Xavier Jais

List of Publications by Year in descending order

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146	16,810	59	125
papers	citations	h-index	g-index
151	151	151	9121
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Survival in Patients With Idiopathic, Familial, and Anorexigen-Associated Pulmonary Arterial Hypertension in the Modern Management Era. Circulation, 2010, 122, 156-163.	1.6	1,264
2	Long-Term Response to Calcium Channel Blockers in Idiopathic Pulmonary Arterial Hypertension. Circulation, 2005, 111, 3105-3111.	1.6	1,040
3	Chronic Thromboembolic Pulmonary Hypertension (CTEPH). Circulation, 2011, 124, 1973-1981.	1.6	860
4	Chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2019, 53, 1801915.	3.1	607
5	Pulmonary Arterial Hypertension in Patients Treated by Dasatinib. Circulation, 2012, 125, 2128-2137.	1.6	548
6	Risk assessment, prognosis and guideline implementation in pulmonary arterial hypertension. European Respiratory Journal, 2017, 50, 1700889.	3.1	527
7	Bosentan for Treatment of Inoperable Chronic Thromboembolic Pulmonary Hypertension. Journal of the American College of Cardiology, 2008, 52, 2127-2134.	1.2	506
8	Long-Term Outcome of Patients With Chronic Thromboembolic Pulmonary Hypertension. Circulation, 2016, 133, 859-871.	1.6	506
9	Complications of Right Heart Catheterization Procedures in Patients With Pulmonary Hypertension in Experienced Centers. Journal of the American College of Cardiology, 2006, 48, 2546-2552.	1.2	498
10	Severe Pulmonary Hypertension during Pregnancy. Anesthesiology, 2005, 102, 1133-1137.	1.3	483
11	Immunosuppressive therapy in lupus―and mixed connective tissue disease–associated pulmonary arterial hypertension: A retrospective analysis of twentyâ€ŧhree cases. Arthritis and Rheumatism, 2008, 58, 521-531.	6.7	321
12	Upfront triple combination therapy in pulmonary arterial hypertension: a pilot study. European Respiratory Journal, 2014, 43, 1691-1697.	3.1	319
13	Immunosuppressive Therapy in Connective Tissue Diseases-Associated Pulmonary Arterial Hypertension. Chest, 2006, 130, 182-189.	0.4	316
14	Pulmonary Veno-Occlusive Disease. Medicine (United States), 2008, 87, 220-233.	0.4	295
15	Pulmonary veno-occlusive disease. European Respiratory Journal, 2016, 47, 1518-1534.	3.1	289
16	ERS statement on chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2021, 57, 2002828.	3.1	287
17	Advances in Therapeutic Interventions for Patients With Pulmonary Arterial Hypertension. Circulation, 2014, 130, 2189-2208.	1.6	278
18	Deleterious Effects of \hat{I}^2 -Blockers on Exercise Capacity and Hemodynamics in Patients With Portopulmonary Hypertension. Gastroenterology, 2006, 130, 120-126.	0.6	277

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19	Long-term outcome with first-line bosentan therapy in idiopathic pulmonary arterial hypertension. European Heart Journal, 2006, 27, 589-595.	1.0	272
20	Clinical Outcomes of Pulmonary Arterial Hypertension in Carriers of <i>BMPR2</i> Mutation. American Journal of Respiratory and Critical Care Medicine, 2008, 177, 1377-1383.	2.5	269
21	Clinical Outcomes of Pulmonary Arterial Hypertension in Patients Carrying an <i>ACVRL1</i> (<i>ALK1</i>) Mutation. American Journal of Respiratory and Critical Care Medicine, 2010, 181, 851-861.	2.5	259
22	Pulmonary arterial hypertension. Orphanet Journal of Rare Diseases, 2013, 8, 97.	1.2	226
23	Pregnancy outcomes in pulmonary arterial hypertension in the modern management era. European Respiratory Journal, 2012, 40, 881-885.	3.1	221
24	Portopulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2008, 178, 637-643.	2.5	220
25	Long-term response to calcium-channel blockers in non-idiopathic pulmonary arterial hypertension. European Heart Journal, 2010, 31, 1898-1907.	1.0	218
26	Criteria for diagnosis of exercise pulmonary hypertension. European Respiratory Journal, 2015, 46, 728-737.	3.1	213
27	Microvascular disease in chronic thromboembolic pulmonary hypertension: a role for pulmonary veins and systemic vasculature. European Respiratory Journal, 2014, 44, 1275-1288.	3.1	201
28	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
29	Targeted therapies in pulmonary arterial hypertension. , 2014, 141, 172-191.		171
30	Pulmonary Arterial Hypertension: A Rare Complication of Primary Sjögren Syndrome. Medicine (United) Tj ETQq	0 0.0 rgBT	/Qyerlock 10
31	Prognostic Value of Follow-Up Hemodynamic Variables After Initial Management in Pulmonary Arterial Hypertension. Circulation, 2018, 137, 693-704.	1.6	155
32	HIV-associated pulmonary arterial hypertension: survival and prognostic factors in the modern therapeutic era. Aids, 2010, 24, 67-75.	1.0	149
33	Chemotherapy-Induced Pulmonary Hypertension. American Journal of Pathology, 2015, 185, 356-371.	1.9	149
34	Outcomes of noncardiac, nonobstetric surgery in patients with PAH: an international prospective survey. European Respiratory Journal, 2013, 41, 1302-1307.	3.1	131
35	Intravenous Epoprostenol in Inoperable Chronic Thromboembolic Pulmonary Hypertension. Journal of Heart and Lung Transplantation, 2007, 26, 357-362.	0.3	126
36	Treatment of pulmonary arterial hypertension with targeted therapies. Nature Reviews Cardiology, 2011, 8, 526-538.	6.1	125

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37	Initial dual oral combination therapy in pulmonary arterial hypertension. European Respiratory Journal, 2016, 47, 1727-1736.	3.1	124
38	Clinical phenotypes and outcomes of heritable and sporadic pulmonary veno-occlusive disease: a population-based study. Lancet Respiratory Medicine, the, 2017, 5, 125-134.	5.2	123
39	Potts Shunt in Children With Idiopathic Pulmonary Arterial Hypertension: Long-Term Results. Annals of Thoracic Surgery, 2012, 94, 817-824.	0.7	116
40	Pharmacokinetic and clinical profile of a novel formulation of bosentan in children with pulmonary arterial hypertension: the FUTUREâ€l study. British Journal of Clinical Pharmacology, 2009, 68, 948-955.	1.1	105
41	Mitomycin-Induced Pulmonary Veno-Occlusive Disease. Circulation, 2015, 132, 834-847.	1.6	103
42	Phosphodiesterase type 5 inhibitors in pulmonary arterial hypertension. Advances in Therapy, 2009, 26, 813-825.	1.3	96
43	Pulmonary veno-occlusive disease: Recent progress and current challenges. Respiratory Medicine, 2010, 104, S23-S32.	1.3	94
44	Association between Initial Treatment Strategy and Long-Term Survival in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2021, 204, 842-854.	2.5	94
45	Efficacy, safety and pharmacokinetics of bosentan in portopulmonary hypertension. European Respiratory Journal, 2013, 41, 96-103.	3.1	92
46	Usefulness of first-line combination therapy with epoprostenol and bosentan in pulmonary arterial hypertension: An observational study. Journal of Heart and Lung Transplantation, 2012, 31, 150-158.	0.3	91
47	Characteristics and outcomes of asthmatic patients with COVID-19 pneumonia who require hospitalisation. European Respiratory Journal, 2020, 56, 2001875.	3.1	90
48	Long-term outcomes of dasatinib-induced pulmonary arterial hypertension: a population-based study. European Respiratory Journal, 2017, 50, 1700217.	3.1	89
49	Genetic counselling in a national referral centre for pulmonary hypertension. European Respiratory Journal, 2016, 47, 541-552.	3.1	87
50	External validation of a refined four-stratum risk assessment score from the French pulmonary hypertension registry. European Respiratory Journal, 2022, 59, 2102419.	3.1	83
51	Pulmonary hypertension in lymphangioleiomyomatosis: characteristics in 20 patients. European Respiratory Journal, 2012, 40, 630-640.	3.1	80
52	Pulmonary arterial hypertension in patients treated with interferon. European Respiratory Journal, 2014, 44, 1627-1634.	3.1	80
53	Occupational exposure to organic solvents: a risk factor for pulmonary veno-occlusive disease. European Respiratory Journal, 2015, 46, 1721-1731.	3.1	80
54	Portopulmonary hypertension in the current era of pulmonary hypertension management. Journal of Hepatology, 2020, 73, 130-139.	1.8	78

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55	Rapid Switch From Intravenous Epoprostenol to Intravenous Treprostinil in Patients With Pulmonary Arterial Hypertension. Journal of Cardiovascular Pharmacology, 2007, 49, 1-5.	0.8	77
56	Pulmonary hypertension associated with benfluorex exposure. European Respiratory Journal, 2012, 40, 1164-1172.	3.1	75
57	<i>BMPR2</i> mutation status influences bronchial vascular changes in pulmonary arterial hypertension. European Respiratory Journal, 2016, 48, 1668-1681.	3.1	68
58	Longâ€term outcome in liver transplantation candidates with portopulmonary hypertension. Hepatology, 2017, 65, 1683-1692.	3.6	68
59	Pulmonary Arterial Hypertension Associated With Systemic Lupus Erythematosus. Chest, 2018, 153, 143-151.	0.4	68
60	Pulmonary Hypertension in Patients With Neurofibromatosis Type I. Medicine (United States), 2011, 90, 201-211.	0.4	60
61	Haemodynamics and serial risk assessment in systemic sclerosis associated pulmonary arterial hypertension. European Respiratory Journal, 2018, 52, 1800678.	3.1	60
62	An Extreme Consequence of Splenectomy in Dehydrated Hereditary Stomatocytosis: Gradual Thromboâ€embolic Pulmonary Hypertension and Lungâ€"Heart Transplantation. Hemoglobin, 2003, 27, 139-147.	0.4	59
63	Screening for pulmonary arterial hypertension in systemic sclerosis. European Respiratory Review, 2019, 28, 190023.	3.0	59
64	Predictors of survival in patients with not-operated chronic thromboembolic pulmonary hypertension. Journal of Heart and Lung Transplantation, 2019, 38, 833-842.	0.3	57
65	Loss of Vascular Distensibility During Exercise Is an Early Hemodynamic Marker of Pulmonary Vascular Disease. Chest, 2016, 149, 353-361.	0.4	55
66	Ventilation/perfusion lung scan in pulmonary veno-occlusive disease. European Respiratory Journal, 2012, 40, 75-83.	3.1	53
67	Pulmonary vascular remodeling patterns and expression of general control nonderepressible 2 (GCN2) in pulmonary veno-occlusive disease. Journal of Heart and Lung Transplantation, 2018, 37, 647-655.	0.3	50
68	Screening for pulmonary arterial hypertension in adults carrying a <i>BMPR2</i> mutation. European Respiratory Journal, 2021, 58, 2004229.	3.1	50
69	Acute decompensated pulmonary hypertension. European Respiratory Review, 2017, 26, 170092.	3.0	48
70	Clinical phenotypes and survival of pre-capillary pulmonary hypertension in systemic sclerosis. PLoS ONE, 2018, 13, e0197112.	1.1	47
71	Characterization of Pulmonary Arterial Hypertension Patients Walking More Than 450 m in 6 Min at Diagnosis. Chest, 2010, 137, 1297-1303.	0.4	46
72	Pulmonary Hypertension Complicating Fibrosing Mediastinitis. Medicine (United States), 2015, 94, e1800.	0.4	46

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73	Deterioration of pulmonary hypertension and pleural effusion with bosutinib following dasatinib lung toxicity. European Respiratory Journal, 2016, 48, 1517-1519.	3.1	44
74	Resting pulmonary artery pressure of 21–24â€mmHg predicts abnormal exercise haemodynamics. European Respiratory Journal, 2016, 47, 1436-1444.	3.1	44
75	Mechanisms of exertional dyspnoea in pulmonary veno-occlusive disease with <i>EIF2AK4</i> mutations. European Respiratory Journal, 2014, 44, 1069-1072.	3.1	43
76	Association Between BMI and Obesity With Survival in Pulmonary Arterial Hypertension. Chest, 2018, 154, 872-881.	0.4	43
77	RV Fractional Area Change and TAPSE as Predictors of Severe Right Ventricular Dysfunction in Pulmonary Hypertension: A CMR Study. Lung, 2018, 196, 157-164.	1.4	42
78	Prevalence of pulmonary embolism in patients with COVID-19 at the time of hospital admission. European Respiratory Journal, 2021, 58, 2100116.	3.1	41
79	Phenotype and outcome of pulmonary arterial hypertension patients carrying a <i>TBX4</i> mutation. European Respiratory Journal, 2020, 55, 1902340.	3.1	40
80	Birth Control and Pregnancy Management in Pulmonary Hypertension. Seminars in Respiratory and Critical Care Medicine, 2013, 34, 681-688.	0.8	39
81	FUTURE-2: Results from an open-label, long-term safety and tolerability extension study using the pediatric FormUlation of bosenTan in pUlmonary arterial hypeRtEnsion. International Journal of Cardiology, 2016, 202, 52-58.	0.8	37
82	Outcome of adults with Eisenmenger syndrome treated with drugs specific to pulmonary arterial hypertension: A French multicentre study. Archives of Cardiovascular Diseases, 2017, 110, 303-316.	0.7	37
83	Dead-space ventilation is linked to exercise capacity and survival in distal chronic thromboembolic pulmonary hypertension. Journal of Heart and Lung Transplantation, 2017, 36, 1234-1242.	0.3	37
84	Comparative Safety and Tolerability of Prostacyclins in Pulmonary Hypertension. Drug Safety, 2016, 39, 287-294.	1.4	35
85	Review: Therapeutic advances in pulmonary arterial hypertension. Therapeutic Advances in Respiratory Disease, 2008, 2, 249-265.	1.0	33
86	Characteristics of Pulmonary Arterial Hypertension in Affected Carriers of a Mutation Located in the Cytoplasmic Tail of Bone Morphogenetic Protein Receptor Type 2. Chest, 2015, 147, 1385-1394.	0.4	33
87	Diagnostic concordance of different criteria for exercise pulmonary hypertension in subjects with normal resting pulmonary artery pressure. European Respiratory Journal, 2016, 48, 254-257.	3.1	31
88	Sex and gender in pulmonary arterial hypertension. European Respiratory Review, 2021, 30, 200330.	3.0	31
89	Impact of High-Priority Allocation on Lung and Heart-Lung Transplantation for Pulmonary Hypertension. Annals of Thoracic Surgery, 2017, 104, 404-411.	0.7	29
90	Factors predicting outcome after pulmonary endarterectomy. PLoS ONE, 2018, 13, e0198198.	1.1	29

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91	Current epoprostenol use in patients with severe idiopathic, heritable or anorexigen-associated pulmonary arterial hypertension: Data from the French pulmonary hypertension registry. International Journal of Cardiology, 2014, 172, 561-567.	0.8	28
92	Pulmonary complications of Bcr-Abl tyrosine kinase inhibitors. European Respiratory Journal, 2020, 56, 2000279.	3.1	28
93	Serum and pulmonary uric acid in pulmonary arterial hypertension. European Respiratory Journal, 2021, 58, 2000332.	3.1	28
94	Clinical Pharmacology of Endothelin Receptor Antagonists Used in the Treatment of Pulmonary Arterial Hypertension. American Journal of Cardiovascular Drugs, 2015, 15, 13-26.	1.0	27
95	Extracorporeal Life Support After Pulmonary Endarterectomy as a Bridge to Recovery or Transplantation: Lessons From 31 Consecutive Patients. Annals of Thoracic Surgery, 2016, 102, 260-268.	0.7	27
96	Left Ventricular Ejection Time in Acute Heart Failure Complicating Precapillary Pulmonary Hypertension. Chest, 2013, 144, 1512-1520.	0.4	26
97	Chronic thromboembolic pulmonary hypertension. Presse Medicale, 2015, 44, e409-e416.	0.8	26
98	Pulmonary veno-occlusive disease: The $b\tilde{A}^a$ te noire of pulmonary hypertension in connective tissue diseases? Presse Medicale, 2011, 40, e87-e100.	0.8	25
99	A Clinical and Echocardiographic Score to Identify Pulmonary Hypertension Due to HFpEF. Journal of Cardiac Failure, 2017, 23, 29-35.	0.7	25
100	Pulmonary hypertension associated with neurofibromatosis type 1. European Respiratory Review, 2018, 27, 180053.	3.0	25
101	Intensity and quality of exertional dyspnoea in patients with stable pulmonary hypertension. European Respiratory Journal, 2020, 55, 1802108.	3.1	24
102	Clinical Challenges in Pulmonary Hypertension. Chest, 2005, 128, 622S-628S.	0.4	23
103	Endothelin receptor antagonists for the treatment of pulmonary arterial hypertension. Expert Opinion on Pharmacotherapy, 2011, 12, 1585-1596.	0.9	23
104	Out-of-Proportion Pulmonary Hypertension and Heart Failure with Preserved Ejection Fraction. Respiration, 2013, 85, 471-477.	1.2	20
105	New pharmacotherapy options for pulmonary arterial hypertension. Expert Opinion on Pharmacotherapy, 2015, 16, 2113-2131.	0.9	20
106	Impact of the initiation of balloon pulmonary angioplasty program on referral of patients with chronic thromboembolic pulmonary hypertension to surgery. Journal of Heart and Lung Transplantation, 2018, 37, 1102-1110.	0.3	20
107	Association between Rheumatoid Arthritis and Pulmonary Hypertension: Data from the French Pulmonary Hypertension Registry. Respiration, 2018, 95, 244-250.	1.2	17
108	Respiratory symptoms and radiological findings in post-acute COVID-19 syndrome. ERJ Open Research, 2022, 8, 00479-2021.	1.1	16

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109	Medical Therapy in Chronic Thromboembolic Pulmonary Hypertension. Annals of the American Thoracic Society, 2016, 13, S248-S254.	1.5	15
110	Long-term outcomes of pulmonary arterial hypertension under specific drug therapy in Eisenmenger syndrome. Journal of Heart and Lung Transplantation, 2017, 36, 386-398.	0.3	15
111	Characteristics and Long-term Outcomes of Pulmonary Venoocclusive Disease Induced by Mitomycin C. Chest, 2021, 159, 1197-1207.	0.4	14
112	EBUS-TBNA in the differential diagnosis of pulmonary artery sarcoma and thromboembolism: Figure $1\hat{a}\in$ ". European Respiratory Journal, 2012, 39, 1549-1550.	3.1	13
113	Direct-Acting Antiviral Medications for Hepatitis C Virus Infection and Pulmonary Arterial Hypertension. Chest, 2016, 150, 256-258.	0.4	12
114	Phenotype and Outcomes of Pulmonary Hypertension Associated with Neurofibromatosis Type 1. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 843-852.	2.5	12
115	Chronic thromboembolic pulmonary hypertension and totally implantable central venous access systems. European Respiratory Journal, 2021, 57, 2002208.	3.1	12
116	Usefulness of Cardiovascular Magnetic Resonance IndicesÂto Rule In or Rule Out Precapillary Pulmonary Hypertension. Canadian Journal of Cardiology, 2015, 31, 1469-1476.	0.8	10
117	Lung capillary blood volume and membrane diffusion in precapillary pulmonary hypertension. Journal of Heart and Lung Transplantation, 2016, 35, 647-656.	0.3	10
118	Clinical and Hemodynamic Correlates of Pulmonary Arterial Stiffness in Incident, Untreated Patients With Idiopathic Pulmonary Arterial Hypertension. Chest, 2018, 154, 882-892.	0.4	10
119	Gas Exchange and Ventilatory Efficiency During Exercise in Pulmonary Vascular Diseases. Archivos De Bronconeumologia, 2020, 56, 578-585.	0.4	10
120	Association between Leflunomide and Pulmonary Hypertension. Annals of the American Thoracic Society, 2021, 18, 1306-1315.	1.5	8
121	The potential for macitentan, a new dual endothelin receptor antagonist, in the treatment of pulmonary arterial hypertension. Therapeutic Advances in Respiratory Disease, 2014, 8, 84-92.	1.0	7
122	A rare case of sarcoidosis-associated pulmonary hypertension in a patient exposed to silica. European Respiratory Review, 2016, 25, 93-96.	3.0	7
123	Preoperative C-reactive protein predicts early postoperative outcomes after pulmonary endarterectomy in patients with chronic thromboembolic pulmonary hypertension. Journal of Thoracic and Cardiovascular Surgery, 2021, 161, 1532-1542.e5.	0.4	7
124	Five-year survival after an acute episode of decompensated pulmonary arterial hypertension in the modern management era of right heart failure. European Respiratory Journal, 2021, 58, 2100466.	3.1	7
125	Pulmonary thromboendarterectomy: The Marie Lannelongue Hospital experience. Annals of Cardiothoracic Surgery, 2022, 11, 143-150.	0.6	6
126	Risk stratification in patients with pulmonary arterial hypertension at the time of listing for lung transplantation. Journal of Heart and Lung Transplantation, 2022, 41, 1285-1293.	0.3	6

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127	Mediastinal Fibrosis Mimicking Proximal Chronic Thromboembolic Disease. Circulation, 2012, 125, 2045-2047.	1.6	5
128	Atypical Vasculitis Mimicking Chronic Thromboembolic Pulmonary Hypertension. American Journal of Medicine, 2015, 128, e47-e49.	0.6	5
129	Pharmacovigilance in a rare disease: example of the VIGIAPATH program in pulmonary arterial hypertension. International Journal of Clinical Pharmacy, 2018, 40, 790-794.	1.0	5
130	Risks and outcomes of gastrointestinal endoscopy with anaesthesia in patients with pulmonary hypertension. British Journal of Anaesthesia, 2020, 125, e466-e468.	1.5	5
131	Response to Letter Regarding Article, "Mitomycin-Induced Pulmonary Veno-Occlusive Disease: Evidence From Human Disease and Animal Model― Circulation, 2016, 133, e592-3.	1.6	4
132	Are indexed values better for defining exercise pulmonary hypertension?. European Respiratory Journal, 2017, 50, 1700240.	3.1	4
133	Evaluation of a collaborative care program for pulmonary hypertension patients: a multicenter randomized trial. International Journal of Clinical Pharmacy, 2020, 42, 1128-1138.	1.0	4
134	Lung Ventilation/Perfusion Scintigraphy for the Screening of Chronic Thromboembolic Pulmonary Hypertension (CTEPH): Which Criteria to Use?. Frontiers in Medicine, 2022, 9, 851935.	1.2	4
135	Chronic thromboembolic pulmonary hypertension complicating long-term cyproterone acetate therapy. European Respiratory Review, 2014, 23, 260-263.	3.0	3
136	Response to Letter Regarding Article, "Advances in Therapeutic Interventions for Patients With Pulmonary Arterial Hypertension― Circulation, 2015, 132, e154.	1.6	3
137	Relation between left ventricular ejection time and pulmonary hemodynamics in pulmonary hypertension. International Journal of Cardiology, 2015, 184, 763-765.	0.8	3
138	Reversible pulmonary hypertension associated with multivisceral Whipple's disease. European Respiratory Journal, 2021, 57, 2003132.	3.1	3
139	Pulmonary Hypertension in Patients with Common Variable Immunodeficiency. Journal of Clinical Immunology, 2021, 41, 1549-1562.	2.0	3
140	Transplantation for pulmonary arterial hypertension with congenital heart disease: Impact on outcomes of the current therapeutic approach including a high-priority allocation program. American Journal of Transplantation, 2021, 21, 3388-3400.	2.6	3
141	Pulmonary hypertension associated with busulfan. Pulmonary Circulation, 2021, 11, 1-12.	0.8	3
142	Sequential combination therapy with parenteral prostacyclin in BMPR2 mutations carriers. Pulmonary Circulation, 2022, 12, e12023.	0.8	2
143	Pulmonary Hypertension Complicating Pulmonary Artery Involvement in Pseudoxanthoma Elasticum. American Journal of Respiratory and Critical Care Medicine, 2020, 202, e90-e91.	2.5	1
144	Pulmonary Vascular Resistance in Pulmonary Arterial Hypertension: La Pièce de Résistance?. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 524-525.	2.5	1

#	ARTICLE	IF	CITATIONS
145	Reply to: Jin et al. and Sun et al American Journal of Respiratory and Critical Care Medicine, 2021, , .	2.5	O
146	Right heart failure., 0,, 32-47.		0