GÃ©rald Simonneau

List of Publications by Year
in descending order

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336	85,082	733 120	340 285
papers	citations	h-index	g-index

350	350	350	26159
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	2015 ESC/ERS Guidelines for the diagnosis and treatment of pulmonary hypertension. European Heart Journal, 2016, 37, 67-119.	1.0	5,074
2	Guidelines for the diagnosis and treatment of pulmonary hypertension: The Task Force for the Diagnosis and Treatment of Pulmonary Hypertension of the European Society of Cardiology (ESC) and the European Respiratory Society (ERS), endorsed by the International Society of Heart and Lung Transplantation (ISHLT). European Heart Journal, 2009, 30, 2493-2537.	1.0	3,108
3	Updated Clinical Classification of Pulmonary Hypertension. Journal of the American College of Cardiology, 2013, 62, D34-D41.	1.2	2,865
4	Haemodynamic definitions and updated clinical classification of pulmonary hypertension. European Respiratory Journal, 2019, 53, 1801913.	3.1	2,583
5	Bosentan Therapy for Pulmonary Arterial Hypertension. New England Journal of Medicine, 2002, 346, 896-903.	13.9	2,545
6	2015 ESC/ERS Guidelines for the diagnosis and treatment of pulmonary hypertension. European Respiratory Journal, 2015, 46, 903-975.	3.1	2,415
7	Sildenafil Citrate Therapy for Pulmonary Arterial Hypertension. New England Journal of Medicine, 2005, 353, 2148-2157.	13.9	2,237
8	Updated Clinical Classification of Pulmonary Hypertension. Journal of the American College of Cardiology, 2009, 54, S43-S54.	1.2	1,919
9	Pulmonary Arterial Hypertension in France. American Journal of Respiratory and Critical Care Medicine, 2006, 173, 1023-1030.	2.5	1,736
10	A Clinical Trial of Vena Caval Filters in the Prevention of Pulmonary Embolism in Patients with Proximal Deep-Vein Thrombosis. New England Journal of Medicine, 1998, 338, 409-416.	13.9	1,676
11	Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2004, 351, 1425-1436.	13.9	1,627
12	Inhaled Iloprost for Severe Pulmonary Hypertension. New England Journal of Medicine, 2002, 347, 322-329.	13.9	1,626
13	Clinical classification of pulmonary hypertension. Journal of the American College of Cardiology, 2004, 43, S5-S12.	1.2	1,542
14	Effects of the dual endothelin-receptor antagonist bosentan in patients with pulmonary hypertension: a randomised placebocontrolled study. Lancet, The, 2001, 358, 1119-1123.	6.3	1,421
15	Long-term intravenous epoprostenol infusion in primary pulmonary hypertension. Journal of the American College of Cardiology, 2002, 40, 780-788.	1.2	1,290
16	Continuous Subcutaneous Infusion of Treprostinil, a Prostacyclin Analogue, in Patients with Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2002, 165, 800-804.	2.5	1,288
17	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
18	Macitentan and Morbidity and Mortality in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2013, 369, 809-818.	13.9	1,168

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19	Riociguat for the Treatment of Chronic Thromboembolic Pulmonary Hypertension. New England Journal of Medicine, 2013, 369, 319-329.	13.9	1,144
20	Appetite-Suppressant Drugs and the Risk of Primary Pulmonary Hypertension. New England Journal of Medicine, 1996, 335, 609-616.	13.9	1,127
21	Treatment of Venous Thrombosis with Intravenous Unfractionated Heparin Administered in the Hospital as Compared with Subcutaneous Low-Molecular-Weight Heparin Administered at Home. New England Journal of Medicine, 1996, 334, 682-687.	13.9	1,108
22	Long-Term Response to Calcium Channel Blockers in Idiopathic Pulmonary Arterial Hypertension. Circulation, 2005, 111, 3105-3111.	1.6	1,040
23	Tadalafil Therapy for Pulmonary Arterial Hypertension. Circulation, 2009, 119, 2894-2903.	1.6	956
24	New Formula for Predicting Mean Pulmonary Artery Pressure Using Systolic Pulmonary Artery Pressure. Chest, 2004, 126, 1313-1317.	0.4	923
25	Initial Use of Ambrisentan plus Tadalafil in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 834-844.	13.9	906
26	Guidelines on diagnosis and treatment of pulmonary arterial hypertension. The Task Force on Diagnosis and Treatment of Pulmonary Arterial Hypertension of the European Society of Cardiology. European Heart Journal, 2004, 25, 2243-2278.	1.0	903
27	Chronic Thromboembolic Pulmonary Hypertension (CTEPH). Circulation, 2011, 124, 1973-1981.	1.6	860
28	Chronic Thromboembolic Pulmonary Hypertension. Circulation, 2006, 113, 2011-2020.	1.6	791
29	Selexipag for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 2522-2533.	13.9	790
30	A Comparison of Low-Molecular-Weight Heparin with Unfractionated Heparin for Acute Pulmonary Embolism. New England Journal of Medicine, 1997, 337, 663-669.	13.9	767
31	Clinical and Molecular Genetic Features of Pulmonary Hypertension in Patients with Hereditary Hemorrhagic Telangiectasia. New England Journal of Medicine, 2001, 345, 325-334.	13.9	676
32	Surgical management and outcome of patients with chronic thromboembolic pulmonary hypertension: Results from an international prospective registry. Journal of Thoracic and Cardiovascular Surgery, 2011, 141, 702-710.	0.4	605
33	Medical Therapy For Pulmonary Arterial Hypertension. Chest, 2004, 126, 35S-62S.	0.4	592
34	Effects of beraprost sodium, an oral prostacyclin analogue, in patients with pulmonary arterial hypertension: a randomized, double-blind, placebo-controlled trial. Journal of the American College of Cardiology, 2002, 39, 1496-1502.	1.2	584
35	Addition of Sildenafil to Long-Term Intravenous Epoprostenol Therapy in Patients with Pulmonary Arterial Hypertension. Annals of Internal Medicine, 2008, 149, 521.	2.0	558
36	Pulmonary Arterial Hypertension in Patients Treated by Dasatinib. Circulation, 2012, 125, 2128-2137.	1.6	548

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37	Pulmonary Hypertension Due to Left Heart Diseases. Journal of the American College of Cardiology, 2013, 62, D100-D108.	1.2	541
38	Risk assessment, prognosis and guideline implementation in pulmonary arterial hypertension. European Respiratory Journal, 2017, 50, 1700889.	3.1	527
39	Pulmonary Hypertension in Chronic Lung Diseases. Journal of the American College of Cardiology, 2013, 62, D109-D116.	1.2	518
40	Bosentan for Treatment of Inoperable Chronic Thromboembolic Pulmonary Hypertension. Journal of the American College of Cardiology, 2008, 52, 2127-2134.	1.2	506
41	Long-Term Outcome of Patients With Chronic Thromboembolic Pulmonary Hypertension. Circulation, 2016, 133, 859-871.	1.6	506
42	Increased plasma serotonin in primary pulmonary hypertension. American Journal of Medicine, 1995, 99, 249-254.	0.6	500
43	Severe Pulmonary Hypertension during Pregnancy. Anesthesiology, 2005, 102, 1133-1137.	1.3	483
44	Imatinib Mesylate as Add-on Therapy for Pulmonary Arterial Hypertension. Circulation, 2013, 127, 1128-1138.	1.6	482
45	Medical Therapy for Pulmonary Arterial Hypertension. Chest, 2007, 131, 1917-1928.	0.4	477
46	Serotonin transporter overexpression is responsible for pulmonary artery smooth muscle hyperplasia in primary pulmonary hypertension. Journal of Clinical Investigation, 2001, 108, 1141-1150.	3.9	446
47	Ambrisentan Therapy for Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2005, 46, 529-535.	1.2	441
48	Endothelial-to-Mesenchymal Transition in Pulmonary Hypertension. Circulation, 2015, 131, 1006-1018.	1.6	441
49	A Hemodynamic Study of Pulmonary Hypertension in Sickle Cell Disease. New England Journal of Medicine, 2011, 365, 44-53.	13.9	432
50	Platelet-derived Growth Factor Expression and Function in Idiopathic Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2008, 178, 81-88.	2.5	405
51	Prevalence of HIV-related Pulmonary Arterial Hypertension in the Current Antiretroviral Therapy Era. American Journal of Respiratory and Critical Care Medicine, 2008, 177, 108-113.	2.5	402
52	Diagnosis and Classification of 17 Diseases from 1404 Subjects via Pattern Analysis of Exhaled Molecules. ACS Nano, 2017, 11, 112-125.	7.3	386
53	Diagnostic strategy for patients with suspected pulmonary embolism: a prospective multicentre outcome study. Lancet, The, 2002, 360, 1914-1920.	6.3	384
54	Mutations of the TGF-Î² type II receptorBMPR2 in pulmonary arterial hypertension. Human Mutation, 2006, 27, 121-132.	1.1	368

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55	EIF2AK4 mutations cause pulmonary veno-occlusive disease, a recessive form of pulmonary hypertension. Nature Genetics, 2014, 46, 65-69.	9.4	351
56	Effects of the oral endothelin-receptorantagonist bosentan on echocardiographicand doppler measures in patients with pulmonary arterial hypertension. Journal of the American College of Cardiology, 2003, 41, 1380-1386.	1.2	334
57	Immunosuppressive therapy in lupusâ€•and mixed connective tissue diseaseâ€“associated pulmonary arterial hypertension: A retrospective analysis of twentyâ€three cases. Arthritis and Rheumatism, 2008, 58, 521-531.	6.7	321
58	Upfront triple combination therapy in pulmonary arterial hypertension: a pilot study. European Respiratory Journal, 2014, 43, 1691-1697.	3.1	319
59	Immunosuppressive Therapy in Connective Tissue Diseases-Associated Pulmonary Arterial Hypertension. Chest, 2006, 130, 182-189.	0.4	316
60	Dysregulated Reninâ€“Angiotensinâ€“Aldosterone System Contributes to Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2012, 186, 780-789.	2.5	309
61	BMPR2 mutations and survival in pulmonary arterial hypertension: an individual participant data meta-analysis. Lancet Respiratory Medicine,the, 2016, 4, 129-137.	5.2	307
62	The pathophysiology of chronic thromboembolic pulmonary hypertension. European Respiratory Review, 2017, 26, 160112.	3.0	307
63	Prognostic Factors for Survival in Human Immunodeficiency Virusâ€“associated Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2003, 167, 1433-1439.	2.5	295
64	Pulmonary Veno-Occlusive Disease. Medicine (United States), 2008, 87, 220-233.	0.4	295
65	Fibrous remodeling of the pulmonary venous system in pulmonary arterial hypertension associated with connective tissue diseases. Human Pathology, 2007, 38, 893-902.	1.1	291
66	Screening for pulmonary arterial hypertension in patients with systemic sclerosis: Clinical characteristics at diagnosis and longâ€term survival. Arthritis and Rheumatism, 2011, 63, 3522-3530.	6.7	291
67	Pulmonary veno-occlusive disease. European Respiratory Journal, 2016, 47, 1518-1534.	3.1	289
68	ERS statement on chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2021, 57, 2002828.	3.1	287
69	Advances in Therapeutic Interventions for Patients With Pulmonary Arterial Hypertension. Circulation, 2014, 130, 2189-2208.	1.6	278
70	Deleterious Effects of Î²-Blockers on Exercise Capacity and Hemodynamics in Patients With Portopulmonary Hypertension. Gastroenterology, 2006, 130, 120-126.	0.6	277
71	Long-term outcome with first-line bosentan therapy in idiopathic pulmonary arterial hypertension. European Heart Journal, 2006, 27, 589-595.	1.0	272
72	Effects of the Dual Endothelin Receptor Antagonist Bosentan in Patients With Pulmonary Arterial Hypertension. Chest, 2003, 124, 247-254.	0.4	271

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73	Prevalence of chronic thromboembolic pulmonary hypertension after acute pulmonary embolism. Thrombosis and Haemostasis, 2014, 112, 598-605.	1.8	271
74	Clinical Outcomes of Pulmonary Arterial Hypertension in Carriers of BMPR2 Mutation. American Journal of Respiratory and Critical Care Medicine, 2008, 177, 1377-1383.	2.5	269
75	Selexipag: an oral, selective prostacyclin receptor agonist for the treatment of pulmonary arterial hypertension. European Respiratory Journal, 2012, 40, 874-880.	3.1	267
76	Clinical Outcomes of Pulmonary Arterial Hypertension in Patients Carrying an ACVRL1 (ALK1) Mutation. American Journal of Respiratory and Critical Care Medicine, 2010, 181, 851-861.	2.5	259
77	Cross Talk Between Endothelial and Smooth Muscle Cells in Pulmonary Hypertension. Circulation, 2006, 113, 1857-1864.	1.6	257
78	Inhibition of Hypoxic Pulmonary Vasoconstriction by Nifedipine. New England Journal of Medicine, 1981, 304, 1582-1585.	13.9	256
79	CX3C Chemokine Fractalkine in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2002, 165, 1419-1425.	2.5	247
80	Riociguat for the treatment of chronic thromboembolic pulmonary hypertension: a long-term extension study (CHEST-2). European Respiratory Journal, 2015, 45, 1293-1302.	3.1	247
81	Chemokine RANTES in Severe Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2002, 165, 534-539.	2.5	239
82	Bosentan for the Treatment of Human Immunodeficiency Virusâ€“associated Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2004, 170, 1212-1217.	2.5	238
83	Long-term Treatment With Sildenafil Citrate in Pulmonary Arterial Hypertension. Chest, 2011, 140, 1274-1283.	0.4	237
84	Pulmonary Hypertension: CT of the Chest in Pulmonary Venoocclusive Disease. American Journal of Roentgenology, 2004, 183, 65-70.	1.0	234
85	Treprostinil, a Prostacyclin Analogue, in Pulmonary Arterial Hypertension Associated With Connective Tissue Disease. Chest, 2004, 126, 420-427.	0.4	232
86	Pulmonary arterial hypertension. Orphanet Journal of Rare Diseases, 2013, 8, 97.	1.2	226
87	Portopulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2008, 178, 637-643.	2.5	220
88	Long-term response to calcium-channel blockers in non-idiopathic pulmonary arterial hypertension. European Heart Journal, 2010, 31, 1898-1907.	1.0	218
89	Dasatinib induces lung vascular toxicity and predisposes to pulmonary hypertension. Journal of Clinical Investigation, 2016, 126, 3207-3218.	3.9	208
90	Comparative analysis of clinical trials and evidence-based treatment algorithm in pulmonary arterial hypertension. Journal of the American College of Cardiology, 2004, 43, S81-S88.	1.2	206

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91	Stress Doppler Echocardiography in Relatives of Patients With Idiopathic and Familial Pulmonary Arterial Hypertension. Circulation, 2009, 119, 1747-1757.	1.6	205
92	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
93	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
94	Systematic Lung Scans Reveal a High Frequency of Silent Pulmonary Embolism in Patients With Proximal Deep Venous Thrombosis. Archives of Internal Medicine, 2000, 160, 159.	4.3	197
95	BMPR2 gene rearrangements account for a significant proportion of mutations in familial and idiopathic pulmonary arterial hypertension. Human Mutation, 2006, 27, 212-213.	1.1	196
96	Role of Endothelium-derived CC Chemokine Ligand 2 in Idiopathic Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2007, 176, 1041-1047.	2.5	196
97	Serotonin-Induced Smooth Muscle Hyperplasia in Various Forms of Human Pulmonary Hypertension. Circulation Research, 2004, 94, 1263-1270.	2.0	187
98	Endothelial-derived FGF2 contributes to the progression of pulmonary hypertension in humans and rodents. Journal of Clinical Investigation, 2009, 119, 512-523.	3.9	177
99	C-Kitâ€“Positive Cells Accumulate in Remodeled Vessels of Idiopathic Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2011, 184, 116-123.	2.5	176
100	French experience of balloon pulmonary angioplasty for chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2019, 53, 1802095.	3.1	173
101	Targeted therapies in pulmonary arterial hypertension. , 2014, 141, 172-191.		171
102	High Prevalence of Detectable Deep Venous Thrombosis in Patients With Acute Pulmonary Embolism. Chest, 1999, 116, 903-908.	0.4	163
103	Pulmonary Edema Complicating Continuous Intravenous Prostacyclin in Pulmonary Capillary Hemangiomatosis. American Journal of Respiratory and Critical Care Medicine, 1998, 157, 1681-1685.	2.5	161
104	Pulmonary Arterial Hypertension: A Rare Complication of Primary SjÃ¶gren Syndrome. Medicine (United) Tj ETQq0 0 0 rgBT /Overlock 10	0.4	160
105	Prognostic Value of Follow-Up Hemodynamic Variables After Initial Management in Pulmonary Arterial Hypertension. Circulation, 2018, 137, 693-704.	1.6	155
106	Primary pulmonary hypertension in a patient with a familial platelet storage pool disease: Role of serotonin. American Journal of Medicine, 1990, 89, 117-120.	0.6	154
107	PATHOBIOLOGY OF PULMONARY HYPERTENSION. Clinics in Chest Medicine, 2001, 22, 451-458.	0.8	153
108	Sildenafil for pulmonary arterial hypertension associated with connective tissue disease. Journal of Rheumatology, 2007, 34, 2417-22.	1.0	152

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109	Pulmonary artery pulse pressure and wave reflection in chronic pulmonary thromboembolism and primary pulmonary hypertension. Journal of the American College of Cardiology, 2001, 37, 1085-1092.	1.2	151
110	HIV-associated pulmonary arterial hypertension: survival and prognostic factors in the modern therapeutic era. Aids, 2010, 24, 67-75.	1.0	149
111	Chemotherapy-Induced Pulmonary Hypertension. American Journal of Pathology, 2015, 185, 356-371.	1.9	149
112	Surgical Treatments/Interventions for Pulmonary Arterial Hypertension. Chest, 2004, 126, 63S-71S.	0.4	144
113	Potassium Channel Subfamily K Member 3 (KCNK3) Contributes to the Development of Pulmonary Arterial Hypertension. Circulation, 2016, 133, 1371-1385.	1.6	141
114	Initial combination therapy with ambrisentan and tadalafil in connective tissue disease-associated pulmonary arterial hypertension (CTD-PAH): subgroup analysis from the AMBITION trial. Annals of the Rheumatic Diseases, 2017, 76, 1219-1227.	0.5	135
115	Long-term outcome of double-lung and heartâ€“lung transplantation for pulmonary hypertension: a comparative retrospective study of 219 patientsâ˜†. European Journal of Cardio-thoracic Surgery, 2010, 38, 277-284.	0.6	130
116	Predictors of long-term outcomes in patients treated with riociguat for chronic thromboembolic pulmonary hypertension: data from the CHEST-2 open-label, randomised, long-term extension trial. Lancet Respiratory Medicine,the, 2016, 4, 372-380.	5.2	130
117	New Treatments for Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2002, 165, 1209-1216.	2.5	129
118	Survival in systemic sclerosis-associated pulmonary arterial hypertension in the modern management era. Annals of the Rheumatic Diseases, 2013, 72, 1940-1946.	0.5	128
119	Intravenous Epoprostenol in Inoperable Chronic Thromboembolic Pulmonary Hypertension. Journal of Heart and Lung Transplantation, 2007, 26, 357-362.	0.3	126
120	Autocrine Fibroblast Growth Factor-2 Signaling Contributes to Altered Endothelial Phenotype in Pulmonary Hypertension. American Journal of Respiratory Cell and Molecular Biology, 2011, 45, 311-322.	1.4	125
121	Treatment of pulmonary arterial hypertension with targeted therapies. Nature Reviews Cardiology, 2011, 8, 526-538.	6.1	125
122	Tadalafil for the Treatment of Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2012, 60, 768-774.	1.2	124
123	Palliative Potts shunt for the treatment of children with drug-refractory pulmonary arterial hypertension: updated data from the first 24 patients. European Journal of Cardio-thoracic Surgery, 2015, 47, e105-e110.	0.6	124
124	Initial dual oral combination therapy in pulmonary arterial hypertension. European Respiratory Journal, 2016, 47, 1727-1736.	3.1	124
125	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
126	Nebivolol for Improving Endothelial Dysfunction, Pulmonary Vascular Remodeling, and Right Heart Function inÂPulmonary Hypertension. Journal of the American College of Cardiology, 2015, 65, 668-680.	1.2	119

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127	Is Pulmonary Arterial Hypertension Really a Late Complication of Systemic Sclerosis?. Chest, 2009, 136, 1211-1219.	0.4	117
128	RISK FACTORS FOR PULMONARY ARTERIAL HYPERTENSION. Clinics in Chest Medicine, 2001, 22, 459-475.	0.8	116
129	Potts Shunt in Children With Idiopathic Pulmonary Arterial Hypertension: Long-Term Results. Annals of Thoracic Surgery, 2012, 94, 817-824.	0.7	116
130	RESPITE: switching to riociguat in pulmonary arterial hypertension patients with inadequate response to phosphodiesterase-5 inhibitors. European Respiratory Journal, 2017, 50, 1602425.	3.1	113
131	Proinflammatory cytokine levels are linked to death in pulmonary arterial hypertension. European Respiratory Journal, 2014, 43, 915-917.	3.1	111
132	Systemic sclerosisâ€“related pulmonary hypertension associated with interstitial lung disease: Impact of pulmonary arterial hypertension therapies. Arthritis and Rheumatism, 2011, 63, 2456-2464.	6.7	109
133	Angiopoietin/Tie2 Pathway Influences Smooth Muscle Hyperplasia in Idiopathic Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2006, 174, 1025-1033.	2.5	106
134	Pulmonary Langerhans Cell Histiocytosis-Associated Pulmonary Hypertension. Chest, 2012, 142, 1150-1157.	0.4	104
135	Long-term safety and efficacy of imatinib in pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2015, 34, 1366-1375.	0.3	103
136	Mitomycin-Induced Pulmonary Veno-Occlusive Disease. Circulation, 2015, 132, 834-847.	1.6	103
137	Nitric Oxide Deficiency in Fenfluramine- and Dexfenfluramine-induced Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 1998, 158, 1061-1067.	2.5	102
138	Evaluation of Various Empirical Formulas for Estimating Mean Pulmonary Artery Pressure by Using Systolic Pulmonary Artery Pressure in Adults. Chest, 2009, 135, 760-768.	0.4	102
139	Vascular and right ventricular remodelling in chronic thromboembolic pulmonary hypertension. European Respiratory Journal, 2013, 41, 224-232.	3.1	100
140	Chronic thromboembolic pulmonary hypertension: role of medical therapy. European Respiratory Journal, 2013, 41, 985-990.	3.1	99
141	Plasma proteome analysis in patients with pulmonary arterial hypertension: an observational cohort study. Lancet Respiratory Medicine,the, 2017, 5, 717-726.	5.2	99
142	Primary Pulmonary Hypertension Associated With the Use of Fenfluramine Derivatives. Chest, 1998, 114, 195S-199S.	0.4	97
143	Selexipag for the treatment of connective tissue disease-associated pulmonary arterial hypertension. European Respiratory Journal, 2017, 50, 1602493.	3.1	97
144	Phosphodiesterase type 5 inhibitors in pulmonary arterial hypertension. Advances in Therapy, 2009, 26, 813-825.	1.3	96

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145	Factors associated with diagnosis and operability of chronic thromboembolic pulmonary hypertension. Thrombosis and Haemostasis, 2013, 110, 83-91.	1.8	96
146	Tadalafil monotherapy and as add-on to background bosentan in patients with pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2011, 30, 632-643.	0.3	95
147	Pulmonary Artery Pressureâ€“Flow Relations after Prostacyclin in Primary Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2002, 165, 338-340.	2.5	94
148	Pulmonary veno-occlusive disease: Recent progress and current challenges. Respiratory Medicine, 2010, 104, S23-S32.	1.3	94
149	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
150	Genome-wide association analysis identifies a susceptibility locus for pulmonary arterial hypertension. Nature Genetics, 2013, 45, 518-521.	9.4	93
151	Efficacy, safety and pharmacokinetics of bosentan in portopulmonary hypertension. European Respiratory Journal, 2013, 41, 96-103.	3.1	92
152	Long-term outcome of systemic sclerosis-associated pulmonary arterial hypertension treated with bosentan as first-line monotherapy followed or not by the addition of prostanoids or sildenafil. Rheumatology, 2010, 49, 490-500.	0.9	91
153	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
154	Three- Versus Two-Drug Therapy for Patients With Newly Diagnosed Pulmonary ArterialÂHypertension. Journal of the American College of Cardiology, 2021, 78, 1393-1403.	1.2	90
155	Long-term outcomes of dasatinib-induced pulmonary arterial hypertension: a population-based study. European Respiratory Journal, 2017, 50, 1700217.	3.1	89
156	Genetic counselling in a national referral centre for pulmonary hypertension. European Respiratory Journal, 2016, 47, 541-552.	3.1	87
157	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.	5.2	85
158	Serotonin Transporter Polymorphisms in Familial and Idiopathic Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2006, 173, 798-802.	2.5	83
159	Key Role of the Endothelial TGF-Î²/ALK1/Endoglin Signaling Pathway in Humans and Rodents Pulmonary Hypertension. PLoS ONE, 2014, 9, e100310.	1.1	83
160	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
161	The CX3C chemokine fractalkine in allergic asthma and rhinitis. Journal of Allergy and Clinical Immunology, 2003, 112, 1139-1146.	1.5	82
162	Controversies, Uncertainties and Future Research on the Treatment of Chronic Thromboembolic Pulmonary Hypertension. Proceedings of the American Thoracic Society, 2006, 3, 608-614.	3.5	82

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163	Validation of two predictive models for survival in pulmonary arterial hypertension. European Respiratory Journal, 2015, 46, 152-164.	3.1	82
164	Pulmonary Arterial Hypertension-Related Morbidity Is Prognostic for Mortality. Journal of the American College of Cardiology, 2018, 71, 752-763.	1.2	82
165	Absence of influence of gender and BMPR2 mutation type on clinical phenotypes of pulmonary arterial hypertension. Respiratory Research, 2010, 11, 73.	1.4	81
166	Drug-induced pulmonary arterial hypertension: a recent outbreak. European Respiratory Review, 2013, 22, 244-250.	3.0	81
167	Pulmonary hypertension in lymphangioleiomyomatosis: characteristics in 20 patients. European Respiratory Journal, 2012, 40, 630-640.	3.1	80
168	Pulmonary arterial hypertension in patients treated with interferon. European Respiratory Journal, 2014, 44, 1627-1634.	3.1	80
169	Occupational exposure to organic solvents: a risk factor for pulmonary veno-occlusive disease. European Respiratory Journal, 2015, 46, 1721-1731.	3.1	80
170	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
171	Pulmonary hypertension associated with benfluorex exposure. European Respiratory Journal, 2012, 40, 1164-1172.	3.1	75
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