Ralph T Schermuly

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.

188 106 12,103 55 h-index g-index citations papers 206 8.5 5.81 14,193 avg, IF L-index ext. citations ext. papers

#	Paper	IF	Citations
188	The effect of long-term doxycycline treatment in a mouse model of cigarette smoke-induced emphysema and pulmonary hypertension. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2021 , 320, L903-L915	5.8	1
187	Targeting Jak-Stat Signaling in Experimental Pulmonary Hypertension. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2021 , 64, 100-114	5.7	10
186	The role of chemokines and chemokine receptors in pulmonary arterial hypertension. <i>British Journal of Pharmacology</i> , 2021 , 178, 72-89	8.6	23
185	Kinases as potential targets for treatment of pulmonary hypertension and right ventricular dysfunction. <i>British Journal of Pharmacology</i> , 2021 , 178, 31-53	8.6	5
184	Genetic Delivery and Gene Therapy in Pulmonary Hypertension. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	3
183	Experimental Models 2021 , 27-52		
182	Pulmonary Hypertension in Acute and Chronic High Altitude Maladaptation Disorders. <i>International Journal of Environmental Research and Public Health</i> , 2021 , 18,	4.6	7
181	Novel Therapeutic Targets for the Treatment of Right Ventricular Remodeling: Insights from the Pulmonary Artery Banding Model. <i>International Journal of Environmental Research and Public Health</i> , 2021 , 18,	4.6	1
180	Deficiency of Axl aggravates pulmonary arterial hypertension via BMPR2. <i>Communications Biology</i> , 2021 , 4, 1002	6.7	
179	Generation of pulmonary arterial hypertension patient-specific induced pluripotent stem cell lines from three unrelated patients with a heterozygous missense mutation in exon 12, a heterozygous in-frame deletion in exon 3 and a missense mutation in exon 11 of the BMPR2 gene. Stem Cell	1.6	0
178	Research, 2021, 55, 102488 Lack of Contribution of p66shc to Pressure Overload-Induced Right Heart Hypertrophy. International Journal of Molecular Sciences, 2020, 21,	6.3	2
177	Effect of p53 activation on experimental right ventricular hypertrophy. <i>PLoS ONE</i> , 2020 , 15, e0234872	3.7	2
176	Influence of gender in monocrotaline and chronic hypoxia induced pulmonary hypertension in obese rats and mice. <i>Respiratory Research</i> , 2020 , 21, 136	7-3	4
175	NADPH oxidase subunit NOXO1 is a target for emphysema treatment in COPD. <i>Nature Metabolism</i> , 2020 , 2, 532-546	14.6	4
174	Cytochrome P450 epoxygenase-derived 5,6-epoxyeicosatrienoic acid relaxes pulmonary arteries in normoxia but promotes sustained pulmonary vasoconstriction in hypoxia. <i>Acta Physiologica</i> , 2020 , 230, e13521	5.6	4
173	is a Promising Therapeutic Option for Treatment of Pulmonary Hypertension due to the Potent Anti-Proliferative and Vasorelaxant Properties. <i>Medicina (Lithuania)</i> , 2020 , 56,	3.1	1
172	Profiling of human lymphocytes reveals a specific network of protein kinases modulated by endurance training status. <i>Scientific Reports</i> , 2020 , 10, 888	4.9	9

(2019-2020)

171	Lung epithelium damage in COPD - An unstoppable pathological event?. <i>Cellular Signalling</i> , 2020 , 68, 109540	4.9	9
170	FHL-1 is not involved in pressure overload-induced maladaptive right ventricular remodeling and dysfunction. <i>Basic Research in Cardiology</i> , 2020 , 115, 17	11.8	14
169	IRAG1 Deficient Mice Develop PKG1 Dependent Pulmonary Hypertension. Cells, 2020, 9,	7.9	2
168	Effects of macitentan and tadalafil monotherapy or their combination on the right ventricle and plasma metabolites in pulmonary hypertensive rats. <i>Pulmonary Circulation</i> , 2020 , 10, 204589402094728	3 3 .7	2
167	Genetic Deficiency and Pharmacological Stabilization of Mast Cells Ameliorate Pressure Overload-Induced Maladaptive Right Ventricular Remodeling in Mice. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	2
166	Bypassing mitochondrial complex III using alternative oxidase inhibits acute pulmonary oxygen sensing. <i>Science Advances</i> , 2020 , 6, eaba0694	14.3	18
165	Update in Pulmonary Vascular Diseases and Right Ventricular Dysfunction 2019. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020 , 202, 22-28	10.2	3
164	Is PKM2 Phosphorylation a Prerequisite for Oligomer Disassembly in Pulmonary Arterial Hypertension?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019 , 200, 1550-1554	10.2	2
163	Pulmonary Vascular Pressure Response to Acute Cold Exposure in Kyrgyz Highlanders. <i>High Altitude Medicine and Biology</i> , 2019 , 20, 375-382	1.9	2
162	Targeting cyclin-dependent kinases for the treatment of pulmonary arterial hypertension. <i>Nature Communications</i> , 2019 , 10, 2204	17.4	39
161	Lipids - two sides of the same coin in lung fibrosis. <i>Cellular Signalling</i> , 2019 , 60, 65-80	4.9	8
160	Circulating Apoptotic Signals During Acute and Chronic Exposure to High Altitude in Kyrgyz Population. <i>Frontiers in Physiology</i> , 2019 , 10, 54	4.6	3
159	LRP1 promotes synthetic phenotype of pulmonary artery smooth muscle cells in pulmonary hypertension. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2019 , 1865, 1604-1616	6.9	10
158	Riociguat for treatment of pulmonary hypertension in COPD: a translational study. <i>European Respiratory Journal</i> , 2019 , 53,	13.6	15
157	Reply to Bogaard: Emphysema Is-at the Most-Only a Mild Phenotype in the Sugen/Hypoxia Rat Model of Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019 , 200, 1450-1452	10.2	4
156	Metformin induces lipogenic differentiation in myofibroblasts to reverse lung fibrosis. <i>Nature Communications</i> , 2019 , 10, 2987	17.4	89
155	Depletion of Bone Marrow-Derived Fibrocytes Attenuates TAA-Induced Liver Fibrosis in Mice. <i>Cells</i> , 2019 , 8,	7.9	9
154	Evidence for the Fucoidan/P-Selectin Axis as a Therapeutic Target in Hypoxia-induced Pulmonary Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019 , 199, 1407-1420	10.2	25

153	Pathology and pathobiology of pulmonary hypertension: state of the art and research perspectives. European Respiratory Journal, 2019 , 53,	13.6	407
152	Nitric Oxide Synthase 2 Induction Promotes Right Ventricular Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2019 , 60, 346-356	5.7	14
151	Impact of the mitochondria-targeted antioxidant MitoQ on hypoxia-induced pulmonary hypertension. <i>European Respiratory Journal</i> , 2018 ,	13.6	30
150	Eplerenone attenuates pathological pulmonary vascular rather than right ventricular remodeling in pulmonary arterial hypertension. <i>BMC Pulmonary Medicine</i> , 2018 , 18, 41	3.5	31
149	ASK1 Inhibition Halts Disease Progression in Preclinical Models of Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2018 , 197, 373-385	10.2	57
148	Inflammatory Mediators Drive Adverse Right Ventricular Remodeling and Dysfunction and Serve as Potential Biomarkers. <i>Frontiers in Physiology</i> , 2018 , 9, 609	4.6	26
147	Is the fibroblast growth factor signaling pathway a victim of receptor tyrosine kinase inhibition in pulmonary parenchymal and vascular remodeling?. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2018 , 315, L248-L252	5.8	8
146	Myeloperoxidase aggravates pulmonary arterial hypertension by activation of vascular Rho-kinase. <i>JCI Insight</i> , 2018 , 3,	9.9	25
145	Response to: Comment on "Effect of Riociguat and Sildenafil on Right Heart Remodeling and Function in Pressure Overload Induced Model of Pulmonary Arterial Banding". <i>BioMed Research International</i> , 2018 , 2018, 7491284	3	
144	The Role of G Protein-Coupled Receptors in the Right Ventricle in Pulmonary Hypertension. <i>Frontiers in Cardiovascular Medicine</i> , 2018 , 5, 179	5.4	6
143	Evaluating Systolic and Diastolic Cardiac Function in Rodents Using Microscopic Computed Tomography. <i>Circulation: Cardiovascular Imaging</i> , 2018 , 11, e007653	3.9	5
142	Nintedanib in Severe Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018 , 198, 808-810	10.2	11
141	Effect of Riociguat and Sildenafil on Right Heart Remodeling and Function in Pressure Overload Induced Model of Pulmonary Arterial Banding. <i>BioMed Research International</i> , 2018 , 2018, 3293584	3	17
140	Hypoxic pulmonary vasoconstriction in isolated mouse pulmonary arterial vessels. <i>Experimental Physiology</i> , 2018 , 103, 1185-1191	2.4	7
139	Riociguat: Mode of Action and Clinical Development in Pulmonary Hypertension. <i>Chest</i> , 2017 , 151, 468-	-4 5 .6	57
138	The Giessen Pulmonary Hypertension Registry: Survival in pulmonary hypertension subgroups. Journal of Heart and Lung Transplantation, 2017 , 36, 957-967	5.8	138
137	Amplified canonical transforming growth factor-Bignalling heat shock protein 90 in pulmonary fibrosis. <i>European Respiratory Journal</i> , 2017 , 49,	13.6	43
136	Enhanced inflammatory cell profiles in schistosomiasis-induced pulmonary vascular remodeling. <i>Pulmonary Circulation</i> , 2017 , 7, 244-252	2.7	5

135	Mitochondrial Complex IV Subunit 4 Isoform 2 Is Essential for Acute Pulmonary Oxygen Sensing. <i>Circulation Research</i> , 2017 , 121, 424-438	15.7	58
134	Plasma MMP2/TIMP4 Ratio at Follow-up Assessment Predicts Disease Progression of Idiopathic Pulmonary Arterial Hypertension. <i>Lung</i> , 2017 , 195, 489-496	2.9	10
133	Long Noncoding RNA MANTIS Facilitates Endothelial Angiogenic Function. <i>Circulation</i> , 2017 , 136, 65-79	16.7	145
132	Oxidative injury of the pulmonary circulation in the perinatal period: Short- and long-term consequences for the human cardiopulmonary system. <i>Pulmonary Circulation</i> , 2017 , 7, 55-66	2.7	17
131	Maintained right ventricular pressure overload induces ventricular-arterial decoupling in mice. <i>Experimental Physiology</i> , 2017 , 102, 180-189	2.4	14
130	Differential Alterations of the Mitochondrial Morphology and Respiratory Chain Complexes during Postnatal Development of the Mouse Lung. <i>Oxidative Medicine and Cellular Longevity</i> , 2017 , 2017, 9169	146	11
129	Developmental vascular remodeling defects and postnatal kidney failure in mice lacking Gpr116 (Adgrf5) and Eltd1 (Adgrl4). <i>PLoS ONE</i> , 2017 , 12, e0183166	3.7	17
128	Pressure overload leads to an increased accumulation and activity of mast cells in the right ventricle. <i>Physiological Reports</i> , 2017 , 5, e13146	2.6	30
127	Lung cancer-associated pulmonary hypertension: Role of microenvironmental inflammation based on tumor cell-immune cell cross-talk. <i>Science Translational Medicine</i> , 2017 , 9,	17.5	50
126	p38 MAPK Inhibition Improves Heart Function in Pressure-Loaded Right Ventricular Hypertrophy. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2017 , 57, 603-614	5.7	48
125	Two-Way Conversion between Lipogenic and Myogenic Fibroblastic Phenotypes Marks the Progression and Resolution of Lung Fibrosis. <i>Cell Stem Cell</i> , 2017 , 20, 261-273.e3	18	118
124	Translating Research into Improved Patient Care in Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2017 , 195, 583-595	10.2	95
123	Chronic intratracheal application of the soluble guanylyl cyclase stimulator BAY 41-8543 ameliorates experimental pulmonary hypertension. <i>Oncotarget</i> , 2017 , 8, 29613-29624	3.3	6
122	The prognostic impact of thyroid function in pulmonary hypertension. <i>Journal of Heart and Lung Transplantation</i> , 2016 , 35, 1427-1434	5.8	14
121	Soluble guanylate cyclase stimulator riociguat and phosphodiesterase 5 inhibitor sildenafil ameliorate pulmonary hypertension due to left heart disease in mice. <i>International Journal of Cardiology</i> , 2016 , 216, 85-91	3.2	20
120	Prolonged vasodilatory response to nanoencapsulated sildenafil in pulmonary hypertension. <i>Nanomedicine: Nanotechnology, Biology, and Medicine</i> , 2016 , 12, 63-8	6	13
119	miR-223-IGF-IR signalling in hypoxia- and load-induced right-ventricular failure: a novel therapeutic approach. <i>Cardiovascular Research</i> , 2016 , 111, 184-93	9.9	42
118	Endothelial actions of atrial natriuretic peptide prevent pulmonary hypertension in mice. <i>Basic Research in Cardiology</i> , 2016 , 111, 22	11.8	12

117	Nestin-expressing vascular wall cells drive development of pulmonary hypertension. <i>European Respiratory Journal</i> , 2016 , 47, 876-88	13.6	24
116	Circulating Angiopoietin-1 Is Not a Biomarker of Disease Severity or Prognosis in Pulmonary Hypertension. <i>PLoS ONE</i> , 2016 , 11, e0165982	3.7	5
115	Relevance of angiopoietin-2 and soluble P-selectin levels in patients with pulmonary arterial hypertension receiving combination therapy with oral treprostinil: a FREEDOM-C2 biomarker substudy. <i>Pulmonary Circulation</i> , 2016 , 6, 516-523	2.7	5
114	Notch1 signalling regulates endothelial proliferation and apoptosis in pulmonary arterial hypertension. <i>European Respiratory Journal</i> , 2016 , 48, 1137-1149	13.6	57
113	Genetic Ablation of PDGF-Dependent Signaling Pathways Abolishes Vascular Remodeling and Experimental Pulmonary Hypertension. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2015 , 35, 123	6 ⁹ 45	31
112	Chymase: a multifunctional player in pulmonary hypertension associated with lung fibrosis. <i>European Respiratory Journal</i> , 2015 , 46, 1084-94	13.6	29
111	Loss of secreted frizzled-related protein-1 leads to deterioration of cardiac function in mice and plays a role in human cardiomyopathy. <i>Circulation: Heart Failure</i> , 2015 , 8, 362-72	7.6	43
110	New potential diagnostic biomarkers for pulmonary hypertension. <i>European Respiratory Journal</i> , 2015 , 46, 1390-6	13.6	22
109	The F-BAR Protein NOSTRIN Dictates the Localization of the Muscarinic M3 Receptor and Regulates Cardiovascular Function. <i>Circulation Research</i> , 2015 , 117, 460-9	15.7	11
108	5-HT2B receptor antagonists inhibit fibrosis and protect from RV heart failure. <i>BioMed Research International</i> , 2015 , 2015, 438403	3	48
107	Cigarette Smoke-Induced Emphysema and Pulmonary Hypertension Can Be Prevented by Phosphodiesterase 4 and 5 Inhibition in Mice. <i>PLoS ONE</i> , 2015 , 10, e0129327	3.7	24
106	Impact of S-adenosylmethionine decarboxylase 1 on pulmonary vascular remodeling. <i>Circulation</i> , 2014 , 129, 1510-23	16.7	17
105	Pro-proliferative and inflammatory signaling converge on FoxO1 transcription factor in pulmonary hypertension. <i>Nature Medicine</i> , 2014 , 20, 1289-300	50.5	183
104	Endothelin-1 driven proliferation of pulmonary arterial smooth muscle cells is c-fos dependent. <i>International Journal of Biochemistry and Cell Biology</i> , 2014 , 54, 137-48	5.6	34
103	Protective effects of 10-nitro-oleic acid in a hypoxia-induced murine model of pulmonary hypertension. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2014 , 51, 155-62	5.7	45
102	Novel and emerging therapies for pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014 , 189, 394-400	10.2	62
101	Hypoxia- or PDGF-BB-dependent paxillin tyrosine phosphorylation in pulmonary hypertension is reversed by HIF-1 depletion or imatinib treatment. <i>Thrombosis and Haemostasis</i> , 2014 , 112, 1288-303	7	15
100	Histological characterization of mast cell chymase in patients with pulmonary hypertension and chronic obstructive pulmonary disease. <i>Pulmonary Circulation</i> , 2014 , 4, 128-36	2.7	26

(2013-2014)

99	Stimulation of soluble guanylate cyclase prevents cigarette smoke-induced pulmonary hypertension and emphysema. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014 , 189, 1359-73	10.2	59
98	Structural and functional prevention of hypoxia-induced pulmonary hypertension by individualized exercise training in mice. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2014 , 306, L986-95	5.8	25
97	Experimental Models 2014 , 45-67		
96	Mitochondrial hyperpolarization in pulmonary vascular remodeling. Mitochondrial uncoupling protein deficiency as disease model. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2013 , 49, 358-67	5.7	50
95	Deletion of Fn14 receptor protects from right heart fibrosis and dysfunction. <i>Basic Research in Cardiology</i> , 2013 , 108, 325	11.8	54
94	Effects of multikinase inhibitors on pressure overload-induced right ventricular remodeling. <i>International Journal of Cardiology</i> , 2013 , 167, 2630-7	3.2	29
93	Relevant issues in the pathology and pathobiology of pulmonary hypertension. <i>Journal of the American College of Cardiology</i> , 2013 , 62, D4-12	15.1	379
92	Function of NADPH oxidase 1 in pulmonary arterial smooth muscle cells after monocrotaline-induced pulmonary vascular remodeling. <i>Antioxidants and Redox Signaling</i> , 2013 , 19, 221	8 3 1	57
91	The role of dimethylarginine dimethylaminohydrolase (DDAH) in pulmonary fibrosis. <i>Journal of Pathology</i> , 2013 , 229, 242-9	9.4	29
90	Cofilin, a hypoxia-regulated protein in murine lungs identified by 2DE: role of the cytoskeletal protein cofilin in pulmonary hypertension. <i>Proteomics</i> , 2013 , 13, 75-88	4.8	14
89	Mast cells and fibroblasts work in concert to aggravate pulmonary fibrosis: role of transmembrane SCF and the PAR-2/PKC-[]Raf-1/p44/42 signaling pathway. <i>American Journal of Pathology</i> , 2013 , 182, 2094-108	5.8	69
88	Inhibition of overactive transforming growth factor-Isignaling by prostacyclin analogs in pulmonary arterial hypertension. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2013 , 48, 733-41	5.7	35
87	Classical transient receptor potential channel 1 in hypoxia-induced pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2013 , 188, 1451-9	10.2	58
86	Anticipated classes of new medications and molecular targets for pulmonary arterial hypertension. <i>Pulmonary Circulation</i> , 2013 , 3, 226-44	2.7	35
85	The peroxisome proliferator-activated receptor Magonist GW0742 has direct protective effects on right heart hypertrophy. <i>Pulmonary Circulation</i> , 2013 , 3, 926-35	2.7	18
84	Sildenafil potentiates bone morphogenetic protein signaling in pulmonary arterial smooth muscle cells and in experimental pulmonary hypertension. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2013 , 33, 34-42	9.4	55
83	A molecular mechanism for therapeutic effects of cGMP-elevating agents in pulmonary arterial hypertension. <i>Journal of Biological Chemistry</i> , 2013 , 288, 16557-16566	5.4	20
82	The role of cGMP in the physiological and molecular responses of the right ventricle to pressure overload. <i>Experimental Physiology</i> , 2013 , 98, 1274-8	2.4	9

81	Mast cell chymase: an indispensable instrument in the pathological symphony of idiopathic pulmonary fibrosis?. <i>Histology and Histopathology</i> , 2013 , 28, 691-9	1.4	13
80	Phosphodiesterase 5 (PDE5) inhibition, ANP and NO rapidly reduce epididymal duct contractions, but long-term PDE5 inhibition in vivo does not. <i>Molecular and Cellular Endocrinology</i> , 2012 , 349, 145-53	4.4	11
79	Immune and inflammatory cell involvement in the pathology of idiopathic pulmonary arterial hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2012 , 186, 897-908	10.2	219
78	BDNF/TrkB signaling augments smooth muscle cell proliferation in pulmonary hypertension. <i>American Journal of Pathology</i> , 2012 , 181, 2018-29	5.8	35
77	Activation of TRPC6 channels is essential for lung ischaemia-reperfusion induced oedema in mice. <i>Nature Communications</i> , 2012 , 3, 649	17.4	137
76	The soluble guanylate cyclase stimulator riociguat ameliorates pulmonary hypertension induced by hypoxia and SU5416 in rats. <i>PLoS ONE</i> , 2012 , 7, e43433	3.7	89
75	Hypoxia induces Kv channel current inhibition by increased NADPH oxidase-derived reactive oxygen species. <i>Free Radical Biology and Medicine</i> , 2012 , 52, 1033-42	7.8	60
74	Role of Src tyrosine kinases in experimental pulmonary hypertension. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2012 , 32, 1354-65	9.4	90
73	Inhibition of microRNA-17 improves lung and heart function in experimental pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2012 , 185, 409-19	10.2	171
72	PAR-2 inhibition reverses experimental pulmonary hypertension. Circulation Research, 2012, 110, 1179-	91 5.7	52
71	Mitochondrial complex II is essential for hypoxia-induced pulmonary vasoconstriction of intra- but not of pre-acinar arteries. <i>Cardiovascular Research</i> , 2012 , 93, 702-10	9.9	16
70	Riociguat for the treatment of pulmonary hypertension. <i>Expert Opinion on Investigational Drugs</i> , 2011 , 20, 567-76	5.9	69
69	Inducible NOS inhibition reverses tobacco-smoke-induced emphysema and pulmonary hypertension in mice. <i>Cell</i> , 2011 , 147, 293-305	56.2	226
68	Phosphodiesterase 10A upregulation contributes to pulmonary vascular remodeling. <i>PLoS ONE</i> , 2011 , 6, e18136	3.7	30
67	Involvement of mast cells in monocrotaline-induced pulmonary hypertension in rats. <i>Respiratory Research</i> , 2011 , 12, 60	7.3	53
66	Therapeutic efficacy of TBC3711 in monocrotaline-induced pulmonary hypertension. <i>Respiratory Research</i> , 2011 , 12, 87	7.3	16
65	Mechanisms of disease: pulmonary arterial hypertension. <i>Nature Reviews Cardiology</i> , 2011 , 8, 443-55	14.8	472
64	Hypoxia enhances platelet-derived growth factor signaling in the pulmonary vasculature by down-regulation of protein tyrosine phosphatases. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2011 , 183, 1092-102	10.2	64

(2009-2011)

63	cAMP phosphodiesterase inhibitors increases nitric oxide production by modulating dimethylarginine dimethylaminohydrolases. <i>Circulation</i> , 2011 , 123, 1194-204	16.7	34
62	Hypoxic pulmonary hypertension in mice with constitutively active platelet-derived growth factor receptor- [Pulmonary Circulation, 2011, 1, 259-68]	2.7	38
61	The role of dimethylarginine dimethylaminohydrolase in idiopathic pulmonary fibrosis. <i>Science Translational Medicine</i> , 2011 , 3, 87ra53	17.5	50
60	A role for coagulation factor Xa in experimental pulmonary arterial hypertension. <i>Cardiovascular Research</i> , 2011 , 92, 159-68	9.9	26
59	Glycogen synthase kinase 3beta contributes to proliferation of arterial smooth muscle cells in pulmonary hypertension. <i>PLoS ONE</i> , 2011 , 6, e18883	3.7	33
58	Targeting non-malignant disorders with tyrosine kinase inhibitors. <i>Nature Reviews Drug Discovery</i> , 2010 , 9, 956-70	64.1	102
57	Role of epidermal growth factor inhibition in experimental pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2010 , 181, 158-67	10.2	99
56	Pulmonary vascular remodeling correlates with lung eggs and cytokines in murine schistosomiasis. American Journal of Respiratory and Critical Care Medicine, 2010, 181, 279-88	10.2	93
55	Dysregulation of the IL-13 receptor system: a novel pathomechanism in pulmonary arterial hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2010 , 182, 805-18	10.2	54
54	Animal models of pulmonary hypertension: role in translational research. <i>Drug Discovery Today:</i> Disease Models, 2010 , 7, 89-97	1.3	8
53	Phosphodiesterase 6 subunits are expressed and altered in idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2010 , 11, 146	7-3	17
52	Identification of right heart-enriched genes in a murine model of chronic outflow tract obstruction. Journal of Molecular and Cellular Cardiology, 2010 , 49, 598-605	5.8	47
51	PDGF receptor and its antagonists: role in treatment of PAH. <i>Advances in Experimental Medicine and Biology</i> , 2010 , 661, 435-46	3.6	47
50	Targeting cancer with phosphodiesterase inhibitors. <i>Expert Opinion on Investigational Drugs</i> , 2010 , 19, 117-31	5.9	105
49	Effects of phosphodiesterase 4 inhibition on bleomycin-induced pulmonary fibrosis in mice. <i>BMC Pulmonary Medicine</i> , 2010 , 10, 26	3.5	29
48	Heme oxygenase-2 and large-conductance Ca2+-activated K+ channels: lung vascular effects of hypoxia. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2009 , 180, 353-64	10.2	34
47	The noncanonical WNT pathway is operative in idiopathic pulmonary arterial hypertension. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2009 , 40, 683-91	5.7	80
46	The soluble guanylate cyclase activator HMR1766 reverses hypoxia-induced experimental pulmonary hypertension in mice. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2009 , 297, L658-65	5.8	32

45	Inflammation, growth factors, and pulmonary vascular remodeling. <i>Journal of the American College of Cardiology</i> , 2009 , 54, S10-S19	15.1	518
44	Inhibition of the soluble epoxide hydrolase attenuates monocrotaline-induced pulmonary hypertension in rats. <i>Journal of Hypertension</i> , 2009 , 27, 322-31	1.9	50
43	Fhl-1, a new key protein in pulmonary hypertension. <i>Circulation</i> , 2008 , 118, 1183-94	16.7	71
42	Combined tyrosine and serine/threonine kinase inhibition by sorafenib prevents progression of experimental pulmonary hypertension and myocardial remodeling. <i>Circulation</i> , 2008 , 118, 2081-90	16.7	121
41	Role of the prostanoid EP4 receptor in iloprost-mediated vasodilatation in pulmonary hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2008 , 178, 188-96	10.2	66
40	Characterization of a murine model of monocrotaline pyrrole-induced acute lung injury. <i>BMC Pulmonary Medicine</i> , 2008 , 8, 25	3.5	31
39	Dysregulated bone morphogenetic protein signaling in monocrotaline-induced pulmonary arterial hypertension. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2007 , 27, 1072-8	9.4	115
38	Iloprost-induced desensitization of the prostacyclin receptor in isolated rabbit lungs. <i>Respiratory Research</i> , 2007 , 8, 4	7.3	31
37	Receptor for activated C-kinase 1, a novel interaction partner of type II bone morphogenetic protein receptor, regulates smooth muscle cell proliferation in pulmonary arterial hypertension. <i>Circulation</i> , 2007 , 115, 2957-68	16.7	43
36	Hypoxia-dependent regulation of nonphagocytic NADPH oxidase subunit NOX4 in the pulmonary vasculature. <i>Circulation Research</i> , 2007 , 101, 258-67	15.7	279
35	Phosphodiesterase 1 upregulation in pulmonary arterial hypertension: target for reverse-remodeling therapy. <i>Circulation</i> , 2007 , 115, 2331-9	16.7	118
34	Hypoxia-induced pulmonary hypertension: different impact of iloprost, sildenafil, and nitric oxide. <i>Respiratory Medicine</i> , 2007 , 101, 2125-32	4.6	24
33	In vivo characterization of the novel imidazopyridine BYK191023 [2-[2-(4-methoxy-pyridin-2-yl)-ethyl]-3H-imidazo[4,5-b]pyridine], a potent and highly selective inhibitor of inducible nitric-oxide synthase. <i>Journal of Pharmacology and Experimental Therapeutics</i> ,	4.7	11
32	2006 , 317, 181-7 Activation of soluble guanylate cyclase reverses experimental pulmonary hypertension and vascular remodeling. <i>Circulation</i> , 2006 , 113, 286-95	16.7	183
31	Classical transient receptor potential channel 6 (TRPC6) is essential for hypoxic pulmonary vasoconstriction and alveolar gas exchange. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006 , 103, 19093-8	11.5	247
30	Comparison of pharmacokinetics and vasodilatory effect of nebulized and infused iloprost in experimental pulmonary hypertension: rapid tolerance development. <i>Journal of Aerosol Medicine and Pulmonary Drug Delivery</i> , 2006 , 19, 353-63		13
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