Olivier Sitbon

List of Publications by Year
in descending order

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356 papers	35,753 citations	4388 86 h-index	3579 181 g-index
377 all docs	377 docs citations	377 times ranked	13290 citing authors

#	Article	IF	Citations
1	Change in REVEAL Lite 2 risk score predicts outcomes in patients with pulmonary arterial hypertension in the PATENT study. Journal of Heart and Lung Transplantation, 2022, 41, 411-420.	0.6	4
2	Long-Term Survival, Safety and Tolerability with Selexipag in Patients with Pulmonary Arterial Hypertension: Results from GRIPHON and its Open-Label Extension. Advances in Therapy, 2022, 39, 796-810.	2.9	12
3	External validation of a refined four-stratum risk assessment score from the French pulmonary hypertension registry. European Respiratory Journal, 2022, 59, 2102419.	6.7	83
4	The impact of comorbidities on selexipag treatment effect in patients with pulmonary arterial hypertension: insights from the <scp>GRIPHON</scp> study. European Journal of Heart Failure, 2022, 24, 205-214.	7.1	22
5	Using the Plasma Proteome for Risk Stratifying Patients with Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2022, 205, 1102-1111.	5.6	35
6	Respiratory symptoms and radiological findings in post-acute COVID-19 syndrome. ERJ Open Research, 2022, 8, 00479-2021.	2.6	16
7	Sequential combination therapy with parenteral prostacyclin in BMPR2 mutations carriers. Pulmonary Circulation, 2022, 12, e12023.	1.7	2
8	ERS statement on chronic thromboembolic pulmonary hypertension. Pulmonologiya, 2022, 32, 13-52.	0.8	0
9	Lung Ventilation/Perfusion Scintigraphy for the Screening of Chronic Thromboembolic Pulmonary Hypertension (CTEPH): Which Criteria to Use?. Frontiers in Medicine, 2022, 9, 851935.	2.6	4
10	Aggressive Afterload Lowering to Improve the Right Ventricle: A New Target for Medical Therapy in Pulmonary Arterial Hypertension?. American Journal of Respiratory and Critical Care Medicine, 2022, 205, 751-760.	5.6	27
11	Mining the Plasma Proteome for Insights into the Molecular Pathology of Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2022, 205, 1449-1460.	5.6	19
12	COVID-19 in Patients with Pulmonary Hypertension: A National Prospective Cohort Study. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 573-583.	5.6	16
13	ERS International Congress 2021: highlights from the Pulmonary Vascular Diseases Assembly. ERJ Open Research, 2022, 8, 00665-2021.	2.6	2
14	An emerging phenotype of pulmonary arterial hypertension patients carryingSOX17variants. European Respiratory Journal, 2022, 60, 2200656.	6.7	15
15	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.	0.6	6
16	To be or not to beâ€¦ treated with initial combination therapy, that is the (PAH) question. European Respiratory Journal, 2022, 59, 2200390.	6.7	2
17	Outcomes of cirrhotic patients with pre-capillary pulmonary hypertension and pulmonary vascular resistance between 2 and 3 Wood Units. European Respiratory Journal, 2022, 60, 2200107.	6.7	5
18	Long-Term Safety, Tolerability and Survival in Patients with Pulmonary Arterial Hypertension Treated with Macitentan: Results from the SERAPHIN Open-Label Extension. Advances in Therapy, 2022, 39, 4374-4390.	2.9	2

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19	Current and future treatments of pulmonary arterial hypertension. British Journal of Pharmacology, 2021, 178, 6-30.	5.4	104
20	Portopulmonary hypertension: An unfolding story. Clinics and Research in Hepatology and Gastroenterology, 2021, 45, 101492.	1.5	3
21	Chronic thromboembolic pulmonary hypertension and totally implantable central venous access systems. European Respiratory Journal, 2021, 57, 2002208.	6.7	12
22	Characteristics and Long-term Outcomes of Pulmonary Venoocclusive Disease Induced by Mitomycin C. Chest, 2021, 159, 1197-1207.	0.8	14
23	Riociguat: Clinical research and evolving role in therapy. British Journal of Clinical Pharmacology, 2021, 87, 2645-2662.	2.4	18
24	Riociguat treatment in patients with chronic thromboembolic pulmonary hypertension: Final safety data from the EXPERT registry. Respiratory Medicine, 2021, 178, 106220.	2.9	23
25	Reply to: â€œManagement of portopulmonary hypertension: What is more important, PAH severity or liver disease severity?â€• Journal of Hepatology, 2021, 74, 238-239.	3.7	0
26	Pulmonary Vascular Resistance in Pulmonary Arterial Hypertension: La PiÃ¨ce de RÃ©sistance?. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 524-525.	5.6	1
27	Reversible pulmonary hypertension associated with multivisceral Whipple's disease. European Respiratory Journal, 2021, 57, 2003132.	6.7	3
28	Initial triple therapy in pulmonary arterial hypertension: coming of age and rejuvenated. European Respiratory Journal, 2021, 57, 2004258.	6.7	0
29	Riociguat treatment in patients with pulmonary arterial hypertension: Final safety data from the EXPERT registry. Respiratory Medicine, 2021, 177, 106241.	2.9	13
30	The isobaric pulmonary arterial compliance in pulmonary hypertension. ERJ Open Research, 2021, 7, 00941-2020.	2.6	5
31	Hypoxemia during sleep and overnight rostral fluid shift in pulmonary arterial hypertension: a pilot study. Pulmonary Circulation, 2021, 11, 1-9.	1.7	5
32	Prevalence of pulmonary embolism in patients with COVID-19 at the time of hospital admission. European Respiratory Journal, 2021, 58, 2100116.	6.7	41
33	Outcomes of patients with decreased arterial oxyhaemoglobin saturation on pulmonary arterial hypertension drugs. European Respiratory Journal, 2021, 58, 2004066.	6.7	14
34	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.	5.6	94
35	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.	6.7	7
36	Pulmonary Hypertension in Patients with Common Variable Immunodeficiency. Journal of Clinical Immunology, 2021, 41, 1549-1562.	3.8	3

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37	French recommendations for the management of systemic sclerosis. Orphanet Journal of Rare Diseases, 2021, 16, 322.	2.7	37
38	Relationship Between Time From Diagnosis and Morbidity/Mortality in Pulmonary Arterial Hypertension. Chest, 2021, 160, 277-286.	0.8	21
39	Association between Leflunomide and Pulmonary Hypertension. Annals of the American Thoracic Society, 2021, 18, 1306-1315.	3.2	8
40	Severe pulmonary hypertension associated with chronic obstructive pulmonary disease: A prospective French multicenter cohort. Journal of Heart and Lung Transplantation, 2021, 40, 1009-1018.	0.6	24
41	Pulmonary hypertension associated with busulfan. Pulmonary Circulation, 2021, 11, 1-12.	1.7	3
42	Three- Versus Two-Drug Therapy for Patients With Newly Diagnosed Pulmonary ArterialÂHypertension. Journal of the American College of Cardiology, 2021, 78, 1393-1403.	2.8	90
43	Serum and pulmonary uric acid in pulmonary arterial hypertension. European Respiratory Journal, 2021, 58, 2000332.	6.7	28
44	ERS statement on chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2021, 57, 2002828.	6.7	287
45	Screening for pulmonary arterial hypertension in adults carrying a BMPR2 mutation. European Respiratory Journal, 2021, 58, 2004229.	6.7	50
46	TORREY, a Phase 2 study to evaluate the efficacy and safety of inhaled seralutinib for the treatment of pulmonary arterial hypertension. Pulmonary Circulation, 2021, 11, 1-7.	1.7	24
47	Reply to: Jin et al. and Sun et al.. American Journal of Respiratory and Critical Care Medicine, 2021, , .	5.6	0
48	Sex and gender in pulmonary arterial hypertension. European Respiratory Review, 2021, 30, 200330.	7.1	31
49	Results of an Expert Consensus Survey on the Treatment of Pulmonary Arterial Hypertension With Oral Prostacyclin Pathway Agents. Chest, 2020, 157, 955-965.	0.8	26
50	Intensity and quality of exertional dyspnoea in patients with stable pulmonary hypertension. European Respiratory Journal, 2020, 55, 1802108.	6.7	24
51	Survival Improved in Patients AgedÂâ‰ 70 Years With Systemic Sclerosis-Associated Pulmonary Arterial Hypertension During the Period 2006 to 2017 in France. Chest, 2020, 157, 945-954.	0.8	13
52	Novel composite clinical endpoints and risk scores used in clinical trials in pulmonary arterial hypertension. Pulmonary Circulation, 2020, 10, 1-11.	1.7	12
53	Risks and outcomes of gastrointestinal endoscopy with anaesthesia in patients with pulmonary hypertension. British Journal of Anaesthesia, 2020, 125, e466-e468.	3.4	5
54	The dangerous and contradictory prognostic significance of PVR<3WU when TAPSE<16mm in postcapillary pulmonary hypertension. ESC Heart Failure, 2020, 7, 2398-2405.	3.1	4

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55	Characteristics and outcomes of asthmatic patients with COVID-19 pneumonia who require hospitalisation. European Respiratory Journal, 2020, 56, 2001875.	6.7	90
56	Exercise Hemodynamics in the Prognosis of Patients with Pulmonary Arterial Hypertension. Respiration, 2020, 99, 678-685.	2.6	2
57	Gas Exchange and Ventilatory Efficiency During Exercise in Pulmonary Vascular Diseases. Archivos De Bronconeumologia, 2020, 56, 578-585.	0.8	5
58	Initial combination therapy of macitentan and tadalafil in pulmonary arterial hypertension. European Respiratory Journal, 2020, 56, 2000673.	6.7	35
59	Pulmonary Hypertension Complicating Pulmonary Artery Involvement in Pseudoxanthoma Elasticum. American Journal of Respiratory and Critical Care Medicine, 2020, 202, e90-e91.	5.6	1
60	Severe Pulmonary Hypertension Management Across Europe (PHAROS): an ERS Clinical Research Collaboration. European Respiratory Journal, 2020, 55, 2001047.	6.7	3
61	Phenotype and Outcomes of Pulmonary Hypertension Associated with Neurofibromatosis Type 1. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 843-852.	5.6	12
62	Pulmonary complications of Bcr-Abl tyrosine kinase inhibitors. European Respiratory Journal, 2020, 56, 2000279.	6.7	28
63	Transition from intravenous epoprostenol to selexipag in pulmonary arterial hypertension: a word of caution. European Respiratory Journal, 2020, 55, 1902418.	6.7	17
64	Macitentan Improves Risk Categorization for Liver Transplant Mortality in Patients With Portopulmonary Hypertension: A PORTICO Study Post Hoc Analysis. Liver Transplantation, 2020, 26, 935-940.	2.4	17
65	Phenotype and outcome of pulmonary arterial hypertension patients carrying a TBX4 mutation. European Respiratory Journal, 2020, 55, 1902340.	6.7	40
66	Portopulmonary hypertension in the current era of pulmonary hypertension management. Journal of Hepatology, 2020, 73, 130-139.	3.7	78
67	Risk assessment in pulmonary arterial hypertension: Insights from the GRIPHON study. Journal of Heart and Lung Transplantation, 2020, 39, 300-309.	0.6	39
68	Hereditary hemorrhagic telangiectasia and liver involvement. Clinics and Research in Hepatology and Gastroenterology, 2020, 44, 426-432.	1.5	5
69	Pulmonary Hypertension Associated with Chronic Lung Diseases: Treatment Considerations. Respiratory Medicine, 2020, , 79-96.	0.1	1
70	Gas Exchange and Ventilatory Efficiency During Exercise in Pulmonary Vascular Diseases. Archivos De Bronconeumologia, 2020, 56, 578-585.	0.8	10
71	Evaluation of a collaborative care program for pulmonary hypertension patients: a multicenter randomized trial. International Journal of Clinical Pharmacy, 2020, 42, 1128-1138.	2.1	4
72	Macitentan for the treatment of portopulmonary hypertension (PORTICO): a multicentre, randomised, double-blind, placebo-controlled, phase 4 trial. Lancet Respiratory Medicine,the, 2019, 7, 594-604.	10.7	119

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73	Golden Ratio and the Proportionality Between Pulmonary Pressure Components in Pulmonary Arterial Hypertension. Chest, 2019, 155, 991-998.	0.8	13
74	Integrating Data From Randomized Controlled Trials and Observational Studies to Assess Survival in Rare Diseases. Circulation: Cardiovascular Quality and Outcomes, 2019, 12, e005095.	2.2	8
75	Predictors of survival in patients with not-operated chronic thromboembolic pulmonary hypertension. Journal of Heart and Lung Transplantation, 2019, 38, 833-842.	0.6	57
76	French experience of balloon pulmonary angioplasty for chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2019, 53, 1802095.	6.7	173
77	Highlights from the ERS International Congress 2018: Assembly 13 â€“ Pulmonary Vascular Diseases. ERJ Open Research, 2019, 5, 00202-2018.	2.6	0
78	Assembly 13: placing the pulmonary circulation in the heart of ERS. Breathe, 2019, 15, 88-89.	1.3	1
79	Association of N-Terminal Pro Brain Natriuretic Peptide and Long-Term Outcome in Patients With Pulmonary Arterial Hypertension. Circulation, 2019, 139, 2440-2450.	1.6	67
80	Understanding the Similarities and Differences between Hepatic and Pulmonary Veno-Occlusive Disease. American Journal of Pathology, 2019, 189, 1159-1175.	3.8	19
81	Pulmonary arterial hypertension registries: past, present and into the future. European Respiratory Review, 2019, 28, 190128.	7.1	8
82	Management of pulmonary arterial hypertension in patients aged over 65 years. European Heart Journal Supplements, 2019, 21, K29-K36.	0.1	9
83	Clinical phenotypes and outcomes of precapillary pulmonary hypertension of sickle cell disease. European Respiratory Journal, 2019, 54, 1900585.	6.7	15
84	Genetic determinants of risk in pulmonary arterial hypertension: international genome-wide association studies and meta-analysis. Lancet Respiratory Medicine,the, 2019, 7, 227-238.	10.7	122
85	Selexipag treatment for pulmonary arterial hypertension associated with congenital heart disease after defect correction: insights from the randomised controlled GRIPHON study. European Journal of Heart Failure, 2019, 21, 352-359.	7.1	40
86	Clinical trial design and new therapies for pulmonary arterial hypertension. European Respiratory Journal, 2019, 53, 1801908.	6.7	142
87	Association between Rheumatoid Arthritis and Pulmonary Hypertension: Data from the French Pulmonary Hypertension Registry. Respiration, 2018, 95, 244-250.	2.6	17
88	Outcome of Portopulmonary Hypertension After Liver Transplantation. Transplantation, 2018, 102, e190-e191.	1.0	1
89	Pulmonary Arterial Hypertension-Related Morbidity Is Prognostic for Mortality. Journal of the American College of Cardiology, 2018, 71, 752-763.	2.8	82
90	RV Fractional Area Change and TAPSE as Predictors of Severe Right Ventricular Dysfunction in Pulmonary Hypertension: A CMR Study. Lung, 2018, 196, 157-164.	3.3	42

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91	The Low-Risk Profile in Pulmonary Arterial Hypertension. Time for a Paradigm Shift to Goal-oriented Clinical Trial Endpoints?. American Journal of Respiratory and Critical Care Medicine, 2018, 197, 860-868.	5.6	45
92	Targeting the Prostacyclin Pathway with Selexipag in Patients with Pulmonary Arterial Hypertension Receiving Double Combination Therapy: Insights from the Randomized Controlled GRIPHON Study. American Journal of Cardiovascular Drugs, 2018, 18, 37-47.	2.2	69
93	Natural History over 8 Years of Pulmonary Vascular Disease in a Patient Carrying Biallelic EIF2AK4 Mutations. American Journal of Respiratory and Critical Care Medicine, 2018, 198, 537-541.	5.6	12
94	Risk assessment in pulmonary arterial hypertension. European Respiratory Journal, 2018, 51, 1800279.	6.7	26
95	Validation of a risk assessment instrument for pulmonary arterial hypertension. European Heart Journal, 2018, 39, 4182-4185.	2.2	16
96	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.	0.6	50
97	Temporary treatment interruptions with oral selexipag in pulmonary arterial hypertension: Insights from the Prostacyclin (PGI 2) Receptor Agonist in Pulmonary Arterial Hypertension (GRIPHON) study. Journal of Heart and Lung Transplantation, 2018, 37, 401-408.	0.6	15
98	Prognostic Value of Follow-Up Hemodynamic Variables After Initial Management in Pulmonary Arterial Hypertension. Circulation, 2018, 137, 693-704.	1.6	155
99	Pulmonary Arterial Hypertension Associated With Systemic Lupus Erythematosus. Chest, 2018, 153, 143-151.	0.8	68
100	Chronic blood exchange transfusions in the management of pre-capillary pulmonary hypertension complicating sickle cell disease. European Respiratory Journal, 2018, 52, 1800272.	6.7	21
101	Haemodynamics and serial risk assessment in systemic sclerosis associated pulmonary arterial hypertension. European Respiratory Journal, 2018, 52, 1800678.	6.7	60
102	Pulmonary hypertension associated with neurofibromatosis type 1. European Respiratory Review, 2018, 27, 180053.	7.1	25
103	Clinical phenotypes and survival of pre-capillary pulmonary hypertension in systemic sclerosis. PLoS ONE, 2018, 13, e0197112.	2.5	47
104	Factors predicting outcome after pulmonary endarterectomy. PLoS ONE, 2018, 13, e0198198.	2.5	29
105	Association Between BMI and Obesity With Survival in Pulmonary Arterial Hypertension. Chest, 2018, 154, 872-881.	0.8	43
106	Clinical and Hemodynamic Correlates of Pulmonary Arterial Stiffness in Incident, Untreated Patients With Idiopathic Pulmonary Arterial Hypertension. Chest, 2018, 154, 882-892.	0.8	10
107	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.	0.6	20
108	Age, risk and outcomes in idiopathic pulmonary arterial hypertension. European Respiratory Journal, 2018, 51, 1800629.	6.7	9

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109	Pharmacovigilance in a rare disease: example of the VIGIAPATH program in pulmonary arterial hypertension. International Journal of Clinical Pharmacy, 2018, 40, 790-794.	2.1	5
110	Risk stratification in pulmonary arterial hypertension. Current Opinion in Pulmonary Medicine, 2018, 24, 407-415.	2.6	18
111	Late Breaking Abstract - Efficacy and safety of macitentan in portopulmonary hypertension: the PORTICO trial. , 2018, , .		1
112	Risk assessment in pulmonary arterial hypertension (PAH): Insights from the GRIPHON study. , 2018, , .		1
113	Impact of initial treatment strategy on long-term survival in pulmonary arterial hypertension (PAH). , 2018, , .		2
114	Initial dual oral combination therapy in inoperable chronic thromboembolic pulmonary hypertension (CTEPH). , 2018, , .		3
115	Reappraising the effects of pulmonary artery wedge pressure on right ventricular pulsatile loading. , 2018, , .		1
116	Association between six-minute walk distance and long-term outcomes in patients with pulmonary arterial hypertension: Data from the randomized SERAPHIN trial. PLoS ONE, 2018, 13, e0193226.	2.5	33
117	Balloon pulmonary angioplasty (BPA) for inoperable chronic thromboembolic pulmonary hypertension (CTEPH). , 2018, , .		0
118	Identifying chronic thromboembolic pulmonary hypertension (CTEPH) through French national hospital database (PMSI). , 2018, , .		0
119	Survival in portopulmonary hypertension (PoPH) in the era of modern PAH-targeted therapy. , 2018, , .		0
120	Factors associated with survival in patients with not-operated chronic thromboembolic pulmonary hypertension (CTEPH) in the modern management era. , 2018, , .		0
121	Efficacy and safety of tadalafil in portopulmonary hypertension. , 2018, , .		0
122	Ambrisentan use for pulmonary arterial hypertension in a post-authorization drug registry: The VOLibris Tracking Study. Journal of Heart and Lung Transplantation, 2017, 36, 399-406.	0.6	13
123	Clinical phenotypes and outcomes of heritable and sporadic pulmonary veno-occlusive disease: a population-based study. Lancet Respiratory Medicine,the, 2017, 5, 125-134.	10.7	123
124	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.	1.6	37
125	Epoprostenol and pulmonary arterial hypertension: 20â€…years of clinical experience. European Respiratory Review, 2017, 26, 160055.	7.1	70
126	Gutâ€“Lung Connection in Pulmonary Arterial Hypertension. American Journal of Respiratory Cell and Molecular Biology, 2017, 56, 402-405.	2.9	34

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127	Pulmonary hypertension due to left heart disease. Archives of Cardiovascular Diseases, 2017, 110, 420-431.	1.6	10
128	Plasma proteome analysis in patients with pulmonary arterial hypertension: an observational cohort study. Lancet Respiratory Medicine,the, 2017, 5, 717-726.	10.7	99
129	Impact of High-Priority Allocation on Lung and Heart-Lung Transplantation for Pulmonary Hypertension. Annals of Thoracic Surgery, 2017, 104, 404-411.	1.3	29
130	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.	0.6	37
131	Longâ€term outcome in liver transplantation candidates with portopulmonary hypertension. Hepatology, 2017, 65, 1683-1692.	7.3	68
132	Medical Treatment of Pulmonary Arterial Hypertension. Seminars in Respiratory and Critical Care Medicine, 2017, 38, 686-700.	2.1	9
133	Portopulmonary Hypertension. Seminars in Respiratory and Critical Care Medicine, 2017, 38, 651-661.	2.1	30
134	Management and long-term outcomes of sarcoidosis-associated pulmonary hypertension. European Respiratory Journal, 2017, 50, 1700465.	6.7	111
135	Selexipag for the treatment of connective tissue disease-associated pulmonary arterial hypertension. European Respiratory Journal, 2017, 50, 1602493.	6.7	97
136	Exertional dyspnoea in pulmonary arterial hypertension. European Respiratory Review, 2017, 26, 170039.	7.1	25
137	Are indexed values better for defining exercise pulmonary hypertension?. European Respiratory Journal, 2017, 50, 1700240.	6.7	4
138	Long-term outcomes of dasatinib-induced pulmonary arterial hypertension: a population-based study. European Respiratory Journal, 2017, 50, 1700217.	6.7	89
139	Risk assessment, prognosis and guideline implementation in pulmonary arterial hypertension. European Respiratory Journal, 2017, 50, 1700889.	6.7	527
140	Acute decompensated pulmonary hypertension. European Respiratory Review, 2017, 26, 170092.	7.1	48
141	A Clinical and Echocardiographic Score to Identify Pulmonary Hypertension Due to HFpEF. Journal of Cardiac Failure, 2017, 23, 29-35.	1.7	25
142	Long-term outcomes of pulmonary arterial hypertension under specific drug therapy in Eisenmenger syndrome. Journal of Heart and Lung Transplantation, 2017, 36, 386-398.	0.6	15
143	Non-invasive diagnosis of pulmonary hypertension from lung Doppler signal: a proof of concept study. Journal of Clinical Monitoring and Computing, 2017, 31, 903-910.	1.6	3
144	Macitentan Improves Health-Related QualityÂof Life for Patients With Pulmonary Arterial Hypertension. Chest, 2017, 151, 106-118.	0.8	46

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145	SERAPHIN haemodynamic substudy: the effect of the dual endothelin receptor antagonist macitentan on haemodynamic parameters and NT-proBNP levels and their association with disease progression in patients with pulmonary arterial hypertension. European Heart Journal, 2017, 38, 1147-1155.	2.2	65
146	Bacterial translocation in pulmonary hypertension. , 2017, , .		0
147	Pulmonary arterial hypertension-related morbidity is prognostic for survival: insights from the SERAPHIN and GRIPHON studies. , 2017, , .		0
148	Outcomes of patients with precapillary pulmonary hypertension admitted to an intensive care unit for acute right heart failure. , 2017, , .		0
149	Using controlled and real-world data in concert to assess survival benefits in pulmonary arterial hypertension: Insights from SERAPHIN and REVEAL. , 2017, , .		0
150	Is there an obesity paradox in pulmonary arterial hypertension?. , 2017, , .		0
151	Risk categories from European guidelines applied to the French Pulmonary Hypertension (PH) registry. , 2017, , .		0
152	Prognostic value of hemodynamics following treatment initiation in pulmonary arterial hypertension. , 2017, , .		0
153	Initial combination of macitentan and tadalafil in newly diagnosed patients with pulmonary arterial hypertension (PAH): results from the OPTIMA study. , 2017, , .		1
154	Kinetics of Cardiac Output at the Onset of Exercise in Precapillary Pulmonary Hypertension. BioMed Research International, 2016, 2016, 1-8.	1.9	7
155	Direct-Acting Antiviral Medications for Hepatitis C Virus Infection and Pulmonary Arterial Hypertension. Chest, 2016, 150, 256-258.	0.8	12
156	Response to Letter Regarding Article, â€œMitomycin-Induced Pulmonary Veno-Occlusive Disease: Evidence From Human Disease and Animal Modelâ€• Circulation, 2016, 133, e592-3.	1.6	4
157	Pulmonary arterial hypertension in idiopathic inflammatory myopathies. Medicine (United States), 2016, 95, e4911.	1.0	40
158	International Liver Transplant Society Practice Guidelines. Transplantation, 2016, 100, 1440-1452.	1.0	309
159	BMPR2mutation status influences bronchial vascular changes in pulmonary arterial hypertension. European Respiratory Journal, 2016, 48, 1668-1681.	6.7	68
160	Beyond a single pathway: combination therapy in pulmonary arterial hypertension. European Respiratory Review, 2016, 25, 408-417.	7.1	53
161	Deterioration of pulmonary hypertension and pleural effusion with bosutinib following dasatinib lung toxicity. European Respiratory Journal, 2016, 48, 1517-1519.	6.7	44
162	Interferon-induced pulmonary hypertension. Current Opinion in Pulmonary Medicine, 2016, 22, 415-420.	2.6	28

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163	Pulmonary veno-occlusive disease. European Respiratory Journal, 2016, 47, 1518-1534.	6.7	289
164	Initial dual oral combination therapy in pulmonary arterial hypertension. European Respiratory Journal, 2016, 47, 1727-1736.	6.7	124
165	Lung capillary blood volume and membrane diffusion in precapillary pulmonary hypertension. Journal of Heart and Lung Transplantation, 2016, 35, 647-656.	0.6	10
166	Diagnostic concordance of different criteria for exercise pulmonary hypertension in subjects with normal resting pulmonary artery pressure. European Respiratory Journal, 2016, 48, 254-257.	6.7	31
167	BMPR2 mutations and survival in pulmonary arterial hypertension: an individual participant data meta-analysis. Lancet Respiratory Medicine,the, 2016, 4, 129-137.	10.7	307
168	Resting pulmonary artery pressure of 21â€“24â€…mmHg predicts abnormal exercise haemodynamics. European Respiratory Journal, 2016, 47, 1436-1444.	6.7	44
169	Loss of Vascular Distensibility During Exercise Is an Early Hemodynamic Marker of Pulmonary Vascular Disease. Chest, 2016, 149, 353-361.	0.8	55
170	Regulatory T Cell Dysfunction in Idiopathic, Heritable and Connective Tissue-Associated Pulmonary Arterial Hypertension. Chest, 2016, 149, 1482-1493.	0.8	63
171	A rare case of sarcoidosis-associated pulmonary hypertension in a patient exposed to silica. European Respiratory Review, 2016, 25, 93-96.	7.1	7
172	Patientsâ€™, relativesâ€™, and practitionersâ€™ views of pulmonary arterial hypertension: A qualitative study. Presse Medicale, 2016, 45, e11-e27.	1.9	18
173	Comparative Safety and Tolerability of Prostacyclins in Pulmonary Hypertension. Drug Safety, 2016, 39, 287-294.	3.2	35
174	Genetic counselling in a national referral centre for pulmonary hypertension. European Respiratory Journal, 2016, 47, 541-552.	6.7	87
175	A prospective study of the 6â€…min walk test as a surrogate marker for haemodynamics in two independent cohorts of treatment-naÃ¯ve systemic sclerosis-associated pulmonary arterial hypertension. Annals of the Rheumatic Diseases, 2016, 75, 1457-1465.	0.9	16
176	Dasatinib induces lung vascular toxicity and predisposes to pulmonary hypertension. Journal of Clinical Investigation, 2016, 126, 3207-3218.	8.2	208
177	Management of selexipag interruptions in the GRIPHON study. , 2016, , .		1
178	Clinical phenotypes and outcomes of pulmonary veno-occlusive disease in carriers of bi-allelicEIF2AK4mutations. , 2016, , .		0
179	Illicit drug use and pulmonary arterial hypertension: Not so frequent. , 2016, , .		0
180	Pulmonary arterial hypertension in patients treated with interferon. European Respiratory Journal, 2015, 46, 1851-1853.	6.7	35

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181	Individualized Dosing of Selexipag Based on Tolerability in the GRIPHON Study Shows Consistent Efficacy and Safety in Patients With Pulmonary Arterial Hypertension (PAH). Chest, 2015, 148, 961A.	0.8	3
182	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.	0.8	33
183	Incident and prevalent cohorts with pulmonary arterial hypertension: insight from SERAPHIN. European Respiratory Journal, 2015, 46, 1711-1720.	6.7	39
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