

## 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.

64  
papers

3,355  
citations

29  
h-index

57  
g-index

79  
ext. papers

4,106  
ext. citations

8.3  
avg, IF

5  
L-index

#	Paper	IF	Citations
64	Dysregulated renin-angiotensin-aldosterone system contributes to pulmonary arterial hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , <b>2012</b> , 186, 780-9	10.2	244
63	Impact of interleukin-6 on hypoxia-induced pulmonary hypertension and lung inflammation in mice. <i>Respiratory Research</i> , <b>2009</b> , 10, 6	7.3	210
62	Role of endothelium-derived CC chemokine ligand 2 in idiopathic pulmonary arterial hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , <b>2007</b> , 176, 1041-7	10.2	173
61	Transgenic mice overexpressing the 5-hydroxytryptamine transporter gene in smooth muscle develop pulmonary hypertension. <i>Circulation Research</i> , <b>2006</b> , 98, 1323-30	15.7	158
60	Endothelial-derived FGF2 contributes to the progression of pulmonary hypertension in humans and rodents. <i>Journal of Clinical Investigation</i> , <b>2009</b> , 119, 512-23	15.9	148
59	Dasatinib induces lung vascular toxicity and predisposes to pulmonary hypertension. <i>Journal of Clinical Investigation</i> , <b>2016</b> , 126, 3207-18	15.9	144
58	RhoA and Rho kinase activation in human pulmonary hypertension: role of 5-HT signaling. <i>American Journal of Respiratory and Critical Care Medicine</i> , <b>2009</b> , 179, 1151-8	10.2	134
57	Increased pericyte coverage mediated by endothelial-derived fibroblast growth factor-2 and interleukin-6 is a source of smooth muscle-like cells in pulmonary hypertension. <i>Circulation</i> , <b>2014</b> , 129, 1586-97	16.7	131
56	Pathogenesis of pulmonary arterial hypertension: lessons from cancer. <i>European Respiratory Review</i> , <b>2013</b> , 22, 543-51	9.8	126
55	Role for interleukin-6 in COPD-related pulmonary hypertension. <i>Chest</i> , <b>2009</b> , 136, 678-687	5.3	126
54	Immune dysregulation and endothelial dysfunction in pulmonary arterial hypertension: a complex interplay. <i>Circulation</i> , <b>2014</b> , 129, 1332-40	16.7	110
53	New molecular targets of pulmonary vascular remodeling in pulmonary arterial hypertension: importance of endothelial communication. <i>Chest</i> , <b>2015</b> , 147, 529-537	5.3	109
52	Proinflammatory Signature of the Dysfunctional Endothelium in Pulmonary Hypertension. Role of the Macrophage Migration Inhibitory Factor/CD74 Complex. <i>American Journal of Respiratory and Critical Care Medicine</i> , <b>2015</b> , 192, 983-97	10.2	108
51	Autocrine fibroblast growth factor-2 signaling contributes to altered endothelial phenotype in pulmonary hypertension. <i>American Journal of Respiratory Cell and Molecular Biology</i> , <b>2011</b> , 45, 311-22	5.7	98
50	Ectopic upregulation of membrane-bound IL6R drives vascular remodeling in pulmonary arterial hypertension. <i>Journal of Clinical Investigation</i> , <b>2018</b> , 128, 1956-1970	15.9	87
49	Leptin and regulatory T-lymphocytes in idiopathic pulmonary arterial hypertension. <i>European Respiratory Journal</i> , <b>2012</b> , 40, 895-904	13.6	84
48	A critical role for p130Cas in the progression of pulmonary hypertension in humans and rodents. <i>American Journal of Respiratory and Critical Care Medicine</i> , <b>2012</b> , 186, 666-76	10.2	68

47	Dichloroacetate treatment partially regresses established pulmonary hypertension in mice with SM22alpha-targeted overexpression of the serotonin transporter. <i>FASEB Journal</i> , <b>2009</b> , 23, 4135-47	0.9	66
46	Bone morphogenetic protein signalling in heritable versus idiopathic pulmonary hypertension. <i>European Respiratory Journal</i> , <b>2009</b> , 34, 1100-10	13.6	61
45	Delayed Microvascular Shear Adaptation in Pulmonary Arterial Hypertension. Role of Platelet Endothelial Cell Adhesion Molecule-1 Cleavage. <i>American Journal of Respiratory and Critical Care Medicine</i> , <b>2016</b> , 193, 1410-20	10.2	60
44	Contribution of Impaired Parasympathetic Activity to Right Ventricular Dysfunction and Pulmonary Vascular Remodeling in Pulmonary Arterial Hypertension. <i>Circulation</i> , <b>2018</b> , 137, 910-924	16.7	60
43	Leptin signalling system as a target for pulmonary arterial hypertension therapy. <i>European Respiratory Journal</i> , <b>2015</b> , 45, 1066-80	13.6	48
42	Selective BMP-9 Inhibition Partially Protects Against Experimental Pulmonary Hypertension. <i>Circulation Research</i> , <b>2019</b> , 124, 846-855	15.7	48
41	Pan-PPAR agonist IVA337 is effective in experimental lung fibrosis and pulmonary hypertension. <i>Annals of the Rheumatic Diseases</i> , <b>2017</b> , 76, 1931-1940	2.4	47
40	Chronic inflammation within the vascular wall in pulmonary arterial hypertension: more than a spectator. <i>Cardiovascular Research</i> , <b>2020</b> , 116, 885-893	9.9	35
39	N-acetylcysteine improves established monocrotaline-induced pulmonary hypertension in rats. <i>Respiratory Research</i> , <b>2014</b> , 15, 65	7.3	34
38	Regulatory T Cell Dysfunction in Idiopathic, Heritable and Connective Tissue-Associated Pulmonary Arterial Hypertension. <i>Chest</i> , <b>2016</b> , 149, 1482-93	5.3	33
37	Dasatinib increases endothelial permeability leading to pleural effusion. <i>European Respiratory Journal</i> , <b>2018</b> , 51,	13.6	29
36	T-cell costimulation blockade is effective in experimental digestive and lung tissue fibrosis. <i>Arthritis Research and Therapy</i> , <b>2018</b> , 20, 197	5.7	29
35	Lineage Tracing Reveals the Dynamic Contribution of Pericytes to the Blood Vessel Remodeling in Pulmonary Hypertension. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , <b>2020</b> , 40, 766-782	9.4	27
34	Renal Denervation Reduces Pulmonary Vascular Remodeling and Right Ventricular Diastolic Stiffness in Experimental Pulmonary Hypertension. <i>JACC Basic To Translational Science</i> , <b>2017</b> , 2, 22-35	8.7	26
33	Neutralization of CXCL12 attenuates established pulmonary hypertension in rats. <i>Cardiovascular Research</i> , <b>2020</b> , 116, 686-697	9.9	25
32	Nintedanib improves cardiac fibrosis but leaves pulmonary vascular remodelling unaltered in experimental pulmonary hypertension. <i>Cardiovascular Research</i> , <b>2019</b> , 115, 432-439	9.9	24
31	Prevention of progression of pulmonary hypertension by the Nur77 agonist 6-mercaptopurine: role of BMP signalling. <i>European Respiratory Journal</i> , <b>2019</b> , 54,	13.6	20
30	Regression of flow-induced pulmonary arterial vasculopathy after flow correction in piglets. <i>Journal of Thoracic and Cardiovascular Surgery</i> , <b>2009</b> , 137, 1538-46	1.5	20

29	Role of Nerve Growth Factor in Development and Persistence of Experimental Pulmonary Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , <b>2015</b> , 192, 342-55	10.2	16
28	Design, Synthesis, and Biological Activity of New N-(Phenylmethyl)-benzoxazol-2-thiones as Macrophage Migration Inhibitory Factor (MIF) Antagonists: Efficacies in Experimental Pulmonary Hypertension. <i>Journal of Medicinal Chemistry</i> , <b>2018</b> , 61, 2725-2736	8.3	14
27	Role of Stromelysin 2 (Matrix Metalloproteinase 10) as a Novel Mediator of Vascular Remodeling Underlying Pulmonary Hypertension Associated With Systemic Sclerosis. <i>Arthritis and Rheumatology</i> , <b>2017</b> , 69, 2209-2221	9.5	14
26	Therapeutic effect of pirfenidone in the sugen/hypoxia rat model of severe pulmonary hypertension. <i>FASEB Journal</i> , <b>2019</b> , 33, 3670-3679	0.9	14
25	The BMP Receptor 2 in Pulmonary Arterial Hypertension: When and Where the Animal Model Matches the Patient. <i>Cells</i> , <b>2020</b> , 9,	7.9	13
24	Macrophage Migration Inhibitory Factor (MIF) Inhibition in a Murine Model of Bleomycin-Induced Pulmonary Fibrosis. <i>International Journal of Molecular Sciences</i> , <b>2018</b> , 19,	6.3	12
23	Switching-Off Adora2b in Vascular Smooth Muscle Cells Halts the Development of Pulmonary Hypertension. <i>Frontiers in Physiology</i> , <b>2018</b> , 9, 555	4.6	11
22	Right lung ischemia induces contralateral pulmonary vasculopathy in an animal model. <i>Journal of Thoracic and Cardiovascular Surgery</i> , <b>2012</b> , 143, 967-73	1.5	11
21	New targets for pulmonary arterial hypertension: going beyond the currently targeted three pathways. <i>Current Opinion in Pulmonary Medicine</i> , <b>2017</b> , 23, 377-385	3	11
20	Angiomatoid fibrous histiocytoma of the pulmonary artery: a multidisciplinary discussion. <i>Histopathology</i> , <b>2014</b> , 65, 278-82	7.3	9
19	Pulmonary hemodynamic responses to inhaled NO in chronic heart failure depend on PDE5 G(-1142)T polymorphism. <i>Pulmonary Circulation</i> , <b>2011</b> , 1, 377-82	2.7	9
18	Connexin-43 is a promising target for pulmonary hypertension due to hypoxaemic lung disease. <i>European Respiratory Journal</i> , <b>2020</b> , 55,	13.6	9
17	An endothelial activin A-bone morphogenetic protein receptor type 2 link is overdriven in pulmonary hypertension. <i>Nature Communications</i> , <b>2021</b> , 12, 1720	17.4	9
16	Altered TGF $\beta$ /SMAD Signaling in Human and Rat Models of Pulmonary Hypertension: An Old Target Needs Attention. <i>Cells</i> , <b>2021</b> , 10,	7.9	9
15	A genome-wide association analysis identifies   locus on chromosome 2 associated with idiopathic pulmonary arterial hypertension in a Japanese population. <i>Oncotarget</i> , <b>2017</b> , 8, 74917-74926	3.3	7
14	Different cardiovascular and pulmonary phenotypes for single- and double-knock-out mice deficient in BMP9 and BMP10. <i>Cardiovascular Research</i> , <b>2021</b> ,	9.9	7
13	The Thousand Faces of Leptin in the Lung. <i>Chest</i> , <b>2021</b> , 159, 239-248	5.3	7
12	Purinergic Dysfunction in Pulmonary Arterial Hypertension. <i>Journal of the American Heart Association</i> , <b>2020</b> , 9, e017404	6	6

11	Therapeutic potential of melatonin and melatonergic drugs on K18-hACE2 mice infected with SARS-CoV-2. <i>Journal of Pineal Research</i> , <b>2021</b> , e12772	10.4	6
10	Serum and pulmonary uric acid in pulmonary arterial hypertension. <i>European Respiratory Journal</i> , <b>2021</b> , 58,	13.6	6
9	Emerging Molecular Targets for Anti-proliferative Strategies in Pulmonary Arterial Hypertension. <i>Handbook of Experimental Pharmacology</i> , <b>2013</b> , 409-436	3.2	5
8	Additive protective effects of sacubitril/valsartan and bosentan on vascular remodelling in experimental pulmonary hypertension. <i>Cardiovascular Research</i> , <b>2021</b> , 117, 1391-1401	9.9	5
7	Emerging molecular targets for anti-proliferative strategies in pulmonary arterial hypertension. <i>Handbook of Experimental Pharmacology</i> , <b>2013</b> , 218, 409-36	3.2	4
6	Phenotypic Diversity of Vascular Smooth Muscle Cells in Pulmonary Arterial Hypertension: Implications for Therapy. <i>Chest</i> , <b>2021</b> ,	5.3	4
5	Lower Plasma Melatonin Levels Predict Worse Long-Term Survival in Pulmonary Arterial Hypertension. <i>Journal of Clinical Medicine</i> , <b>2020</b> , 9,	5.1	3
4	Response by Guignabert et al to Letter Regarding Article, "Selective BMP-9 Inhibition Partially Protects Against Experimental Pulmonary Hypertension". <i>Circulation Research</i> , <b>2019</b> , 124, e82-e83	15.7	2
3	Acazicolcept (ALPN-101), a dual ICOS/CD28 antagonist, demonstrates efficacy in systemic sclerosis preclinical mouse models.. <i>Arthritis Research and Therapy</i> , <b>2022</b> , 24, 13	5.7	1
2	Platelet-Derived Growth Factor Receptor Type 1 Activation Drives Pulmonary Vascular Remodeling Via Progenitor Cell Proliferation and Induces Pulmonary Hypertension.. <i>Journal of the American Heart Association</i> , <b>2022</b> , e023021	6	1
1	Pulmonary hypertension associated with neurofibromatosis type 2. <i>Pulmonary Circulation</i> , <b>2021</b> , 11, 2045894021102955	15.7	1