

A Phase 3 Trial of Pirfenidone in Patients with Idiopathic

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Citation Report

#	ARTICLE	IF	CITATIONS
2	Upregulation of activin-B and follistatin in pulmonary fibrosis – a translational study using human biopsies and a specific inhibitor in mouse fibrosis models. BMC Pulmonary Medicine, 2014, 14, 170.	0.8	17
3	Hypoxia-sensitive pathways in inflammation-driven fibrosis. American Journal of Physiology - Regulatory Integrative and Comparative Physiology, 2014, 307, R1369-R1380.	0.9	40
4	Molecular Mechanism and Treatment of Viral Hepatitis-Related Liver Fibrosis. International Journal of Molecular Sciences, 2014, 15, 10578-10604.	1.8	60
5	Open-Access Biorepository for Idiopathic Pulmonary Fibrosis. The Way Forward. Annals of the American Thoracic Society, 2014, 11, 1171-1175.	1.5	15
7	Measuring diffusion limitation with a perfusion-limited gas – Hyperpolarized ¹²⁹ Xe gas-transfer spectroscopy in patients with idiopathic pulmonary fibrosis. Journal of Applied Physiology, 2014, 117, 577-585.	1.2	77
9	Novel approaches to pulmonary fibrosis. Clinical Medicine, 2014, 14, s45-s49.	0.8	9
10	A new era of drug therapy for idiopathic pulmonary fibrosis. Lancet Respiratory Medicine, the, 2014, 2, 964-966.	5.2	1
11	Parenchymal cryobiopsies for interstitial lung diseases: A step <i>forward</i> in disease management. Respiriology, 2014, 19, 773-774.	1.3	6
12	Improving care for patients with idiopathic pulmonary fibrosis (IPF) in the UK: a round table discussion. Thorax, 2014, 69, 1136-1140.	2.7	31
13	ALK and ROS1 non-small-cell lung cancer: two molecular subgroups sensitive to targeted therapy. Lancet Respiratory Medicine, the, 2014, 2, 966-968.	5.2	14
14	Moving stem cell therapy to patients with idiopathic pulmonary fibrosis. Respiriology, 2014, 19, 950-951.	1.3	8
15	Computed Tomography: Revolutionizing the Practice of Medicine for 40 Years. Radiology, 2014, 273, S45-S74.	3.6	128
16	Pulmonary Hypertension due to Fibrotic Lung Disease: Hidden Value in a Neutral Trial. American Journal of Respiratory and Critical Care Medicine, 2014, 190, 131-132.	2.5	3
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19	Treatment with pirfenidone for two years decreases fibrosis, cytokine levels and enhances CB2 gene expression in patients with chronic hepatitis C. BMC Gastroenterology, 2014, 14, 131.	0.8	65
20	Clinical significance of epithelial mesenchymal transition (EMT) in chronic obstructive pulmonary disease (COPD): potential target for prevention of airway fibrosis and lung cancer. Clinical and Translational Medicine, 2014, 3, 33.	1.7	65
21	IPF clinical trial design and endpoints. Current Opinion in Pulmonary Medicine, 2014, 20, 463-471.	1.2	58
22	Treatment strategies for idiopathic interstitial pneumonias. Current Opinion in Pulmonary Medicine, 2014, 20, 442-448.	1.2	19

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23	A New Hope for Idiopathic Pulmonary Fibrosis. <i>New England Journal of Medicine</i> , 2014, 370, 2142-2143.	13.9	50
24	Combination therapy: the future of management for idiopathic pulmonary fibrosis?. <i>Lancet Respiratory Medicine</i> , 2014, 2, 933-942.	5.2	128
26	Hope for Disease-Modifying Treatment of Systemic Sclerosis/Scleroderma. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2014, 350, 480-482.	1.3	0
27	Beyond TGF β 2: Novel ways to target airway and parenchymal fibrosis. <i>Pulmonary Pharmacology and Therapeutics</i> , 2014, 29, 166-180.	1.1	14
30	Fibrosis pulmonar idiopática. <i>Medicine</i> , 2014, 11, 3799-3807.	0.0	0
31	Treatments for Idiopathic Pulmonary Fibrosis. <i>New England Journal of Medicine</i> , 2014, 371, 781-784.	13.9	31
32	New Therapeutic Targets in Idiopathic Pulmonary Fibrosis. Aiming to Rein in Runaway Wound-Healing Responses. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014, 190, 867-878.	2.5	209
33	Pirfenidone for Idiopathic Pulmonary Fibrosis, Thrombocytosis in Chronic Obstructive Pulmonary Disease Exacerbations, and a Longitudinal Study on E-Cigarettes. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014, 190, 699-700.	2.5	0
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47	Management of Idiopathic Pulmonary Fibrosis in the Elderly Patient. <i>Chest</i> , 2015, 148, 242-252.	0.4	36
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50	Which Patients With ARDS Benefit From Lung Biopsy?. <i>Chest</i> , 2015, 148, 1073-1082.	0.4	32
51	Recent Advances in Dyspnea. <i>Chest</i> , 2015, 147, 232-241.	0.4	110
52	The need for patient-centred clinical research in idiopathic pulmonary fibrosis. <i>BMC Medicine</i> , 2015, 13, 240.	2.3	30
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54	17(R)-resolvin D1 ameliorates bleomycin-induced pulmonary fibrosis in mice. <i>Physiological Reports</i> , 2015, 3, e12628.	0.7	35
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56	Molecular classification of idiopathic pulmonary fibrosis: Personalized medicine, genetics and biomarkers. <i>Respirology</i> , 2015, 20, 1010-1022.	1.3	44
57	Mesenchymal Stromal Cells in Animal Bleomycin Pulmonary Fibrosis Models: A Systematic Review. <i>Stem Cells Translational Medicine</i> , 2015, 4, 1500-1510.	1.6	94
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61	Pharmacological treatment of idiopathic pulmonary fibrosis â€“ preclinical and clinical studies of pirfenidone, nintedanib, and N-acetylcysteine. <i>European Clinical Respiratory Journal</i> , 2015, 2, 26385.	0.7	78
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63	CD248/Endosialin: A Novel Pericyte Target in Renal Fibrosis. <i>Nephron</i> , 2015, 131, 262-264.	0.9	6
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65	Idiopathic Pulmonary Fibrosis: Treatment and Prognosis. <i>Clinical Medicine Insights: Circulatory, Respiratory and Pulmonary Medicine</i> , 2015, 9s1, CCRPM.S23321.	0.5	42
67	Current and Future Therapies for Idiopathic Pulmonary Fibrosis. <i>Pulmonary Therapy</i> , 2015, 1, 1-18.	1.1	2
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79	Biomarkers for Staging and Evaluating the Therapy for Idiopathic Pulmonary Fibrosis. <i>Clinical Pulmonary Medicine</i> , 2015, 22, 165-171.	0.3	1
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81	Double-Blind Randomized Trial of Pirfenidone in Chinese Idiopathic Pulmonary Fibrosis Patients. <i>Medicine (United States)</i> , 2015, 94, e1600.	0.4	27
82	Honeycomb Lung: Time for a Change. <i>Archives of Pathology and Laboratory Medicine</i> , 2015, 139, 1398-1399.	1.2	5
83	Taking the "œœ-out of IPF. <i>European Respiratory Journal</i> , 2015, 45, 1539-1541.	3.1	3
84	Clinical highlights: messages from Munich. <i>ERJ Open Research</i> , 2015, 1, 00002-2015.	1.1	0

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104	Update on therapeutic management of idiopathic pulmonary fibrosis. <i>Therapeutics and Clinical Risk Management</i> , 2015, 11, 359.	0.9	51

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105	Exploring Clinical and Epidemiological Characteristics of Interstitial Lung Diseases: Rationale, Aims, and Design of a Nationwide Prospective Registry—The EXCITING-ILD Registry. <i>BioMed Research International</i> , 2015, 2015, 1-9.	0.9	42
106	Fibrosis Related Inflammatory Mediators: Role of the IL-10 Cytokine Family. <i>Mediators of Inflammation</i> , 2015, 2015, 1-15.	1.4	206
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108	Inflammatory Response Mechanisms Exacerbating Hypoxemia in Coexistent Pulmonary Fibrosis and Sleep Apnea. <i>Mediators of Inflammation</i> , 2015, 2015, 1-13.	1.4	27
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140	Pneumopathies interstitielles diffuses. <i>Revue Des Maladies Respiratoires Actualites</i> , 2015, 7, S54-S64.	0.0	0
141	Lung-Dominant Connective Tissue Disease. <i>Chest</i> , 2015, 148, 1438-1446.	0.4	49
142	Cough in interstitial lung disease. <i>Pulmonary Pharmacology and Therapeutics</i> , 2015, 35, 122-128.	1.1	13
143	Personalizing Therapy in Idiopathic Pulmonary Fibrosis: A Glimpse of the Future?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2015, 192, 1409-1411.	2.5	3

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145	MMP-9 >T Gene Polymorphism and Efficacy of Glucocorticoid Therapy in Idiopathic Pulmonary Fibrosis Patients. Genetic Testing and Molecular Biomarkers, 2015, 19, 591-597.	0.3	10
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148	New Treatments for Idiopathic Pulmonary Fibrosis: 'Die Another Day' if Diagnosed Early?. Respiration, 2015, 90, 352-352.	1.2	9
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157	Nintedanib and Pirfenidone. New Antifibrotic Treatments Indicated for Idiopathic Pulmonary Fibrosis Offer Hopes and Raises Questions. American Journal of Respiratory and Critical Care Medicine, 2015, 191, 252-254.	2.5	135
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162	Pharmacological treatment of idiopathic pulmonary fibrosis: an update. Drug Discovery Today, 2015, 20, 514-524.	3.2	26

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165	Quantitative Proteomics of Bronchoalveolar Lavage Fluid in Idiopathic Pulmonary Fibrosis. <i>Journal of Proteome Research</i> , 2015, 14, 1238-1249.	1.8	79
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169	Disruption of Calcium Signaling in Fibroblasts and Attenuation of Bleomycin-Induced Fibrosis by Nifedipine. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2015, 53, 450-458.	1.4	42
170	Principles of Rehabilitation and Reactivation: Interstitial Lung Disease, Sarcoidosis and Rheumatoid Disease with Respiratory Involvement. <i>Respiration</i> , 2015, 89, 89-99.	1.2	42
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172	Nintedanib modulates surfactant protein-D expression in A549 human lung epithelial cells via the c-Jun N-terminal kinase-activator protein-1 pathway. <i>Pulmonary Pharmacology and Therapeutics</i> , 2015, 32, 29-36.	1.1	20
173	La criobiopsia transbronquial en la enfermedad pulmonar intersticial: excelente relación coste/beneficio. <i>Archivos De Bronconeumología</i> , 2015, 51, 257-258.	0.4	2
174	Update on New Treatments for Idiopathic Pulmonary Fibrosis. <i>Current Emergency and Hospital Medicine Reports</i> , 2015, 3, 134-138.	0.6	0
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177	A novel segmental challenge model for bleomycin-induced pulmonary fibrosis in sheep. <i>Experimental Lung Research</i> , 2015, 41, 115-134.	0.5	26
178	Characteristic Patterns in the Fibrotic Lung. Comparing Idiopathic Pulmonary Fibrosis with Chronic Lung Allograft Dysfunction. <i>Annals of the American Thoracic Society</i> , 2015, 12, S34-S41.	1.5	16
179	Liposomal hydrogel formulation for transdermal delivery of pirfenidone. <i>Journal of Liposome Research</i> , 2016, 26, 1-9.	1.5	17
180	Pulmonary Resection for Lung Cancer in Patients With Idiopathic Interstitial Pneumonia. <i>Annals of Thoracic Surgery</i> , 2015, 100, 954-960.	0.7	66
181	FIBROSIS PULMONAR IDIOPÁTICA. <i>Revista Médica Clínica Las Condes</i> , 2015, 26, 292-301.	0.2	2

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183	Early Experience of Pirfenidone in Daily Clinical Practice in Belgium and The Netherlands: a Retrospective Cohort Analysis. <i>Advances in Therapy</i> , 2015, 32, 691-704.	1.3	31
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875	Diagnostic criteria for idiopathic pulmonary fibrosis: a Fleischner Society White Paper. <i>Lancet Respiratory Medicine</i> , 2018, 6, 138-153.	5.2	739
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878	Clinical experience in idiopathic pulmonary fibrosis: a retrospective study. <i>Acta Clinica Belgica</i> , 2018, 73, 139-143.	0.5	9
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885	Lung transplantation: indications and contraindications. <i>Journal of Thoracic Disease</i> , 2018, 10, 4574-4587.	0.6	76
886	Expression of mutant Sftpc in murine alveolar epithelia drives spontaneous lung fibrosis. <i>Journal of Clinical Investigation</i> , 2018, 128, 4008-4024.	3.9	152
887	Idiopathic Pulmonary Fibrosis-Unknown Cause, Global Occurrence and New Medical Possibilities. <i>Internal Medicine: Open Access</i> , 2018, 08, .	0.0	0
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889	Palliative and End-of-Life Care in Idiopathic Pulmonary Fibrosis. , 2018, , 97-106.		0

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914	Targeting interleukin-13 in idiopathic pulmonary fibrosis: from promising path to dead end. <i>European Respiratory Journal</i> , 2018, 52, 1802111.	3.1	23
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916	Impact of novel antifibrotic therapy on patient outcomes in idiopathic pulmonary fibrosis: patient selection and perspectives. <i>Patient Related Outcome Measures</i> , 2018, Volume 9, 321-328.	0.7	33
917	A Volumetric Computed Tomography Analysis of the Normal Lung in Idiopathic Pulmonary Fibrosis: The Relationship with the Survival. <i>Internal Medicine</i> , 2018, 57, 929-937.	0.3	5
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919	PD-1 up-regulation on CD4 ⁺ T cells promotes pulmonary fibrosis through STAT3-mediated IL-17A and TGF- β 1 production. <i>Science Translational Medicine</i> , 2018, 10, .	5.8	225
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922	Pirfenidone in patients with unclassifiable progressive fibrosing interstitial lung disease: design of a double-blind, randomised, placebo-controlled phase II trial. <i>BMJ Open Respiratory Research</i> , 2018, 5, e000289.	1.2	48
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929	Long-term safety of pirfenidone: results of the prospective, observational PASSPORT study. <i>ERJ Open Research</i> , 2018, 4, 00084-2018.	1.1	78
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932	Cytokine and Anti-Cytokine Agents as Future Therapeutics for Fibrostenosing IBD. , 2018, , 59-75.		0
933	Connective Tissue Related Interstitial Lung Disease. <i>Current Pulmonology Reports</i> , 2018, 7, 133-148.	0.5	0
934	Imaging biomarkers and staging in IPF. <i>Current Opinion in Pulmonary Medicine</i> , 2018, 24, 445-452.	1.2	16
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944	Ambulatory oxygen and quality of life in interstitial lung disease. <i>Lancet Respiratory Medicine</i> , 2018, 6, 730-731.	5.2	1
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947	Identification of MMP28 as a biomarker for the differential diagnosis of idiopathic pulmonary fibrosis. <i>PLoS ONE</i> , 2018, 13, e0203779.	1.1	19
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949	Drug and chemical induced photosensitivity from a clinical perspective. <i>Photochemical and Photobiological Sciences</i> , 2018, 17, 1885-1903.	1.6	33
950	Interstitial Lung Disease in the Elderly: A Review of Pathogenesis and Clinical Management. <i>Clinical Pulmonary Medicine</i> , 2018, 25, 157-165.	0.3	1
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952	Recommendations for evaluating and managing idiopathic pulmonary fibrosis. <i>JAAPA: Official Journal of the American Academy of Physician Assistants</i> , 2018, 31, 21-26.	0.1	7
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954	Targeting TGF- β 2 Signaling in Kidney Fibrosis. <i>International Journal of Molecular Sciences</i> , 2018, 19, 2532.	1.8	164
955	Real-World Practice Patterns for Prevention and Management of Potential Adverse Events with Pirfenidone in Patients with Idiopathic Pulmonary Fibrosis. <i>Pulmonary Therapy</i> , 2018, 4, 103-114.	1.1	4
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962	Optimising experimental research in respiratory diseases: an ERS statement. <i>European Respiratory Journal</i> , 2018, 51, 1702133.	3.1	98
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966	Fibrotic microtissue array to predict anti-fibrosis drug efficacy. <i>Nature Communications</i> , 2018, 9, 2066.	5.8	102
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969	Clinical Trials in Idiopathic Pulmonary Fibrosis in the "Posttreatment Era". <i>JAMA - Journal of the American Medical Association</i> , 2018, 319, 2275.	3.8	7
970	Safety, tolerability, pharmacokinetics, and pharmacodynamics of GLPG1690, a novel autotaxin inhibitor, to treat idiopathic pulmonary fibrosis (FLORA): a phase 2a randomised placebo-controlled trial. <i>Lancet Respiratory Medicine</i> , 2018, 6, 627-635.	5.2	173
971	Heterogeneity in Unclassifiable Interstitial Lung Disease. A Systematic Review and Meta-Analysis. <i>Annals of the American Thoracic Society</i> , 2018, 15, 854-863.	1.5	74
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973	Economic consequences of idiopathic pulmonary fibrosis in Denmark. <i>ERJ Open Research</i> , 2018, 4, 00045-2017.	1.1	4
974	Abnormal Bolus Reflux Is Associated With Poor Pulmonary Outcome in Patients With Idiopathic Pulmonary Fibrosis. <i>Journal of Neurogastroenterology and Motility</i> , 2018, 24, 395-402.	0.8	12
975	Targeting the Myofibroblast in Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018, 198, 834-835.	2.5	13
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978	Safety of nintedanib added to pirfenidone treatment for idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2018, 52, 1800230.	3.1	95
979	Therapeutic effects of telomerase in mice with pulmonary fibrosis induced by damage to the lungs and short telomeres. <i>ELife</i> , 2018, 7, .	2.8	88
980	Effect of pirfenidone on gastric emptying in a rat model. <i>Pulmonary Pharmacology and Therapeutics</i> , 2018, 51, 41-47.	1.1	3
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984	Diagnosis and management of idiopathic pulmonary fibrosis: Thoracic Society of Australia and New Zealand and Lung Foundation Australia position statements summary. <i>Medical Journal of Australia</i> , 2018, 208, 82-88.	0.8	13
985	Improvement in Patient-Reported Outcomes and Forced Vital Capacity during Nintedanib Treatment of Idiopathic Pulmonary Fibrosis. <i>Tohoku Journal of Experimental Medicine</i> , 2018, 245, 107-114.	0.5	5
986	Fibroblast senescence in the pathology of idiopathic pulmonary fibrosis. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2018, 315, L162-L172.	1.3	114
987	Casein Kinase 1 γ Inhibitor, PF670462 Attenuates the Fibrogenic Effects of Transforming Growth Factor- β ² in Pulmonary Fibrosis. <i>Frontiers in Pharmacology</i> , 2018, 9, 738.	1.6	28
988	New treatment paradigms for connective tissue disease-associated interstitial lung disease. <i>Current Opinion in Pulmonary Medicine</i> , 2018, 24, 453-460.	1.2	10
989	Unclassifiable interstitial lung disease. <i>Current Opinion in Pulmonary Medicine</i> , 2018, 24, 461-468.	1.2	15
990	The European IPF registry (eurIPFreg): baseline characteristics and survival of patients with idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2018, 19, 141.	1.4	199
991	Treating heart failure with preserved ejection fraction: learning from pulmonary fibrosis. <i>European Journal of Heart Failure</i> , 2018, 20, 1385-1391.	2.9	38
992	Amifostine Analog, DRDE-30, Attenuates Bleomycin-Induced Pulmonary Fibrosis in Mice. <i>Frontiers in Pharmacology</i> , 2018, 9, 394.	1.6	13
993	Proton Pump Inhibitors in IPF: A Call for Clinical Trials. <i>Frontiers in Pharmacology</i> , 2018, 9, 499.	1.6	15
994	Interstitial Pneumonia With Autoimmune Features. <i>Arthritis and Rheumatology</i> , 2018, 70, 1901-1913.	2.9	38
995	Prolonged treatment with open-label pirfenidone in Hermansky-Pudlak syndrome pulmonary fibrosis. <i>Molecular Genetics and Metabolism</i> , 2018, 125, 168-173.	0.5	24
996	The relationship between complement C3 expression and the MUC5B genotype in pulmonary fibrosis. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2018, 315, L1-L10.	1.3	28
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1001	Human alveolar epithelial cells type II are capable of TGF β ² -dependent epithelial-mesenchymal-transition and collagen-synthesis. <i>Respiratory Research</i> , 2018, 19, 138.	1.4	52
1002	Tissue and Bronchoalveolar Lavage Biomarkers in Idiopathic Pulmonary Fibrosis Patients on Pirfenidone. <i>Lung</i> , 2018, 196, 543-552.	1.4	13
1003	Idiopathic pulmonary fibrosis: time for greater expectations?. <i>European Respiratory Journal</i> , 2018, 52, 1801312.	3.1	2
1004	S100a4 Is Secreted by Alternatively Activated Alveolar Macrophages and Promotes Activation of Lung Fibroblasts in Pulmonary Fibrosis. <i>Frontiers in Immunology</i> , 2018, 9, 1216.	2.2	64
1005	The Role of Immune and Inflammatory Cells in Idiopathic Pulmonary Fibrosis. <i>Frontiers in Medicine</i> , 2018, 5, 43.	1.2	216
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1007	Mesenchymal Stem Cells for the Treatment of Idiopathic Pulmonary Fibrosis. <i>Frontiers in Medicine</i> , 2018, 5, 142.	1.2	60
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1011	Effect of pirfenidone on wound healing in lung transplant patients. <i>Multidisciplinary Respiratory Medicine</i> , 2018, 13, 16.	0.6	20
1012	SIS3, a specific inhibitor of smad3, attenuates bleomycin-induced pulmonary fibrosis in mice. <i>Biochemical and Biophysical Research Communications</i> , 2018, 503, 757-762.	1.0	21
1013	Japanese guideline for the treatment of idiopathic pulmonary fibrosis. <i>Respiratory Investigation</i> , 2018, 56, 268-291.	0.9	72
1014	Macitentan reduces progression of TGF- β ¹ -induced pulmonary fibrosis and pulmonary hypertension. <i>European Respiratory Journal</i> , 2018, 52, 1701857.	3.1	46
1015	Reducing protein oxidation reverses lung fibrosis. <i>Nature Medicine</i> , 2018, 24, 1128-1135.	15.2	88
1016	The Role of the Mammalian Target of Rapamycin (mTOR) in Pulmonary Fibrosis. <i>International Journal of Molecular Sciences</i> , 2018, 19, 778.	1.8	129
1017	Novel formononetin-7-sal ester ameliorates pulmonary fibrosis via MEF2c signaling pathway. <i>Toxicology and Applied Pharmacology</i> , 2018, 356, 15-24.	1.3	10
1018	Treatment of Interstitial Lung Disease Associated Cough. <i>Chest</i> , 2018, 154, 904-917.	0.4	50

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1019	Metformin Does Not Affect Clinically Relevant Outcomes in Patients with Idiopathic Pulmonary Fibrosis. <i>Respiration</i> , 2018, 96, 314-322.	1.2	41
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1021	The Management of Patients With Idiopathic Pulmonary Fibrosis. <i>Frontiers in Medicine</i> , 2018, 5, 148.	1.2	42
1022	Idiopathic pulmonary fibrosis: pathogenesis and management. <i>Respiratory Research</i> , 2018, 19, 32.	1.4	339
1023	Barriers to timely diagnosis of interstitial lung disease in the real world: the INTENSITY survey. <i>BMC Pulmonary Medicine</i> , 2018, 18, 9.	0.8	112
1024	Disease progression in idiopathic pulmonary fibrosis with mild physiological impairment: analysis from the Australian IPF registry. <i>BMC Pulmonary Medicine</i> , 2018, 18, 19.	0.8	58
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1027	N-acetylcysteine exposure is associated with improved survival in anti-nuclear antibody seropositive patients with usual interstitial pneumonia. <i>BMC Pulmonary Medicine</i> , 2018, 18, 30.	0.8	5
1028	Idiopathic Pulmonary Fibrosis. <i>New England Journal of Medicine</i> , 2018, 378, 1811-1823.	13.9	1,177
1029	Associated Pulmonary Hypertension Is an Independent Contributor to Exercise Intolerance in Chronic Fibrosing Interstitial Pneumonias. <i>Respiration</i> , 2018, 96, 543-551.	1.2	7
1030	Comprehensive gene expression profiling identifies distinct and overlapping transcriptional profiles in non-specific interstitial pneumonia and idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2018, 19, 153.	1.4	66
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1189	The Effectiveness, Safety, and Tolerability of Pirfenidone in Idiopathic Pulmonary Fibrosis: A Retrospective Study. <i>Advances in Therapy</i> , 2019, 36, 1126-1131.	1.3	14
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1205	Risk of anastomotic dehiscence in patients with pulmonary fibrosis transplanted while receiving anti-fibrotics: Experience of the Australian Lung Transplant Collaborative. <i>Journal of Heart and Lung Transplantation</i> , 2019, 38, 553-559.	0.3	24
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1283	Lung Transplantation for Interstitial Lung Disease. , 2019, , 131-149.		0
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1307	Pericytes and T Cells in Lung Injury and Fibroproliferation. <i>Molecular and Translational Medicine</i> , 2019, , 175-195.	0.4	2
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1328	Towards a refined definition of combined pulmonary fibrosis and emphysema. <i>Respirology</i> , 2019, 24, 9-10.	1.3	4
1329	Efficacy and Safety of Pirfenidone in Advanced Idiopathic Pulmonary Fibrosis. <i>Respiration</i> , 2019, 97, 242-251.	1.2	20
1330	Implications of the diagnostic criteria of idiopathic pulmonary fibrosis in clinical practice: Analysis from the Australian Idiopathic Pulmonary Fibrosis Registry. <i>Respirology</i> , 2019, 24, 361-368.	1.3	24
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1334	The Antifibrotic Activity of Prostacyclin Receptor Agonism Is Mediated through Inhibition of YAP/TAZ. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2019, 60, 578-591.	1.4	34
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1347	Effectiveness and safety of pirfenidone for idiopathic pulmonary fibrosis. <i>European Journal of Hospital Pharmacy</i> , 2020, 27, 350-354.	0.5	6
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1352	Pamrevlumab, an anti-connective tissue growth factor therapy, for idiopathic pulmonary fibrosis (PRAISE): a phase 2, randomised, double-blind, placebo-controlled trial. <i>Lancet Respiratory Medicine</i> , 2020, 8, 25-33.	5.2	165
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1509	Shedding light on developmental drugs for idiopathic pulmonary fibrosis. <i>Expert Opinion on Investigational Drugs</i> , 2020, 29, 797-808.	1.9	8
1510	The extended utility of antifibrotic therapy in progressive fibrosing interstitial lung disease. <i>Expert Review of Respiratory Medicine</i> , 2020, 14, 1001-1008.	1.0	6
1511	Cluster analysis based clinical profiling of Idiopathic Pulmonary Fibrosis patients according to comorbidities evident prior to diagnosis: a single-center observational study. <i>European Journal of Internal Medicine</i> , 2020, 80, 18-23.	1.0	5
1512	Comprehensive analysis of lncRNA-associated competing endogenous RNA network and immune infiltration in idiopathic pulmonary fibrosis. <i>Journal of Thoracic Disease</i> , 2020, 12, 1856-1865.	0.6	9
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1516	Asbestos-related diseases. <i>International Journal of Tuberculosis and Lung Disease</i> , 2020, 24, 562-567.	0.6	18
1517	Radiation-Induced Lung Fibrosis: Preclinical Animal Models and Therapeutic Strategies. <i>Cancers</i> , 2020, 12, 1561.	1.7	56
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1519	Tolerability of nintedanib-related diarrhea in patients with idiopathic pulmonary fibrosis. <i>Pulmonary Pharmacology and Therapeutics</i> , 2020, 62, 101917.	1.1	9
1520	Circulating matrix metalloproteinases and tissue metalloproteinase inhibitors in patients with idiopathic pulmonary fibrosis in the multicenter IPF-PRO Registry cohort. <i>BMC Pulmonary Medicine</i> , 2020, 20, 64.	0.8	59
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1523	Prominence of IL6, IGF, TLR, and Bioenergetics Pathway Perturbation in Lung Tissues of Scleroderma Patients With Pulmonary Fibrosis. <i>Frontiers in Immunology</i> , 2020, 11, 383.	2.2	40
1524	Perioperative pirfenidone treatment as prophylaxis against acute exacerbation of idiopathic pulmonary fibrosis: a single-center analysis. <i>Surgery Today</i> , 2020, 50, 905-911.	0.7	14
1525	NADPH Oxidase Inhibition in Fibrotic Pathologies. <i>Antioxidants and Redox Signaling</i> , 2020, 33, 455-479.	2.5	20
1526	Emerging biomarkers in chronic lung allograft dysfunction. <i>Expert Review of Molecular Diagnostics</i> , 2020, 20, 467-475.	1.5	6
1527	Contemporary Concise Review 2019: Interstitial lung disease. <i>Respirology</i> , 2020, 25, 756-763.	1.3	2
1528	Disease progression across the spectrum of idiopathic pulmonary fibrosis: A multicentre study. <i>Respirology</i> , 2020, 25, 1144-1151.	1.3	6
1529	Therapeutic Targets for the Treatment of Cardiac Fibrosis and Cancer: Focusing on TGF- β 2 Signaling. <i>Frontiers in Cardiovascular Medicine</i> , 2020, 7, 34.	1.1	85
1530	Efficacy and safety of pirfenidone in systemic sclerosis-related interstitial lung disease—a randomised controlled trial. <i>Rheumatology International</i> , 2020, 40, 703-710.	1.5	64
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1536	Genetic Variation in CCL18 Gene Influences CCL18 Expression and Correlates with Survival in Idiopathic Pulmonary Fibrosisâ€”Part B. <i>Journal of Clinical Medicine</i> , 2020, 9, 1993.	1.0	10
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1542	Metabolic Activation of Pirfenidone Mediated by Cytochrome P450s and Sulfotransferases. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 8059-8068.	2.9	16
1543	The myofibroblast at a glance. <i>Journal of Cell Science</i> , 2020, 133, .	1.2	167
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1546	A Unique Protein Self-Assembling Nanoparticle with Significant Advantages in Vaccine Development and Production. <i>Journal of Nanomaterials</i> , 2020, 2020, 1-10.	1.5	20
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1549	Probable usual interstitial pneumonia pattern on chest CT: is it sufficient for a diagnosis of idiopathic pulmonary fibrosis?. <i>European Respiratory Journal</i> , 2020, 55, 1802465.	3.1	25
1550	Antibody-based therapies for idiopathic pulmonary fibrosis. <i>Expert Opinion on Biological Therapy</i> , 2020, 20, 779-786.	1.4	13

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1552	Parametric Time-to-Event Model for Acute Exacerbations in Idiopathic Pulmonary Fibrosis. <i>CPT: Pharmacometrics and Systems Pharmacology</i> , 2020, 9, 87-95.	1.3	4
1553	Early decrease in erector spinae muscle area and future risk of mortality in idiopathic pulmonary fibrosis. <i>Scientific Reports</i> , 2020, 10, 2312.	1.6	38
1554	Reference values for high attenuation areas on chest CT in a healthy, never-smoker, multi-ethnic sample: The MESA study. <i>Respirology</i> , 2020, 25, 855-862.	1.3	13
1555	Patient gender bias on the diagnosis of idiopathic pulmonary fibrosis. <i>Thorax</i> , 2020, 75, 407-412.	2.7	30
1556	Pirfenidone in idiopathic pulmonary fibrosis: real-life experience in the referral centre of Siena. <i>Therapeutic Advances in Respiratory Disease</i> , 2020, 14, 175346662090632.	1.0	22
1557	Three Steps to Cure Pulmonary Fibrosis. Step 1: The Runaway Train or Groundhog Day?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 201, 1172-1174.	2.5	4
1558	Stereotactic ablative radiotherapy in patients with early-stage non-small cell lung cancer and co-existing interstitial lung disease. <i>Acta Oncologica</i> , 2020, 59, 569-573.	0.8	11
1559	Development of small molecule inhibitors targeting TGF- β 2 ligand and receptor: Structures, mechanism, preclinical studies and clinical usage. <i>European Journal of Medicinal Chemistry</i> , 2020, 191, 112154.	2.6	27
1560	Efficacy and Safety of Traditional Chinese Medicine in Idiopathic Pulmonary Fibrosis: A Meta-Analysis. <i>Evidence-based Complementary and Alternative Medicine</i> , 2020, 2020, 1-11.	0.5	15
1562	Systemic sclerosis-associated interstitial lung disease. <i>Lancet Respiratory Medicine</i> , 2020, 8, 304-320.	5.2	202
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1564	An Open-label Study With Pirfenidone on Chronic Hypersensitivity Pneumonitis. <i>Archivos De Bronconeumologia</i> , 2020, 56, 163-169.	0.4	7
1565	A post-transcriptional program of chemoresistance by AU-rich elements and TTP in quiescent leukemic cells. <i>Genome Biology</i> , 2020, 21, 33.	3.8	22
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1567	Screening for YAP Inhibitors Identifies Statins as Modulators of Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2020, 62, 479-492.	1.4	36
1568	Idiopathic Pulmonary Fibrosis: Pathogenesis and the Emerging Role of Long Non-Coding RNAs. <i>International Journal of Molecular Sciences</i> , 2020, 21, 524.	1.8	41
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1571	Surfactant protein A as a biomarker of outcomes of anti-fibrotic drug therapy in patients with idiopathic pulmonary fibrosis. <i>BMC Pulmonary Medicine</i> , 2020, 20, 27.	0.8	26
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1573	The dietary antioxidant quercetin reduces hallmarks of bleomycin-induced lung fibrogenesis in mice. <i>BMC Pulmonary Medicine</i> , 2020, 20, 112.	0.8	34
1574	Interstitial lung disease. <i>Canadian Journal of Respiratory, Critical Care, and Sleep Medicine</i> , 2020, 4, S21-S24.	0.2	0
1575	Clinical Decision-Making in Hypersensitivity Pneumonitis: Diagnosis and Management. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2020, 41, 214-228.	0.8	11
1576	Nonspecific Interstitial Pneumonia. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2020, 41, 184-201.	0.8	7
1577	Home Monitoring in Patients with Idiopathic Pulmonary Fibrosis. A Randomized Controlled Trial. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 202, 393-401.	2.5	79
1578	Conditional deletion of Nedd4-2 in lung epithelial cells causes progressive pulmonary fibrosis in adult mice. <i>Nature Communications</i> , 2020, 11, 2012.	5.8	52
1579	Pulse oximetry saturation can predict prognosis of idiopathic pulmonary fibrosis. <i>Respiratory Investigation</i> , 2020, 58, 190-195.	0.9	6
1580	Practical Considerations for the Diagnosis and Treatment of Fibrotic Interstitial Lung Disease During the Coronavirus Disease 2019 Pandemic. <i>Chest</i> , 2020, 158, 1069-1078.	0.4	57
1581	Safety and Efficacy of Pirfenidone in Advanced Idiopathic Pulmonary Fibrosis: A Nationwide Post-Marketing Surveillance Study in Korean Patients. <i>Advances in Therapy</i> , 2020, 37, 2303-2316.	1.3	20
1582	Pirfenidone for the treatment of interstitial lung disease associated to rheumatoid arthritis: a new scenario is coming?. <i>Respiratory Medicine Case Reports</i> , 2020, 30, 101051.	0.2	4
1583	Lung cryobiopsy and interstitial lung disease: What is its role in the era of multidisciplinary meetings and antifibrotics?. <i>Respirology</i> , 2020, 25, 987-996.	1.3	9
1584	CXXC5 Attenuates Pulmonary Fibrosis in a Bleomycin-Induced Mouse Model and MLFs by Suppression of the CD40/CD40L Pathway. <i>BioMed Research International</i> , 2020, 2020, 1-15.	0.9	4
1585	Patient-reported outcome measures in systemic sclerosis-related interstitial lung disease for clinical practice and clinical trials. <i>Journal of Scleroderma and Related Disorders</i> , 2020, 5, 48-60.	1.0	13
1586	Pirfenidone Inhibits Cell Proliferation and Collagen I Production of Primary Human Intestinal Fibroblasts. <i>Cells</i> , 2020, 9, 775.	1.8	31
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1590	Idiopathic pulmonary fibrosis and pulmonary hypertension: Heracles meets the Hydra. <i>British Journal of Pharmacology</i> , 2021, 178, 172-186.	2.7	20
1591	Refining the unilateral ureteral obstruction mouse model: No sham, no shame. <i>Laboratory Animals</i> , 2021, 55, 21-29.	0.5	3
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1595	Pirfenidone plus inhaled N-acetylcysteine for idiopathic pulmonary fibrosis: a randomised trial. <i>European Respiratory Journal</i> , 2021, 57, 2000348.	3.1	31
1596	A novel tree shrew model of pulmonary fibrosis. <i>Laboratory Investigation</i> , 2021, 101, 116-124.	1.7	9
1597	Chemokine Receptor 2—targeted Molecular Imaging in Pulmonary Fibrosis. A Clinical Trial. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021, 203, 78-89.	2.5	61
1598	Pharmacological management of Idiopathic Pulmonary Fibrosis: current and emerging options. <i>Expert Opinion on Pharmacotherapy</i> , 2021, 22, 191-204.	0.9	16
1599	Using forced vital capacity (FVC) in the clinic to monitor patients with idiopathic pulmonary fibrosis (IPF): pros and cons. <i>Expert Review of Respiratory Medicine</i> , 2021, 15, 175-181.	1.0	7
1600	Mechanisms of progressive fibrosis in connective tissue disease (CTD)-associated interstitial lung diseases (ILDs). <i>Annals of the Rheumatic Diseases</i> , 2021, 80, 143-150.	0.5	120
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1607	Rictor-targeting exosomal microRNA-16 ameliorates lung fibrosis by inhibiting the mTORC2-SPARC axis. <i>Experimental Cell Research</i> , 2021, 398, 112416.	1.2	12
1608	Transcriptome analysis of IPF fibroblastic foci identifies key pathways involved in fibrogenesis. <i>Thorax</i> , 2021, 76, 73-82.	2.7	25
1609	Target inhibition of galectin-3 by inhaled TD139 in patients with idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2021, 57, 2002559.	3.1	106
1610	Safety and efficacy of pirfenidone and nintedanib in patients with idiopathic pulmonary fibrosis and carrying a telomere-related gene mutation. <i>European Respiratory Journal</i> , 2021, 57, 2003198.	3.1	36
1612	Betulinic acid attenuated bleomycin-induced pulmonary fibrosis by effectively intervening Wnt/ β^2 -catenin signaling. <i>Phytomedicine</i> , 2021, 81, 153428.	2.3	14
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1614	Extended Delivery of Pirfenidone with Novel, Soft Contact Lenses <i>In Vitro</i> and <i>In Vivo</i> . <i>Journal of Ocular Pharmacology and Therapeutics</i> , 2021, 37, 75-83.	0.6	7
1615	Effectiveness of pirfenidone for idiopathic pulmonary fibrosis associated with pleuroparenchymal fibroelastosis-like lesions and nonspecific interstitial pneumonia. <i>Clinical Respiratory Journal</i> , 2021, 15, 272-279.	0.6	2
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1617	Real-World Study Analysing Progression and Survival of Patients with Idiopathic Pulmonary Fibrosis with Preserved Lung Function on Antifibrotic Treatment. <i>Advances in Therapy</i> , 2021, 38, 268-277.	1.3	13
1618	Leukocyte telomere length and mycophenolate therapy in chronic hypersensitivity pneumonitis. <i>European Respiratory Journal</i> , 2021, 57, 2002872.	3.1	32
1619	Associations of ω -3 Fatty Acids With Interstitial Lung Disease and Lung Imaging Abnormalities Among Adults. <i>American Journal of Epidemiology</i> , 2021, 190, 95-108.	1.6	11
1620	Clinical Molecular Imaging of Pulmonary CXCR4 Expression to Predict Outcome of Pirfenidone Treatment in Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2021, 159, 1094-1106.	0.4	23
1621	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. <i>Lancet Respiratory Medicine</i> , 2021, 9, 85-95.	5.2	96
1622	A disease progression model of longitudinal lung function decline in idiopathic pulmonary fibrosis patients. <i>Journal of Pharmacokinetics and Pharmacodynamics</i> , 2021, 48, 55-67.	0.8	4
1623	Priorities and expectations of patients attending a multidisciplinary interstitial lung disease clinic. <i>Respirology</i> , 2021, 26, 80-86.	1.3	12
1624	Effectiveness of Proton Pump Inhibitors in Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2021, 159, 673-682.	0.4	31

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1626	Anti-fibrotic effects of different sources of MSC in bleomycin-induced lung fibrosis in C57BL/6 male mice. <i>Respirology</i> , 2021, 26, 161-170.	1.3	24
1627	Progressive fibrosing interstitial lung diseases: A new concept and indication of nintedanib. <i>Modern Rheumatology</i> , 2021, 31, 13-19.	0.9	20
1628	Cause of mortality and sarcopenia in patients with idiopathic pulmonary fibrosis receiving antifibrotic therapy. <i>Respirology</i> , 2021, 26, 171-179.	1.3	24
1629	Deep Learning of Computed Tomography Virtual Wedge Resection for Prediction of Histologic Usual Interstitial Pneumonitis. <i>Annals of the American Thoracic Society</i> , 2021, 18, 51-59.	1.5	22
1630	Spectrum of Pulmonary Fibrosis from Interstitial Lung Abnormality to Usual Interstitial Pneumonia: Importance of Identification and Quantification of Traction Bronchiectasis in Patient Management. <i>Korean Journal of Radiology</i> , 2021, 22, 811.	1.5	20
1631	Treatment of systemic sclerosis-associated interstitial lung disease: a work in progress. <i>Lancet Respiratory Medicine</i> , 2021, 9, 5-7.	5.2	4
1632	Lung organoids and other preclinical models of pulmonary fibrosis. <i>QJM - Monthly Journal of the Association of Physicians</i> , 2021, 114, 167-173.	0.2	9
1633	Update in Diagnosis of Idiopathic Pulmonary Fibrosis and Interstitial Lung Abnormality. <i>Journal of the Korean Society of Radiology</i> , 2021, 82, 770.	0.1	0
1634	Antifibrotic Therapy: Is There a Role in Myositis-Interstitial Lung Disease?. <i>Respiration</i> , 2021, 100, 923-932.	1.2	1
1635	Current trends in candidate selection, contraindications, and indications for lung transplantation. <i>Journal of Thoracic Disease</i> , 2021, 13, 6514-6527.	0.6	9
1636	Clinical Significance of Continuable Treatment with Nintedanib Over 12 Months for Idiopathic Pulmonary Fibrosis in a Real-World Setting. <i>Drug Design, Development and Therapy</i> , 2021, Volume 15, 223-230.	2.0	10
1637	Idiopathic pulmonary fibrosis patients with severe physiologic impairment: characteristics and outcomes. <i>Respiratory Research</i> , 2021, 22, 5.	1.4	10
1638	Monitoring and management of fibrosing interstitial lung diseases: a narrative review for practicing clinicians. <i>Therapeutic Advances in Respiratory Disease</i> , 2021, 15, 175346662110397.	1.0	11
1639	Idiopathic pulmonary fibrosis "A challenging disease with new horizons and changing therapeutic landscape. <i>Clinical Respiratory Journal</i> , 2021, 15, 369-373.	0.6	0
1640	Establishing content-validity of a disease-specific health-related quality of life instrument for patients with chronic hypersensitivity pneumonitis. <i>Journal of Patient-Reported Outcomes</i> , 2021, 5, 9.	0.9	4
1641	Fibrosis on a Chip for Screening of Anti-Fibrosis Drugs. <i>Methods in Molecular Biology</i> , 2021, 2299, 263-274.	0.4	3
1642	Fibrotic interstitial lung disease occurring as sequelae of COVID-19 pneumonia despite concomitant steroids. <i>Lung India</i> , 2021, 38, 61.	0.3	15

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1644	Weiterentwicklung in der Therapie rheumatischer Erkrankungen bei Kindern und Jugendlichen. <i>Springer Reference Medizin</i> , 2021, , 1-19.	0.0	0
1645	Interstitial lung abnormality (ILA) and nonspecific interstitial pneumonia (NSIP). <i>European Journal of Radiology Open</i> , 2021, 8, 100336.	0.7	8
1646	Inhibition of TGF β 2 improves hematopoietic stem cell niche and ameliorates cancer-related anemia. <i>Stem Cell Research and Therapy</i> , 2021, 12, 65.	2.4	6
1647	ERS International Congress 2020: highlights from the Thoracic Surgery and Transplantation Assembly. <i>ERJ Open Research</i> , 2021, 7, 00743-2020.	1.1	0
1649	Targeting Molecular and Cellular Mechanisms in Idiopathic Pulmonary Fibrosis. , 2021, , 287-310.		0
1651	Hesperetin attenuates silica-induced lung injury by reducing oxidative damage and inflammatory response. <i>Experimental and Therapeutic Medicine</i> , 2021, 21, 297.	0.8	11
1652	Mouse Models of Lung Fibrosis. <i>Methods in Molecular Biology</i> , 2021, 2299, 291-321.	0.4	8
1653	Co-delivery of siPTPN13 and siNOX4 <i>via</i> (myo)fibroblast-targeting polymeric micelles for idiopathic pulmonary fibrosis therapy. <i>Theranostics</i> , 2021, 11, 3244-3261.	4.6	14
1654	Resident Interstitial Lung Fibroblasts and their Role in Alveolar Stem Cell Niche Development, Homeostasis, Injury, and Regeneration. <i>Stem Cells Translational Medicine</i> , 2021, 10, 1021-1032.	1.6	44
1655	Use of nintedanib in interstitial lung disease other than idiopathic pulmonary fibrosis: much caution is warranted. <i>Pulmonary Pharmacology and Therapeutics</i> , 2021, 66, 101987.	1.1	6
1656	Baseline characteristics and survival of patients of idiopathic pulmonary fibrosis: a longitudinal analysis of the Swedish IPF Registry. <i>Respiratory Research</i> , 2021, 22, 40.	1.4	36
1657	Chronic Lung Allograft Dysfunction: Review of CT and Pathologic Findings. <i>Radiology: Cardiothoracic Imaging</i> , 2021, 3, e200314.	0.9	15
1658	Pulmonary rehabilitation for interstitial lung disease. <i>The Cochrane Library</i> , 2021, 2021, CD006322.	1.5	67
1659	Transcriptional and Proteomic Characterization of Telomere-Induced Senescence in a Human Alveolar Epithelial Cell Line. <i>Frontiers in Medicine</i> , 2021, 8, 600626.	1.2	8
1661	Hermansky-Pudlak syndrome pulmonary fibrosis: a rare inherited interstitial lung disease. <i>European Respiratory Review</i> , 2021, 30, 200193.	3.0	18
1662	Long non-coding RNAs: Promising new targets in pulmonary fibrosis. <i>Journal of Gene Medicine</i> , 2021, 23, e3318.	1.4	25
1663	Hermansky-Pudlak syndrome-2 alters mitochondrial homeostasis in the alveolar epithelium of the lung. <i>Respiratory Research</i> , 2021, 22, 49.	1.4	5

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1767	Reliability of histopathologic diagnosis of fibrotic interstitial lung disease: An international collaborative standardization project. <i>BMC Pulmonary Medicine</i> , 2021, 21, 184.	0.8	0
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1769	Diagnosis and Management of Fibrotic Interstitial Lung Diseases. <i>Clinics in Chest Medicine</i> , 2021, 42, 321-335.	0.8	7
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1778	Idiopathic pulmonary fibrosis complications: what a radiologist should know. <i>Journal of Radiological Review</i> , 2021, 8, .	0.1	1
1779	Mesenchymal growth hormone receptor deficiency leads to failure of alveolar progenitor cell function and severe pulmonary fibrosis. <i>Science Advances</i> , 2021, 7, .	4.7	10
1780	Clinical Features and Outcomes of Combined Pulmonary Fibrosis and Emphysema After Lung Transplantation. <i>Chest</i> , 2021, 160, 1743-1750.	0.4	12
1781	Idiopathic pulmonary fibrosis and systemic sclerosis: pathogenic mechanisms and therapeutic interventions. <i>Cellular and Molecular Life Sciences</i> , 2021, 78, 5527-5542.	2.4	22
1782	Myofibroblast fate plasticity in tissue repair and fibrosis: Deactivation, apoptosis, senescence and reprogramming. <i>Wound Repair and Regeneration</i> , 2021, 29, 678-691.	1.5	20
1783	Unclassifiable, or simply unclassified interstitial lung disease?. <i>Current Opinion in Pulmonary Medicine</i> , 2021, 27, 405-413.	1.2	5
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1785	The Role of miRNAs in Extracellular Matrix Repair and Chronic Fibrotic Lung Diseases. <i>Cells</i> , 2021, 10, 1706.	1.8	13
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1788	Anti-fibrotic treatments for chronic liver diseases: The present and the future. <i>Clinical and Molecular Hepatology</i> , 2021, 27, 413-424.	4.5	23
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1795	Questions of ethical regulation of immunobiological therapy of some occupational lung diseases. <i>Perm Medical Journal</i> , 2021, 38, 131-140.	0.0	1
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1803	Fine crackles on chest auscultation in the early diagnosis of idiopathic pulmonary fibrosis: a prospective cohort study. <i>BMJ Open Respiratory Research</i> , 2021, 8, e000815.	1.2	12
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1806	Efficacy of early antifibrotic treatment for idiopathic pulmonary fibrosis. <i>BMC Pulmonary Medicine</i> , 2021, 21, 218.	0.8	9
1807	Co-trimoxazole to reduce mortality, transplant, or unplanned hospitalisation in people with moderate to very severe idiopathic pulmonary fibrosis: the EME-TIPAC RCT. <i>Efficacy and Mechanism Evaluation</i> , 2021, 8, 1-110.	0.9	1
1808	Calycosin attenuates pulmonary fibrosis by the epithelial-mesenchymal transition repression upon inhibiting the AKT/GSK3 β / β -catenin signaling pathway. <i>Acta Histochemica</i> , 2021, 123, 151746.	0.9	14
1809	Management and support of patients with fibrosing interstitial lung diseases. <i>Nurse Practitioner</i> , 2021, 46, 39-44.	0.2	2
1810	Outcomes for hospitalized patients with idiopathic pulmonary fibrosis treated with antifibrotic medications. <i>BMC Pulmonary Medicine</i> , 2021, 21, 239.	0.8	6
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1814	Adoption of the Antifibrotic Medications Pirfenidone and Nintedanib for Patients with Idiopathic Pulmonary Fibrosis. <i>Annals of the American Thoracic Society</i> , 2021, 18, 1121-1128.	1.5	37
1815	Efficacy of lower dose pirfenidone for idiopathic pulmonary fibrosis in real practice: a retrospective cohort study. <i>Korean Journal of Internal Medicine</i> , 2022, 37, 366-376.	0.7	7
1816	Current Status of Pharmacologic and Nonpharmacologic Therapy in Heart Failure with Preserved Ejection Fraction. <i>Heart Failure Clinics</i> , 2021, 17, 463-482.	1.0	4
1817	The attitudes and practices of physicians caring for patients with rheumatoid arthritis-associated interstitial lung disease: an international survey. <i>Rheumatology</i> , 2022, 61, 1459-1467.	0.9	9
1818	Switching antifibrotics in patients with idiopathic pulmonary fibrosis: a multi-center retrospective cohort study. <i>BMC Pulmonary Medicine</i> , 2021, 21, 221.	0.8	15
1819	Impact of Antifibrotic Therapy on Mortality and Acute Exacerbation in Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2021, 160, 1751-1763.	0.4	88
1820	Patient-centered Outcomes Research in Interstitial Lung Disease: An Official American Thoracic Society Research Statement. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021, 204, e3-e23.	2.5	41
1821	Pterostilbene alleviates pulmonary fibrosis by regulating ASIC2. <i>Chinese Medicine</i> , 2021, 16, 66.	1.6	6
1822	An update on interstitial lung disease. <i>British Journal of Hospital Medicine (London, England: 2005)</i> , 2021, 82, 1-14.	0.2	2
1823	Lung transplantation for interstitial lung disease. <i>European Respiratory Review</i> , 2021, 30, 210017.	3.0	36
1824	Fibrometabolism—An emerging therapeutic frontier in pulmonary fibrosis. <i>Science Signaling</i> , 2021, 14, .	1.6	31
1826	Human bronchial epithelial cell-derived extracellular vesicle therapy for pulmonary fibrosis via inhibition of TGF β 2-WNT crosstalk. <i>Journal of Extracellular Vesicles</i> , 2021, 10, e12124.	5.5	74
1827	Clusters of comorbidities in idiopathic pulmonary fibrosis. <i>Respiratory Medicine</i> , 2021, 185, 106490.	1.3	13
1828	Protective function of interleukin-22 in pulmonary fibrosis. <i>Clinical and Translational Medicine</i> , 2021, 11, e509.	1.7	18
1829	Modeling Extracellular Matrix-Cell Interactions in Lung Repair and Chronic Disease. <i>Cells</i> , 2021, 10, 2145.	1.8	16
1830	Pharmacological Interactions of Nintedanib and Pirfenidone in Patients with Idiopathic Pulmonary Fibrosis in Times of COVID-19 Pandemic. <i>Pharmaceuticals</i> , 2021, 14, 819.	1.7	12

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1832	Riociguat for Sarcoidosis-Associated Pulmonary Hypertension. <i>Chest</i> , 2022, 161, 448-457.	0.4	24
1833	Surgical treatment for patients with idiopathic pulmonary fibrosis and lung cancer: postoperative acute exacerbation of idiopathic pulmonary fibrosis and outcomes. <i>Surgery Today</i> , 2022, 52, 736-744.	0.7	12
1834	Small Airway Reduction and Fibrosis Is an Early Pathologic Feature of Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021, 204, 1048-1059.	2.5	31
1835	Pleuroparenchymal fibroelastosis in idiopathic pulmonary fibrosis: Survival analysis using visual and computer-based computed tomography assessment. <i>EClinicalMedicine</i> , 2021, 38, 101009.	3.2	6
1836	3-Carbamoyl-proxyl nitroxide radicals attenuate bleomycin-induced pulmonary fibrosis in mice. <i>Free Radical Biology and Medicine</i> , 2021, 171, 135-142.	1.3	7
1837	Global research hotspots and research trends on idiopathic pulmonary fibrosis: a bibliometric and visualization analysis. <i>Annals of Palliative Medicine</i> , 2021, 10, 9057-9068.	0.5	9
1838	Clinical Impact of Surgical Lung Biopsy for Interstitial Lung Disease in a Reference Center. <i>Annals of Thoracic Surgery</i> , 2021, , .	0.7	0
1839	Inspiratory muscle training in interstitial lung disease: a systematic scoping review. <i>Jornal Brasileiro De Pneumologia</i> , 2021, 47, e20210089.	0.4	6
1840	An updated approach to determine minimal clinically important differences in idiopathic pulmonary fibrosis. <i>ERJ Open Research</i> , 2021, 7, 00142-2021.	1.1	4
1841	Diagnostic Accuracy of Endobronchial Optical Coherence Tomography for the Microscopic Diagnosis of Usual Interstitial Pneumonia. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021, 204, 1164-1179.	2.5	32
1842	Serial Measurements of Circulating KL-6, SP-D, MMP-7, CA19-9, CA-125, CCL18, and Periostin in Patients with Idiopathic Pulmonary Fibrosis Receiving Antifibrotic Therapy: An Exploratory Study. <i>Journal of Clinical Medicine</i> , 2021, 10, 3864.	1.0	14
1843	Novel Artificial Intelligence-based Technology for Chest Computed Tomography Analysis of Idiopathic Pulmonary Fibrosis. <i>Annals of the American Thoracic Society</i> , 2022, 19, 399-406.	1.5	29
1844	COPD, Pulmonary Fibrosis and ILAs in Aging Smokers: The Paradox of Striking Different Responses to the Major Risk Factors. <i>International Journal of Molecular Sciences</i> , 2021, 22, 9292.	1.8	14
1845	Identification of Hub Genes and Pathways Associated With Idiopathic Pulmonary Fibrosis via Bioinformatics Analysis. <i>Frontiers in Molecular Biosciences</i> , 2021, 8, 711239.	1.6	21
1846	Pulmonary rehabilitation in patients with interstitial lung diseases: Correlates of success. <i>Respiratory Medicine</i> , 2021, 185, 106473.	1.3	7
1847	Pirfenidone in heart failure with preserved ejection fraction: a randomized phase 2 trial. <i>Nature Medicine</i> , 2021, 27, 1477-1482.	15.2	92
1848	Diagnostic and prognostic implications of 2018 guideline for the diagnosis of idiopathic pulmonary fibrosis in clinical practice. <i>Scientific Reports</i> , 2021, 11, 16481.	1.6	8

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1850	Cost drivers in the pharmacological treatment of interstitial lung disease. <i>Respiratory Research</i> , 2021, 22, 218.	1.4	5
1851	Endothelial-Specific Loss of Sphingosine-1-Phosphate Receptor 1 Increases Vascular Permeability and Exacerbates Bleomycin-induced Pulmonary Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2022, 66, 38-52.	1.4	21
1852	Interstitial lung disease and obstructive sleep apnea. <i>Sleep Medicine Reviews</i> , 2021, 58, 101442.	3.8	22
1853	A phase 1, randomized study to evaluate safety, tolerability, and pharmacokinetics of GDC-3280, a potential novel anti-fibrotic small molecule, in healthy subjects. <i>Pulmonary Pharmacology and Therapeutics</i> , 2021, 69, 102051.	1.1	2
1855	The impact of SARS-COV2 pandemic on the management of IPF patients: Our narrative experience. <i>Pulmonary Pharmacology and Therapeutics</i> , 2021, 69, 102038.	1.1	2
1856	Physical activity decline is disproportionate to decline in pulmonary physiology in IPF. <i>Respirology</i> , 2021, 26, 1152-1159.	1.3	6
1857	Targeting Cpt1a-Bcl-2 interaction modulates apoptosis resistance and fibrotic remodeling. <i>Cell Death and Differentiation</i> , 2022, 29, 118-132.	5.0	18
1858	Spatially Resolved Identification of Transglutaminase Substrates by Proteomics in Pulmonary Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2021, 65, 319-330.	1.4	7
1859	Myeloperoxidase and associated lung disease: Review of the latest developments. <i>International Journal of Rheumatic Diseases</i> , 2021, 24, 1460-1466.	0.9	8
1860	Computed Tomography Findings Suggestive of Connective Tissue Disease in the Setting of Usual Interstitial Pneumonia. <i>Journal of Computer Assisted Tomography</i> , 2021, 45, 776-781.	0.5	5
1861	Effect of antifibrotic therapy in patients with idiopathic pulmonary fibrosis undergoing lung transplant in the peri and post-operative period. <i>Respiratory Medicine</i> , 2021, 190, 106599.	1.3	5
1862	Design of phase 2 study of TAS-115, a novel oral multi-kinase inhibitor, in patients with idiopathic pulmonary fibrosis. <i>Contemporary Clinical Trials Communications</i> , 2021, 23, 100832.	0.5	2
1863	The effect of heated humidified nasal high flow oxygen supply on exercise tolerance in patients with interstitial lung disease: A pilot study. <i>Respiratory Medicine</i> , 2021, 186, 106523.	1.3	5
1864	Progression of Idiopathic Pulmonary Fibrosis Is Associated with Silica/Silicate Inhalation. <i>Environmental Science and Technology Letters</i> , 2021, 8, 903-910.	3.9	8
1865	A systematic review of blood biomarkers with individual participant data meta-analysis of matrix metalloproteinase-7 in idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2022, 59, 2101612.	3.1	20
1866	Special considerations for pulmonary rehabilitation in conditions other than COPD. , 2021, , 145-164.		3
1867	Therapeutically Targeting Cancers That Overexpress FOXC1: A Transcriptional Driver of Cell Plasticity, Partial EMT, and Cancer Metastasis. <i>Frontiers in Oncology</i> , 2021, 11, 721959.	1.3	15

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1868	A New Tool to Assess Quality of Life in Patients with Idiopathic Pulmonary Fibrosis or Non-specific Interstitial Pneumonia. <i>Pneumologie</i> , 2022, 76, 25-34.	0.1	0
1869	Progressive Fibrosing Interstitial Lung Diseases: A Current Perspective. <i>Biomedicines</i> , 2021, 9, 1237.	1.4	10
1870	Clinical Utility of Home versus Hospital Spirometry in Fibrotic Interstitial Lung Disease: Evaluation after INJUSTIS Interim Analysis. <i>Annals of the American Thoracic Society</i> , 2022, 19, 506-509.	1.5	12
1871	Lung cancer and interstitial lung diseases: the lack of prognostic impact of lung cancer in IPF. <i>Internal and Emergency Medicine</i> , 2022, 17, 457-464.	1.0	3
1872	Effect of sildenafil added to antifibrotic treatment in idiopathic pulmonary fibrosis. <i>Scientific Reports</i> , 2021, 11, 17824.	1.6	5
1873	Early diagnosis of fibrotic interstitial lung disease: challenges and opportunities. <i>Lancet Respiratory Medicine</i> , 2021, 9, 1065-1076.	5.2	55
1874	GED-0507 attenuates lung fibrosis by counteracting myofibroblast transdifferentiation in vivo and in vitro. <i>PLoS ONE</i> , 2021, 16, e0257281.	1.1	5
1875	Therapeutic targets in lung tissue remodelling and fibrosis. , 2021, 225, 107839.		98
1876	Comparison of Palliative Care Models in Idiopathic Pulmonary Fibrosis. <i>Applied Sciences (Switzerland)</i> , 2021, 11, 9028.	1.3	2
1877	Cross-species integration of single-cell RNA-seq resolved alveolar-epithelial transitional states in idiopathic pulmonary fibrosis. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2021, 321, L491-L506.	1.3	13
1879	Combined assessment of the GAP index and body mass index at antifibrotic therapy initiation for prognosis of idiopathic pulmonary fibrosis. <i>Scientific Reports</i> , 2021, 11, 18579.	1.6	4
1880	Global and regional burden of interstitial lung disease and pulmonary sarcoidosis from 1990 to 2019: results from the Global Burden of Disease study 2019. <i>Thorax</i> , 2022, 77, 596-605.	2.7	13
1881	Therapeutic Approaches to Systemic Sclerosis: Recent Approvals and Future Candidate Therapies. <i>Clinical Reviews in Allergy and Immunology</i> , 2023, 64, 239-261.	2.9	20
1882	An IPF-like disease course in disorders other than IPF: how can this be anticipated, recognized, and managed?. <i>Expert Review of Clinical Immunology</i> , 2021, 17, 1091-1101.	1.3	4
1883	The Aggregate Index of Systemic Inflammation (AISI): A Novel Prognostic Biomarker in Idiopathic Pulmonary Fibrosis. <i>Journal of Clinical Medicine</i> , 2021, 10, 4134.	1.0	32
1884	Associations of sleep duration and sleep-wake rhythm with lung parenchymal abnormalities on computed tomography: TheAMESA study. <i>Journal of Sleep Research</i> , 2022, 31, e13475.	1.7	5
1885	Pulmonary fibrosis in dyskeratosis congenita: a case report with a PRISMA-compliant systematic review. <i>BMC Pulmonary Medicine</i> , 2021, 21, 279.	0.8	10
1886	Interstitial Lung Diseases and the Impact of Gender. <i>Clinics in Chest Medicine</i> , 2021, 42, 531-541.	0.8	5

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1887	From ARDS to pulmonary fibrosis: the next phase of the COVID-19 pandemic?. <i>Translational Research</i> , 2022, 241, 13-24.	2.2	68
1888	Ageing mechanisms that contribute to tissue remodeling in lung disease. <i>Ageing Research Reviews</i> , 2021, 70, 101405.	5.0	22
1889	Antifibrotics in systemic sclerosis. <i>Best Practice and Research in Clinical Rheumatology</i> , 2021, 35, 101671.	1.4	4
1890	Development of antifibrotic therapy for stricturing Crohn's disease: lessons from randomized trials in other fibrotic diseases. <i>Physiological Reviews</i> , 2022, 102, 605-652.	13.1	31
1891	Treatment of fibrotic interstitial lung disease: current approaches and future directions. <i>Lancet</i> , The, 2021, 398, 1450-1460.	6.3	47
1892	Improved Radiolytic Stability of a ⁶⁸ Ga-labelled Collagelin Analogue for the Imaging of Fibrosis. <i>Pharmaceuticals</i> , 2021, 14, 990.	1.7	3
1893	Impact of smoking on the development of idiopathic pulmonary fibrosis: results from a nationwide population-based cohort study. <i>Thorax</i> , 2022, 77, 470-476.	2.7	15
1894	Synthesis and biological evaluation of selenogefitinib for reducing bleomycin-induced pulmonary fibrosis. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 48, 128238.	1.0	2
1895	The effect of additional antimicrobial therapy on the outcomes of patients with idiopathic pulmonary fibrosis: a systematic review and meta-analysis. <i>Respiratory Research</i> , 2021, 22, 243.	1.4	2
1896	Diagnosis Yield and Safety of Surgical Biopsy in Interstitial Lung Diseases: A Prospective Study. <i>Annals of Thoracic Surgery</i> , 2022, 114, 1911-1917.	0.7	2
1897	Targeting Alveolar Repair in Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2021, 65, 347-365.	1.4	29
1898	Ion therapy of pulmonary fibrosis by inhalation of ionic solution derived from silicate bioceramics. <i>Bioactive Materials</i> , 2021, 6, 3194-3206.	8.6	15
1899	Hyperpolarized ¹²⁹ Xe MRI and Spectroscopy of Gas-Exchange Abnormalities in Nonspecific Interstitial Pneumonia. <i>Radiology</i> , 2021, 301, 211-220.	3.6	11
1900	Mesoporous silica nanoparticles for pulmonary drug delivery. <i>Advanced Drug Delivery Reviews</i> , 2021, 177, 113953.	6.6	64
1901	Beyond TGF β 1 - novel treatment strategies targeting lung fibrosis. <i>International Journal of Biochemistry and Cell Biology</i> , 2021, 141, 106090.	1.2	2
1902	Molecular Signatures of Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2021, 65, 430-441.	1.4	23
1903	Tetraethylthiuram disulphide alleviates pulmonary fibrosis through modulating transforming growth factor- β signalling. <i>Pharmacological Research</i> , 2021, 174, 105923.	3.1	3
1904	Incidence of acute exacerbation of idiopathic pulmonary fibrosis in patients receiving antifibrotic agents: Real-world experience. <i>Respiratory Medicine</i> , 2021, 187, 106551.	1.3	8

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1906	Interstitial Lung Abnormalities: State of the Art. <i>Radiology</i> , 2021, 301, 19-34.	3.6	63
1907	Mortality differences in pulmonary fibrosis among the most populated states in the United States. <i>Respiratory Medicine</i> , 2021, 187, 106565.	1.3	0
1908	The Role of Surgical Lung Biopsy in the Diagnosis of Fibrotic Interstitial Lung Disease: Perspective from the Pulmonary Fibrosis Foundation. <i>Annals of the American Thoracic Society</i> , 2021, 18, 1601-1609.	1.5	8
1909	Pirfenidone is a renal protective drug: Mechanisms, signalling pathways, and preclinical evidence. <i>European Journal of Pharmacology</i> , 2021, 911, 174503.	1.7	16
1910	Icariin attenuates bleomycin-induced pulmonary fibrosis by targeting Hippo/YAP pathway. <i>Biomedicine and Pharmacotherapy</i> , 2021, 143, 112152.	2.5	16
1911	The phosphodiesterase 4 inhibitor AA6216 suppresses activity of fibrosis-specific macrophages. <i>Biochemistry and Biophysics Reports</i> , 2021, 28, 101118.	0.7	1
1912	Scleroderma Associated Interstitial Lung Disease. , 2022, , 319-325.		0
1913	Rheumatoid Arthritis Interstitial Lung Disease. , 2022, , 307-318.		0
1914	Interstitial Pneumonia With Autoimmune Features. , 2022, , 298-306.		1
1916	A Comprehensive Guide to Lung Transplantation for the Recipient With Pulmonary Fibrosis. , 2022, , 661-675.		0
1917	Other Idiopathic Interstitial Pneumonias and Unclassifiable Interstitial Lung Disease. , 2022, , 257-274.		0
1918	Comprehensive Care of Interstitial Lung Disease. , 2022, , 64-78.		0
1919	Idiopathic Pulmonary Fibrosis-Treatment and Management. , 2022, , 218-233.		0
1920	Common Co-Morbidities in Fibrosing Interstitial Lung Disease. , 2022, , 79-87.		0
1921	Palliative Care in Interstitial Lung Disease. <i>Respiratory Medicine</i> , 2021, , 189-207.	0.1	0
1922	An Introduction to Advanced Lung Disease. <i>Respiratory Medicine</i> , 2021, , 11-25.	0.1	0
1923	<i>Hirsutiella sinensis</i> mycelium regulates autophagy of alveolar macrophages via TLR4/NF- κ B signaling pathway. <i>International Journal of Medical Sciences</i> , 2021, 18, 1810-1823.	1.1	7

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1925	Real-Time Evaluation of Hydrogen Peroxide Injuries in Pulmonary Fibrosis Mice Models with a Mitochondria-Targeted Near-Infrared Fluorescent Probe. ACS Sensors, 2021, 6, 1228-1239.	4.0	70
1926	Pulmonary and Critical Care Medicine. , 2021, , 325-338.		0
1927	Post-COVID lung fibrosis: The tsunami that will follow the earthquake. Lung India, 2021, 38, 41.	0.3	69
1928	Immune dysregulation as a driver of idiopathic pulmonary fibrosis. Journal of Clinical Investigation, 2021, 131, .	3.9	114
1929	Molecular Imaging of Fibrosis. , 2021, , 1447-1468.		0
1930	S2K Guideline for Diagnosis of Idiopathic Pulmonary Fibrosis. Respiration, 2021, 100, 238-271.	1.2	19
1931	Antifibrotic drugs in connective tissue disease-related interstitial lung disease (CTD-ILD): from mechanistic insights to therapeutic applications. Drugs in Context, 2021, 10, 1-13.	1.0	5
1933	The Promise (and Pitfalls) of Administrative Data for Idiopathic Pulmonary Fibrosis. Chest, 2021, 159, 9-10.	0.4	0
1934	Controllable release of pirfenidone by polyvinyl alcohol film embedded soft contact lenses <i>in vitro</i> and <i>in vivo</i> . Drug Delivery, 2021, 28, 634-641.	2.5	8
1935	Post-COVID-19 pulmonary fibrosis: A case series and review of literature. Journal of Family Medicine and Primary Care, 2021, 10, 2028.	0.3	11
1936	Combined Pulmonary Fibrosis and Emphysema (CPFE) Clinical Features and Management. International Journal of COPD, 2021, Volume 16, 167-177.	0.9	16
1937	Pulmonary Fibrosis Progression Prediction Using Image Processing and Machine Learning. Advances in Science, Technology and Innovation, 2021, , 159-177.	0.2	1
1938	Monocyte Count as a Prognostic Biomarker in Patients with Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2021, 204, 74-81.	2.5	107
1939	Genetic Risk Factors for Idiopathic Pulmonary Fibrosis: Insights into Immunopathogenesis. Journal of Inflammation Research, 2020, Volume 13, 1305-1318.	1.6	29
1940	Potential treatment for vocal fold scar with pirfenidone. Laryngoscope, 2018, 128, E171-E177.	1.1	10
1941	p38 Gamma MAPK. , 2016, , 1-11.		1
1942	Stem Cell Based Therapy for Lung Disease Preclinical evidence for the role of stem/stromal cells Clinical application of stem/stromal cells in lung fibrosis. , 2019, , 119-130.		1

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1944	Mesenchymal Stromal Cell-Based Therapies for Lung Disease. <i>Pancreatic Islet Biology</i> , 2015, , 225-242.	0.1	1
1945	p38 Gamma MAPK. , 2018, , 3718-3727.		1
1946	Risk Factors and Biomarkers of RA-ILD. <i>Respiratory Medicine</i> , 2018, , 59-72.	0.1	3
1947	Clinical Trials in IPF: What Are the Best Endpoints?. <i>Respiratory Medicine</i> , 2019, , 433-453.	0.1	6
1948	Pharmacotherapy of IPF Using Antifibrotic Compounds. , 2016, , 147-159.		1
1949	Circulating Plasma Biomarkers of Survival in Antifibrotic-Treated Patients With Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2020, 158, 1526-1534.	0.4	31
1950	CPAP Adherence, Mortality, and Progression-Free Survival in Interstitial Lung Disease and OSA. <i>Chest</i> , 2020, 158, 1701-1712.	0.4	19
1951	Evolution and treatment of idiopathic pulmonary fibrosis. <i>Presse Medicale</i> , 2020, 49, 104025.	0.8	15
1952	Self DNA perpetuates IPF lung fibroblast senescence in a cGAS-dependent manner. <i>Clinical Science</i> , 2020, 134, 889-905.	1.8	28
1953	The K-Ras effector p38 ^β MAPK confers intrinsic resistance to tyrosine kinase inhibitors by stimulating EGFR transcription and EGFR dephosphorylation. <i>Journal of Biological Chemistry</i> , 2017, 292, 15070-15079.	1.6	10
1954	Pamrevlumab for the treatment of idiopathic pulmonary fibrosis. <i>Expert Opinion on Investigational Drugs</i> , 2020, 29, 771-777.	1.9	40
1955	Forced Vital Capacity (FVC) decline, mortality and healthcare resource utilization in idiopathic pulmonary fibrosis. <i>European Clinical Respiratory Journal</i> , 2020, 7, 1702618.	0.7	16
1956	Telomerase treatment prevents lung profibrotic pathologies associated with physiological aging. <i>Journal of Cell Biology</i> , 2020, 219, .	2.3	36
1957	New Developments in Imaging Idiopathic Pulmonary Fibrosis With Hyperpolarized Xenon Magnetic Resonance Imaging. <i>Journal of Thoracic Imaging</i> , 2019, 34, 136-150.	0.8	43
1963	Safety and effectiveness of pirfenidone combined with carboplatin-based chemotherapy in patients with idiopathic pulmonary fibrosis and non-small cell lung cancer: A retrospective cohort study. <i>Thoracic Cancer</i> , 2020, 11, 3317-3325.	0.8	20
1964	Tissue-resident mesenchymal stromal cells: Implications for tissue-specific antifibrotic therapies. <i>Science Translational Medicine</i> , 2018, 10, .	5.8	95
1965	Ambulatory oxygen for treatment of exertional hypoxaemia in pulmonary fibrosis (PFOX trial): a randomised controlled trial. <i>BMJ Open</i> , 2020, 10, e040798.	0.8	9

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1967	Impact of an interstitial lung disease service in the diagnosis and management of interstitial lung disease in Singapore. <i>Singapore Medical Journal</i> , 2020, 61, 302-307.	0.3	6
1968	Combination Therapy and the Start of a New Epoch for Idiopathic Pulmonary Fibrosis?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018, 197, 283-284.	2.5	7
1969	Interstitial Lung Disease in Relatives of Patients with Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 201, 1240-1248.	2.5	65
1970	CCR10+ epithelial cells from idiopathic pulmonary fibrosis lungs drive remodeling. <i>JCI Insight</i> , 2018, 3, .	2.3	30
1971	Fibulin-1c regulates transforming growth factor α 2 activation in pulmonary tissue fibrosis. <i>JCI Insight</i> , 2019, 4, .	2.3	42
1972	TAZ is required for lung alveolar epithelial cell differentiation after injury. <i>JCI Insight</i> , 2019, 4, .	2.3	54
1973	Soluble Thy-1 reverses lung fibrosis via its integrin-binding motif. <i>JCI Insight</i> , 2019, 4, .	2.3	20
1974	Transcriptional regulatory model of fibrosis progression in the human lung. <i>JCI Insight</i> , 2019, 4, .	2.3	113
1975	Brd4-p300 inhibition downregulates Nox4 and accelerates lung fibrosis resolution in aged mice. <i>JCI Insight</i> , 2020, 5, .	2.3	45
1976	Vaccinia vaccine-based immunotherapy arrests and reverses established pulmonary fibrosis. <i>JCI Insight</i> , 2016, 1, e83116.	2.3	22
1977	Attenuation of lung fibrosis in mice with a clinically relevant inhibitor of glutathione-S-transferase α . <i>JCI Insight</i> , 2016, 1, .	2.3	32
1978	The antifibrotic drug pirfenidone promotes pulmonary cavitation and drug resistance in a mouse model of chronic tuberculosis. <i>JCI Insight</i> , 2016, 1, e86017.	2.3	10
1979	Three-dimensional characterization of fibroblast foci in idiopathic pulmonary fibrosis. <i>JCI Insight</i> , 2016, 1, .	2.3	73
1980	Uncoupling of the profibrotic and hemostatic effects of thrombin in lung fibrosis. <i>JCI Insight</i> , 2017, 2, .	2.3	67
1981	Cannabinoid CB1 receptor overactivity contributes to the pathogenesis of idiopathic pulmonary fibrosis. <i>JCI Insight</i> , 2017, 2, .	2.3	59
1982	Molecular imaging of fibrosis: recent advances and future directions. <i>Journal of Clinical Investigation</i> , 2019, 129, 24-33.	3.9	86
1983	Immunometabolism of pro-repair cells. <i>Journal of Clinical Investigation</i> , 2019, 129, 2597-2607.	3.9	30

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1985	Contributions of alveolar epithelial cell quality control to pulmonary fibrosis. <i>Journal of Clinical Investigation</i> , 2020, 130, 5088-5099.	3.9	135
1986	FVC variability in patients with idiopathic pulmonary fibrosis and role of 6-min walk test to predict further change. <i>European Respiratory Journal</i> , 2020, 55, 1902151.	3.1	19
1987	The therapy of idiopathic pulmonary fibrosis: what is next?. <i>European Respiratory Review</i> , 2019, 28, 190021.	3.0	157
1988	A feasibility, randomised controlled trial of a complex breathlessness intervention in idiopathic pulmonary fibrosis (BREEZE-IPF): study protocol. <i>ERJ Open Research</i> , 2019, 5, 00186-2019.	1.1	3
1989	Idiopathic pulmonary fibrosis in the UK: analysis of the British Thoracic Society electronic registry between 2013 and 2019. <i>ERJ Open Research</i> , 2021, 7, 00187-2020.	1.1	17
1991	Pharmacological management. , 0, , 196-217.		1
1992	Progression of fibrosing interstitial lung disease. <i>Respiratory Research</i> , 2020, 21, 32.	1.4	94
1993	Patient and site characteristics associated with pirfenidone and nintedanib use in the United States; an analysis of idiopathic pulmonary fibrosis patients enrolled in the Pulmonary Fibrosis Foundation Patient Registry. <i>Respiratory Research</i> , 2020, 21, 48.	1.4	31
1994	Application of Isobaric Tags for Relative and Absolute Quantification (iTRAQ) Coupled with Two-Dimensional Liquid Chromatography/Tandem Mass Spectrometry in Quantitative Proteomic Analysis for Discovery of Serum Biomarkers for Idiopathic Pulmonary Fibrosis. <i>Medical Science Monitor</i> , 2018, 24, 4146-4153.	0.5	14
1995	Recent advances in managing idiopathic pulmonary fibrosis. <i>F1000Research</i> , 2017, 6, 2052.	0.8	12
1996	BRD4 inhibition for the treatment of pathological organ fibrosis. <i>F1000Research</i> , 2017, 6, 1015.	0.8	47
1997	Developing better drugs for pulmonary sarcoidosis: determining indications for treatment and endpoints to assess therapy based on patient and clinician concerns. <i>F1000Research</i> , 2019, 8, 2149.	0.8	8
1998	Therapeutic potential of targeting kinase inhibition in patients with idiopathic pulmonary fibrosis. <i>Yeungnam University Journal of Medicine</i> , 2020, 37, 269-276.	0.7	4
1999	The hospitalized patient with interstitial lung disease: a hospitalist primer. <i>Journal of Hospital Medicine</i> , 2017, 12, 580-584.	0.7	1
2000	Promising new treatment targets in patients with fibrosing lung disorders. <i>World Journal of Clinical Cases</i> , 2014, 2, 668.	0.3	4
2001	A Critical Role for the mTORC2 Pathway in Lung Fibrosis. <i>PLoS ONE</i> , 2014, 9, e106155.	1.1	49
2002	Validation of the 2nd Generation Proteasome Inhibitor Oprozomib for Local Therapy of Pulmonary Fibrosis. <i>PLoS ONE</i> , 2015, 10, e0136188.	1.1	11

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2004	Immune Inflammation and Disease Progression in Idiopathic Pulmonary Fibrosis. <i>PLoS ONE</i> , 2016, 11, e0154516.	1.1	87
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2028	Effectiveness of support groups to improve the quality of life of people with idiopathic pulmonary fibrosis a pre-post test pilot study. <i>Acta Biomedica</i> , 2017, 88, 5-12.	0.2	10
2029	Lung Disease in Rheumatic Disorders. <i>Mediterranean Journal of Rheumatology</i> , 2019, 30, 147.	0.3	4
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2031	Lung Ultrasound Imaging, a Technical Review. <i>Applied Sciences (Switzerland)</i> , 2020, 10, 462.	1.3	34
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2073	Interstitial Lung Diseases in Small Lung Biopsies. , 2015, , 39-49.		0
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2078	An image case report of a complex pirfenidone skin rash in a patient with idiopathic pulmonary fibrosis. International Journal of Case Reports and Images, 2015, 6, 758.	0.0	1
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2093	Current pharmacotherapy of idiopathic pulmonary fibrosis. Klinicka Farmakologie A Farmacie, 2016, 30, 30-34.	0.1	0
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2113	Lung Allograft Dysfunction (LAD) and Bronchiolitis Obliterans Syndrome. , 2018, , 263-278.		0
2114	Respiratory Organ Aging and Cancer. , 2018, , 1-30.		0
2115	Gastroesophageal Reflux and Idiopathic Pulmonary Fibrosis. , 2018, , 195-204.		0
2117	Interstitielle Lungenerkrankungen. , 2018, , 268-275.		0
2118	4. Recent Progress in Diagnosis and Treatment of Interstitial Pneumonias. The Journal of the Japanese Society of Internal Medicine, 2018, 107, 505-509.	0.0	1
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2120	Chronic Lung Allograft Dysfunction: Phenotypes and the Future. , 2019, , 119-129.		0
2121	Cardiovascular risk in Idiopathic Pulmonary Fibrosis. International Journal of Pulmonary & Respiratory Sciences, 2018, 3, .	0.1	0
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2124	Future Directions for IPF Research. Respiratory Medicine, 2019, , 455-467.	0.1	0
2125	The Role of Pulmonary Rehabilitation and Supplemental Oxygen Therapy in the Treatment of Patients with Idiopathic Pulmonary Fibrosis. Respiratory Medicine, 2019, , 389-399.	0.1	0
2126	Pharmacologic Treatment of IPF. Respiratory Medicine, 2019, , 325-364.	0.1	1
2128	Gastroesophageal Reflux and IPF. Respiratory Medicine, 2019, , 379-387.	0.1	0
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2132	Hermansky-Pudlak syndrome with interstitial lung disease: A holistically worked up couplet. Lung India, 2019, 36, 345.	0.3	1
2134	Complications of Transbronchial Cryobiopsy. , 2019, , 59-65.		0
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2149	Lung transplantation for idiopathic pulmonary fibrosis. Presse Medicale, 2020, 49, 104026.	0.8	9
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2153	Idiopathic Interstitial Pneumonias: A Review of the Past and Emerging Therapies. <i>Journal of Pharmacy Practice</i> , 2021, , 089719002110532.	0.5	0
2154	Antifibrotic drugs for idiopathic pulmonary fibrosis: What we should know?. <i>Indian Journal of Medical Research</i> , 2020, 152, 177.	0.4	5
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2159	Interstitial lung abnormalities – current knowledge and future directions. <i>European Clinical Respiratory Journal</i> , 2021, 8, 1994178.	0.7	7
2160	Interstitial Lung Disease. , 2020, , 93-131.		0
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2164	Pharmacological treatment of idiopathic pulmonary fibrosis: time to step out of the comfort zone?. <i>Jornal Brasileiro De Pneumologia</i> , 2020, 46, e20200193-e20200193.	0.4	0
2165	Use of biologics in the treatment of asthma, COPD, ACOS, and idiopathic pulmonary fibrosis. , 2020, , 97-115.		0
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2168	Antifibrotic Therapy for Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2021, 160, 1589-1591.	0.4	1
2169	Impact of Lung Biopsy Information on Treatment Strategy of Patients with Interstitial Lung Diseases. <i>Annals of the American Thoracic Society</i> , 2022, 19, 737-745.	1.5	9
2170	Acute exacerbations of interstitial lung disease. , 0, , 117-131.		0
2171	IPF: treatment and prevention of pulmonary exacerbations. , 0, , 199-223.		0
2174	The evaluation of disease severity/staging for prognosis. , 0, , 97-105.		1

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2175	Acute exacerbations. , 0 , 143-150.		1
2176	CPFE: distinctive and non-distinctive features. , 0 , 175-185.		1
2177	Symptom management: dyspnoea and cough. , 0 , 218-229.		0
2178	Key ongoing issues in trial design. , 0 , 253-259.		0
2179	Perspectives for the future. , 0 , 260-274.		1
2182	IPF: definition, severity and impact of pulmonary exacerbations. , 0 , 58-65.		0
2183	Coexistent COPD and ILD. , 0 , 109-120.		1
2184	Patterns of cardiopulmonary response to exercise in fibrotic ILD. , 0 , 128-145.		0
2185	Multiple Choice Questions with explanations. , 0 , 1-544.		0
2186	Question 13. , 0 , 25-26.		0
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2190	Interstitial lung diseases with progressive pulmonary fibrosis: pathogenetic features and approaches to therapy. <i>Meditinskiy Sovet</i> , 2020, , 99-106.	0.1	1
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2192	Efficacy study of edaravone and acetylcysteine towards bleomycin-induced rat pulmonary fibrosis. <i>International Journal of Clinical and Experimental Medicine</i> , 2015, 8, 8730-9.	1.3	4
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2194	CCAAT/enhancer binding protein delta (C/EBP δ) deficiency does not affect bleomycin-induced pulmonary fibrosis. <i>Journal of Clinical and Translational Research</i> , 2018, 3, 358-365.	0.3	3
2195	Metformin ameliorates bleomycin-induced pulmonary fibrosis in mice by suppressing IGF-1. <i>American Journal of Translational Research (discontinued)</i> , 2020, 12, 940-949.	0.0	9
2196	Combined pirfenidone, azithromycin and prednisolone in post-H1N1 ARDS pulmonary fibrosis. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2018, 35, 85-90.	0.2	13

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2199	Pirfenidone for primary Sjögren's syndrome-related fibrotic interstitial pneumonia. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2017, 34, 91-96.	0.2	2
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2201	Identification of transcriptomic markers for developing idiopathic pulmonary fibrosis: an integrative analysis of gene expression profiles. <i>International Journal of Clinical and Experimental Pathology</i> , 2020, 13, 1698-1706.	0.5	4
2202	Management of interstitial lung diseases: A consensus statement of the Indian Chest Society (ICS) and National College of Chest Physicians (NCCP). <i>Lung India</i> , 2020, 37, 359-378.	0.3	2
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2204	A real-world study of the dosing and tolerability of pirfenidone and its effect on survival in idiopathic pulmonary fibrosis. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2020, 37, 148-157.	0.2	7
2205	Real world experience of response to pirfenidone in patients with idiopathic pulmonary fibrosis: a two centre retrospective study. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2020, 37, 218-224.	0.2	5
2206	Thoracic Ultrasound in Idiopathic Pulmonary Fibrosis Evolution (TOUPIE): research protocol of a multicentric prospective study. <i>BMJ Open</i> , 2021, 11, e039078.	0.8	0
2207	Does 1-minute walk test predict results of 6-minute walk test in patients with idiopathic pulmonary fibrosis?. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2021, 38, e2021005.	0.2	2
2209	The Extent of Inflammatory Cell Infiltrate and Fibrosis in Lungs of Telomere- and Surfactant-Related Familial Pulmonary Fibrosis. <i>Frontiers in Medicine</i> , 2021, 8, 736485.	1.2	0
2210	The clinical relevance of lymphocyte to monocyte ratio in patients with Idiopathic Pulmonary Fibrosis (IPF). <i>Respiratory Medicine</i> , 2022, 191, 106686.	1.3	4
2211	Ageing Immune System and Its Correlation With Liability to Severe Lung Complications. <i>Frontiers in Public Health</i> , 2021, 9, 735151.	1.3	12
2212	Tetrandrine Modulates Rheb-mTOR Signaling-Mediated Selective Autophagy and Protects Pulmonary Fibrosis. <i>Frontiers in Pharmacology</i> , 2021, 12, 739220.	1.6	9
2213	Neighborhood-Level Disadvantage Impacts on Patients with Fibrotic Interstitial Lung Disease. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2022, 205, 459-467.	2.5	25
2214	Longitudinal serological assessment of type VI collagen turnover is related to progression in a real-world cohort of idiopathic pulmonary fibrosis. <i>BMC Pulmonary Medicine</i> , 2021, 21, 382.	0.8	5
2215	Pirfenidone increases IL-10 and improves acute pancreatitis in multiple clinically relevant murine models. <i>JCI Insight</i> , 2022, 7, .	2.3	10

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2217	Impact of concomitant idiopathic pulmonary fibrosis on prognosis in lung cancer patients: A meta-analysis. <i>PLoS ONE</i> , 2021, 16, e0259784.	1.1	3
2218	Variables Associated With Response to Therapy in Patients With Interstitial Pneumonia With Autoimmune Features. <i>Journal of Clinical Rheumatology</i> , 2022, 28, 84-88.	0.5	9
2219	Interstitial lung disease on the acute take for the non-respiratory physician. <i>Clinical Medicine</i> , 2021, 21, e584-e590.	0.8	2
2220	Survival of patients with idiopathic pulmonary fibrosis and pulmonary hypertension under therapy with nintedanib or pirfenidone. <i>Internal and Emergency Medicine</i> , 2022, 17, 815-822.	1.0	3
2221	Naringenin: A Promising Therapeutic Agent against Organ Fibrosis. <i>Oxidative Medicine and Cellular Longevity</i> , 2021, 2021, 1-13.	1.9	23
2222	A Randomized Phase 1 Evaluation of Deupirfenidone, a Novel Deuterium-Containing Drug Candidate for Interstitial Lung Disease and Other Inflammatory and Fibrotic Diseases. <i>Clinical Pharmacology in Drug Development</i> , 2022, 11, 220-234.	0.8	6
2223	Advances in glioma-associated oncogene (GLI) inhibitors for cancer therapy. <i>Investigational New Drugs</i> , 2022, 40, 370-388.	1.2	10
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2225	Antifibrotic therapies reduce mortality and hospitalization among Medicare beneficiaries with idiopathic pulmonary fibrosis. <i>Journal of Managed Care & Specialty Pharmacy</i> , 2021, 27, 1724-1733.	0.5	6
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2227	Management of interstitial lung diseases: A consensus statement of the Indian Chest Society (ICS) and National College of Chest Physicians (NCCP). <i>Lung India</i> , 2020, 37, 359.	0.3	9
2228	Role of antifibrotics in the management of idiopathic inflammatory myopathy associated interstitial lung disease. <i>Therapeutic Advances in Musculoskeletal Disease</i> , 2021, 13, 1759720X2110609.	1.2	7
2229	Inhibiting TGF- β 1-Mediated Cellular Processes as an Effective Strategy for the Treatment of Pulmonary Fibrosis with Chinese Herbal Medicines. <i>The American Journal of Chinese Medicine</i> , 2021, 49, 1965-1999.	1.5	3
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2231	Differential Discontinuation Profiles between Pirfenidone and Nintedanib in Patients with Idiopathic Pulmonary Fibrosis. <i>Cells</i> , 2022, 11, 143.	1.8	13
2232	Impact of Concomitant Medication Burden on Tolerability of Disease-targeted Therapy and Survival in Interstitial Lung Disease. <i>Annals of the American Thoracic Society</i> , 2022, 19, 962-970.	1.5	5
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2237	Retrospective analysis of skin photosensitivity induced by pirfenidone. <i>Journal of Clinical Pharmacy and Therapeutics</i> , 2022, 47, 194-199.	0.7	2
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2242	Wenfei Buqi Tongluo Formula Against Bleomycin-Induced Pulmonary Fibrosis by Inhibiting TGF- β 2/Smad3 Pathway. <i>Frontiers in Pharmacology</i> , 2021, 12, 762998.	1.6	4
2243	Overexpression of bone morphogenetic protein receptor type 2 suppresses transforming growth factor β -induced profibrotic responses in lung fibroblasts. <i>Experimental Lung Research</i> , 2022, 48, 35-51.	0.5	1
2244	The use of exhaled air analysis in discriminating interstitial lung diseases: a pilot study. <i>Respiratory Research</i> , 2022, 23, 12.	1.4	10
2245	Disparities in Lung Transplant among Patients with Idiopathic Pulmonary Fibrosis: An Analysis of the IPF-PRO Registry. <i>Annals of the American Thoracic Society</i> , 2022, 19, 981-990.	1.5	14
2248	An 8-ferroptosis-related genes signature from Bronchoalveolar Lavage Fluid for prognosis in patients with idiopathic pulmonary fibrosis. <i>BMC Pulmonary Medicine</i> , 2022, 22, 15.	0.8	16
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2252	Induced pluripotent stem cells. , 2022, , 1-58.		0
2253	Long-Term Effects of COVID-19. <i>Mayo Clinic Proceedings</i> , 2022, 97, 579-599.	1.4	49

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2255	Challenges for Clinical Drug Development in Pulmonary Fibrosis. <i>Frontiers in Pharmacology</i> , 2022, 13, 823085.	1.6	20
2256	Systemic Sclerosis: From Pathophysiology to Novel Therapeutic Approaches. <i>Biomedicines</i> , 2022, 10, 163.	1.4	16
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2258	Mortality and survival in idiopathic pulmonary fibrosis: a systematic review and meta-analysis. <i>ERJ Open Research</i> , 2022, 8, 00591-2021.	1.1	25
2259	Demystifying idiopathic interstitial pneumonia: time for more etiology-focused nomenclature in interstitial lung disease. <i>Expert Review of Respiratory Medicine</i> , 2022, 16, 235-245.	1.0	2
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2262	Idiopathic pulmonary fibrosis in the practice of a family doctor. <i>Spravočnik Vraća Obćej Praktiki</i> , 2022, , 34-43.	0.0	0
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2264	Novedades diagnósticas y terapéuticas en fibrosis pulmonar progresiva. <i>Archivos De Bronconeumología</i> , 2022, , .	0.4	7
2265	CC-90001, a c-Jun N-terminal kinase (JNK) inhibitor, in patients with pulmonary fibrosis: design of a phase 2, randomised, placebo-controlled trial. <i>BMJ Open Respiratory Research</i> , 2022, 9, e001060.	1.2	17
2266	Inhibition of RUNX1 blocks the differentiation of lung fibroblasts to myofibroblasts. <i>Journal of Cellular Physiology</i> , 2022, 237, 2169-2182.	2.0	16
2267	Analysis of Forced Vital Capacity (FVC) Trajectories in Idiopathic Pulmonary Fibrosis (IPF) Identifies Four Distinct Clusters of Disease Behaviour. <i>SSRN Electronic Journal</i> , 0, , .	0.4	0
2268	The safety and tolerability of pirfenidone in Indian patients with idiopathic pulmonary fibrosis. <i>Current Medical Issues</i> , 2022, 20, 27.	0.1	0
2269	Pemafibrate Attenuates Pulmonary Fibrosis by Inhibiting Myofibroblast Differentiation. <i>SSRN Electronic Journal</i> , 0, , .	0.4	0
2270	Longitudinal and Comparative Measures of Serum Chitotriosidase and YKL-40 in Patients With Idiopathic Pulmonary Fibrosis. <i>Frontiers in Immunology</i> , 2022, 13, 760776.	2.2	11
2271	The Anti-fibrosis drug Pirfenidone modifies the immunosuppressive tumor microenvironment and prevents the progression of renal cell carcinoma by inhibiting tumor autocrine TGF-β ² . <i>Cancer Biology and Therapy</i> , 2022, 23, 150-162.	1.5	13
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2275	A single injection of CM1021, a long half-life hepatocyte growth factor mimetic, increases liver mass in mice. <i>Biochemistry and Biophysics Reports</i> , 2021, 28, 101186.	0.7	0
2276	Vardenafil Activity in Lung Fibrosis and In Vitro Synergy with Nintedanib. <i>Cells</i> , 2021, 10, 3502.	1.8	6
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2279	The role of antifibrotics in the treatment of rheumatoid arthritis-associated interstitial lung disease. <i>Therapeutic Advances in Musculoskeletal Disease</i> , 2022, 14, 1759720X2210744.	1.2	7
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2282	Usual interstitial pneumonia: a clinically significant pattern, but not the final word. <i>Modern Pathology</i> , 2022, 35, 589-593.	2.9	4
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2289	Investigation of clinical predictors of survival in idiopathic pulmonary fibrosis patients: A cohort study in Taiwan. <i>Journal of the Chinese Medical Association</i> , 2022, 85, 578-583.	0.6	2
2290	Caveolin-1-Derived Peptide Reduces ER Stress and Enhances Gelatinolytic Activity in IPF Fibroblasts. <i>International Journal of Molecular Sciences</i> , 2022, 23, 3316.	1.8	1
2291	Editorial: Mechanisms of Lung Fibrosis: Is Immunity Back in the Game?. <i>Frontiers in Immunology</i> , 2022, 13, 882979.	2.2	0
2292	Study on the Mechanism of Astragalus Polysaccharide in Treating Pulmonary Fibrosis Based on "Drug-Target-Pathway" Network. <i>Frontiers in Pharmacology</i> , 2022, 13, 865065.	1.6	13
2293	Responsiveness and meaningful change thresholds of the Living with Pulmonary Fibrosis (L-PF) questionnaire Dyspnoea and Cough scores in patients with progressive fibrosing interstitial lung diseases. <i>BMJ Open Respiratory Research</i> , 2022, 9, e001167.	1.2	6
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2297	BMP1 is not required for lung fibrosis in mice. <i>Scientific Reports</i> , 2022, 12, 5466.	1.6	3
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2302	Insights into the Pathogenesis of Pulmonary Fibrosis from Genetic Diseases. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2022, , .	1.4	6
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2305	Pre-exposure to Aerosolized Polyvalent Bacterial Lysates Protects Against Bleomycin-Induced Pulmonary Fibrosis in Mice. <i>Inflammation</i> , 2022, 45, 1692-1699.	1.7	1
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2307	Arsenic trioxide inhibits the functions of lung fibroblasts derived from patients with idiopathic pulmonary fibrosis. <i>Toxicology and Applied Pharmacology</i> , 2022, 441, 115972.	1.3	6
2308	Disparities in Rural Populations With Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2022, 162, 630-634.	0.4	10
2309	Risk Factors of Silicosis Progression: A Retrospective Cohort Study in China. <i>Frontiers in Medicine</i> , 2022, 9, 832052.	1.2	9
2311	Chronic Obstructive Pulmonary Disease Combined with Interstitial Lung Disease. <i>Tuberculosis and Respiratory Diseases</i> , 2022, 85, 122-136.	0.7	3
2312	Screening for Inhibitors of YAP Nuclear Localization Identifies Aurora Kinase A as a Modulator of Lung Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2022, , .	1.4	6
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2321	Efficacy of antifibrotic drugs, nintedanib and pirfenidone, in treatment of progressive pulmonary fibrosis in both idiopathic pulmonary fibrosis (IPF) and non-IPF: a systematic review and meta-analysis. <i>BMC Pulmonary Medicine</i> , 2021, 21, 411.	0.8	47
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2323	Validation of the risk stratification score in idiopathic pulmonary fibrosis: study protocol of a prospective, multi-centre, observational, 3-year clinical trial. <i>BMC Pulmonary Medicine</i> , 2021, 21, 396.	0.8	0
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2380	The Extent of Inflammatory Cell Infiltrate and Fibrosis in Lungs of Telomere- and Surfactant-Related Familial Pulmonary Fibrosis. Frontiers in Medicine, 2021, 8, 736485.	1.2	4
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2401	Inhalational exposures in patients with fibrotic interstitial lung disease: Presentation, pulmonary function and survival in the <scp>Canadian Registry</scp> for <scp>Pulmonary Fibrosis</scp>. Respirology, 2022, 27, 635-644.	1.3	12
2402	Cardiac Fibrosis in the Pressure Overloaded Left and Right Ventricle as a Therapeutic Target. Frontiers in Cardiovascular Medicine, 2022, 9, .	1.1	30
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2413	Orphan Drug Use in Patients With Rare Diseases: A Population-Based Cohort Study. <i>Frontiers in Pharmacology</i> , 2022, 13, .	1.6	4
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