

James R Klinger

List of Publications by Year in descending order

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Version: 2024-02-01

81
papers

3,729
citations

159585

30
h-index

133252

59
g-index

81
all docs

81
docs citations

81
times ranked

4136
citing authors

#	ARTICLE	IF	CITATIONS
1	Remote 6-Minute-Walk Testing in Patients with Pulmonary Hypertension: A Pilot Study. American Journal of Respiratory and Critical Care Medicine, 2022, 205, 851-854.	5.6	8
2	Targeting RUNX1 as a novel treatment modality for pulmonary arterial hypertension. Cardiovascular Research, 2022, 118, 3211-3224.	3.8	16
3	Hispanic Ethnicity and Social Determinants of Health in Pulmonary Arterial Hypertension: The Pulmonary Hypertension Association Registry. Annals of the American Thoracic Society, 2022, 19, 1459-1468.	3.2	13
4	Delphi consensus recommendation for optimization of pulmonary hypertension therapy focusing on switching from a phosphodiesterase 5 inhibitor to riociguat. Pulmonary Circulation, 2022, 12, e12055.	1.7	6
5	Mesenchymal Stromal Cell Extracellular Vesicles - A New Approach for Preventing Bronchopulmonary Dysplasia?. American Journal of Respiratory and Critical Care Medicine, 2022, , .	5.6	1
6	Riociguat: Clinical research and evolving role in therapy. British Journal of Clinical Pharmacology, 2021, 87, 2645-2662.	2.4	18
7	EmPHasis-10 as a measure of health-related quality of life in pulmonary arterial hypertension: data from PHAR. European Respiratory Journal, 2021, 57, 2000414.	6.7	24
8	Prediction of Health-related Quality of Life and Hospitalization in Pulmonary Arterial Hypertension: The Pulmonary Hypertension Association Registry. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 761-764.	5.6	12
9	Insights from the Menstrual Cycle in Pulmonary Arterial Hypertension. Annals of the American Thoracic Society, 2021, 18, 218-228.	3.2	15
10	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.	10.7	85
11	Health disparities and treatment approaches in portopulmonary hypertension and idiopathic pulmonary arterial hypertension: an analysis of the Pulmonary Hypertension Association Registry. Pulmonary Circulation, 2021, 11, 1-10.	1.7	17
12	Treatment of Pulmonary Hypertension Associated With COPD. Chest, 2021, 160, 409-410.	0.8	2
13	Effect of dose, dosing intervals, and hypoxic stress on the reversal of pulmonary hypertension by mesenchymal stem cell extracellular vesicles. Pulmonary Circulation, 2021, 11, 1-11.	1.7	3
14	Novel Pharmacological Targets for Pulmonary Arterial Hypertension. , 2021, 11, 2297-2349.		5
15	Mesenchymal Stem Cell Extracellular Vesicles Reverse Sugen/Hypoxia Pulmonary Hypertension in Rats. American Journal of Respiratory Cell and Molecular Biology, 2020, 62, 577-587.	2.9	54
16	Guidelines for the Treatment of Pulmonary Arterial Hypertension. Lung, 2020, 198, 581-596.	3.3	37
17	Prevalence and risk factors of pulmonary hypertension among adult patients with HIV infection in Ethiopia. Pulmonary Circulation, 2020, 10, 204589402097151.	1.7	4
18	Residence at moderately high altitude and its relationship with WHO Group 1 pulmonary arterial hypertension symptom severity and clinical characteristics: the Pulmonary Hypertension Association Registry. Pulmonary Circulation, 2020, 10, 1-8.	1.7	5

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19	Identifying potential parameters associated with response to switching from a PDE5i to riociguat in RESPITE. <i>International Journal of Cardiology</i> , 2020, 317, 188-192.	1.7	5
20	Rapid development of pulmonary hypertension and right ventricular failure due to large vessel intravascular microcrystalline cellulosis in an intravenous drug user. <i>Pulmonary Circulation</i> , 2020, 10, 1-3.	1.7	4
21	Alternative Splicing of the Cardiac Sodium Channel in Pulmonary Arterial Hypertension. <i>Chest</i> , 2020, 158, 735-738.	0.8	6
22	Culture of pulmonary artery endothelial cells from pulmonary artery catheter balloon tips: considerations for use in pulmonary vascular disease. <i>European Respiratory Journal</i> , 2020, 55, 1901313.	6.7	10
23	Late Breaking Abstract - Switching from PDE5i to riociguat in patients with PAH: The REPLACE study. , 2020, , .		2
24	Response. <i>Chest</i> , 2019, 156, 187-188.	0.8	0
25	Low dose 100â€%cGy irradiation as a potential therapy for pulmonary hypertension. <i>Journal of Cellular Physiology</i> , 2019, 234, 21193-21198.	4.1	9
26	Genetic determinants of risk in pulmonary arterial hypertension: international genome-wide association studies and meta-analysis. <i>Lancet Respiratory Medicine</i> , the, 2019, 7, 227-238.	10.7	122
27	Therapy for Pulmonary Arterial Hypertension in Adults. <i>Chest</i> , 2019, 155, 565-586.	0.8	216
28	Pathology and pathobiology of pulmonary hypertension: state of the art and research perspectives. <i>European Respiratory Journal</i> , 2019, 53, 1801887.	6.7	776
29	Chronic Thromboembolic Pulmonary Hypertension. <i>Heart Failure Clinics</i> , 2018, 14, 339-351.	2.1	19
30	Anastrozole in Pulmonary Arterial Hypertension. A Randomized, Double-Blind, Placebo-controlled Trial. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2017, 195, 360-368.	5.6	88
31	Riociguat: Mode of Action and Clinical Development in Pulmonary Hypertension. <i>Chest</i> , 2017, 151, 468-480.	0.8	79
32	Rationale and study design of RESPITE: An open-label, phase 3b study of riociguat in patients with pulmonary arterial hypertension who demonstrate an insufficient response to treatment with phosphodiesterase-5 inhibitors. <i>Respiratory Medicine</i> , 2017, 122, S18-S22.	2.9	15
33	The Nitric Oxide Pathway in Pulmonary Vascular Disease. <i>American Journal of Cardiology</i> , 2017, 120, S71-S79.	1.6	79
34	RESPITE: switching to riociguat in pulmonary arterial hypertension patients with inadequate response to phosphodiesterase-5 inhibitors. <i>European Respiratory Journal</i> , 2017, 50, 1602425.	6.7	113
35	Ask The Expert: What Are Some Pitfalls and Promises of the Current PAH Treatment Guidelines?. <i>Advances in Pulmonary Hypertension</i> , 2017, 15, 182-183.	0.1	0
36	Plasma nitrite/nitrate levels: a new biomarker for pulmonary arterial hypertension?. <i>European Respiratory Journal</i> , 2016, 48, 1265-1267.	6.7	5

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37	Group III Pulmonary Hypertension. <i>Cardiology Clinics</i> , 2016, 34, 413-433.	2.2	70
38	Socioeconomic Status Affects Pulmonary Hypertension Disease Severity at Time of First Evaluation. <i>Pulmonary Circulation</i> , 2016, 6, 191-195.	1.7	27
39	Exosomes induce and reverse monocrotaline-induced pulmonary hypertension in mice. <i>Cardiovascular Research</i> , 2016, 110, 319-330.	3.8	196
40	Modulation of cGMP Synthesis and Metabolism. <i>Respiratory Medicine</i> , 2015, , 355-375.	0.1	1
41	Sepsis and Pulmonary Arterial Hypertension in the ICU. <i>Advances in Pulmonary Hypertension</i> , 2015, 13, 188-196.	0.1	4
42	Pharmacologic Therapy for Pulmonary Arterial Hypertension in Adults. <i>Chest</i> , 2014, 146, 449-475.	0.8	237
43	Cost-effectiveness of Dalteparin vs Unfractionated Heparin for the Prevention of Venous Thromboembolism in Critically Ill Patients. <i>JAMA - Journal of the American Medical Association</i> , 2014, 312, 2135.	7.4	50
44	Effects of Dose and Age on Adverse Events Associated with Tadalafil in the Treatment of Pulmonary Arterial Hypertension. <i>Pulmonary Circulation</i> , 2014, 4, 45-52.	1.7	5
45	Economic evaluation of the prophylaxis for thromboembolism in critical care trial (E-PROTECT): study protocol for a randomized controlled trial. <i>Trials</i> , 2014, 15, 502.	1.6	10
46	Nitric Oxide Deficiency and Endothelial Dysfunction in Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2013, 188, 639-646.	5.6	165
47	Oral Therapies for Pulmonary Arterial Hypertension. <i>Clinics in Chest Medicine</i> , 2013, 34, 811-824.	2.1	6
48	Atrial natriuretic peptide attenuates agonist-induced pulmonary edema in mice with targeted disruption of the gene for natriuretic peptide receptor-A. <i>Journal of Applied Physiology</i> , 2013, 114, 307-315.	2.5	8
49	WHO Group 1 pulmonary arterial hypertension: Current and investigative therapies. <i>Progress in Cardiovascular Diseases</i> , 2012, 55, 89-103.	3.1	27
50	Initial Risk Assessment for Pulmonary Hypertension in Patients with COPD. <i>Lung</i> , 2012, 190, 83-89.	3.3	17
51	Transfer of Monocrotaline-Induced Pulmonary Hypertension to Healthy Mice Via Microparticles. <i>Blood</i> , 2012, 120, 5190-5190.	1.4	0
52	C-type natriuretic peptide does not attenuate the development of pulmonary hypertension caused by hypoxia and VEGF receptor blockade. <i>Life Sciences</i> , 2011, 89, 460-466.	4.3	10
53	Tadalafil for the treatment of pulmonary arterial hypertension. <i>Expert Review of Respiratory Medicine</i> , 2011, 5, 315-328.	2.5	7
54	Long-Term Pulmonary Hemodynamic Effects of Ambrisentan in Pulmonary Arterial Hypertension. <i>American Journal of Cardiology</i> , 2011, 108, 302-307.	1.6	44

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55	Echocardiographic Evidence of Pulmonary Hypertension is Associated with Increased 1-year Mortality in Patients Admitted with Chronic Obstructive Pulmonary Disease. <i>Lung</i> , 2011, 189, 207-212.	3.3	33
56	Pulmonary Hypertension in a Stable Community-Based COPD Population. <i>Lung</i> , 2011, 189, 377-382.	3.3	35
57	Tadalafil in Geriatric Patients With Pulmonary Arterial Hypertension. <i>Chest</i> , 2010, 138, 367A.	0.8	3
58	Cardiac atria are the primary source of ANP release in hypoxia-adapted rats. <i>Life Sciences</i> , 2010, 87, 382-389.	4.3	11
59	Brain natriuretic peptide in pulmonary arterial hypertension: biomarker and potential therapeutic agent. <i>Drug Design, Development and Therapy</i> , 2009, 3, 269.	4.3	48
60	Marrow Cell Infusion Attenuates Vascular Remodeling in a Murine Model of Monocrotaline-Induced Pulmonary Hypertension. <i>Stem Cells and Development</i> , 2009, 18, 773-781.	2.1	16
61	Pulmonary Arterial Hypertension in Pregnancy. , 2009, , 285-312.		0
62	Diagnosis and Management of Pulmonary Hypertension Associated With Pulmonary Fibrosis. <i>Advances in Pulmonary Hypertension</i> , 2009, 8, 141-147.	0.1	0
63	Pulmonary Arterial Hypertension: An Overview. <i>Seminars in Cardiothoracic and Vascular Anesthesia</i> , 2007, 11, 96-103.	1.0	3
64	Pulmonary hypertension in the intensive care unit: Critical role of the right ventricle*. <i>Critical Care Medicine</i> , 2007, 35, 2210-2211.	0.9	9
65	The Nitric Oxide/cGMP Signaling Pathway in Pulmonary Hypertension. <i>Clinics in Chest Medicine</i> , 2007, 28, 143-167.	2.1	74
66	Rottlerin causes pulmonary edema in vivo: a possible role for PKC δ . <i>Journal of Applied Physiology</i> , 2007, 103, 2084-2094.	2.5	25
67	Natriuretic peptides differentially attenuate thrombin-induced barrier dysfunction in pulmonary microvascular endothelial cells. <i>Experimental Cell Research</i> , 2006, 312, 401-410.	2.6	34
68	Pulmonary Hemodynamic Responses to Brain Natriuretic Peptide and Sildenafil in Patients With Pulmonary Arterial Hypertension. <i>Chest</i> , 2006, 129, 417-425.	0.8	90
69	Acute Cardiopulmonary Hemodynamic Effects of Brain Natriuretic Peptide in Patients With Pulmonary Arterial Hypertension. <i>Chest</i> , 2005, 128, 618S-619S.	0.8	8
70	Pulmonary hypertension: inhaled nitric oxide, sildenafil and natriuretic peptides. <i>Current Opinion in Pharmacology</i> , 2005, 5, 245-250.	3.5	47
71	Acute and chronic effects of sildenafil in patients with pulmonary arterial hypertension. <i>Respiratory Medicine</i> , 2005, 99, 1501-1510.	2.9	79
72	Synergistic Effects of ANP and Sildenafil on cGMP Levels and Amelioration of Acute Hypoxic Pulmonary Hypertension. <i>Experimental Biology and Medicine</i> , 2004, 229, 920-925.	2.4	38

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73	Pulmonary Edema Caused by Inhaled Nitric Oxide Therapy in Two Patients With Pulmonary Hypertension Associated With the CREST Syndrome. <i>Chest</i> , 2002, 121, 656-659.	0.8	48
74	Inhaled nitric oxide in ARDS. <i>Critical Care Clinics</i> , 2002, 18, 45-68.	2.6	27
75	Targeted disruption of the gene for natriuretic peptide receptor-A worsens hypoxia-induced cardiac hypertrophy. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2002, 282, H58-H65.	3.2	36
76	Vasoresponsiveness of Sarcoidosis-Associated Pulmonary Hypertension. <i>Chest</i> , 2001, 120, 866-872.	0.8	121
77	Genetic disruption of atrial natriuretic peptide causes pulmonary hypertension in normoxic and hypoxic mice. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 1999, 276, L868-L874.	2.9	38
78	Brain natriuretic peptide inhibits hypoxic pulmonary hypertension in rats. <i>Journal of Applied Physiology</i> , 1998, 84, 1646-1652.	2.5	50
79	C-Receptor Ligand Blocks Pulmonary Clearance of Atrial Natriuretic Peptide in Isolated Rat Lungs. <i>Experimental Biology and Medicine</i> , 1992, 201, 154-158.	2.4	11
80	Right Ventricular Dysfunction in Chronic Obstructive Pulmonary Disease*. <i>Chest</i> , 1991, 99, 715-723.	0.8	88
81	Reply to: Remote 6-minute Walk Testing in Patients with Pulmonary Hypertension: Further Validation Needed?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 0, , .	5.6	0