

Ly Tu

List of Publications by Year in Descending Order

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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	Acazicolcept (ALPN-101), a dual ICOS/CD28 antagonist, demonstrates efficacy in systemic sclerosis preclinical mouse models.. <i>Arthritis Research and Therapy</i> , 2022 , 24, 13	5.7	1
63	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> , 2022 , e023021	6	1
62	An endothelial activin A-bone morphogenetic protein receptor type 2 link is overdriven in pulmonary hypertension. <i>Nature Communications</i> , 2021 , 12, 1720	17.4	9
61	Pulmonary hypertension associated with neurofibromatosis type 2. <i>Pulmonary Circulation</i> , 2021 , 11, 2045894021102955	17.4	9
60	Different cardiovascular and pulmonary phenotypes for single- and double-knock-out mice deficient in BMP9 and BMP10. <i>Cardiovascular Research</i> , 2021 ,	9.9	7
59	Additive protective effects of sacubitril/valsartan and bosentan on vascular remodelling in experimental pulmonary hypertension. <i>Cardiovascular Research</i> , 2021 , 117, 1391-1401	9.9	5
58	The Thousand Faces of Leptin in the Lung. <i>Chest</i> , 2021 , 159, 239-248	5.3	7
57	Altered TGF β /SMAD Signaling in Human and Rat Models of Pulmonary Hypertension: An Old Target Needs Attention. <i>Cells</i> , 2021 , 10,	7.9	9
56	Phenotypic Diversity of Vascular Smooth Muscle Cells in Pulmonary Arterial Hypertension: Implications for Therapy. <i>Chest</i> , 2021 ,	5.3	4
55	Therapeutic potential of melatonin and melatonergic drugs on K18-hACE2 mice infected with SARS-CoV-2. <i>Journal of Pineal Research</i> , 2021 , e12772	10.4	6
54	Serum and pulmonary uric acid in pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2021 , 58,	13.6	6
53	The BMP Receptor 2 in Pulmonary Arterial Hypertension: When and Where the Animal Model Matches the Patient. <i>Cells</i> , 2020 , 9,	7.9	13
52	Lineage Tracing Reveals the Dynamic Contribution of Pericytes to the Blood Vessel Remodeling in Pulmonary Hypertension. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2020 , 40, 766-782	9.4	27
51	Lower Plasma Melatonin Levels Predict Worse Long-Term Survival in Pulmonary Arterial Hypertension. <i>Journal of Clinical Medicine</i> , 2020 , 9,	5.1	3
50	Chronic inflammation within the vascular wall in pulmonary arterial hypertension: more than a spectator. <i>Cardiovascular Research</i> , 2020 , 116, 885-893	9.9	35
49	Connexin-43 is a promising target for pulmonary hypertension due to hypoxaemic lung disease. <i>European Respiratory Journal</i> , 2020 , 55,	13.6	9
48	Purinergic Dysfunction in Pulmonary Arterial Hypertension. <i>Journal of the American Heart Association</i> , 2020 , 9, e017404	6	6

47	Neutralization of CXCL12 attenuates established pulmonary hypertension in rats. <i>Cardiovascular Research</i> , 2020 , 116, 686-697	9.9	25
46	Response by Guignabert et al to Letter Regarding Article, "Selective BMP-9 Inhibition Partially Protects Against Experimental Pulmonary Hypertension". <i>Circulation Research</i> , 2019 , 124, e82-e83	15.7	2
45	Nintedanib improves cardiac fibrosis but leaves pulmonary vascular remodelling unaltered in experimental pulmonary hypertension. <i>Cardiovascular Research</i> , 2019 , 115, 432-439	9.9	24
44	Prevention of progression of pulmonary hypertension by the Nur77 agonist 6-mercaptopurine: role of BMP signalling. <i>European Respiratory Journal</i> , 2019 , 54,	13.6	20
43	Therapeutic effect of pirfenidone in the sugen/hypoxia rat model of severe pulmonary hypertension. <i>FASEB Journal</i> , 2019 , 33, 3670-3679	0.9	14
42	Selective BMP-9 Inhibition Partially Protects Against Experimental Pulmonary Hypertension. <i>Circulation Research</i> , 2019 , 124, 846-855	15.7	48
41	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> , 2018 , 61, 2725-2736	8.3	14
40	Dasatinib increases endothelial permeability leading to pleural effusion. <i>European Respiratory Journal</i> , 2018 , 51,	13.6	29
39	Switching-Off Adora2b in Vascular Smooth Muscle Cells Halts the Development of Pulmonary Hypertension. <i>Frontiers in Physiology</i> , 2018 , 9, 555	4.6	11
38	Ectopic upregulation of membrane-bound IL6R drives vascular remodeling in pulmonary arterial hypertension. <i>Journal of Clinical Investigation</i> , 2018 , 128, 1956-1970	15.9	87
37	Contribution of Impaired Parasympathetic Activity to Right Ventricular Dysfunction and Pulmonary Vascular Remodeling in Pulmonary Arterial Hypertension. <i>Circulation</i> , 2018 , 137, 910-924	16.7	60
36	Macrophage Migration Inhibitory Factor (MIF) Inhibition in a Murine Model of Bleomycin-Induced Pulmonary Fibrosis. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	12
35	T-cell costimulation blockade is effective in experimental digestive and lung tissue fibrosis. <i>Arthritis Research and Therapy</i> , 2018 , 20, 197	5.7	29
34	Renal Denervation Reduces Pulmonary Vascular Remodeling and Right Ventricular Diastolic Stiffness in Experimental Pulmonary Hypertension. <i>JACC Basic To Translational Science</i> , 2017 , 2, 22-35	8.7	26
33	A genome-wide association analysis identifies a locus on chromosome 2 associated with idiopathic pulmonary arterial hypertension in a Japanese population. <i>Oncotarget</i> , 2017 , 8, 74917-74926	3.3	7
32	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> , 2017 , 69, 2209-2221	9.5	14
31	Pan-PPAR agonist IVA337 is effective in experimental lung fibrosis and pulmonary hypertension. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 1931-1940	2.4	47
30	New targets for pulmonary arterial hypertension: going beyond the currently targeted three pathways. <i>Current Opinion in Pulmonary Medicine</i> , 2017 , 23, 377-385	3	11

29	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> , 2016 , 193, 1410-20	10.2	60
28	Regulatory T Cell Dysfunction in Idiopathic, Heritable and Connective Tissue-Associated Pulmonary Arterial Hypertension. <i>Chest</i> , 2016 , 149, 1482-93	5.3	33
27	Dasatinib induces lung vascular toxicity and predisposes to pulmonary hypertension. <i>Journal of Clinical Investigation</i> , 2016 , 126, 3207-18	15.9	144
26	New molecular targets of pulmonary vascular remodeling in pulmonary arterial hypertension: importance of endothelial communication. <i>Chest</i> , 2015 , 147, 529-537	5.3	109
25	Leptin signalling system as a target for pulmonary arterial hypertension therapy. <i>European Respiratory Journal</i> , 2015 , 45, 1066-80	13.6	48
24	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> , 2015 , 192, 983-97	10.2	108
23	Role of Nerve Growth Factor in Development and Persistence of Experimental Pulmonary Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2015 , 192, 342-55	10.2	16
22	N-acetylcysteine improves established monocrotaline-induced pulmonary hypertension in rats. <i>Respiratory Research</i> , 2014 , 15, 65	7.3	34
21	Immune dysregulation and endothelial dysfunction in pulmonary arterial hypertension: a complex interplay. <i>Circulation</i> , 2014 , 129, 1332-40	16.7	110
20	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> , 2014 , 129, 1586-97	16.7	131
19	Angiomatoid fibrous histiocytoma of the pulmonary artery: a multidisciplinary discussion. <i>Histopathology</i> , 2014 , 65, 278-82	7.3	9
18	Pathogenesis of pulmonary arterial hypertension: lessons from cancer. <i>European Respiratory Review</i> , 2013 , 22, 543-51	9.8	126
17	Emerging molecular targets for anti-proliferative strategies in pulmonary arterial hypertension. <i>Handbook of Experimental Pharmacology</i> , 2013 , 218, 409-36	3.2	4
16	Emerging Molecular Targets for Anti-proliferative Strategies in Pulmonary Arterial Hypertension. <i>Handbook of Experimental Pharmacology</i> , 2013 , 409-436	3.2	5
15	Right lung ischemia induces contralateral pulmonary vasculopathy in an animal model. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2012 , 143, 967-73	1.5	11
14	Dysregulated renin-angiotensin-aldosterone system contributes to pulmonary arterial hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2012 , 186, 780-9	10.2	244
13	Leptin and regulatory T-lymphocytes in idiopathic pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2012 , 40, 895-904	13.6	84
12	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> , 2012 , 186, 666-76	10.2	68

11	Autocrine fibroblast growth factor-2 signaling contributes to altered endothelial phenotype in pulmonary hypertension. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2011 , 45, 311-22	5.7	98
10	Pulmonary hemodynamic responses to inhaled NO in chronic heart failure depend on PDE5 G(-1142)T polymorphism. <i>Pulmonary Circulation</i> , 2011 , 1, 377-82	2.7	9
9	Dichloroacetate treatment partially regresses established pulmonary hypertension in mice with SM22alpha-targeted overexpression of the serotonin transporter. <i>FASEB Journal</i> , 2009 , 23, 4135-47	0.9	66
8	Bone morphogenetic protein signalling in heritable versus idiopathic pulmonary hypertension. <i>European Respiratory Journal</i> , 2009 , 34, 1100-10	13.6	61
7	Role for interleukin-6 in COPD-related pulmonary hypertension. <i>Chest</i> , 2009 , 136, 678-687	5.3	126
6	RhoA and Rho kinase activation in human pulmonary hypertension: role of 5-HT signaling. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2009 , 179, 1151-8	10.2	134
5	Impact of interleukin-6 on hypoxia-induced pulmonary hypertension and lung inflammation in mice. <i>Respiratory Research</i> , 2009 , 10, 6	7.3	210
4	Regression of flow-induced pulmonary arterial vasculopathy after flow correction in piglets. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2009 , 137, 1538-46	1.5	20
3	Endothelial-derived FGF2 contributes to the progression of pulmonary hypertension in humans and rodents. <i>Journal of Clinical Investigation</i> , 2009 , 119, 512-23	15.9	148
2	Role of endothelium-derived CC chemokine ligand 2 in idiopathic pulmonary arterial hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2007 , 176, 1041-7	10.2	173
1	Transgenic mice overexpressing the 5-hydroxytryptamine transporter gene in smooth muscle develop pulmonary hypertension. <i>Circulation Research</i> , 2006 , 98, 1323-30	15.7	158