Steven D Nathan

List of Publications by Year
in descending order

273 papers	15,415 citations	26567 56 h-index	19136 118 g-index
293 all docs	293 docs citations	293 times ranked	10849 citing authors

#	Article	IF	Citations
1	Derivation and validation of a simple multidimensional index incorporating exercise capacity parameters for survival prediction in idiopathic pulmonary fibrosis. Thorax, 2023, 78, 368-375.	2.7	10
2	Riociguat for Sarcoidosis-Associated Pulmonary Hypertension. Chest, 2022, 161, 448-457.	0.4	24
3	Lung Transplantation for Patients With COVID-19. Chest, 2022, 161, 169-178.	0.4	54
4	Fostamatinib for the Treatment of Hospitalized Adults With Coronavirus Disease 2019: A Randomized Trial. Clinical Infectious Diseases, 2022, 75, e491-e498.	2.9	34
5	A Phase-2 Exploratory Randomized Controlled Trial of INOpulse in Patients with Fibrotic Interstitial Lung Disease Requiring Oxygen. Annals of the American Thoracic Society, 2022, 19, 594-602.	1.5	17
6	Lung Diseaseâ€“Related Pulmonary Hypertension. Cardiology Clinics, 2022, 40, 77-88.	0.9	2
7	Efficacy of Inhaled Treprostinil on Multiple Disease Progression Events in Patients with Pulmonary Hypertension due to Parenchymal Lung Disease in the INCREASE Trial. American Journal of Respiratory and Critical Care Medicine, 2022, 205, 198-207.	2.5	32
8	Piecing together the bigger picture: Idiopathic pulmonary fibrosis in Australia and beyond. Respirology, 2022, , .	1.3	0
9	Biological Variation of Donor-Derived Cell-Free DNA in Stable Lung Transplant Recipients. journal of applied laboratory medicine, The, 2022, , .	0.6	4
10	WASOG statement on the diagnosis and management of sarcoidosis-associated pulmonary hypertension. European Respiratory Review, 2022, 31, 210165.	3.0	28
11	Screening Strategies for Pulmonary Hypertension in Patients With Interstitial Lung Disease. Chest, 2022, 162, 145-155.	0.4	24
12	Reply to: Inhaled Treprostinil after Initial Clinical Worsening: To Continue or Not to Continue, That's the Question. American Journal of Respiratory and Critical Care Medicine, 2022, , .	2.5	0
13	Integrating Clinical Probability into the Diagnostic Approach to Idiopathic Pulmonary Fibrosis: An International Working Group Perspective. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 247-259.	2.5	15
14	Donor-derived cell-free DNA as a composite marker of acute lung allograft dysfunction in clinical care. Journal of Heart and Lung Transplantation, 2022, 41, 458-466.	0.3	20
15	The six-minute walk test in sarcoidosis associated pulmonary hypertension: Results from an international registry. Respiratory Medicine, 2022, 196, 106801.	1.3	15
16	Relative environmental and social disadvantage in patients with idiopathic pulmonary fibrosis. Thorax, 2022, 77, 1237-1242.	2.7	14
17	Echocardiographic estimate of pulmonary artery pressure in sarcoidosis patients - real world data from a multi-national study.. Sarcoidosis Vasculitis and Diffuse Lung Diseases, 2022, 38, e2021032.	0.2	2
18	Elevated cell-free DNA in respiratory viral infection and associated lung allograft dysfunction. American Journal of Transplantation, 2022, 22, 2560-2570.	2.6	7

#	Article	IF	Citations
19	The Antifibrotic Effects of Inhaled Treprostinil: An Emerging Option for ILD. Advances in Therapy, 2022, 39, 3881-3895.	1.3	15
20	Utility of a Molecular Classifier as a Complement to High-Resolution Computed Tomography to Identify Usual Interstitial Pneumonia. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 211-220.	2.5	55
21	Using forced vital capacity (FVC) in the clinic to monitor patients with idiopathic pulmonary fibrosis (IPF): pros and cons. Expert Review of Respiratory Medicine, 2021, 15, 175-181.	1.0	7
22	Association Between Anticoagulation and Survival in Interstitial Lung Disease. Chest, 2021, 159, 1507-1516.	0.4	10
23	Standardization of the 6-min walk test in clinical trials of idiopathic pulmonary fibrosis. Contemporary Clinical Trials, 2021, 100, 106227.	0.8	4
24	High-Flow Nasal Cannula Therapy in COVID-19: Using the ROX Index to Predict Success. Respiratory Care, 2021, 66, 909-919.	0.8	119
25	Efficacy and safety of sildenafil added to pirfenidone in patients with advanced idiopathic pulmonary fibrosis and risk of pulmonary hypertension: a double-blind, randomised, placebo-controlled, phase 2b trial. Lancet Respiratory Medicine,the, 2021, 9, 85-95.	5.2	96
26	Inhaled Nitric Oxide via High-Flow Nasal Cannula in Patients with Acute Respiratory Failure Related to COVID-19. Clinical Medicine Insights: Circulatory, Respiratory and Pulmonary Medicine, 2021, 15, 117954842110470.	0.5	9
27	Idiopathic pulmonary fibrosis patients with severe physiologic impairment: characteristics and outcomes. Respiratory Research, 2021, 22, 5.	1.4	10
28	Atraumatic forearm swelling in a patient with poorly controlled asthma. Respiratory Medicine Case Reports, 2021, 33, 101454.	0.2	0
29	Impact of the new definition for pulmonary hypertension in patients with lung disease: an analysis of the United Network for Organ Sharing database. Pulmonary Circulation, 2021, 11, 1-7.	0.8	13
30	Changes in Neutrophilâ€“Lymphocyte or Plateletâ€“Lymphocyte Ratios and Their Associations with Clinical Outcomes in Idiopathic Pulmonary Fibrosis. Journal of Clinical Medicine, 2021, 10, 1427.	1.0	17
31	Development and Validation of a Clinical Diagnostic Scoring System for the Diagnosis of IPF. Annals of the American Thoracic Society, 2021, 18, 1803-1810.	1.5	2
32	Incidence and prognostic significance of pleural effusions in pulmonary arterial hypertension. Pulmonary Circulation, 2021, 11, 1-10.	0.8	5
33	Cell-free DNA maps COVID-19 tissue injury and risk of death and can cause tissue injury. JCI Insight, 2021, 6, .	2.3	86
34	Pulmonary hypertension due to interstitial lung disease or chronic obstructive pulmonary disease: a patient experience study of symptoms and their impact on quality of life. Pulmonary Circulation, 2021, 11, 1-9.	0.8	8
35	Donor derived cell free DNA% is elevated with pathogens that are risk factors for acute and chronic lung allograft injury. Journal of Heart and Lung Transplantation, 2021, 40, 1454-1462.	0.3	13
36	Effect of Antimicrobial Therapy on Respiratory Hospitalization or Death in Adults With Idiopathic Pulmonary Fibrosis. JAMA - Journal of the American Medical Association, 2021, 325, 1841.	3.8	43

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37	Automated Digital Quantification of Pulmonary Fibrosis in Human Histopathology Specimens. Frontiers in Medicine, 2021, 8, 607720.	1.2	13
38	Pulmonary hypertension in interstitial lung disease: screening, diagnosis and treatment. Current Opinion in Pulmonary Medicine, 2021, 27, 396-404.	1.2	16
39	Use of donor-derived-cell-free DNA as a marker of early allograft injury in primary graft dysfunction (PGD) to predict the risk of chronic lung allograft dysfunction (CLAD). Journal of Heart and Lung Transplantation, 2021, 40, 488-493.	0.3	26
40	Impact of lung morphology on clinical outcomes with riociguat in patients with pulmonary hypertension and idiopathic interstitial pneumonia: A post hoc subgroup analysis of the RISE-IIP study. Journal of Heart and Lung Transplantation, 2021, 40, 494-503.	0.3	20
41	Differentiation of Idiopathic Pulmonary Fibrosis from Connective Tissue Disease-Related Interstitial Lung Disease Using Quantitative Imaging. Journal of Clinical Medicine, 2021, 10, 2663.	1.0	11
42	Inhaled treprostinil and forced vital capacity in patients with interstitial lung disease and associated pulmonary hypertension: a post-hoc analysis of the INCREASE study. Lancet Respiratory Medicine,the, 2021, 9, 1266-1274.	5.2	62
43	Serum levels of small HDL particles are negatively correlated with death or lung transplantation in an observational study of idiopathic pulmonary fibrosis. European Respiratory Journal, 2021, 58, 2004053.	3.1	10
44	Experience of Treating COVID-19 With Remdesivir and Convalescent Plasma in a Resource-Limited Setting: A Prospective, Observational Study. Open Forum Infectious Diseases, 2021, 8, ofab391.	0.4	11
45	Donor-derived cell-free DNA accurately detects acute rejection in lung transplant patients, a multicenter cohort study. Journal of Heart and Lung Transplantation, 2021, 40, 822-830.	0.3	34
46	Computed Tomography Findings Suggestive of Connective Tissue Disease in the Setting of Usual Interstitial Pneumonia. Journal of Computer Assisted Tomography, 2021, 45, 776-781.	0.5	5
47	Inhaled Treprostinil in Pulmonary Hypertension Due to Interstitial Lung Disease. New England Journal of Medicine, 2021, 384, 325-334.	13.9	292
48	CHARACTERIZATION OF PATIENTS WITH PULMONARY HYPERTENSION DUE TO COPD: A REAL-WORLD DATA ANALYSIS. Chest, 2021, 160, A1784-A1785.	0.4	0
49	COMPARISON OF IDIOPATHIC VS CONNECTIVE TISSUE DISEASE-ASSOCIATED PULMONARY ARTERIAL HYPERTENSION GROUPS IN US CLINICAL PRACTICE. Chest, 2021, 160, A2301-A2303.	0.4	0
50	SCREENING FOR PULMONARY HYPERTENSION IN PATIENTS WITH INTERSTITIAL LUNG DISEASE: RECOMMENDATIONS FROM A DELPHI CONSENSUS PANEL. Chest, 2021, 160, A1239-A1242.	0.4	0
51	INCIDENCE AND IMPACT OF POST-OPERATIVE ACUTE KIDNEY INJURY REQUIRING RENAL REPLACEMENT THERAPY DURING INDEX LUNG TRANSPLANT HOSPITALIZATION. Chest, 2021, 160, A2500-A2501.	0.4	0
52	TRANSTHORACIC ECHOCARDIOGRAM (ECHO) AND RIGHT HEART CATHETERIZATION (RHC) AS DISEASE MANAGEMENT TOOLS FOR PATIENTS WITH PULMONARY ARTERIAL HYPERTENSION (PAH) IN US CLINICAL PRACTICE. Chest, 2021, 160, A2331-A2333.	0.4	0
53	DOSE RESPONSE ANALYSIS OF INHALED TREPROSTINIL IN PULMONARY HYPERTENSION ASSOCIATED WITH INTERSTITIAL LUNG DISEASE AND ITS EFFECTS ON CLINICAL WORSENING. Chest, 2021, 160, A2279-A2280.	0.4	0
54	SEVERE THROMBOCYTOPENIA DUE TO IV EPOPROSTENOL: DON'T MUCK WITH THE PLATELETS. Chest, 2021, 160, A2180-A2181.	0.4	1

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55	CARE AND CHARACTERISTICS OF PATIENTS WITH PULMONARY ARTERIAL HYPERTENSION (PAH) IN US CLINICAL PRACTICE. Chest, 2021, 160, A2287-A2289.	0.4	0
56	COMPARISON OF EFFECTS OF INHALED TREPROSTINIL ON LUNG FUNCTION IN PATIENTS WITH PULMONARY HYPERTENSION ASSOCIATED WITH INTERSTITIAL LUNG DISEASE AND PULMONARY ARTERIAL HYPERTENSION. Chest, 2021, 160, A2244-A2246.	0.4	1
57	THE IMPACT OF HEMODYNAMIC PARAMETERS ON INHALED TREPROSTINIL TREATMENT RESPONSE: A SUBGROUP ANALYSIS FROM THE INCREASE TRIAL. Chest, 2021, 160, A2265-A2266.	0.4	0
58	Does 1-minute walk test predict results of 6-minute walk test in patients with idiopathic pulmonary fibrosis?. Sarcoidosis Vasculitis and Diffuse Lung Diseases, 2021, 38, e2021005.	0.2	2
59	Lung nodules due to Candida parapsilosis in a person with cystic fibrosis. BMJ Case Reports, 2021, 14, e245441.	0.2	1
60	ISCHEMIC COLITIS WITH NINTEDANIB USE: THE CONUNDRUM OF A COMMON SYMPTOM DUE TO A RARE CAUSE. Chest, 2020, 158, A1110-A1111.	0.4	2
61	THE IMPACT OF INHALED TREPROSTINIL ON PATIENT LUNG FUNCTION: RESULTS FROM THE INCREASE STUDY. Chest, 2020, 158, A2179-A2180.	0.4	2
62	SUCCESSFUL CALCINEURIN-INHIBITOR-FREE IMMUNOSUPPRESSION REGIMEN WITH SIROLIMUS AND PREDNISONE IN LUNG TRANSPLANT RECIPIENTS: A CASE SERIES. Chest, 2020, 158, A2383.	0.4	0
63	HIGHER DONOR PAO2/FIO2 RATIO APPEARS TO BE ASSOCIATED WITH INCREASED INCIDENCE OF PRIMARY GRAFT DYSFUNCTION IN LUNG TRANSPLANT RECIPIENTS. Chest, 2020, 158, A2399-A2400.	0.4	0
64	RELATIONSHIP BETWEEN ENVISIA GENOMIC CLASSIFIER AND AN HRCT-DERIVED FIBROTIC INDEX FROM DATA DRIVEN TEXTURE ANALYSIS ON 50 ILD PATIENTS. Chest, 2020, 158, A1064-A1065.	0.4	0
65	The Pulmonary Fibrosis Foundation Patient Registry. Rationale, Design, and Methods. Annals of the American Thoracic Society, 2020, 17, 1620-1628.	1.5	27
66	Expert consensus on the management of adverse events and prescribing practices associated with the treatment of patients taking pirfenidone for idiopathic pulmonary fibrosis: a Delphi consensus study. BMC Pulmonary Medicine, 2020, 20, 191.	0.8	6
67	HRCT evaluation of patients with interstitial lung disease: comparison of the 2018 and 2011 diagnostic guidelines. Therapeutic Advances in Respiratory Disease, 2020, 14, 175346662096849.	1.0	12
68	A RETROSPECTIVE DESCRIPTIVE ANALYSIS OF SYSTEMIC SCLEROSIS-RELATED INTERSTITIAL LUNG DISEASE AND PULMONARY HYPERTENSION. Chest, 2020, 158, A1878-A1879.	0.4	0
69	DOES SURGICAL LUNG BIOPSY CHANGE MANAGEMENT IN HOSPITALIZED PATIENTS WITH SUSPECTED INTERSTITIAL LUNG DISEASE?. Chest, 2020, 158, A1068-A1069.	0.4	0
70	OUTCOMES IN PATIENTS WITH IDIOPATHIC PULMONARY FIBROSIS AND PULMONARY HYPERTENSION. Chest, 2020, 158, A1071-A1072.	0.4	0
71	Lung transplantation in China: a firm foundation for a solid future. Annals of Translational Medicine, 2020, 8, 265-265.	0.7	3
72	POINT: Should Every Patient With Idiopathic Pulmonary Fibrosis Be Referred for Transplant Evaluation? Yes. Chest, 2020, 157, 1411-1412.	0.4	2

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73	A 48-Year-Old South African Woman with Rheumatoid Arthritis and Lung Nodules. Chest, 2020, 157, e151-e155.	0.4	2
74	A Molecular Classifier That Identifies Usual Interstitial Pneumonia in Transbronchial Biopsy Specimens of Patients With Interstitial Lung Disease. Chest, 2020, 157, 1391-1392.	0.4	1
75	The association between white blood cell count and outcomes in patients with idiopathic pulmonary fibrosis. Respiratory Medicine, 2020, 170, 106068.	1.3	16
76	Rebuttal From Dr Nathan. Chest, 2020, 157, 1415.	0.4	0
77	Physiological predictors of survival in patients with sarcoidosis-associated pulmonary hypertension: results from an international registry. European Respiratory Journal, 2020, 55, 1901747.	3.1	67
78	Targeting the Wnt signaling pathway through R-spondin 3 identifies an anti-fibrosis treatment strategy for multiple organs. PLoS ONE, 2020, 15, e0229445.	1.1	23
79	A Randomized, Double-Blind, Placebo-Controlled Study of Pulsed, Inhaled Nitric Oxide in Subjects at Risk ofÂPulmonary Hypertension Associated With Pulmonary Fibrosis. Chest, 2020, 158, 637-645.	0.4	62
80	FVC variability in patients with idiopathic pulmonary fibrosis and role of 6-min walk test to predict further change. European Respiratory Journal, 2020, 55, 1902151.	3.1	19
81	Outcomes of mechanically ventilated patients with COVID-19 associated respiratory failure. PLoS ONE, 2020, 15, e0242651.	1.1	67
82	IPF in Saudi Arabia: Lessons for all. Annals of Thoracic Medicine, 2020, 15, 183.	0.7	0
83	Outcomes of mechanically ventilated patients with COVID-19 associated respiratory failure. , 2020, 15, e0242651.		0
84	Outcomes of mechanically ventilated patients with COVID-19 associated respiratory failure. , 2020, 15, e0242651.		0
85	Outcomes of mechanically ventilated patients with COVID-19 associated respiratory failure. , 2020, 15, e0242651.		0
86	Outcomes of mechanically ventilated patients with COVID-19 associated respiratory failure. , 2020, 15, e0242651.		0
87	Cardiovascular Risks, Bleeding Risks, and Clinical Events from 3 Phase III Trials of Pirfenidone in Patients with Idiopathic Pulmonary Fibrosis. Advances in Therapy, 2019, 36, 2910-2926.	1.3	18
88	Contemporary optimized practice in the management of pulmonary sarcoidosis. Therapeutic Advances in Respiratory Disease, 2019, 13, 175346661986893.	1.0	13
89	Nintedanib and Sildenafil in Patients with Idiopathic Pulmonary Fibrosis. Echoes of the Past, Lessons for the Future. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 1459-1461.	2.5	4
90	Multimodal noninvasive prediction of pulmonary hypertension in IPF. Clinical Respiratory Journal, 2019, 13, 567-573.	0.6	15

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91	COMBINING RADIOLOGY AND ENVISIA, A MOLECULAR CLASSIFIER, TO IMPROVE USUAL INTERSTITIAL PNEUMONIA (UIP) DIAGNOSIS. Chest, 2019, 156, A253-A256.	0.4	0
92	WBC COUNT AS A PROGNOSTIC INDICATOR IN THE TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS. Chest, 2019, 156, A1071-A1072.	0.4	0
93	CATEGORIZATION OF GROUP 3 PULMONARY HYPERTENSION BY THE 2018 DEFINITION: WHO IS IN, WHO IS OUT?. Chest, 2019, 156, A872-A873.	0.4	1
94	OPEN-LABEL DOSE-ESCALATION DATA FROM THE RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY TO ASSESS THE SAFETY AND EFFICACY OF PULSED, INHALED NITRIC OXIDE (INO) IN SUBJECTS AT RISK OF PULMONARY HYPERTENSION ASSOCIATED WITH PULMONARY FIBROSIS (PH-PF) ON LONG TERM OXYGEN THERAPY. Chest, 2019, 156, A2273-A2275.	0.4	1
95	Singleâ€center experience with use of letermovir for CMV prophylaxis or treatment in thoracic organ transplant recipients. Transplant Infectious Disease, 2019, 21, e13166.	0.7	40
96	Riociguat for idiopathic interstitial pneumonia-associated pulmonary hypertension (RISE-IIP): a randomised, placebo-controlled phase 2b study. Lancet Respiratory Medicine,the, 2019, 7, 780-790.	5.2	139
97	Donor-derived cell-free DNA predicts allograft failure and mortality after lung transplantation. EBioMedicine, 2019, 40, 541-553.	2.7	83
98	Connective tissue disease-associated interstitial lung disease and outcomes after hospitalization: A cohort study. Respiratory Medicine, 2019, 154, 1-5.	1.3	11
99	Pirfenidone in patients with idiopathic pulmonary fibrosis and more advanced lung function impairment. Respiratory Medicine, 2019, 153, 44-51.	1.3	54
100	Resequencing Study Confirms That Host Defense and Cell Senescence Gene Variants Contribute to the Risk of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 199-208.	2.5	90
101	Use of a molecular classifier to identify usual interstitial pneumonia in conventional transbronchial lung biopsy samples: a prospective validation study. Lancet Respiratory Medicine,the, 2019, 7, 487-496.	5.2	119
102	EVALUATING CLINICAL UTILITY OF A UIP GENOMIC CLASSIFIER IN SUBJECTS WITH AND WITHOUT A HRCT PATTERN OF UIP. Chest, 2019, 156, A175-A178.	0.4	0
103	Pulmonary hypertension due to interstitial lung disease. Current Opinion in Pulmonary Medicine, 2019, 25, 459-467.	1.2	22
104	Critical Care of the Adult Patient With Cystic Fibrosis. Chest, 2019, 155, 202-214.	0.4	28
105	Genetic determinants of risk in pulmonary arterial hypertension: international genome-wide association studies and meta-analysis. Lancet Respiratory Medicine,the, 2019, 7, 227-238.	5.2	122
106	Efficacy of Pirfenidone in the Context of Multiple Disease Progression Events in Patients With Idiopathic Pulmonary Fibrosis. Chest, 2019, 155, 712-719.	0.4	24
107	Pulmonary hypertension in chronic lung disease and hypoxia. European Respiratory Journal, 2019, 53, 1801914.	3.1	428
108	Extracorporeal Membrane Oxygenation as a Bridge to Initial Medical Therapy in a Patient With Decompensated Pulmonary Arterial Hypertension Presenting With Biventricular Failure. Journal of Medical Cases, 2019, 10, 260-263.	0.4	0

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109	A 24-Year-Old Woman With Precipitous Respiratory Failure After Lung Transplantation. Chest, 2018, 153, e53-e56.	0.4	8
110	Late manifestation of alloantibody-associated injury and clinical pulmonary antibody-mediated rejection: Evidence from cell-free DNA analysis. Journal of Heart and Lung Transplantation, 2018, 37, 925-932.	0.3	69
111	Sildenafil added to pirfenidone in patients with advanced idiopathic pulmonary fibrosis and risk of pulmonary hypertension: A Phase IIb, randomised, double-blind, placebo-controlled study â€“ Rationale and study design. Respiratory Medicine, 2018, 138, 13-20.	1.3	27
112	Single vs. bilateral lung transplantation. Current Opinion in Organ Transplantation, 2018, 23, 316-323.	0.8	18
113	RNAseq analysis of bronchial epithelial cells to identify COPD-associated genes and SNPs. BMC Pulmonary Medicine, 2018, 18, 42.	0.8	20
114	The Value and Application of the 6-Minute-Walk Test in Idiopathic Pulmonary Fibrosis. Annals of the American Thoracic Society, 2018, 15, 3-10.	1.5	54
115	RISING INCIDENCE OF PULMONARY EMBOLISM POST-LUNG TRANSPLANTATION: A SINGLE CENTER EXPERIENCE. Chest, 2018, 154, 1103A-1104A.	0.4	0
116	CONNECTIVE TISSUE DISEASE-ASSOCIATED INTERSTITIAL LUNG DISEASE AND OUTCOMES AFTER HOSPITALIZATION: A COHORT STUDY. Chest, 2018, 154, 418A-419A.	0.4	0
117	EFFECT OF PIRFENIDONE ON EXERCISE CAPACITY AND DYSPNEA IN PATIENTS WITH IDIOPATHIC PULMONARY FIBROSIS (IPF) AND MORE ADVANCED LUNG FUNCTION IMPAIRMENT. Chest, 2018, 154, 432A-433A.	0.4	1
118	Prevalence and impact of WHO group 3 pulmonary hypertension in advanced idiopathic nonspecific interstitial pneumonia. European Respiratory Journal, 2018, 52, 1800545.	3.1	19
119	Identifying Barriers to Idiopathic Pulmonary Fibrosis Treatment: A Survey of Patient and Physician Views. Respiration, 2018, 96, 514-524.	1.2	54
120	Dose modification and dose intensity during treatment with pirfenidone: analysis of pooled data from three multinational phase III trials. BMJ Open Respiratory Research, 2018, 5, e000323.	1.2	35
121	Novel management strategies for idiopathic pulmonary fibrosis. Expert Review of Respiratory Medicine, 2018, 12, 831-842.	1.0	9
122	Exercise pulmonary haemodynamic response predicts outcomes in fibrotic lung disease. European Respiratory Journal, 2018, 52, 1801015.	3.1	6
123	Clinical management and outcomes of patients with Hermansky-Pudlak syndrome pulmonary fibrosis evaluated for lung transplantation. PLoS ONE, 2018, 13, e0194193.	1.1	29
124	The Diagnosis and Management of AirwayÂComplications Following LungÂTransplantation. Chest, 2017, 152, 627-638.	0.4	67
125	Early postoperative management after lung transplantation: Results of an international survey. Clinical Transplantation, 2017, 31, e12985.	0.8	8
126	Enhancing Insights into Pulmonary Vascular Disease through a Precision Medicine Approach. A Joint NHLBIâ€“Cardiovascular Medical Research and Education Fund Workshop Report. American Journal of Respiratory and Critical Care Medicine, 2017, 195, 1661-1670.	2.5	59

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127	sGC stimulators: Evidence for riociguat beyond groups 1 and 4 pulmonary hypertension. Respiratory Medicine, 2017, 122, S28-S34.	1.3	6
128	A randomized, placebo-controlled, double-blinded, crossover trial of pioglitazone for severe asthma. Journal of Allergy and Clinical Immunology, 2017, 140, 1716-1718.	1.5	17
129	Ventricular Diastolic Pressure Ratio as a Marker of Treatment Response in Pulmonary Hypertension. Chest, 2017, 152, 980-989.	0.4	2
130	Pirfenidone safety and adverse event management in idiopathic pulmonary fibrosis. European Respiratory Review, 2017, 26, 170057.	3.0	162
131	Does 1-Minute Walk Test Predict Results of 6-Minute Walk Test in Patients With Idiopathic Pulmonary Fibrosis?. Chest, 2017, 152, A486.	0.4	4
132	Tolerability and Efficacy of Selexipag in Real Life Clinical Setting. Chest, 2017, 152, A998.	0.4	0
133	Pitfalls in developing new compounds for idiopathic pulmonary fibrosis. Current Opinion in Pulmonary Medicine, 2017, 23, 426-431.	1.2	2
134	Pulmonary Hypertension in Diffuse Parenchymal Lung Diseases. Chest, 2017, 151, 204-214.	0.4	15
135	Idiopathic pulmonary fibrosis: effects and optimal management of comorbidities. Lancet Respiratory Medicine,the, 2017, 5, 72-84.	5.2	137
136	Idiopathic interstitial pneumonia-associated pulmonary hypertension: A target for therapy?. Respiratory Medicine, 2017, 122, S10-S13.	1.3	15
137	Effect of pirfenidone on mortality: pooled analyses and meta-analyses of clinical trials in idiopathic pulmonary fibrosis. Lancet Respiratory Medicine,the, 2017, 5, 33-41.	5.2	240
138	Incidence and Impact of Gastroparesis After Lung Transplantation. Chest, 2017, 152, A1101.	0.4	2
139	Tracheobronchial Tree Size as a Predictor of Disease Severity and Outcomes in Idiopathic Pulmonary Fibrosis. Chest, 2017, 152, A487.	0.4	1
140	Survival in Idiopathic Pulmonary Fibrosis: Perspectives from Pulmonary Arterial Hypertension. Journal of Managed Care & Specialty Pharmacy, 2017, 23, S3-S4.	0.5	1
141	Predicting Life Expectancy for Pirfenidone in Idiopathic Pulmonary Fibrosis. Journal of Managed Care & Specialty Pharmacy, 2017, 23, S17-S24.	0.5	65
142	Evaluating new treatment options. American Journal of Managed Care, 2017, 23, S183-S190.	0.8	4
143	Effect of continued treatment with pirfenidone following clinically meaningful declines in forced vital capacity: analysis of data from three phase 3 trials in patients with idiopathic pulmonary fibrosis. Thorax, 2016, 71, 429-435.	2.7	151
144	A safety evaluation of pirfenidone for the treatment of idiopathic pulmonary fibrosis. Expert Opinion on Drug Safety, 2016, 15, 975-982.	1.0	3

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145	Antacid therapy and idiopathic pulmonary fibrosis: cause for heartburn?. Lancet Respiratory Medicine,the, 2016, 4, 340-341.	5.2	2
146	Efficacy of pirfenidone in patients with idiopathic pulmonary fibrosis with more preserved lung function. European Respiratory Journal, 2016, 48, 843-851.	3.1	134
147	Ambrisentan response in connective tissue disease-associated pulmonary arterial hypertension (CTD-PAH) â€“ A subgroup analysis of the ARIES-E clinical trial. Respiratory Medicine, 2016, 117, 254-263.	1.3	26
148	Hemothorax following lung transplantation: incidence, risk factors, and effect on morbidity and mortality. Multidisciplinary Respiratory Medicine, 2016, 11, 40.	0.6	21
149	POINT: Should All Patients With Idiopathic Pulmonary Fibrosis, Even Those With More Than Moderate Impairment, Be Treated With Nintedanib or Pirfenidone? Yes. Chest, 2016, 150, 273-275.	0.4	13
150	Rebuttal From Drs King andÂNathan. Chest, 2016, 150, 278.	0.4	0
151	Sildenafil for pulmonary hypertension complicating idiopathic pulmonary fibrosis: a rationale grounded in basic science. European Respiratory Journal, 2016, 47, 1615-1617.	3.1	7
152	Pirfenidone for idiopathic pulmonary fibrosis: analysis of pooled data from three multinational phase 3 trials. European Respiratory Journal, 2016, 47, 243-253.	3.1	349
153	Lung transplantation in <scp>IIP</scp>: A review. Respirology, 2016, 21, 1173-1184.	1.3	31
154	Safety of pirfenidone in patients with idiopathic pulmonary fibrosis: integrated analysis of cumulative data from 5 clinical trials. BMJ Open Respiratory Research, 2016, 3, e000105.	1.2	96
155	Pulmonary artery size as a predictor of outcomes in idiopathic pulmonary fibrosis. European Respiratory Journal, 2016, 47, 1445-1451.	3.1	49
156	Upfront combination therapy: does the AMBITION study herald a new era in the treatment of pulmonary arterial hypertension?. Thorax, 2016, 71, 107-109.	2.7	5
157	Effect of Pirfenidone on All-Cause Mortality in Patients With Idiopathic Pulmonary Fibrosis (IPF): Comparison of Pooled Analysis With Meta-analysis From the ASCEND and CAPACITY Trials. Chest, 2015, 148, 363A.	0.4	0
158	Effect of Pirfenidone on IPF-Related Mortality Outcome Measures in Patients With Idiopathic Pulmonary Fibrosis (IPF): Pooled Data Analysis From the ASCEND and CAPACITY Trials. Chest, 2015, 148, 391A.	0.4	2
159	Management of Idiopathic Pulmonary Fibrosis in the Elderly Patient. Chest, 2015, 148, 242-252.	0.4	36
160	The Future of Lung Transplantation. Chest, 2015, 147, 309-316.	0.4	37
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