-671.	3.1	24
285	A Case Series of Patients with Severe Pulmonary Hypertension Receiving an Implantable Pump for Intravenous Prostanoid Therapy. American Journal of Respiratory and Critical Care Medicine, 2012, 186, 1196-1198.	2.5	24
286	Plasma MMP2/TIMP4 Ratio at Follow-up Assessment Predicts Disease Progression of Idiopathic Pulmonary Arterial Hypertension. Lung, 2017, 195, 489-496.	1.4	24
287	TORREY, a Phase 2 study to evaluate the efficacy and safety of inhaled seralutinib for the treatment of pulmonary arterial hypertension. Pulmonary Circulation, 2021, 11, 1-7.	0.8	24
288	Alveolar epithelial barrier functions in ventilated perfused rabbit lungs. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2001, 280, L896-L904.	1.3	23

#	Article	IF	Citations
289	Pharmacokinetics and Metabolism of Infused versus Inhaled Iloprost in Isolated Rabbit Lungs. Journal of Pharmacology and Experimental Therapeutics, 2002, 303, 741-745.	1.3	23
290	Impact of S-Adenosylmethionine Decarboxylase 1 on Pulmonary Vascular Remodeling. Circulation, 2014, 129, 1510-1523.	1.6	23
291	NADPH oxidase subunit NOXO1 is a target for emphysema treatment in COPD. Nature Metabolism, 2020, 2, 532-546.	5.1	23
292	Evaluation of the prognostic value of electrocardiography parameters and heart rhythm in patients with pulmonary hypertension. Cardiology Journal, 2016, 23, 465-472.	0.5	23
293	Urodilatin, a Natriuretic Peptide Stimulating Particulate Guanylate Cyclase, and the Phosphodiesterase 5 Inhibitor Dipyridamole Attenuate Experimental Pulmonary Hypertension. American Journal of Respiratory Cell and Molecular Biology, 2001, 25, 219-225.	1.4	22
294	An Evaluation of Fracture Stabilization Comparing Kyphoplasty and Titanium Mesh Repair Techniques for Vertebral Compression Fractures. Spine, 2010, 35, E768-E773.	1.0	22
295	Phosphodiesterase 6 subunits are expressed and altered in idiopathic pulmonary fibrosis. Respiratory Research, 2010, 11, 146.	1.4	22
296	Cardiovocal Syndrome (Ortner's Syndrome) Associated with Chronic Thromboembolic Pulmonary Hypertension and Giant Pulmonary Artery Aneurysm: Case Report and Review of the Literature. Case Reports in Medicine, 2012, 2012, 1-5.	0.3	22
297	Updating Clinical Endpoint Definitions. Pulmonary Circulation, 2013, 3, 206-216.	0.8	22
298	Comparison of hemodynamic parameters in treatment-na $\tilde{A}$ -ve and pre-treated patients with pulmonary arterial hypertension in the randomized phase III PATENT-1 study. Journal of Heart and Lung Transplantation, 2017, 36, 509-519.	0.3	22
299	A novel non-invasive and echocardiography-derived method for quantification of right ventricular pressure–volume loops. European Heart Journal Cardiovascular Imaging, 2022, 23, 498-507.	0.5	22
300	Validity of echocardiographic tricuspid regurgitation gradient to screen for new definition of pulmonary hypertension. EClinicalMedicine, 2021, 34, 100822.	3.2	22
301	Air Travel Can Be Safe and Well Tolerated in Patients with Clinically Stable Pulmonary Hypertension. Pulmonary Circulation, 2011, 1, 239-243.	0.8	21
302	Riociguat for Pulmonary Hypertension. New England Journal of Medicine, 2013, 369, 2266-2268.	13.9	21
303	Pulmonary Hemodynamic Response to Exercise in Chronic Thromboembolic Pulmonary Hypertension before and after Pulmonary Endarterectomy. Respiration, 2015, 90, 63-73.	1.2	21
304	Impaired right ventricular lusitropy is associated with ventilatory inefficiency in pulmonary arterial hypertension. European Respiratory Journal, 2019, 54, 1900342.	3.1	21
305	Pulmonary Hypertension in Adults with Congenital Heart Disease: Real-World Data from the International COMPERA-CHD Registry. Journal of Clinical Medicine, 2020, 9, 1456.	1.0	21
306	Right ventricular pressure-volume loop shape and systolic pressure change in pulmonary hypertension. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2021, 320, L715-L725.	1.3	21

#	Article	IF	Citations
307	SPARC, a Novel Regulator of Vascular Cell Function in Pulmonary Hypertension. Circulation, 2022, 145, 916-933.	1.6	21
308	Novel soluble guanylyl cyclase stimulator BAY 41-2272 attenuates ischemia-reperfusion-induced lung injury. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2009, 296, L462-L469.	1.3	20
309	Effects of hypercapnia and NO synthase inhibition in sustained hypoxic pulmonary vasoconstriction. Respiratory Research, 2012, 13, 7.	1.4	20
310	Shortâ€Term Improvement in Pulmonary Hemodynamics is Strongly Predictive of Longâ€Term Survival in Patients with Pulmonary Arterial Hypertension. Pulmonary Circulation, 2013, 3, 523-532.	0.8	20
311	Biomarkers of tissue remodeling predict survival in patients with pulmonary hypertension. International Journal of Cardiology, 2016, 223, 821-826.	0.8	20
312	The safety and pharmacokinetics of rapid iloprost aerosol delivery via the BREELIB nebulizer in pulmonary arterial hypertension. Pulmonary Circulation, 2017, 7, 505-513.	0.8	20
313	Severe Emphysema in the SU5416/Hypoxia Rat Model of Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 515-518.	2.5	20
314	Nitric Oxide Synthase 2 Induction Promotes Right Ventricular Fibrosis. American Journal of Respiratory Cell and Molecular Biology, 2019, 60, 346-356.	1.4	20
315	Zardaverine and aerosolised iloprost in a model of acute respiratoryfailure. European Respiratory Journal, 2003, 22, 342-347.	3.1	19
316	Prolonged vasodilatory response to nanoencapsulated sildenafil in pulmonary hypertension. Nanomedicine: Nanotechnology, Biology, and Medicine, 2016, 12, 63-68.	1.7	19
317	Initial combination therapy with ambrisentan + tadalafil on pulmonary arterial hypertensionâ€'related hospitalization in the AMBITION trial. Journal of Heart and Lung Transplantation, 2019, 38, 194-202.	0.3	19
318	Intravenous treprostinil infusion via a fully implantable pump for pulmonary arterial hypertension. Clinical Research in Cardiology, 2017, 106, 776-783.	1.5	18
319	Maintained right ventricular pressure overload induces ventricular–arterial decoupling in mice. Experimental Physiology, 2017, 102, 180-189.	0.9	18
320	Conebulization of surfactant and urokinase restores gas exchange in perfused lungs with alveolar fibrin formation. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2001, 280, L792-L800.	1.3	17
321	Comparison of Pharmacokinetics and Vasodilatory Effect of Nebulized and Infused lloprost in Experimental Pulmonary Hypertension: Rapid Tolerance Development. Journal of Aerosol Medicine and Pulmonary Drug Delivery, 2006, 19, 353-363.	1.2	17
322	Therapeutic efficacy of TBC3711 in monocrotaline-induced pulmonary hypertension. Respiratory Research, 2011, 12, 87.	1.4	17
323	Effects of Dimethylarginine Dimethylaminohydrolase–1 Overexpression on the Response of the Pulmonary Vasculature to Hypoxia. American Journal of Respiratory Cell and Molecular Biology, 2013, 49, 491-500.	1.4	17
324	Nintedanib in Severe Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2018, 198, 808-810.	2.5	17

#	Article	IF	Citations
325	Right ventricular function inÂpulmonary (arterial) hypertension. Herz, 2019, 44, 509-516.	0.4	17
326	FHL-1 is not involved in pressure overload-induced maladaptive right ventricular remodeling and dysfunction. Basic Research in Cardiology, 2020, 115, 17.	2.5	17
327	Amelioration of elastaseâ€induced lung emphysema and reversal of pulmonary hypertension by pharmacological iNOS inhibition in mice. British Journal of Pharmacology, 2021, 178, 152-171.	2.7	17
328	Apical, But Not Basolateral, Endotoxin Preincubation Protects Alveolar Epithelial Cells Against Hydrogen Peroxide-Induced Loss of Barrier Function: The Role of Nitric Oxide Synthesis. Journal of Immunology, 2002, 169, 1474-1481.	0.4	16
329	Long-term effects of intravenous iloprost in patients with idiopathic pulmonary arterial hypertension deteriorating on non-parenteral therapy. BMC Pulmonary Medicine, 2011, 11, 56.	0.8	16
330	Cofilin, a hypoxiaâ€regulated protein in murine lungs identified by 2 <scp>DE</scp> : Role of the cytoskeletal protein cofilin in pulmonary hypertension. Proteomics, 2013, 13, 75-88.	1.3	16
331	The Prognostic Significance of Inspiratory Capacity in Pulmonary Arterial Hypertension. Respiration, 2014, 88, 24-30.	1.2	16
332	Hemodynamic phenotyping based on exercise catheterization predicts outcome in patients with heart failure and reduced ejection fraction. Journal of Heart and Lung Transplantation, 2017, 36, 880-889.	0.3	16
333	Procedural safety of a fully implantable intravenous prostanoid pump for pulmonary hypertension. Clinical Research in Cardiology, 2017, 106, 174-182.	1.5	16
334	Clinical outcomes stratified by baseline functional class after initial combination therapy for pulmonary arterial hypertension. Respiratory Research, 2019, 20, 208.	1.4	16
335	Protection against pressure overload-induced right heart failure by uncoupling protein 2 silencing. Cardiovascular Research, 2019, 115, 1217-1227.	1.8	16
336	Association of right atrial conduit phase with right ventricular lusitropic function in pulmonary hypertension. International Journal of Cardiovascular Imaging, 2020, 36, 633-642.	0.7	16
337	Comparison of MRI and VQ-SPECT as a Screening Test for Patients With Suspected CTEPH: CHANGE-MRI Study Design and Rationale. Frontiers in Cardiovascular Medicine, 2020, 7, 51.	1.1	16
338	Impairment of hypoxic pulmonary vasoconstriction in acute respiratory distress syndrome. European Respiratory Review, 2021, 30, 210059.	3.0	16
339	AMBRISENTAN HAS NO CLINICALLY RELEVANT EFFECT ON THE PHARMACOKINETICS OR PHARMACODYNAMICS OF WARFARIN. Chest, 2006, 130, 256S.	0.4	15
340	Acute hemodynamic response to single oral doses of BAY 60-4552, a soluble guanylate cyclase stimulator, in patients with biventricular heart failure. BMC Pharmacology, 2009, 9, .	0.4	15
341	Update on Pulmonary Hypertension 2009. American Journal of Respiratory and Critical Care Medicine, 2010, 181, 1020-1026.	2.5	15
342	Sildenafil versus Nitric Oxide for Acute Vasodilator Testing in Pulmonary Arterial Hypertension. Pulmonary Circulation, 2015, 5, 305-312.	0.8	15

#	Article	IF	CITATIONS
343	Temporary treatment interruptions with oral selexipag in pulmonary arterial hypertension: Insights from the Prostacyclin (PGI 2) Receptor Agonist in Pulmonary Arterial Hypertension (GRIPHON) study. Journal of Heart and Lung Transplantation, 2018, 37, 401-408.	0.3	15
344	Impact of SARSâ€CoVâ€2 pandemic on pulmonary hypertension outâ€patient clinics in Germany: a multiâ€centre study. Pulmonary Circulation, 2020, 10, 1-3.	0.8	15
345	Risk assessment in severe pulmonary hypertension due to interstitial lung disease. Journal of Heart and Lung Transplantation, 2020, 39, 1118-1125.	0.3	15
346	CILP1 as a biomarker for right ventricular maladaptation in pulmonary hypertension. European Respiratory Journal, 2021, 57, 1901192.	3.1	15
347	Mast cell chymase: an indispensable instrument in the pathological symphony of idiopathic pulmonary fibrosis?. Histology and Histopathology, 2013, 28, 691-9.	0.5	15
348	2015 ESC/ERS GUIDELINES FOR THE DIAGNOSIS AND TREATMENT OF PULMONARY HYPERTENSION. Russian Journal of Cardiology, 2016, , 5-64.	0.4	15
349	Increase in alveolar antioxidant levels in hyperoxic and anoxic ventilated rabbit lungs during ischemia. Free Radical Biology and Medicine, 2004, 36, 78-89.	1.3	14
350	Heart Rate Variability is Related to Disease Severity in Children and Young Adults with Pulmonary Hypertension. Frontiers in Pediatrics, 2015, 3, 63.	0.9	14
351	Hypoxic pulmonary vasoconstriction in isolated mouse pulmonary arterial vessels. Experimental Physiology, 2018, 103, 1185-1191.	0.9	14
352	Advanced risk stratification of intermediate risk group in pulmonary arterial hypertension. Pulmonary Circulation, 2020, 10, 1-5.	0.8	14
353	Physical Activity and Mental Health of Patients with Pulmonary Hypertension during the COVID-19 Pandemic. Journal of Clinical Medicine, 2020, 9, 4023.	1.0	14
354	Modulating cGMP to Treat Lung Diseases. Handbook of Experimental Pharmacology, 2009, , 469-483.	0.9	14
355	Nebulization of the acidified sodium nitrite formulation attenuates acute hypoxic pulmonary vasoconstriction. Respiratory Research, 2010, 11, 81.	1.4	13
356	Electrophysiological Studies in Patients with Pulmonary Hypertension: A Retrospective Investigation. BioMed Research International, 2014, 2014, 1-6.	0.9	13
357	Use of responder threshold criteria to evaluate the response to treatment in the phase III CHEST-1 study. Journal of Heart and Lung Transplantation, 2015, 34, 348-355.	0.3	13
358	Ventricular tachycardias in patients with pulmonary hypertension: an underestimated prevalence? A prospective clinical study. Herzschrittmachertherapie Und Elektrophysiologie, 2015, 26, 155-162.	0.3	13
359	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― Circulation: Cardiovascular Imaging, 2019, 12, e010059.	1.3	13
360	Acute Hemodynamic Effects of Nebulized Iloprost via the Iâ€Neb Adaptive Aerosol Delivery System in Pulmonary Hypertension. Pulmonary Circulation, 2015, 5, 162-170.	0.8	12

#	Article	IF	CITATIONS
361	Prognostic Relevance of Nonsustained Ventricular Tachycardia in Patients with Pulmonary Hypertension. BioMed Research International, 2016, 2016, 1-7.	0.9	12
362	Heart rate response during 6-minute walking testing predicts outcome in operable chronic thromboembolic pulmonary hypertension. BMC Pulmonary Medicine, 2016, 16, 96.	0.8	12
363	Survival with sildenafil and inhaled iloprost in a cohort with pulmonary hypertension: an observational study. BMC Pulmonary Medicine, 2016, 16, 5.	0.8	12
364	Hypoxic Pulmonary Vasoconstriction-Triggered by an Increase in Reactive Oxygen Species?. Novartis Foundation Symposium, 0, , 196-213.	1.2	12
365	Risk assessment in pulmonary hypertension based on routinely measured laboratory parameters. Journal of Heart and Lung Transplantation, 2022, 41, 400-410.	0.3	12
366	Unmasking right ventricular-arterial uncoupling during fluid challenge in pulmonary hypertension. Journal of Heart and Lung Transplantation, 2022, 41, 345-355.	0.3	12
367	Animal models of pulmonary hypertension: role in translational research. Drug Discovery Today: Disease Models, 2010, 7, 89-97.	1.2	11
368	A retrospective review of long anterior fusions to the sacrum. Spine Journal, 2011, 11, 290-294.	0.6	11
369	Individual dose adjustment of riociguat in patients with pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension. Respiratory Medicine, 2017, 129, 124-129.	1.3	11
370	Enhanced circulating levels of CD3 cellsâ€derived extracellular vesicles in different forms of pulmonary hypertension. Pulmonary Circulation, 2019, 9, 1-4.	0.8	11
371	SPARCL1 as a biomarker of maladaptive right ventricular remodelling in pulmonary hypertension. Biomarkers, 2020, 25, 290-295.	0.9	11
372	Protein expression profiling suggests relevance of noncanonical pathways in isolated pulmonary embolism. Blood, 2021, 137, 2681-2693.	0.6	11
373	Stem cellâ€mediated natural tissue engineering. Journal of Cellular and Molecular Medicine, 2011, 15, 52-62.	1.6	10
374	Use of clinically relevant responder threshold criteria to evaluate the response to treatment in the Phase III PATENT-1 study. Journal of Heart and Lung Transplantation, 2015, 34, 338-347.	0.3	10
375	Evaluating Systolic and Diastolic Cardiac Function in Rodents Using Microscopic Computed Tomography. Circulation: Cardiovascular Imaging, 2018, 11, e007653.	1.3	10
376	Cologne consensus conference on pulmonary hypertension – Update 2018. International Journal of Cardiology, 2018, 272, 1-3.	0.8	10
377	Circulating Angiopoietin-1 Is Not a Biomarker of Disease Severity or Prognosis in Pulmonary Hypertension. PLoS ONE, 2016, 11, e0165982.	1.1	10
378	Inhaled Iloprost Improves Right Ventricular Load–Independent Contractility in Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 111-114.	2.5	10

#	Article	IF	Citations
379	Synergism of alveolar endotoxin "priming" and intravascular exotoxin challenge in lung injury American Journal of Respiratory and Critical Care Medicine, 1996, 154, 460-468.	2.5	9
380	Measurement of exhaled hydrogen peroxide from rabbit lungs. Biological Chemistry, 2004, 385, 259-264.	1.2	9
381	Prostanoids and Phosphodiesterase Inhibitors in Experimental Pulmonary Hypertension. Current Topics in Developmental Biology, 2005, 67, 251-284.	1.0	9
382	Effects of carbon monoxide-releasing molecules on pulmonary vasoreactivity in isolated perfused lungs. Journal of Applied Physiology, 2016, 120, 271-281.	1.2	9
383	Altered proteasome function in right ventricular hypertrophy. Cardiovascular Research, 2019, 116, 406-415.	1.8	9
384	Circulating Apoptotic Signals During Acute and Chronic Exposure to High Altitude in Kyrgyz Population. Frontiers in Physiology, 2019, 10, 54.	1.3	9
385	Effects of macitentan and tadalafil monotherapy or their combination on the right ventricle and plasma metabolites in pulmonary hypertensive rats. Pulmonary Circulation, 2020, 10, 1-16.	0.8	9
386	Evaluation of pulmonary hypertension by right heart catheterisation: does timing matter?. European Respiratory Journal, 2020, 56, 1901892.	3.1	9
387	Cytochrome P450 epoxygenaseâ€derived 5,6â€epoxyeicosatrienoic acid relaxes pulmonary arteries in normoxia but promotes sustained pulmonary vasoconstriction in hypoxia. Acta Physiologica, 2020, 230, e13521.	1.8	9
388	Impact of SARS-CoV-2-Pandemic on Mental Disorders and Quality of Life in Patients With Pulmonary Arterial Hypertension. Frontiers in Psychiatry, 2021, 12, 668647.	1.3	9
389	PINK1-mediated Mitophagy Contributes to Pulmonary Vascular Remodeling in Pulmonary Hypertension. American Journal of Respiratory Cell and Molecular Biology, 2021, 65, 226-228.	1.4	9
390	Macitentan for inoperable chronic thromboembolic pulmonary hypertension (CTEPH): results from the randomised controlled MERIT study., 2017,,.		9
391	Chronic intratracheal application of the soluble guanylyl cyclase stimulator BAY 41-8543 ameliorates experimental pulmonary hypertension. Oncotarget, 2017, 8, 29613-29624.	0.8	9
392	Is PKM2 Phosphorylation a Prerequisite for Oligomer Disassembly in Pulmonary Arterial Hypertension?. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 1550-1554.	2.5	8
393	Integrating Data From Randomized Controlled Trials and Observational Studies to Assess Survival in Rare Diseases. Circulation: Cardiovascular Quality and Outcomes, 2019, 12, e005095.	0.9	8
394	Multibeat Right Ventricular–Arterial Coupling during a Positive Acute Vasoreactivity Test. American Journal of Respiratory and Critical Care Medicine, 2019, 199, e41-e42.	2.5	8
395	Genetic deletion of p66shc and/or cyclophilin D results in decreased pulmonary vascular tone. Cardiovascular Research, 2022, 118, 305-315.	1.8	8
396	Medical treatment of pulmonary hypertension in adults with congenital heart disease: updated and extended results from the International COMPERA-CHD Registry. Cardiovascular Diagnosis and Therapy, 2021, 11, 1255-1268.	0.7	8

#	Article	IF	CITATIONS
397	The Clinical Significance of HbA1c in Operable Chronic Thromboembolic Pulmonary Hypertension. PLoS ONE, 2016, 11, e0152580.	1.1	8
398	Relevance of Cor Pulmonale in COPD With and Without Pulmonary Hypertension: A Retrospective Cohort Study. Frontiers in Cardiovascular Medicine, 2022, 9, 826369.	1.1	8
399	Subthreshold doses of nebulized prostacyclin and rolipram synergistaically protect against lung ischemia-reperfusion. Transplantation, 2003, 75, 814-821.	0.5	7
400	Resistant Hypertension. Cardiology Clinics, 2015, 33, 75-87.	0.9	7
401	Relevance of Angiopoietinâ€2 and Soluble Pâ€Selectin Levels in Patients with Pulmonary Arterial Hypertension Receiving Combination Therapy with Oral Treprostinil: A FREEDOMâ€C2 Biomarker Substudy. Pulmonary Circulation, 2016, 6, 516-523.	0.8	7
402	The prognostic relevance of oxygen uptake in inoperable chronic thromboembolic pulmonary hypertension. Clinical Respiratory Journal, 2017, 11, 682-690.	0.6	7
403	IRAG1 Deficient Mice Develop PKG1β Dependent Pulmonary Hypertension. Cells, 2020, 9, 2280.	1.8	7
404	Refined risk stratification in pulmonary arterial hypertension and timing of lung transplantation. European Respiratory Journal, 2022, 60, 2103087.	3.1	7
405	Effect of nitric oxide synthase (NOS) inhibition on macro- and microcirculation in a model of rat endotoxic shock. Thrombosis and Haemostasis, 2006, 95, 720-7.	1.8	7
406	Hypoxic pulmonary vasoconstriction–triggered by an increase in reactive oxygen species?. Novartis Foundation Symposium, 2006, 272, 196-208; discussion 208-17.	1.2	7
407	Prevalence of Mental Disorders in Patients With Chronic Thromboembolic Pulmonary Hypertension. Frontiers in Psychiatry, 2022, 13, 821466.	1.3	7
408	Mitochondrial Respiration in Peripheral Blood Mononuclear Cells Negatively Correlates with Disease Severity in Pulmonary Arterial Hypertension. Journal of Clinical Medicine, 2022, 11, 4132.	1.0	7
409	Interaction of ambrisentan and phenprocoumon in patients with pulmonary hypertension. Pulmonary Pharmacology and Therapeutics, 2014, 28, 87-89.	1.1	6
410	Poor sleep quality is associated with exercise limitation in precapillary pulmonary hypertension. BMC Pulmonary Medicine, 2015, 15, 11.	0.8	6
411	Pulmonary function and diffusion capacity are associated with pulmonary arterial systolic pressure in the general population: The Rotterdam Study. Respiratory Medicine, 2017, 132, 50-55.	1.3	6
412	Switching to riociguat: a potential treatment strategy for the management of CTEPH and PAH. Pulmonary Circulation, 2020, 10, 1-12.	0.8	6
413	Flow rate variance of a fully implantable pump for the delivery of intravenous treprostinil in pulmonary arterial hypertension. Pulmonary Circulation, 2020, 10, 1-5.	0.8	6
414	Osteopontin and galectin-3 as biomarkers of maladaptive right ventricular remodelingÂin pulmonary hypertension. Biomarkers in Medicine, 2021, 15, 1021-1034.	0.6	6

#	Article	IF	CITATIONS
415	Effects Of Inhaled Aviptadil (Vasoactive Intestinal Peptide) In Patients With Pulmonary Arterial Hypertension (PAH)., 2010,,.		5
416	Riociguat for the Treatment of Chronic Thromboembolic Pulmonary Hypertension (CTEPH): 1-Year Results from the CHEST-2 Long-term Extension Study. Chest, 2013, 144, 1023A.	0.4	5
417	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)― Circulation, 2014, 130, e110-2.	1.6	5
418	COPD-associated pulmonary hypertension: clinical implications and current methods for treatment. Expert Review of Respiratory Medicine, 2016, 10, 755-766.	1.0	5
419	Switching inhaled iloprost formulations in patients with pulmonary arterial hypertension: the VENTASWITCH Trial. Pulmonary Circulation, 2018, 8, 1-7.	0.8	5
420	Genetic Deficiency and Pharmacological Stabilization of Mast Cells Ameliorate Pressure Overload-Induced Maladaptive Right Ventricular Remodeling in Mice. International Journal of Molecular Sciences, 2020, 21, 9099.	1.8	5
421	Influence of gender in monocrotaline and chronic hypoxia induced pulmonary hypertension in obese rats and mice. Respiratory Research, 2020, 21, 136.	1.4	5
422	Yarsagumba is a Promising Therapeutic Option for Treatment of Pulmonary Hypertension due to the Potent Anti-Proliferative and Vasorelaxant Properties. Medicina (Lithuania), 2020, 56, 131.	0.8	5
423	Right ventricular dyssynchrony: from load-independent right ventricular function to wall stress in severe pulmonary arterial hypertension. Pulmonary Circulation, 2020, 10, 204589402092575.	0.8	5
424	Exercise hemodynamics in heart failure patients with preserved and mid-range ejection fraction: key role of the right heart. Clinical Research in Cardiology, 2022, 111, 393-405.	1.5	5
425	Targeting peptidyl-prolyl isomerase 1 in experimental pulmonary arterial hypertension. European Respiratory Journal, 2022, 60, 2101698.	3.1	5
426	Impact of Pulmonary Arterial Hypertension on Employment, Work Productivity, and Quality of Life - Results of a Cross-Sectional Multi-Center Study. Frontiers in Psychiatry, 2021, 12, 781532.	1.3	5
427	Profiles and treatment patterns of patients with pulmonary arterial hypertension on monotherapy at experienced centres. ESC Heart Failure, 2022, 9, 2873-2885.	1.4	5
428	Cologne Consensus Conference on pulmonary hypertension. International Journal of Cardiology, 2011, 154, S1-S2.	0.8	4
429	Riociguat for the Treatment of Pulmonary Arterial Hypertension (PAH): 1-Year Results from the PATENT-2 Long-term Extension Study. Chest, 2013, 144, 1024A.	0.4	4
430	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. Chest, 2014, 145, 535B.	0.4	4
431	Reply to Bogaard et al.: Emphysema Is—at the Most—Only a Mild Phenotype in the Sugen/Hypoxia Rat Model of Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 1450-1452.	2.5	4
432	Acute response to rapid iloprost inhalation using the Breelibâ,,¢ nebulizer in pulmonary arterial hypertension: the Breelibâ,,¢ acute study. Pulmonary Circulation, 2019, 9, 1-3.	0.8	4

#	Article	IF	CITATIONS
433	Therapeutic Potential of Regorafenib—A Multikinase Inhibitor in Pulmonary Hypertension. International Journal of Molecular Sciences, 2021, 22, 1502.	1.8	4
434	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. Chest, 2014, 145, 513A.	0.4	3
435	Individualized Dosing of Selexipag Based on Tolerability in the GRIPHON Study Shows Consistent Efficacy and Safety in Patients With Pulmonary Arterial Hypertension (PAH). Chest, 2015, 148, 961A.	0.4	3
436	Selexipag for the treatment of pulmonary arterial hypertension. Expert Opinion on Pharmacotherapy, 2016, 17, 1825-1834.	0.9	3
437	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. European Journal of Epidemiology, 2018, 33, 403-413.	2.5	3
438	Pulmonary Vascular Pressure Response to Acute Cold Exposure in Kyrgyz Highlanders. High Altitude Medicine and Biology, 2019, 20, 375-382.	0.5	3
439	Cardiopulmonary haemodynamics in portopulmonary hypertension. Lancet Respiratory Medicine, the, 2019, 7, 556-558.	5.2	3
440	Deficiency of Axl aggravates pulmonary arterial hypertension via BMPR2. Communications Biology, 2021, 4, 1002.	2.0	3
441	Exercise Hemodynamic Profiling Is Associated With Outcome in Patients Undergoing Percutaneous Mitral Valve Repair. Circulation: Cardiovascular Interventions, 2021, 14, e010453.	1.4	3
442	Clinical Relevance of Right Atrial Functional Response to Treatment in Pulmonary Arterial Hypertension. Frontiers in Cardiovascular Medicine, 2021, 8, 775039.	1.1	3
443	Riociguat dose titration in patients with chronic thromboembolic pulmonary hypertension (CTEPH) or pulmonary arterial hypertension (PAH). BMC Pharmacology, 2009, 9, .	0.4	2
444	Long Term Use Of Imatinib In Patients With Severe Pulmonary Arterial Hypertension. , 2010, , .		2
445	Riociguat in Combination With Prostacyclin Analogs for the Treatment of Pulmonary Arterial Hypertension (PAH): A Subgroup Analysis of the PATENT Studies. Chest, 2015, 148, 922A.	0.4	2
446	Effect of Selexipag on Morbidity/Mortality in Pulmonary Arterial Hypertension: Results of the GRIPHON Study. Journal of Heart and Lung Transplantation, 2015, 34, S163.	0.3	2
447	Inspiratory capacity is not altered in operable chronic thromboembolic pulmonary hypertension. Pulmonary Circulation, 2017, 7, 543-546.	0.8	2
448	Beyond interleukin-6 in right ventricular function: Evidence for another biomarker. Journal of Heart and Lung Transplantation, 2018, 37, 674-675.	0.3	2
449	Stem/Progenitor Cells in Cardiopulmonary Health, Disease, and Treatment. Stem Cells International, 2019, 2019, 1-4.	1.2	2
450	Metabolic Reprogramming in Congenital Cyanotic Heart Disease: Another Fight in Puberty?. Circulation, 2021, 143, 2273-2276.	1.6	2

#	Article	IF	CITATIONS
451	Initial combination therapy with ambrisentan (AMB) and tadalafil (TAD) in treatment $na\tilde{A}^-$ ve patients with pulmonary arterial hypertension (PAH): Efficacy and safety in the AMBITION study intent to treat (ITT) population., 2015,,.		2
452	Acute hemodynamic effects of riociguat in pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension. , $2018,  ,  .$		2
453	Metacognitions in Patients With Frequent Mental Disorders After Diagnosis of Pulmonary Arterial Hypertension. Frontiers in Psychiatry, 2022, 13, 812812.	1.3	2
454	IMATINIB MESYLATE TREATMENT FOR SEVERE PULMONARY ARTERIAL HYPERTENSION: A PROPOSED PHASE III 24-WEEK DOUBLE-BLIND PLACEBO-CONTROLLED RANDOMIZED CLINICAL TRIAL. Chest, 2009, 136, 64S.	0.4	1
455	Effects Of Riociguat On Pulmonary Vascular Remodeling In Severe Experimental Pulmonary Hypertension., 2011,,.		1
456	Improvement of right heart structure and function by BAY 41-8543 in pulmonary artery banded mice. BMC Pharmacology, 2011, 11, .	0.4	1
457	Association Between WHO Functional Class and Long-term Prognosis in Patients With Pulmonary Arterial Hypertension: Data From SERAPHIN, A Randomized Controlled Study of Macitentan. Chest, 2013, 144, 879A.	0.4	1
458	Effects of Riociguat in Treatment-Naive vs Pretreated Patients With Pulmonary Arterial Hypertension: 2-Year Efficacy Results From the PATENT-2 Study. Chest, 2016, 150, 1162A.	0.4	1
459	Follow-Up (Measurement) of Corrected QT Interval in Adult Patients before and after Lung Transplantation. BioMed Research International, 2017, 2017, 1-5.	0.9	1
460	Reply to "a pediatric perspective on the TAPSE/PASP ratio in pulmonary arterial hypertension― International Journal of Cardiology, 2019, 278, 240-241.	0.8	1
461	Subcutaneous treprostinil: a new treatment for chronic thromboembolic pulmonary hypertension?. Lancet Respiratory Medicine, the, 2019, 7, 191-193.	5.2	1
462	Phosphodiesterase-5 Inhibitors in Pulmonary Arterial Hypertension., 0,, 105-125.		1
463	Childhood Maltreatment, Mental Well-Being, and Healthy Lifestyle in Patients With Chronic Thromboembolic Pulmonary Hypertension. Frontiers in Psychiatry, 2022, 13, 821468.	1.3	1
464	Schwere pulmonale Hypertonie - Vasodilative Therapie in der Lungenstrombahn. Pneumologie, 2000, 54, 160-169.	0.1	0
465	Sildenafil for lung fibrosis and pulmonary hypertension. Lancet, The, 2003, 361, 263.	6.3	0
466	Prevention of pulmonary vascular and myocardial remodeling by the combined tyrosine and serine-/threonine kinase inhibitor, sorafenib, in pulmonary hypertension and right heart failure. European Respiratory Review, 2008, 17, 72-73.	3.0	0
467	Response to Letter Regarding Article, "Pulmonary Vascular Disease in the Developing World― Circulation, 2009, 120, .	1.6	0
468	Phosphodiesterase 10A Upregulation Contributes To Pulmonary Vascular Remodeling. , 2010, , .		0

#	Article	IF	Citations
469	Role of Reactive Oxygen Species in Acute Pulmonary Oxygen Sensing. Free Radical Biology and Medicine, 2010, 49, S48.	1.3	О
470	Exercise Induced Pulmonary Hypertension Associated With Systemic Sclerosis: Four Distinct Entities. , 2010, , .		0
471	Effects Of The Multikinase Inhibitor Sunitinib On Right Ventricular Remodeling In An Experimental Model Of Right Heart Hypertrophy. , 2010, , .		O
472	Does Sitaxsentan Therapy Offer Benefit To Functional Class II Patients With Pulmonary Arterial Hypertension. , 2010, , .		0
473	Reversal Of Experimental Pulmonary Hypertension By The Multi-kinase Inhibitor Sunitinib., 2010,,.		0
474	Combined Inhibition Of 5-HT2a And 5-HT2b-receptor Ameliorates Myocardial Hypertrophy And Diastolic Dysfunction In The Pressure Overloaded Right Heart. , 2010, , .		0
475	A Highly Selective Endothelin-A Receptor Antagonist TBC3711 Reverses Monocrotaline Induced Pulmonary Hypertension. , 2010, , .		0
476	Combination Therapy With Sildenafil And Sitaxentan In Patients With Severe Pulmonary Hypertension., 2011,,.		0
477	Role Of Notch Signaling In Pulmonary Arterial Hypertension. , 2011, , .		0
478	Prevention Of Experimental Hypoxia-Induced Pulmonary Hypertension By Moderate Exercise Training In Mice. , $2011$ , , .		0
479	Mitochondrial Alterations Of Pulmonary Arterial Smooth Muscle Cells In Pulmonary Hypertension. , 2011, , .		O
480	Contribution Of Progenitor Cells In Experimental Right Heart Hypertrophy Induced By Pulmonary Artery Ligation. , $2011, \ldots$		0
481	Effects Of The Soluble Guanylate Cyclase Stimulator Riociguat On Emphysema Development In Tobacco-Smoke Exposed Mice. , $2011, \ldots$		0
482	Effects of riociguat in severe experimental pulmonary hypertension. BMC Pharmacology, 2011, 11, .	0.4	0
483	The soluble guanylate cyclase stimulator riociguat ameliorates pulmonary hypertension induced by hypoxia and SU5416 in rats. BMC Pharmacology, $2011, 11, \ldots$	0.4	0
484	Do parameters of cardiac function predict long-term outcomes in patients with pulmonary arterial hypertension? Data from SERAPHIN, a randomized controlled study of macitentan. Chest, 2013, 144, 870B.	0.4	0
485	Application of REVEAL Risk Score to Patients With PAH Receiving Riociguat in the PATENT-2 Study. Chest, 2015, 148, 930A.	0.4	0
486	The Clinical Impact of Referral Time in Patients With Pulmonary Hypertension on Disease Progression: A Register Analysis of a Tertiary Referral Center. Chest, 2016, 150, 1169A.	0.4	0

#	Article	IF	Citations
487	Tidal Carbon Dioxide as a Prognostic Feature in Inoperable Chronic Thromboembolic Pulmonary Hypertension. Annals of the American Thoracic Society, 2017, 14, 1603-1604.	1.5	O
488	Effect of Riociguat on Pulmonary Arterial Compliance in Patients With Pulmonary Arterial Hypertension (PAH) in the Respite Study. Chest, 2017, 152, A1005-A1006.	0.4	0
489	Response to: Comment on "Effect of Riociguat and Sildenafil on Right Heart Remodeling and Function in Pressure Overload Induced Model of Pulmonary Arterial Banding†BioMed Research International, 2018, 2018, 1-2.	0.9	0
490	EFFECT OF RIOCIGUAT ON RIGHT VENTRICULAR FUNCTION IN PATIENTS WITH CHRONIC THROMBOEMBOLIC PULMONARY HYPERTENSION. Chest, 2018, 154, 1065A-1066A.	0.4	0
491	EFFECT OF RIOCIGUAT ON RIGHT VENTRICULAR FUNCTION IN PATIENTS WITH PULMONARY ARTERIAL HYPERTENSION. Chest, 2018, 154, 1062A-1064A.	0.4	0
492	Suppressor of Tumorigenicity 2 as a Biomarker in Pulmonary Arterial Hypertension and its Association with REVEAL Risk Score in Riociguat-Treated Patients in the RESPITE Study. Journal of Heart and Lung Transplantation, 2019, 38, S96.	0.3	0
493	Reply to "Risk stratification in PH associated with interstitial lung disease: The Holy Grail?― Journal of Heart and Lung Transplantation, 2021, 40, 317.	0.3	O
494	NO pathway and phosphodiesterase inhibitors in pulmonary arterial hypertension., 2004,, 163-168.		0
495	Classical transient receptor potential channel 6 (TRPC6) is essential for ischemiaâ€reperfusion injury of the lung. FASEB Journal, 2010, 24, 591.2.	0.2	O
496	Definition, Klassifikation und Diagnose der pulmonalen Hypertonie (PH). Atemwegs- Und Lungenkrankheiten, 2015, 41, 166-170.	0.0	0
497	Chronisch thromboembolische pulmonale Hypertonie. Atemwegs- Und Lungenkrankheiten, 2015, 41, 198-203.	0.0	0
498	Therapie der pulmonal-arteriellen Hypertonie. Atemwegs- Und Lungenkrankheiten, 2015, 41, 171-178.	0.0	0
499	Effects of apoptosis signal-regulating kinase $1$ (ASK1) inhibition in experimental pressure overload-induced right ventricular dysfunction. , $2015, \dots$		0
500	Procedural safety of a fully implantable pump for delivery of intravenous prostanoids in severe pulmonary hypertension: A two-center observational study. , 2016, , .		0
501	Anti-fibrotic effects of pirfenidone on pulmonary arterial vascular smooth muscle cells. , 2016, , .		0
502	Pulmonary arterial hypertension-related morbidity is prognostic for survival: insights from the SERAPHIN and GRIPHON studies. , $2017$ , , .		0
503	VENTASWITCH study: Switching from Ventavis (iloprost) V10 to V20 improves inhalation behaviour in patients with pulmonary arterial hypertension (PAH). , 2017, , .		0
504	Effects on Right Ventricular size and function by Riociguat in Pulmonary Arterial Hypertension and Chronic Thromboembolic Pulmonary Hypertension (The RIVER Study)., 2018,,.		0

#	Article	IF	CITATIONS
505	PEGASUS - the effects of commercial air travel on patients suffering from pulmonary hypertension - a prospective, multicenter, multinational study. , $2018$ , , .		O
506	Acute response of iloprost inhalation using the Breelib nebulizer in pulmonary arterial hypertension: the Breelib acute study. , $2018,  ,  .$		0
507	Effects of electronic cigarette aerosol on isolated murine lung cells and bronchoalveolar lavage fluid. , 2018, , .		O
508	Diagnostic accuracy of echocardiography in pulmonary hypertension due to interstitial lung disease. , 2018, , .		0
509	EARLIER: End tidal carbon dioxide for earlier detection of pulmonary hypertension. , 2018, , .		0
510	Childhood Trauma in Patients With PAHâ€"Prevalence, Impact on QoL, and Mental Healthâ€"A Preliminary Report. Frontiers in Psychiatry, 2022, 13, 812862.	1.3	0