List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	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
2	The physiological basis of pulmonary arterial hypertension. European Respiratory Journal, 2022, 59, 2102334.	6.7	61
3	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
4	Comparative bioavailability of inhaled treprostinil administered as LIQ861 and Tyvaso® in healthy subjects. Vascular Pharmacology, 2021, 138, 106840.	2.1	4
5	Relationship Between Time From Diagnosis and Morbidity/Mortality in Pulmonary Arterial Hypertension. Chest, 2021, 160, 277-286.	0.8	21
6	lt's Time to Put the Term "Pulmonary Vasodilators―to Rest. JACC Basic To Translational Science, 2021, 6, 870-871.	4.1	1
7	Risk assessment in pulmonary arterial hypertension: Insights from the GRIPHON study. Journal of Heart and Lung Transplantation, 2020, 39, 300-309.	0.6	39
8	Intravascular Ultrasound Pulmonary Artery Denervation to Treat Pulmonary Arterial Hypertension (TROPHY1). JACC: Cardiovascular Interventions, 2020, 13, 989-999.	2.9	47
9	Patients with pulmonary arterial hypertension with and without cardiovascular risk factors: Results from the AMBITION trial. Journal of Heart and Lung Transplantation, 2019, 38, 1286-1295.	0.6	62
10	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
11	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
12	Initial combination therapy with ambrisentan + tadalafil on pulmonary arterial hypertension‒related hospitalization in the AMBITION trial. Journal of Heart and Lung Transplantation, 2019, 38, 194-202.	0.6	19
13	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
14	Clinical trial design and new therapies for pulmonary arterial hypertension. European Respiratory Journal, 2019, 53, 1801908.	6.7	142
15	An overview of the 6th World Symposium on Pulmonary Hypertension. European Respiratory Journal, 2019, 53, 1802148.	6.7	345
16	Cor Pulmonale Revisited. From Ferrer and Harvey to the Present. Annals of the American Thoracic Society, 2018, 15, S42-S44.	3.2	8
17	Pulmonary Arterial Hypertension-Related Morbidity Is Prognostic for Mortality. Journal of the American College of Cardiology, 2018, 71, 752-763.	2.8	82
18	Assessing Prognosis of Pulmonary Arterial Hypertension in the Therapeutic Era. Circulation, 2018, 137, 705-706.	1.6	5

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19	Macitentan in pulmonary hypertension due to left ventricular dysfunction. European Respiratory Journal, 2018, 51, 1701886.	6.7	139
20	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
21	Overdue to understand anticoagulation in pulmonary arterial hypertension. Pulmonary Circulation, 2018, 8, 1-1.	1.7	1
22	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
23	Risk-stratified outcomes with initial combination therapy in pulmonary arterial hypertension: Application of the REVEAL risk score. Journal of Heart and Lung Transplantation, 2018, 37, 1410-1417.	0.6	15
24	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
25	Comparison of hemodynamic parameters in treatment-naÃ <sup>-</sup> ve and pre-treated patients with pulmonary arterial hypertension in the randomized phase III PATENT-1 study. Journal of Heart and Lung Transplantation, 2017, 36, 509-519.	0.6	22
26	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.	0.9	135
27	Selexipag for the treatment of connective tissue disease-associated pulmonary arterial hypertension. European Respiratory Journal, 2017, 50, 1602493.	6.7	97
28	Macitentan Improves Health-Related QualityÂof Life for Patients With Pulmonary Arterial Hypertension. Chest, 2017, 151, 106-118.	0.8	46
29	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
30	Metabolic dysfunction in pulmonary hypertension: from basic science to clinical practice. European Respiratory Review, 2017, 26, 170094.	7.1	60
31	Treprostinil Administered to Treat Pulmonary Arterial Hypertension Using a Fully Implantable Programmable Intravascular Delivery System. Chest, 2016, 150, 27-34.	0.8	48
32	Are Animal Models in PulmonaryÂHypertension Relevant toÂtheÂClinicalÂDisease? â^—. Journal of the American College of Cardiology, 2016, 67, 2047-2049.	2.8	3
33	Initial combination therapy with ambrisentan and tadalafil and mortality in patients with pulmonary arterial hypertension: a secondary analysis of the results from the randomised, controlled AMBITION study. Lancet Respiratory Medicine,the, 2016, 4, 894-901.	10.7	59
34	Pulmonary Hypertension OverlapÂSyndromes. Journal of the American College of Cardiology, 2016, 68, 379-381.	2.8	5
35	Incident and prevalent cohorts with pulmonary arterial hypertension: insight from SERAPHIN. European Respiratory Journal, 2015, 46, 1711-1720.	6.7	39
36	Pulmonary Artery Denervation for Pulmonary Artery Hypertension. JACC: Cardiovascular Interventions, 2015, 8, 2024-2025.	2.9	5

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37	Selexipag for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 2522-2533.	27.0	790
38	Improving patient outcomes in pulmonary hypertension. European Respiratory Review, 2015, 24, 550-551.	7.1	5
39	Riociguat for the treatment of pulmonary arterial hypertension: a long-term extension study (PATENT-2). European Respiratory Journal, 2015, 45, 1303-1313.	6.7	174
40	The Beta-Adrenergic Receptor in Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2015, 65, 681-683.	2.8	12
41	Use of clinically relevant responder threshold criteria to evaluate the response to treatment in the Phase III PATENT-1 study. Journal of Heart and Lung Transplantation, 2015, 34, 338-347.	0.6	10
42	Navigating the uncharted waters of combination therapy in pulmonary arterial hypertension: COMPASS or dead-reckoning. European Respiratory Journal, 2015, 46, 297-298.	6.7	4
43	Initial Use of Ambrisentan plus Tadalafil in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 834-844.	27.0	906
44	Effect of Macitentan on Hospitalizations. JACC: Heart Failure, 2015, 3, 1-8.	4.1	51
45	New horizons in pulmonary arterial hypertension management. European Respiratory Review, 2014, 23, 408-409.	7.1	3
46	Long-term sildenafil added to intravenous epoprostenol in patients with pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2014, 33, 689-697.	0.6	23
47	Two formulations of epoprostenol sodium in the treatment of pulmonary arterial hypertension: EPITOME-1 (epoprostenol for injection in pulmonary arterial hypertension), a phase IV, open-label, randomized study. American Heart Journal, 2014, 167, 218-225.e1.	2.7	27
48	Long-term results from the EARLY study of bosentan in WHO functional class II pulmonary arterial hypertension patients. International Journal of Cardiology, 2014, 172, 332-339.	1.7	47
49	Efficacy, safety and clinical pharmacology of macitentan in comparison to other endothelin receptor antagonists in the treatment of pulmonary arterial hypertension. Expert Opinion on Drug Safety, 2014, 13, 391-405.	2.4	60
50	Chronic thromboembolic pulmonary hypertension. Lancet Respiratory Medicine,the, 2014, 2, 573-582.	10.7	146
51	Riociguat for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2013, 369, 330-340.	27.0	1,120
52	Macitentan and Morbidity and Mortality in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2013, 369, 809-818.	27.0	1,168
53	Updated Treatment Algorithm of Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2013, 62, D60-D72.	2.8	596
54	New Trial Designs and Potential Therapies for Pulmonary Artery Hypertension. Journal of the American College of Cardiology, 2013, 62, D82-D91.	2.8	113

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55	Riociguat for Pulmonary Hypertension. New England Journal of Medicine, 2013, 369, 2266-2268.	27.0	21
56	Cor Pulmonale Revisited. Journal of the American College of Cardiology, 2013, 62, 1112-1113.	2.8	12
57	Rapid Transition from Inhaled Iloprost to Inhaled Treprostinil in Patients with Pulmonary Arterial Hypertension. Cardiovascular Therapeutics, 2013, 31, 38-44.	2.5	29
58	Treprostinil for the treatment of pulmonary arterial hypertension. Expert Review of Cardiovascular Therapy, 2013, 11, 13-25.	1.5	25
59	Oral Treprostinil for the Treatment of Pulmonary Arterial Hypertension in Patients Receiving Background Endothelin Receptor Antagonist and Phosphodiesterase Type 5 Inhibitor Therapy (The) Tj ETQq1 1 (	).7 <b>&amp;48</b> 14 r	gBīd©verloc
60	Treatment of Pulmonary Hypertension Caused by Left Heart Failure With Pulmonary Arterial Hypertension–Specific Therapies. Circulation, 2013, 128, 475-476.	1.6	6
61	Efficacy and Safety of Oral Treprostinil Monotherapy for the Treatment of Pulmonary Arterial Hypertension. Circulation, 2013, 127, 624-633.	1.6	291
62	A paradigm shift in pulmonary arterial hypertension management. European Respiratory Review, 2013, 22, 423-426.	7.1	3
63	Contemporary Trends in the Diagnosis and Management of Pulmonary Arterial Hypertension. Chest, 2013, 143, 324-332.	0.8	122
64	Pulmonary Hypertension: Current Management and Future Directions. Handbook of Experimental Pharmacology, 2013, 218, 551-555.	1.8	1
65	Pulmonary Hypertension: Current Management and Future Directions. Handbook of Experimental Pharmacology, 2013, , 551-555.	1.8	1
66	Exercise training for pulmonary hypertension: another prescription to write?. European Respiratory Journal, 2012, 40, 7-8.	6.7	10
67	Safety and efficacy evaluation of ambrisentan in pulmonary hypertension. Expert Opinion on Drug Safety, 2012, 11, 1003-1011.	2.4	19
68	Future Perspectives in Pulmonary Arterial Hypertension. Progress in Respiratory Research, 2012, , 276-279.	0.1	1
69	ARIESâ€3: Ambrisentan Therapy in a Diverse Population of Patients with Pulmonary Hypertension. Cardiovascular Therapeutics, 2012, 30, 93-99.	2.5	85
70	The pulmonary arterial hypertension quality enhancement research initiative: comparison of patients with idiopathic PAH to patients with systemic sclerosis-associated PAH. Annals of the Rheumatic Diseases, 2012, 71, 249-252.	0.9	63
71	Safety and Efficacy of Transition from Systemic Prostanoids to Inhaled Treprostinil in Pulmonary Arterial Hypertension. American Journal of Cardiology, 2012, 110, 1546-1550.	1.6	34
72	Endothelin receptor antagonists for the treatment of pulmonary artery hypertension. Life Sciences, 2012, 91, 517-521.	4.3	56

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73	Clinical trials with endothelin receptor antagonists: What went wrong and where can we improve?. Life Sciences, 2012, 91, 528-539.	4.3	76
74	The study of risk in pulmonary arterial hypertension. European Respiratory Review, 2012, 21, 234-238.	7.1	15
75	The 6-Minute Walk Test in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2012, 186, 396-397.	5.6	25
76	Inhaled treprostinil: a therapeutic review. Drug Design, Development and Therapy, 2012, 6, 19.	4.3	43
77	Systemic sclerosis-associated pulmonary hypertension: why disease-specific composite endpoints are needed. Arthritis Research and Therapy, 2011, 13, 114.	3.5	8
78	Long-term effects of inhaled treprostinil in patients with pulmonary arterial hypertension: The TReprostinil sodium Inhalation Used in the Management of Pulmonary arterial Hypertension (TRIUMPH) study open-label extension. Journal of Heart and Lung Transplantation, 2011, 30, 1327-1333.	0.6	98
79	Analysis of the Validation Status of Quality of Life and Functional Disability Measures in Pulmonary Arterial Hypertension Related to Systemic Sclerosis: Results of a Systematic Literature Analysis by the Expert Panel on Outcomes Measures in Pulmonary Arterial Hypertension Related to Systemic Sclerosis (EPOSS), Journal of Rheumatology, 2011, 38, 2419-2427.	2.0	4
80	Combination Therapy for Pulmonary Artery Hypertension: What Is the Evidence?. Cardiology, 2011, 120, 172-173.	1.4	3
81	Long-term Treatment With Sildenafil Citrate in Pulmonary Arterial Hypertension. Chest, 2011, 140, 1274-1283.	0.8	237
82	Validation of the 6 min walk test according to the OMERACT filter: a systematic literature review by the EPOSS-OMERACT group. Annals of the Rheumatic Diseases, 2010, 69, 1360-1363.	0.9	34
83	Echocardiography as an Outcome Measure in Scleroderma-related Pulmonary Arterial Hypertension: A Systematic Literature Analysis by the EPOSS Group. Journal of Rheumatology, 2010, 37, 105-115.	2.0	37
84	Worldwide Physician Education and Training in Pulmonary Hypertension. Chest, 2010, 137, 85S-94S.	0.8	26
85	Addition of Inhaled Treprostinil to Oral Therapy for Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2010, 55, 1915-1922.	2.8	484
86	Editor's response to "Ethical issues associated with globalization of placebo-controlled trials in pulmonary arterial hypertension― Journal of Heart and Lung Transplantation, 2010, 29, 827-828.	0.6	1
87	Metabolic Dysfunction in the Pathogenesis of Pulmonary Hypertension. Cell Metabolism, 2010, 12, 313-314.	16.2	9
88	Pulmonary Vasodilators in COPD. , 2009, , 699-704.		0
89	ACCF/AHA 2009 Expert Consensus Document on Pulmonary Hypertension. Circulation, 2009, 119, 2250-2294.	1.6	992
90	Borderline Pulmonary Arterial Pressure Is Associated with Decreased Exercise Capacity in Scleroderma. American Journal of Respiratory and Critical Care Medicine, 2009, 180, 881-886.	5.6	141

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91	Medical Treatment of Pulmonary Arterial Hypertension. Seminars in Respiratory and Critical Care Medicine, 2009, 30, 484-492.	2.1	6
92	A Functional Single-Nucleotide Polymorphism in the <i>TRPC6</i> Gene Promoter Associated With Idiopathic Pulmonary Arterial Hypertension. Circulation, 2009, 119, 2313-2322.	1.6	173
93	Identification of putative endothelial progenitor cells (CD34 <sup>+</sup> CD133 <sup>+</sup> Flk-1 <sup>+</sup> ) in endarterectomized tissue of patients with chronic thromboembolic pulmonary hypertension. American Journal of Physiology - Lung Cellular and Molecular Physiology. 2009. 296. L870-L878.	2.9	77
94	Longterm Survival Among Patients with Scleroderma-associated Pulmonary Arterial Hypertension Treated with Intravenous Epoprostenol. Journal of Rheumatology, 2009, 36, 2244-2249.	2.0	77
95	Metered dose inhaler delivery of treprostinil for the treatment of pulmonary hypertension. Pulmonary Pharmacology and Therapeutics, 2009, 22, 50-56.	2.6	43
96	ACCF/AHA 2009 Expert Consensus Document on Pulmonary Hypertension. Journal of the American College of Cardiology, 2009, 53, 1573-1619.	2.8	1,797
97	Endothelin and the Systemic Circulation. Journal of the American College of Cardiology, 2009, 53, 1318-1319.	2.8	2
98	Future Perspectives for the Treatment of Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2009, 54, S108-S117.	2.8	62
99	Updated Evidence-Based Treatment Algorithm in Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2009, 54, S78-S84.	2.8	463
100	Long-Term Ambrisentan Therapy for the Treatment of Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2009, 54, 1971-1981.	2.8	227
101	Ambrisentan Therapy in Patients With Pulmonary Arterial Hypertension Who Discontinued Bosentan or Sitaxsentan Due to Liver Function Test Abnormalities. Chest, 2009, 135, 122-129.	0.8	167
102	Hemodynamics and Epoprostenol Use Are Associated With Thrombocytopenia in Pulmonary Arterial Hypertension. Chest, 2009, 135, 130-136.	0.8	49
103	Defining appropriate outcome measures in pulmonary arterial hypertension related to systemic sclerosis: A Delphi consensus study with cluster analysis. Arthritis and Rheumatism, 2008, 59, 867-875.	6.7	56
104	Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2008, 51, 1527-1538.	2.8	269
105	Bosentan for Treatment of Inoperable Chronic Thromboembolic Pulmonary Hypertension. Journal of the American College of Cardiology, 2008, 52, 2127-2134.	2.8	506
106	Acute effects of the combination of sildenafil and inhaled treprostinil on haemodynamics and gas exchange in pulmonary hypertension. Pulmonary Pharmacology and Therapeutics, 2008, 21, 824-832.	2.6	64
107	Treatment of Pulmonary Arterial Hypertension Due to Scleroderma: Challenges for the Future. Rheumatic Disease Clinics of North America, 2008, 34, 191-197.	1.9	7
108	Exercise-Induced Pulmonary Arterial Hypertension. Circulation, 2008, 118, 2120-2121.	1.6	20

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109	Pulmonary Arterial Hypertension. , 2008, , 759-771.		Ο
110	Prednisolone inhibits PDGF-induced nuclear translocation of NF-κB in human pulmonary artery smooth muscle cells. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2008, 295, L648-L657.	2.9	28
111	Ambrisentan for the Treatment of Pulmonary Arterial Hypertension. Circulation, 2008, 117, 3010-3019.	1.6	967
112	<i>BMPR2</i> Mutation and Outcome in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2008, 177, 1300-1301.	5.6	7
113	Addition of Sildenafil to Long-Term Intravenous Epoprostenol Therapy in Patients with Pulmonary Arterial Hypertension. Annals of Internal Medicine, 2008, 149, 521.	3.9	558
114	Upregulation of Na+/Ca2+ exchanger contributes to the enhanced Ca2+ entry in pulmonary artery smooth muscle cells from patients with idiopathic pulmonary arterial hypertension. American Journal of Physiology - Cell Physiology, 2007, 292, C2297-C2305.	4.6	79
115	Hemodynamic and Functional Assessment of Patients with Sickle Cell Disease and Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2007, 175, 1272-1279.	5.6	227
116	Function of Kv1.5 channels and genetic variations ofKCNA5in patients with idiopathic pulmonary arterial hypertension. American Journal of Physiology - Cell Physiology, 2007, 292, C1837-C1853.	4.6	141
117	Central Venous Blood Oxygen Saturation Monitoring in Patients With Chronic Pulmonary Arterial Hypertension Treated With Continuous IV Epoprostenol. Chest, 2007, 132, 786-792.	0.8	10
118	Clinical trials and basic research: defining mechanisms and improving treatment in connective tissue disease. Arthritis Research and Therapy, 2007, 9, S10.	3.5	3
119	Medical Therapy for Pulmonary Arterial Hypertension. Chest, 2007, 131, 1917-1928.	0.8	477
120	Sildenafil for pulmonary arterial hypertension associated with connective tissue disease. Journal of Rheumatology, 2007, 34, 2417-22.	2.0	152
121	Safety and Efficacy of Inhaled Treprostinil as Add-On Therapy to Bosentan in Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2006, 48, 1433-1437.	2.8	115
122	Favorable Effects of Inhaled Treprostinil in Severe Pulmonary Hypertension. Journal of the American College of Cardiology, 2006, 48, 1672-1681.	2.8	135
123	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.	2.8	498
124	Temporal trends and drug exposures in pulmonary hypertension: An American experience. American Heart Journal, 2006, 152, 521-526.	2.7	78
125	ARIES-1: A PLACEBO-CONTROLLED, EFFICACY AND SAFETY STUDY OF AMBRISENTAN IN PATIENTS WITH PULMONARY ARTERIAL HYPERTENSION. Chest, 2006, 130, 121S.	0.8	27
126	Is Methamphetamine Use Associated With Idiopathic Pulmonary Arterial Hypertension?. Chest, 2006, 130, 1657-1663.	0.8	173

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127	Update in Pulmonary Hypertension 2005. American Journal of Respiratory and Critical Care Medicine, 2006, 173, 499-505.	5.6	40
128	Randomized Study of Adding Inhaled Iloprost to Existing Bosentan in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2006, 174, 1257-1263.	5.6	565
129	Chronic Thromboembolic Pulmonary Hypertension. Circulation, 2006, 113, 2011-2020.	1.6	791
130	Pulmonary Arterial Hypertension. Proceedings of the American Thoracic Society, 2006, 3, 111-115.	3.5	80
131	Current and Future Management of Chronic Thromboembolic Pulmonary Hypertension: From Diagnosis to Treatment Responses. Proceedings of the American Thoracic Society, 2006, 3, 601-607.	3.5	48
132	Endothelin-1 and the Pulmonary Vascular Response to Altitude. Circulation, 2006, 114, 1350-1351.	1.6	21
133	Thyrotoxicosis as a Risk Factor for Pulmonary Arterial Hypertension. Annals of Internal Medicine, 2006, 144, 222.	3.9	1
134	Management following pulmonary thromboendarterectomy: Experience versus evidence*. Critical Care Medicine, 2005, 33, 2132-2133.	0.9	2
135	The right ventricle in pulmonary hypertension. Coronary Artery Disease, 2005, 16, 13-18.	0.7	373
136	Ambrisentan for pulmonary arterial hypertension. Future Cardiology, 2005, 1, 425-432.	1.2	17
137	Pathogenesis of Pulmonary Arterial Hypertension. Circulation, 2005, 111, 534-538.	1.6	186
138	lloprost inhalation solution for the treatment of pulmonary arterial hypertension. Expert Opinion on Pharmacotherapy, 2005, 6, 1921-1930.	1.8	18
139	Evaluation and Management of the Patient with Pulmonary Arterial Hypertension. Annals of Internal Medicine, 2005, 143, 282.	3.9	121
140	Sildenafil Citrate Therapy for Pulmonary Arterial Hypertension. New England Journal of Medicine, 2005, 353, 2148-2157.	27.0	2,237
141	Ambrisentan Therapy for Pulmonary Arterial Hypertension. Journal of the American College of Cardiology, 2005, 46, 529-535.	2.8	441
142	Bosentan Inhibits Transient Receptor Potential Channel Expression in Pulmonary Vascular Myocytes. American Journal of Respiratory and Critical Care Medicine, 2004, 170, 1101-1107.	5.6	91
143	Preoperative Partitioning of Pulmonary Vascular Resistance Correlates With Early Outcome After Thromboendarterectomy for Chronic Thromboembolic Pulmonary Hypertension. Circulation, 2004, 109, 18-22.	1.6	377
144	Enhanced expression of transient receptor potential channels in idiopathic pulmonary arterial hypertension. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 13861-13866.	7.1	395

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145	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.	2.2	903
146	Pulmonary arterial hypertension: a look to the future. Journal of the American College of Cardiology, 2004, 43, S89-S90.	2.8	34
147	Clinical classification of pulmonary hypertension. Journal of the American College of Cardiology, 2004, 43, S5-S12.	2.8	1,542
148	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.	2.8	206
149	Endothelin receptor antagonists in pulmonary arterial hypertension. Journal of the American College of Cardiology, 2004, 43, S62-S67.	2.8	153
150	Cellular and molecular mechanisms of pulmonary vascular remodeling: role in the development of pulmonary hypertension. Microvascular Research, 2004, 68, 75-103.	2.5	263
151	Introduction. Chest, 2004, 126, 7S-10S.	0.8	244
152	Treprostinil, a Prostacyclin Analogue, in Pulmonary Arterial Hypertension Associated With Connective Tissue Disease. Chest, 2004, 126, 420-427.	0.8	232
153	Executive Summary. Chest, 2004, 126, 4S-6S.	0.8	60
154	Bosentan: profile report. Drugs and Therapy Perspectives, 2003, 19, 5-6.	0.6	0
155	Effects of the Dual Endothelin Receptor Antagonist Bosentan in Patients With Pulmonary Arterial Hypertension. Chest, 2003, 124, 247-254.	0.8	271
156	Successful Withdrawal of Long-term Epoprostenol Therapy for Pulmonary Arterial Hypertension. Chest, 2003, 124, 1612-1615.	0.8	43
157	Efficacy and Safety of Treprostinil: An Epoprostenol Analog for Primary Pulmonary Hypertension. Journal of Cardiovascular Pharmacology, 2003, 41, 293-299.	1.9	219
158	Epoprostenol Therapy as a Bridge to Pulmonary Thromboendarterectomy for Chronic Thromboembolic Pulmonary Hypertension. Chest, 2003, 123, 319-320.	0.8	32
159	High prevalence of elevated clotting factor VIII in chronic thromboembolic pulmonary hypertension. Thrombosis and Haemostasis, 2003, 90, 372-376.	3.4	221
160	Bone morphogenetic proteins induce apoptosis in human pulmonary vascular smooth muscle cells. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2003, 285, L740-L754.	2.9	237
161	Bosentan: a dual endothelin receptor antagonist. Expert Opinion on Investigational Drugs, 2002, 11, 991-1002.	4.1	65
162	Inhaled Iloprost for Severe Pulmonary Hypertension. New England Journal of Medicine, 2002, 347, 322-329.	27.0	1,626

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163	New Treatments for Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2002, 165, 1209-1216.	5.6	129
164	Differential Effects of Stable Prostacyclin Analogs on Smooth Muscle Proliferation and Cyclic AMP Generation in Human Pulmonary Artery. American Journal of Respiratory Cell and Molecular Biology, 2002, 26, 194-201.	2.9	211
165	Therapy of Pulmonary Hypertension. American Journal of Respiratory and Critical Care Medicine, 2002, 166, 1308-1309.	5.6	57
166	Endothelin in Health and Disease: Endothelin Receptor Antagonists in the Management of Pulmonary Artery Hypertension. Journal of Cardiovascular Pharmacology and Therapeutics, 2002, 7, 9-19.	2.0	41
167	Bosentan Therapy for Pulmonary Arterial Hypertension. New England Journal of Medicine, 2002, 346, 896-903.	27.0	2,545
168	Bosentan. American Journal of Cardiovascular Drugs, 2002, 2, 343.	2.2	1
169	Complete results of the first randomized, placebo-controlled study of bosentan, a dual endothelin receptor antagonist, in pulmonary arterial hypertension. Current Therapeutic Research, 2002, 63, 227-246.	1.2	32
170	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.	5.6	1,288
171	Chronic thromboembolic pulmonary hypertension. Progress in Cardiovascular Diseases, 2002, 45, 203-212.	3.1	14
172	Pulmonary Vasodilators. , 2002, , 605-609.		0
173	Effects of the dual endothelin-receptor antagonist bosentan in patients with pulmonary hypertension: a randomised placebocontrolled study. Lancet, The, 2001, 358, 1119-1123.	13.7	1,421
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