

Efficacy and Safety of Nintedanib in Idiopathic Pulmonary Fibrosis

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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. <i>BMC Pulmonary Medicine</i> , 2014, 14, 170.	0.8	17
3	Hypoxia-sensitive pathways in inflammation-driven fibrosis. <i>American Journal of Physiology - Regulatory Integrative and Comparative Physiology</i> , 2014, 307, R1369-R1380.	0.9	40
4	Molecular Mechanism and Treatment of Viral Hepatitis-Related Liver Fibrosis. <i>International Journal of Molecular Sciences</i> , 2014, 15, 10578-10604.	1.8	60
5	Open-Access Biorepository for Idiopathic Pulmonary Fibrosis. The Way Forward. <i>Annals of the American Thoracic Society</i> , 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. <i>Journal of Applied Physiology</i> , 2014, 117, 577-585.	1.2	77
9	Novel approaches to pulmonary fibrosis. <i>Clinical Medicine</i> , 2014, 14, s45-s49.	0.8	9
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11	Improving care for patients with idiopathic pulmonary fibrosis (IPF) in the UK: a round table discussion. <i>Thorax</i> , 2014, 69, 1136-1140.	2.7	31
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15	Pulmonary Hypertension due to Fibrotic Lung Disease: Hidden Value in a Neutral Trial. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014, 190, 131-132.	2.5	3
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19	Anti-fibrotic effects of nintedanib in lung fibroblasts derived from patients with idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2014, 15, 157.	1.4	199
20	Clinical significance of epithelial mesenchymal transition (EMT) in chronic obstructive pulmonary disease (COPD): potential target for prevention of airway fibrosis and lung cancer. <i>Clinical and Translational Medicine</i> , 2014, 3, 33.	1.7	65
21	Acute exacerbations complicating interstitial lung disease. <i>Current Opinion in Pulmonary Medicine</i> , 2014, 20, 436-441.	1.2	29
22	IPF clinical trial design and endpoints. <i>Current Opinion in Pulmonary Medicine</i> , 2014, 20, 463-471.	1.2	58

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24	Idiopathic and Autoimmune Interstitial Lung Disease. , 2014, , 1105-1123.		0
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37	Review: Interstitial Lung Disease Associated With Systemic Sclerosis and Idiopathic Pulmonary Fibrosis: How Similar and Distinct?. <i>Arthritis and Rheumatology</i> , 2014, 66, 1967-1978.	2.9	162
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373	Cough in idiopathic pulmonary fibrosis. <i>European Respiratory Review</i> , 2016, 25, 278-286.	3.0	82
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474	Therapeutic advances in idiopathic pulmonary fibrosis. <i>Clinical Pharmacology and Therapeutics</i> , 2016, 99, 30-32.	2.3	6
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486	Treatment of idiopathic pulmonary fibrosis: a network meta-analysis. <i>BMC Medicine</i> , 2016, 14, 18.	2.3	79
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#	ARTICLE	IF	CITATIONS
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508	Exercise training in idiopathic pulmonary fibrosis. <i>Expert Review of Respiratory Medicine</i> , 2016, 10, 69-77.	1.0	20
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526	The effect of mTOR inhibitors on respiratory infections in lymphangioleiomyomatosis. <i>European Respiratory Review</i> , 2017, 26, 160004.	3.0	13
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537	The fibrogenic actions of lung fibroblast-derived urokinase: a potential drug target in IPF. <i>Scientific Reports</i> , 2017, 7, 41770.	1.6	26
538	Targeting coagulation factor receptors â€“ proteaseâ€activated receptors in idiopathic pulmonary fibrosis. <i>Journal of Thrombosis and Haemostasis</i> , 2017, 15, 597-607.	1.9	42
539	Plasma membrane wounding and repair in pulmonary diseases. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2017, 312, L371-L391.	1.3	34
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557	The use of pretest probability increases the value of high-resolution CT in diagnosing usual interstitial pneumonia. <i>Thorax</i> , 2017, 72, 424-429.	2.7	103
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646	Understanding and overcoming metformin gastrointestinal intolerance. <i>Diabetes, Obesity and Metabolism</i> , 2017, 19, 473-481.	2.2	141
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661	Airway Complications After Lung Transplantation. <i>Clinics in Chest Medicine</i> , 2017, 38, 693-706.	0.8	33
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682	MicroRNA-29b inhibits supernatants from silica-treated macrophages from inducing extracellular matrix synthesis in lung fibroblasts. <i>Toxicology Research</i> , 2017, 6, 878-888.	0.9	17
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690	Therapeutic targets in idiopathic pulmonary fibrosis. <i>Respiratory Medicine</i> , 2017, 131, 49-57.	1.3	92
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705	Mitochondrial Dysfunction in Pulmonary Fibrosis. <i>Annals of the American Thoracic Society</i> , 2017, 14, S383-S388.	1.5	72
706	An epithelial biomarker signature for idiopathic pulmonary fibrosis: an analysis from the multicentre PROFILE cohort study. <i>Lancet Respiratory Medicine</i> , 2017, 5, 946-955.	5.2	190
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708	Clubbing in patients with fibrotic interstitial lung diseases. <i>Respiratory Medicine</i> , 2017, 132, 226-231.	1.3	18
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1055	Pirfenidone suppresses polarization to M2 phenotype macrophages and the fibrogenic activity of rat lung fibroblasts. <i>Journal of Clinical Biochemistry and Nutrition</i> , 2018, 63, 58-65.	0.6	31
1056	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
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1073	Epithelial cell-derived cytokines CST3 and GDF15 as potential therapeutics for pulmonary fibrosis. <i>Cell Death and Disease</i> , 2018, 9, 506.	2.7	27
1074	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
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1078	Mechanisms and consequences of oxidative stress in lung disease: therapeutic implications for an aging populace. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2018, 314, L642-L653.	1.3	95
1079	Feifukang ameliorates pulmonary fibrosis by inhibiting JAK-STAT signaling pathway. <i>BMC Complementary and Alternative Medicine</i> , 2018, 18, 234.	3.7	20
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1082	Novel management strategies for idiopathic pulmonary fibrosis. <i>Expert Review of Respiratory Medicine</i> , 2018, 12, 831-842.	1.0	9
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1084	ILDgenDB: integrated genetic knowledge resource for interstitial lung diseases (ILDs). <i>Database: the Journal of Biological Databases and Curation</i> , 2018, 2018, .	1.4	2
1085	Cell Therapy in Idiopathic Pulmonary Fibrosis. <i>Medical Sciences (Basel, Switzerland)</i> , 2018, 6, 64.	1.3	17
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1087	Lung cancer in idiopathic pulmonary fibrosis: A systematic review and meta-analysis. <i>PLoS ONE</i> , 2018, 13, e0202360.	1.1	47

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1093	Clinical behaviour of patients exposed to organic dust and diagnosed with idiopathic pulmonary fibrosis. <i>Respirology</i> , 2018, 23, 1160-1165.	1.3	19
1094	Use of zonal distribution of lung crackles during inspiration and expiration to assess disease severity in idiopathic pulmonary fibrosis. <i>Postgraduate Medical Journal</i> , 2018, 94, 381-385.	0.9	5
1095	Clinical spectrum and prognostic factors of possible UIP pattern on high-resolution CT in patients who underwent surgical lung biopsy. <i>PLoS ONE</i> , 2018, 13, e0193608.	1.1	13
1096	An expert consensus to standardise definitions, diagnosis and treatment targets for anti-fibrotic stricture therapies in Crohn's disease. <i>Alimentary Pharmacology and Therapeutics</i> , 2018, 48, 347-357.	1.9	157
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1099	Assessment of Interstitial Lung Disease Using Lung Ultrasound Surface Wave Elastography. <i>Journal of Thoracic Imaging</i> , 2019, 34, 313-319.	0.8	28
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1103	Effects of emphysema on physiological and prognostic characteristics of lung function in idiopathic pulmonary fibrosis. <i>Respirology</i> , 2019, 24, 55-62.	1.3	24
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1108	Current and Emerging Drug Therapies for Connective Tissue Disease-Interstitial Lung Disease (CTD-ILD). <i>Drugs</i> , 2019, 79, 1511-1528.	4.9	35
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1112	Choice of Methodology Impacts Outcome in Indirect Comparisons of Drugs for Idiopathic Pulmonary Fibrosis. <i>Medicina (Lithuania)</i> , 2019, 55, 443.	0.8	1
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1117	Nintedanib and Sildenafil in Patients with Idiopathic Pulmonary Fibrosis and Right Heart Dysfunction. A Prespecified Subgroup Analysis of a Double-Blind Randomized Clinical Trial (INSTAGE). <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 1505-1512.	2.5	50
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1119	Mobile Health Monitoring in Patients with Idiopathic Pulmonary Fibrosis. <i>Annals of the American Thoracic Society</i> , 2019, 16, 1327-1329.	1.5	26
1120	Novel plasma peptide markers involved in the pathology of CKD identified using mass spectrometric approach. <i>Journal of Molecular Medicine</i> , 2019, 97, 1451-1463.	1.7	10
1121	Interstitial Pneumonia with Autoimmune Features. <i>Clinics in Chest Medicine</i> , 2019, 40, 609-616.	0.8	5
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1123	Gastrointestinal adverse effects of nintedanib and the associated risk factors in patients with idiopathic pulmonary fibrosis. <i>Scientific Reports</i> , 2019, 9, 12062.	1.6	45
1124	Morphological and molecular motifs of fibrosing pulmonary injury patterns. <i>Journal of Pathology: Clinical Research</i> , 2019, 5, 256-271.	1.3	16
1125	Patient journey and treatment patterns in adults with IPF based on health care data in Sweden from 2001 to 2015. <i>Respiratory Medicine</i> , 2019, 155, 72-78.	1.3	18
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1128	Longitudinal prediction of outcome in idiopathic pulmonary fibrosis using automated CT analysis. <i>European Respiratory Journal</i> , 2019, 54, 1802341.	3.1	22
1129	Outcome of lung transplantation in non-idiopathic pulmonary fibrosis interstitial lung disease. <i>Clinical Transplantation</i> , 2019, 33, e13661.	0.8	4
1130	Management of Idiopathic Pulmonary Fibrosis. <i>Annals of Pharmacotherapy</i> , 2019, 53, 1238-1248.	0.9	27
1131	Potential of nintedanib in treatment of progressive fibrosing interstitial lung diseases. <i>European Respiratory Journal</i> , 2019, 54, 1900161.	3.1	164
1133	The Role of Surgical Lung Biopsy in Antifibrotic Therapy for Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 1084-1085.	2.5	0
1134	Nintedanib ameliorates experimental pulmonary arterial hypertension via inhibition of endothelial mesenchymal transition and smooth muscle cell proliferation. <i>PLoS ONE</i> , 2019, 14, e0214697.	1.1	31
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1136	<p></p>Differences in tolerability of pirfenidone between elderly and younger patients with idiopathic pulmonary fibrosis</p>. <i>Drug Design, Development and Therapy</i> , 2019, Volume 13, 2295-2303.	2.0	13
1137	Assessing quality of life of idiopathic pulmonary fibrosis patients: the INSTAGE study. <i>Breathe</i> , 2019, 15, 144-146.	0.6	2
1138	The Role of Telomerase and Telomeres in Interstitial Lung Diseases: From Molecules to Clinical Implications. <i>International Journal of Molecular Sciences</i> , 2019, 20, 2996.	1.8	28
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1145	Real-World Comprehensive Disease Management of Patients With Idiopathic Pulmonary Fibrosis. <i>Current Respiratory Medicine Reviews</i> , 2019, 15, 4-15.	0.1	1

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1148	Epithelial Alarmins in Serum and Exhaled Breath in Patients with Idiopathic Pulmonary Fibrosis: A Prospective One-Year Follow-Up Cohort Study. <i>Journal of Clinical Medicine</i> , 2019, 8, 1590.	1.0	10
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1150	Rheumatoid Arthritis-Associated Interstitial Lung Disease: Clinical Characteristics and Predictors of Mortality. <i>Respiration</i> , 2019, 98, 455-460.	1.2	47
1151	Methotrexate-Associated Pneumonitis and Rheumatoid Arthritis-Interstitial Lung Disease: Current Concepts for the Diagnosis and Treatment. <i>Frontiers in Medicine</i> , 2019, 6, 238.	1.2	73
1152	Peripheral blood proteomic profiling of idiopathic pulmonary fibrosis biomarkers in the multicentre IPF-PRO Registry. <i>Respiratory Research</i> , 2019, 20, 227.	1.4	59
1153	Phase II study of nab-paclitaxel+carboplatin for patients with non-small cell lung cancer and interstitial lung disease. <i>Cancer Science</i> , 2019, 110, 3738-3745.	1.7	49
1154	MUC5B variant is associated with visually and quantitatively detected preclinical pulmonary fibrosis. <i>Thorax</i> , 2019, 74, 1131-1139.	2.7	43
1156	Real-life comparison of pirfenidone and nintedanib in patients with idiopathic pulmonary fibrosis: A 24-month assessment. <i>Respiratory Medicine</i> , 2019, 159, 105803.	1.3	50
1158	A prospective phase II study of carboplatin and nab-paclitaxel in patients with advanced non-small cell lung cancer and concomitant interstitial lung disease (HOT1302). <i>Lung Cancer</i> , 2019, 138, 65-71.	0.9	32
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1162	Gas6/TAM System: A Key Modulator of the Interplay between Inflammation and Fibrosis. <i>International Journal of Molecular Sciences</i> , 2019, 20, 5070.	1.8	59
1163	Idiopathic Pulmonary Fibrosis and Lung Transplantation: When it is Feasible. <i>Medicina (Lithuania)</i> , 2019, 55, 702.	0.8	16
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1165	Tackling MARCKS-PIP3 circuit attenuates fibroblast activation and fibrosis progression. <i>FASEB Journal</i> , 2019, 33, 14354-14369.	0.2	13

#	ARTICLE	IF	CITATIONS
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1167	Regulatory network of two circRNAs and an miRNA with their targeted genes under astilbin treatment in pulmonary fibrosis. <i>Journal of Cellular and Molecular Medicine</i> , 2019, 23, 6720-6729.	1.6	19
1168	Temporal echocardiographic assessment of pulmonary hypertension in idiopathic pulmonary fibrosis patients treated with nintedanib with or without oxygen therapy. <i>BMC Pulmonary Medicine</i> , 2019, 19, 157.	0.8	3
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1171	Role of dual-specificity protein phosphatase DUSP10/MKP-5 in pulmonary fibrosis. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2019, 317, L678-L689.	1.3	15
1172	Mitochondrial calcium uniporter regulates PGC-1 α expression to mediate metabolic reprogramming in pulmonary fibrosis. <i>Redox Biology</i> , 2019, 26, 101307.	3.9	56
1173	Antifibrotic therapy for idiopathic pulmonary fibrosis: time to treat. <i>Respiratory Research</i> , 2019, 20, 205.	1.4	166
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1178	In-Hospital Mortality in Patients with Idiopathic Pulmonary Fibrosis: A US Cohort Study. <i>Lung</i> , 2019, 197, 699-707.	1.4	22
1179	Bullous pemphigoid associated with nintedanib. <i>JAAD Case Reports</i> , 2019, 5, 821-823.	0.4	1
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1182	Physician characteristics associated with treatment initiation patterns in idiopathic pulmonary fibrosis. <i>Chronic Respiratory Disease</i> , 2019, 16, 147997311987967.	1.0	4
1183	Understanding Progressive Fibrosing Interstitial Lung Disease through Therapeutic Trials. <i>New England Journal of Medicine</i> , 2019, 381, 1775-1777.	13.9	9
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1186	Current and future perspectives on management of systemic sclerosis-associated interstitial lung disease. <i>Expert Review of Clinical Immunology</i> , 2019, 15, 1009-1017.	1.3	42
1187	Modelling Forced Vital Capacity in Idiopathic Pulmonary Fibrosis: Optimising Trial Design. <i>Advances in Therapy</i> , 2019, 36, 3059-3070.	1.3	4
1188	Sodium Arsenite Inhibits Lung Fibroblast Differentiation and Pulmonary Fibrosis. <i>Pharmacology</i> , 2019, 104, 368-376.	0.9	4
1189	Lung Cryobiopsy for the Diagnosis of Interstitial Lung Diseases: A Series Contribution to a Debated Procedure. <i>Medicina (Lithuania)</i> , 2019, 55, 606.	0.8	13
1190	tPA promotes the proliferation of lung fibroblasts and activates the Wnt/ β 2-catenin signaling pathway in idiopathic pulmonary fibrosis. <i>Cell Cycle</i> , 2019, 18, 3137-3146.	1.3	17
1191	Treatment of idiopathic pulmonary fibrosis with Nintedanib: an update. <i>Expert Review of Respiratory Medicine</i> , 2019, 13, 1139-1146.	1.0	13
1192	Interstitial Pneumonia With Autoimmune Features (IPAF). <i>Frontiers in Medicine</i> , 2019, 6, 209.	1.2	47
1193	High-Resolution CT Change over Time in Patients with Idiopathic Pulmonary Fibrosis on Antifibrotic Treatment. <i>Journal of Clinical Medicine</i> , 2019, 8, 1469.	1.0	17
1194	Autotaxin and chronic inflammatory diseases. <i>Journal of Autoimmunity</i> , 2019, 104, 102327.	3.0	68
1195	Medical Treatments of Peyronie's Disease: Past, Present, and Future. <i>Urology</i> , 2019, 125, 1-5.	0.5	9
1196	Advances in CT Diagnosis of UIP and IPF. <i>Seminars in Roentgenology</i> , 2019, 54, 6-14.	0.2	3
1197	Scleroderma-related interstitial lung disease: principles of management. <i>Expert Review of Respiratory Medicine</i> , 2019, 13, 357-367.	1.0	7
1198	Progress in Understanding and Treating Idiopathic Pulmonary Fibrosis. <i>Annual Review of Medicine</i> , 2019, 70, 211-224.	5.0	75
1199	Frailty and geriatric conditions in older patients with idiopathic pulmonary fibrosis. <i>Respiratory Medicine</i> , 2019, 148, 6-12.	1.3	31
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1205	Natural History of Idiopathic Pulmonary Fibrosis and Disease Monitoring. , 2019, , 89-97.		0
1206	Vitamin D prevents experimental lung fibrosis and predicts survival in patients with idiopathic pulmonary fibrosis. <i>Pulmonary Pharmacology and Therapeutics</i> , 2019, 55, 17-24.	1.1	62
1207	Obstacles to early treatment of idiopathic pulmonary fibrosis: current perspectives. <i>Therapeutics and Clinical Risk Management</i> , 2019, Volume 15, 73-81.	0.9	12
1208	<p>Collecting patient preference information using a Clinical Data Research Network: demonstrating feasibility with idiopathic pulmonary fibrosis<p>. <i>Patient Preference and Adherence</i> , 2019, Volume 13, 795-804.	0.8	2
1209	Design and synthesis of novel pyrazolo[4,3- <i>d</i>]pyrimidines as potential therapeutic agents for acute lung injury. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019, 34, 1121-1130.	2.5	16
1210	Experimental pulmonary fibrosis was suppressed by microRNA-506 through NF-kappa-mediated apoptosis and inflammation. <i>Cell and Tissue Research</i> , 2019, 378, 255-265.	1.5	19
1211	Modeling of Fibrotic Lung Disease Using 3D Organoids Derived from Human Pluripotent Stem Cells. <i>Cell Reports</i> , 2019, 27, 3709-3723.e5.	2.9	175
1212	Application of nintedanib and other potential anti-fibrotic agents in fibrotic diseases. <i>Clinical Science</i> , 2019, 133, 1309-1320.	1.8	26
1213	Diagnosing complications and co-morbidities of fibrotic interstitial lung disease. <i>Expert Review of Respiratory Medicine</i> , 2019, 13, 645-658.	1.0	12
1214	Novel idiopathic pulmonary fibrosis susceptibility variants revealed by deepÂsequencing. <i>ERJ Open Research</i> , 2019, 5, 00071-2019.	1.1	24
1215	Interstitial Lung Disease and ANCA-Associated Vasculitis. <i>Current Treatment Options in Rheumatology</i> , 2019, 5, 213-229.	0.6	1
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1362	Lung Transplantation for Interstitial Lung Disease. , 2019, , 131-149.		0
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1375	The immunomodulatory role of interleukin-35 in fibrotic diseases. <i>Expert Review of Clinical Immunology</i> , 2019, 15, 431-439.	1.3	13
1376	Tissue Continues to Be the Issue: Role of Histopathology in the Context of Recent Updates in the Radiologic Classification of Interstitial Lung Diseases. <i>Archives of Pathology and Laboratory Medicine</i> , 2019, 143, 30-33.	1.2	0
1377	The mTORC1/4E-BP1 axis represents a critical signaling node during fibrogenesis. <i>Nature Communications</i> , 2019, 10, 6.	5.8	159
1378	The Role of Immunity and Inflammation in IPF Pathogenesis. <i>Respiratory Medicine</i> , 2019, , 97-131.	0.1	7
1379	Prognostic significance of early pulmonary function changes after onset of chronic lung allograft dysfunction. <i>Journal of Heart and Lung Transplantation</i> , 2019, 38, 184-193.	0.3	12
1380	Interstitial Lung Abnormality: Recognition and Perspectives. <i>Radiology</i> , 2019, 291, 1-3.	3.6	70
1381	Hyperpolarised xenon magnetic resonance spectroscopy for the longitudinal assessment of changes in gas diffusion in IPF. <i>Thorax</i> , 2019, 74, 500-502.	2.7	53
1382	No relevant pharmacokinetic drug-drug interaction between nintedanib and pirfenidone. <i>European Respiratory Journal</i> , 2019, 53, 1801060.	3.1	22
1383	Nicotine Modulates Growth Factors and MicroRNA to Promote Inflammatory and Fibrotic Processes. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2019, 368, 169-178.	1.3	23
1384	Attenuation of pulmonary fibrosis in type I collagen-targeted reporter mice with ALK-5 inhibitors. <i>Pulmonary Pharmacology and Therapeutics</i> , 2019, 54, 31-38.	1.1	16
1385	Fibrotic Signaling in the Lung. <i>Molecular and Translational Medicine</i> , 2019, , 91-119.	0.4	0

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1387	Fibrosis and Immune Dysregulation in Systemic Sclerosis. <i>Molecular and Translational Medicine</i> , 2019, , 25-60.	0.4	0
1388	The Role of Mast Cells in the Pathophysiology of Pulmonary Fibrosis. <i>Molecular and Translational Medicine</i> , 2019, , 135-173.	0.4	0
1389	Dynamic Reciprocity: The Role of the Extracellular Matrix Microenvironment in Amplifying and Sustaining Pathological Lung Fibrosis. <i>Molecular and Translational Medicine</i> , 2019, , 239-270.	0.4	1
1390	Lung function outcomes in the INPULSIS [®] trials of nintedanib in idiopathic pulmonary fibrosis. <i>Respiratory Medicine</i> , 2019, 146, 42-48.	1.3	34
1391	Telomerase-related monogenic lung fibrosis presenting with subacute onset: a case report and review of literature. <i>Acta Clinica Belgica</i> , 2019, 74, 445-450.	0.5	0
1392	LOX/LOXL in pulmonary fibrosis: potential therapeutic targets. <i>Journal of Drug Targeting</i> , 2019, 27, 790-796.	2.1	58
1393	Tissue-informed engineering strategies for modeling human pulmonary diseases. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2019, 316, L303-L320.	1.3	24
1394	Diagnosis of asbestos-related lung diseases. <i>Expert Review of Respiratory Medicine</i> , 2019, 13, 241-249.	1.0	12
1395	Federated electronic health records research technology to support clinical trial protocol optimization: Evidence from EHR4CR and the InSite platform. <i>Journal of Biomedical Informatics</i> , 2019, 90, 103090.	2.5	29
1396	Randomised, double-blind, placebo-controlled pilot trial of omeprazole in idiopathic pulmonary fibrosis. <i>Thorax</i> , 2019, 74, 346-353.	2.7	45
1397	Senolytics in idiopathic pulmonary fibrosis: Results from a first-in-human, open-label, pilot study. <i>EBioMedicine</i> , 2019, 40, 554-563.	2.7	746
1398	Assessing Patterns of Palliative Care Referral and Location of Death in Patients with Idiopathic Pulmonary Fibrosis: A Sixteen-Year Single-Center Retrospective Cohort Study. <i>Journal of Palliative Medicine</i> , 2019, 22, 538-544.	0.6	14
1399	Therapeutic effects of nintedanib are not influenced by emphysema in the INPULSIS trials. <i>European Respiratory Journal</i> , 2019, 53, 1801655.	3.1	22
1400	Inflammation and immunity in IPF pathogenesis and treatment. <i>Respiratory Medicine</i> , 2019, 147, 79-91.	1.3	259
1401	Existing and emerging treatments for idiopathic pulmonary fibrosis. <i>Expert Review of Respiratory Medicine</i> , 2019, 13, 229-239.	1.0	10
1402	Prolonged Survival in a Patient with Idiopathic Pulmonary Fibrosis Receiving Acupuncture and DHEA-Promoting Herbs with Conventional Management: A Case Report. , 2019, 23, 18-074.		1
1403	Determination of a novel antifibrotic small molecule GDC-03280 in human plasma and urine by liquid chromatography tandem mass spectrometry to support its first-in-human clinical trial. <i>Biomedical Chromatography</i> , 2019, 33, e4482.	0.8	1

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1405	Pirfenidone for Treating Idiopathic Pulmonary Fibrosis: An Evidence Review Group Perspective of a NICE Single Technology Appraisal. <i>Pharmacoeconomics</i> , 2019, 37, 763-775.	1.7	3
1406	Validity of the Patient Experiences and Satisfaction with Medications (PESaM) Questionnaire. <i>Patient</i> , 2019, 12, 149-162.	1.1	10
1407	The Hypoxic Adenosine Response and Inflammation in Lung Disease. , 2019, , 23-41.		0
1408	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
1409	Another Weapon in the Battle against Idiopathic Pulmonary Fibrosis?. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2019, 60, 386-387.	1.4	3
1410	Nintedanib in Idiopathic Pulmonary Fibrosis: Practical Management Recommendations for Potential Adverse Events. <i>Respiration</i> , 2019, 97, 173-184.	1.2	39
1411	The Tyrosine Kinase Inhibitor TAS-115 Attenuates Bleomycin-induced Lung Fibrosis in Mice. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2019, 60, 478-487.	1.4	15
1412	Long-term outcomes in chronic hypersensitivity pneumonitis. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2019, 74, 944-952.	2.7	55
1413	Antifibrotics: Shrinking the Box of Therapeutic Uncertainty. <i>Respiration</i> , 2019, 97, 202-204.	1.2	0
1414	Smoking-Related Lung Disease. <i>Seminars in Ultrasound, CT and MRI</i> , 2019, 40, 229-238.	0.7	9
1415	Determinants of survival in lung transplantation patients with idiopathic pulmonary fibrosis: a retrospective cohort study. <i>Transplant International</i> , 2019, 32, 399-409.	0.8	7
1416	IL-9 Blockade Suppresses Silica-induced Lung Inflammation and Fibrosis in Mice. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2019, 60, 232-243.	1.4	41
1417	Nintedanib-cyclodextrin complex to improve bio-activity and intestinal permeability. <i>Carbohydrate Polymers</i> , 2019, 204, 68-77.	5.1	47
1418	Immunological Lung Diseases. , 2019, , 967-980.e1.		0
1419	P311 in Scar Wars: Myofibroblasts Lost without Transforming Growth Factor β 2 Translation. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2019, 60, 139-140.	1.4	5
1420	The Future of Lung Transplantation. <i>Chest</i> , 2019, 155, 465-473.	0.4	67
1421	Understanding the patient's experience of care in idiopathic pulmonary fibrosis. <i>Respirology</i> , 2019, 24, 270-277.	1.3	31

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1423	Effectiveness and Safety of Chinese Medicine for Idiopathic Pulmonary Fibrosis: A Systematic Review and Meta-Analysis. <i>Chinese Journal of Integrative Medicine</i> , 2019, 25, 778-784.	0.7	21
1424	A Randomized, Double-Blinded, Placebo-Controlled, Dose-Escalation Phase 1 Study of Aerosolized Pirfenidone Delivered via the PARI Investigational eFlow Nebulizer in Volunteers and Patients with Idiopathic Pulmonary Fibrosis. <i>Journal of Aerosol Medicine and Pulmonary Drug Delivery</i> , 2020, 33, 15-20.	0.7	29
1425	What Patients With Idiopathic Pulmonary Fibrosis and Caregivers Want: Filling the Gaps With Patient Reported Outcomes and Experience Measures. <i>American Journal of Medicine</i> , 2020, 133, 281-289.	0.6	36
1426	Usual interstitial pneumonia in ANCA-associated vasculitis: A poor prognostic factor. <i>Journal of Autoimmunity</i> , 2020, 106, 102338.	3.0	43
1427	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
1428	Pamrevlumab in idiopathic pulmonary fibrosis. <i>Lancet Respiratory Medicine</i> , 2020, 8, 2-3.	5.2	14
1429	First-in-human high-cumulative-dose stem cell therapy in idiopathic pulmonary fibrosis with rapid lung function decline. <i>Stem Cells Translational Medicine</i> , 2020, 9, 6-16.	1.6	71
1430	Twenty percent of secondary publications of randomized controlled trials of drugs did not provide new results relative to the primary publication. <i>Journal of Clinical Epidemiology</i> , 2020, 117, 20-28.	2.4	6
1431	Ventilatory Support and Oxygen Therapy in Elder, Palliative and End-of-Life Care Patients. , 2020, , .		3
1432	Synthesis and discovery of new compounds bearing coumarin scaffold for the treatment of pulmonary fibrosis. <i>European Journal of Medicinal Chemistry</i> , 2020, 185, 111790.	2.6	7
1433	Interstitial lung disease: perhaps unclassifiable, but not untreatable. <i>Lancet Respiratory Medicine</i> , 2020, 8, 126-127.	5.2	2
1434	Pirfenidone in patients with unclassifiable progressive fibrosing interstitial lung disease: a double-blind, randomised, placebo-controlled, phase 2 trial. <i>Lancet Respiratory Medicine</i> , 2020, 8, 147-157.	5.2	410
1435	In search for a predictive marker of acute exacerbations of idiopathic pulmonary fibrosis. <i>Respirology</i> , 2020, 25, 234-235.	1.3	1
1436	Old but Gold: Tracking the New Guise of Histone Deacetylase 6 (HDAC6) Enzyme as a Biomarker and Therapeutic Target in Rare Diseases. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 23-39.	2.9	69
1437	Identifying "Real-World" Patients in the Real World. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 201, 256-257.	2.5	0
1438	Long-term treatment with nintedanib in Asian patients with idiopathic pulmonary fibrosis: Results from INPULSIS®. <i>Respirology</i> , 2020, 25, 410-416.	1.3	23
1439	Prediction of idiopathic pulmonary fibrosis progression using early quantitative changes on CT imaging for a short term of clinical 18-24-month follow-ups. <i>European Radiology</i> , 2020, 30, 726-734.	2.3	38

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1441	Idiopathic interstitial pneumonias. , 2020, , 29-140.		1
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1443	Secretory IgA accumulated in the airspaces of idiopathic pulmonary fibrosis and promoted VEGF, TGF- β ² and IL-8 production by A549 cells. Clinical and Experimental Immunology, 2020, 199, 326-336.	1.1	11
1444	Idiopathic and Autoimmune Interstitial Lung Disease. , 2020, , 1335-1354.		0
1445	Mitochondrial quality control in pulmonary fibrosis. Redox Biology, 2020, 33, 101426.	3.9	66
1446	The Role of Palliative Care in Reducing Symptoms and Improving Quality of Life for Patients with Idiopathic Pulmonary Fibrosis: A Review. Pulmonary Therapy, 2020, 6, 35-46.	1.1	24
1447	Ongoing challenges in pulmonary fibrosis and insights from the nintedanib clinical programme. Respiratory Research, 2020, 21, 7.	1.4	19
1448	Idiopathic Pulmonary Fibrosis: A Review of Disease, Pharmacological, and Nonpharmacological Strategies With a Focus on Symptoms, Function, and Health-Related Quality of Life. Journal of Pain and Symptom Management, 2020, 59, 1362-1378.	0.6	20
1449	Metabolic profiling of tyrosine kinase inhibitor nintedanib using metabolomics. Journal of Pharmaceutical and Biomedical Analysis, 2020, 180, 113045.	1.4	11
1450	Lung cancer in patients with Idiopathic Pulmonary Fibrosis. A retrospective multicenter study in Greece. Pulmonary Pharmacology and Therapeutics, 2020, 60, 101880.	1.1	31
1451	Magnetic Resonance in Crohn Disease. Magnetic Resonance Imaging Clinics of North America, 2020, 28, 45-53.	0.6	2
1452	Comparative survival analysis between idiopathic pulmonary fibrosis and chronic hypersensitivity pneumonitis. Pulmonology, 2020, 26, 3-9.	1.0	19
1453	Targeting c-Src Reverses Accelerated GPX-1 mRNA Decay in Chronic Obstructive Pulmonary Disease Airway Epithelial Cells. American Journal of Respiratory Cell and Molecular Biology, 2020, 62, 598-607.	1.4	8
1454	Idiopathic Pulmonary Fibrosis: Best Practice in Monitoring and Managing a Relentless Fibrotic Disease. Respiration, 2020, 99, 73-82.	1.2	58
1455	Corticosteroid use is not associated with improved outcomes in acute exacerbation of IPF. Respiriology, 2020, 25, 629-635.	1.3	47
1456	Nintedanib: New indication for systemic sclerosis-associated interstitial lung disease. Modern Rheumatology, 2020, 30, 225-231.	0.9	29
1457	Ca ²⁺ signalling in fibroblasts and the therapeutic potential of KCa3.1 channel blockers in fibrotic diseases. British Journal of Pharmacology, 2020, 177, 1003-1024.	2.7	23

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1460	Quantitative CT Analysis of Diffuse Lung Disease. <i>Radiographics</i> , 2020, 40, 28-43.	1.4	90
1461	Exosome-Derived microRNA-22 Ameliorates Pulmonary Fibrosis by Regulating Fibroblast-to-Myofibroblast Differentiation <i>in Vitro</i> and <i>in Vivo</i> . <i>Journal of Nippon Medical School</i> , 2020, 87, 118-128.	0.3	34
1462	An Open-label Study With Pirfenidone on Chronic Hypersensitivity Pneumonitis. <i>Archivos De Bronconeumologia</i> , 2020, 56, 163-169.	0.4	20
1463	Restrictive allograft dysfunction after lung transplantation: is there a place for nintedanib? a case report. <i>Fundamental and Clinical Pharmacology</i> , 2020, 34, 408-411.	1.0	5
1464	Microenvironment-Responsive Small-Molecule Probe for Pulmonary Fibrosis Detection. <i>Analytical Chemistry</i> , 2020, 92, 699-706.	3.2	12
1465	Anti-fibrotic mechanisms of exogenously-expanded mesenchymal stromal cells for fibrotic diseases. <i>Seminars in Cell and Developmental Biology</i> , 2020, 101, 87-103.	2.3	31
1466	Traditional Chinese medicine in the treatment of idiopathic pulmonary fibrosis based on syndrome differentiation: Study protocol of an exploratory trial. <i>Journal of Integrative Medicine</i> , 2020, 18, 163-168.	1.4	14
1467	Pathology of Idiopathic Interstitial Pneumonias. <i>Surgical Pathology Clinics</i> , 2020, 13, 91-118.	0.7	9
1468	Occupational exposures and idiopathic pulmonary fibrosis. <i>Current Opinion in Allergy and Clinical Immunology</i> , 2020, 20, 103-111.	1.1	10
1469	Multiple Traditional Chinese Medicine interventions for idiopathic pulmonary fibrosis. <i>Medicine (United States)</i> , 2020, 99, e22396.	0.4	7
1470	Older Idiopathic Pulmonary Fibrosis Male Patients Are at a Higher Risk of Nintedanib Dose Reduction. <i>Respiration</i> , 2020, 99, 646-648.	1.2	5
1471	Interstitial Lung Disease in Patients With Systemic Sclerosis: Toward Personalized-Medicine-Based Prediction and Drug Screening Models of Systemic Sclerosis-Related Interstitial Lung Disease (SSc-ILD). <i>Frontiers in Immunology</i> , 2020, 11, 1990.	2.2	9
1472	Long-Term Follow-Up of Patients With Idiopathic Pulmonary Fibrosis Treated With Pirfenidone or Nintedanib: A Real-Life Comparison Study. <i>Frontiers in Molecular Biosciences</i> , 2020, 7, 581828.	1.6	48
1473	STRATUS: A Phase II Study of Abituzumab in Patients With Systemic Sclerosis-associated Interstitial Lung Disease. <i>Journal of Rheumatology</i> , 2021, 48, 1295-1298.	1.0	12
1474	Healthcare use and costs among Medicare enrollees on pirfenidone versus nintedanib for idiopathic pulmonary fibrosis. <i>Journal of Comparative Effectiveness Research</i> , 2020, 9, 933-943.	0.6	7
1475	Prognostic significance of forced vital capacity decline prior to and following antifibrotic therapy in idiopathic pulmonary fibrosis. <i>Therapeutic Advances in Respiratory Disease</i> , 2020, 14, 175346662095378.	1.0	3
1476	Drug discovery and development in idiopathic pulmonary fibrosis: challenges and opportunities. <i>Drug Discovery Today</i> , 2020, 25, 2277-2283.	3.2	8

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1478	Baicalein alleviated TGF β 1-induced type I collagen production in lung fibroblasts via downregulation of connective tissue growth factor. <i>Biomedicine and Pharmacotherapy</i> , 2020, 131, 110744.	2.5	22
1479	Fibroblast Growth Factor Inhibitors in Lung Fibrosis: Friends or Foes?. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2020, 63, 273-274.	1.4	7
1480	The Pulmonary Fibrosis Foundation Patient Registry. Rationale, Design, and Methods. <i>Annals of the American Thoracic Society</i> , 2020, 17, 1620-1628.	1.5	27
1481	Australasian interstitial lung disease registry (AILDR): objectives, design and rationale of a bi-national prospective database. <i>BMC Pulmonary Medicine</i> , 2020, 20, 257.	0.8	9
1482	Investigation of telomere related gene mutations in idiopathic pulmonary fibrosis. <i>Molecular Biology Reports</i> , 2020, 47, 7851-7860.	1.0	0
1483	A novel phosphodiesterase 4 inhibitor, AA6216, reduces macrophage activity and fibrosis in the lung. <i>European Journal of Pharmacology</i> , 2020, 885, 173508.	1.7	20
1484	Occupational and environmental risk factors for idiopathic pulmonary fibrosis in Australia: case-control study. <i>Thorax</i> , 2020, 75, 864-869.	2.7	48
1485	<p>Inhibitors of the Autotaxin-Lysophosphatidic Acid Axis and Their Potential in the Treatment of Interstitial Lung Disease: Current Perspectives<p>. <i>Clinical Pharmacology: Advances and Applications</i> , 2020, Volume 12, 97-108.	0.8	13
1486	Idiopathic Nonspecific Interstitial Pneumonia. A Case Series and Literature Review. <i>Open Respiratory Archives</i> , 2020, 2, 192-193.	0.0	0
1487	A near-infrared fluorescent probe for evaluating glutamyl transpeptidase fluctuation in idiopathic pulmonary fibrosis cell and mice models. <i>Sensors and Actuators B: Chemical</i> , 2020, 322, 128565.	4.0	17
1488	Clinical relevance of endpoints in clinical trials for acid sphingomyelinase deficiency enzyme replacement therapy. <i>Molecular Genetics and Metabolism</i> , 2020, 131, 116-123.	0.5	18
1489	The adoption of nintedanib in systemic sclerosis: the SENSICIS study. <i>Breathe</i> , 2020, 16, 200005.	0.6	5
1490	Treatment patterns, healthcare resource utilization, and costs among patients with idiopathic pulmonary fibrosis treated with antifibrotic medications in US-based commercial and Medicare Supplemental claims databases: a retrospective cohort study. <i>BMC Pulmonary Medicine</i> , 2020, 20, 188.	0.8	14
1491	Antifibrotic Drug Use in Patients with Idiopathic Pulmonary Fibrosis. Data from the IPF-PRO Registry. <i>Annals of the American Thoracic Society</i> , 2020, 17, 1413-1423.	1.5	32
1492	Imatinib a Tyrosine Kinase Inhibitor: a potential treatment for SARS- COV-2 induced pneumonia. <i>Alexandria Journal of Medicine</i> , 2020, 56, 68-72.	0.4	8
1493	Pan-transcriptome-based candidate therapeutic discovery for idiopathic pulmonary fibrosis. <i>Therapeutic Advances in Respiratory Disease</i> , 2020, 14, 175346662097114.	1.0	7
1494	Transbronchial cryobiopsy in the diagnosis of interstitial lung diseases: methodologies and perspectives from the Cryo-PID and COLDICE studies. <i>Annals of Translational Medicine</i> , 2020, 8, 1330-1330.	0.7	5

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1496	Analysis of body mass index, weight loss and progression of idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2020, 21, 312.	1.4	44
1497	Comparison of reversal of rat pulmonary fibrosis of nintedanib, pirfenidone, and human umbilical mesenchymal stem cells from Wharton's jelly. <i>Stem Cell Research and Therapy</i> , 2020, 11, 513.	2.4	17
1498	Heparanase and the hallmarks of cancer. <i>Journal of Translational Medicine</i> , 2020, 18, 453.	1.8	78
1499	Engeletin ameliorates pulmonary fibrosis through endoplasmic reticulum stress depending on Inc949-mediated TGF- β 1-Smad2/3 and JNK signalling pathways. <i>Pharmaceutical Biology</i> , 2020, 58, 1114-1123.	1.3	12
1500	Facing the Challenge of Post COVID-19 Pulmonary fibrosis: What is so unique about it?. <i>Bangladesh Critical Care Journal</i> , 2020, 8, 102-107.	0.1	1
1501	Dynamics in diagnoses and pharmacotherapy before and after diagnosing idiopathic pulmonary fibrosis. <i>ERJ Open Research</i> , 2020, 6, 00479-2020.	1.1	3
1503	Clinical experience with antifibrotics in fibrotic hypersensitivity pneumonitis: a 3-year real-life observational study. <i>ERJ Open Research</i> , 2020, 6, 00152-2020.	1.1	15
1504	Fibrosis and cancer: shared features and mechanisms suggest common targeted therapeutic approaches. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 1024-1032.	0.4	18
1505	Is operation safe for lung cancer patients with interstitial lung disease on computed tomography?. <i>Therapeutic Advances in Respiratory Disease</i> , 2020, 14, 175346662097113.	1.0	2
1506	In-hospital mortality trends among patients with idiopathic pulmonary fibrosis in the United States between 2013-2017: a comparison of academic and non-academic programs. <i>BMC Pulmonary Medicine</i> , 2020, 20, 289.	0.8	13
1507	Transbronchial cryobiopsy for diffuse parenchymal lung diseases: evidence that demands a (favorable) verdict. <i>Annals of Translational Medicine</i> , 2020, 8, 1324-1324.	0.7	0
1508	Acute exacerbations of idiopathic pulmonary fibrosis and the role of corticosteroids. <i>Breathe</i> , 2020, 16, 200086.	0.6	5
1509	Antifibrotic and Regenerative Effects of Treamid in Pulmonary Fibrosis. <i>International Journal of Molecular Sciences</i> , 2020, 21, 8380.	1.8	9
1510	Functional parameters of small airways can guide bronchodilator use in idiopathic pulmonary fibrosis. <i>Scientific Reports</i> , 2020, 10, 18633.	1.6	4
1511	Hypersensitivity pneumonitis. <i>Nature Reviews Disease Primers</i> , 2020, 6, 65.	18.1	75
1512	Safety of Nintedanib in Patients with Idiopathic Pulmonary Fibrosis: Global Pharmacovigilance Data. <i>Advances in Therapy</i> , 2020, 37, 4209-4219.	1.3	21
1513	Inhalation: A means to explore and optimize nintedanib's pharmacokinetic/pharmacodynamic relationship. <i>Pulmonary Pharmacology and Therapeutics</i> , 2020, 63, 101933.	1.1	9

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1515	Systemic sclerosis-associated interstitial lung disease. <i>Current Opinion in Pulmonary Medicine</i> , 2020, 26, 487-495.	1.2	3
1516	Recent advances in rheumatoid arthritis-associated interstitial lung disease. <i>Current Opinion in Pulmonary Medicine</i> , 2020, 26, 477-486.	1.2	31
1517	Progressive fibrosing interstitial lung disease: treatable traits and therapeutic strategies. <i>Current Opinion in Pulmonary Medicine</i> , 2020, 26, 436-442.	1.2	18
1518	Safety and tolerability of nintedanib in patients with systemic sclerosis-associated interstitial lung disease: data from the SENSIS trial. <i>Annals of the Rheumatic Diseases</i> , 2020, 79, 1478-1484.	0.5	46
1519	Exhaled Biomarkers in Idiopathic Pulmonary Fibrosis—A Six-Month Follow-up Study in Patients Treated with Pirfenidone. <i>Journal of Clinical Medicine</i> , 2020, 9, 2523.	1.0	4
1520	Impact of angiotensin II type 1 and G-protein-coupled Mas receptor expression on the pulmonary performance of patients with idiopathic pulmonary fibrosis. <i>Peptides</i> , 2020, 133, 170384.	1.2	7
1521	Developments in lung transplantation over the past decade. <i>European Respiratory Review</i> , 2020, 29, 190132.	3.0	71
1522	Bi-directional communication: Conversations between fibroblasts and immune cells in systemic sclerosis. <i>Journal of Autoimmunity</i> , 2020, 113, 102526.	3.0	29
1523	Predictors of acute exacerbation in biopsy-proven idiopathic pulmonary fibrosis. <i>Respiratory Investigation</i> , 2020, 58, 177-184.	0.9	6
1524	Type I Collagen Signaling Regulates Opposing Fibrotic Pathways through $\alpha 2(\text{I})\beta 1$ Integrin. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2020, 63, 613-622.	1.4	12
1525	A Comprehensive Review of Clinical Cardiotoxicity Incidence of FDA-Approved Small-Molecule Kinase Inhibitors. <i>Frontiers in Pharmacology</i> , 2020, 11, 891.	1.6	48
1526	Exhaled breath analysis by use of eNose technology: a novel diagnostic tool for interstitial lung disease. <i>European Respiratory Journal</i> , 2021, 57, 2002042.	3.1	28
1527	Anticoagulant Use and Bleeding Risk in Central European Patients with Idiopathic Pulmonary Fibrosis (IPF) Treated with Antifibrotic Therapy: Real-World Data from EMPIRE. <i>Drug Safety</i> , 2020, 43, 971-980.	1.4	13
1528	Imidazolium-based ionic liquid functionalized mesoporous silica nanoparticles as a promising nano-carrier: response surface strategy to investigate and optimize loading and release process for Lapatinib delivery. <i>Pharmaceutical Development and Technology</i> , 2020, 25, 1150-1161.	1.1	14
1529	Acute exacerbation of idiopathic interstitial pneumonias related to chemotherapy for lung cancer: nationwide surveillance in Japan. <i>ERJ Open Research</i> , 2020, 6, 00184-2019.	1.1	18
1530	Prognosis of idiopathic pulmonary fibrosis without anti-fibrotic therapy: a systematic review. <i>European Respiratory Review</i> , 2020, 29, 190158.	3.0	48
1531	Patient expectations, experiences and satisfaction with nintedanib and pirfenidone in idiopathic pulmonary fibrosis: a quantitative study. <i>Respiratory Research</i> , 2020, 21, 196.	1.4	16

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1532	Patient-reported Outcomes for Clinical Trials in Idiopathic Pulmonary Fibrosis: New Opportunities to Understand How Patients Feel and Function. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 202, 1620-1622.	2.5	2
1533	The need for a holistic approach for SSc-ILD â€œ achievements and ambiguity in a devastating disease. <i>Respiratory Research</i> , 2020, 21, 197.	1.4	33
1534	Palliation of chronic breathlessness with morphine in patients with fibrotic interstitial lung disease â€œ a randomised placebo-controlled trial. <i>Respiratory Research</i> , 2020, 21, 195.	1.4	22
1535	A long-acting isomer of Acâ€œSDKP attenuates pulmonary fibrosis through SRPK1 â€œmediated PI3K / AKT and Smad2 pathway inhibition. <i>IUBMB Life</i> , 2020, 72, 2611-2626.	1.5	5
1536	New Perspectives on the Aberrant Alveolar Repair of Idiopathic Pulmonary Fibrosis. <i>Frontiers in Cell and Developmental Biology</i> , 2020, 8, 580026.	1.8	12
1537	Response to: â€œCorrespondence on â€œSafety and tolerability of nintedanib in patients with systemic sclerosis-associated interstitial lung disease: data from the SENSICIS trialâ€™â€™ by Bredemeier. <i>Annals of the Rheumatic Diseases</i> , 2022, 81, e251-e251.	0.5	0
1538	Early diagnosis of idiopathic pulmonary fibrosis: Closer to the goal?. <i>European Journal of Internal Medicine</i> , 2020, 80, 12-13.	1.0	1
1539	New developments in respiratory medicine: a primary immunodeficiency perspective. <i>Current Opinion in Allergy and Clinical Immunology</i> , 2020, 20, 549-556.	1.1	0
1540	Clinical features of patients with small cell lung cancer and idiopathic pulmonary fibrosis treated with chemotherapy or chemoradiotherapy. <i>Therapeutic Advances in Respiratory Disease</i> , 2020, 14, 175346662096386.	1.0	5
1541	Low-Frequency Intrapulmonary Percussive Ventilation Increases Aerosol Penetration in a 2-Compartment Physical Model of Fibrotic Lung Disease. <i>Frontiers in Bioengineering and Biotechnology</i> , 2020, 8, 1022.	2.0	1
1542	Progressive fibrosing interstitial lung disease: clinical uncertainties, consensus recommendations, and research priorities. <i>Lancet Respiratory Medicine</i> , the, 2020, 8, 925-934.	5.2	198
1543	Novel Multitarget Therapies for Lung Cancer and Respiratory Disease. <i>Molecules</i> , 2020, 25, 3987.	1.7	14
1544	Rheumatoid Arthritisâ€œAssociated Interstitial Lung Disease: Current Update on Prevalence, Risk Factors, and Pharmacologic Treatment. <i>Current Treatment Options in Rheumatology</i> , 2020, 6, 337-353.	0.6	35
1545	â€œA Chain Only as Strong as Its Weakest Linkâ€œ An Up-to-Date Literature Review on the Bidirectional Interaction of Pulmonary Fibrosis and COVID-19. <i>Journal of Proteome Research</i> , 2020, 19, 4327-4338.	1.8	33
1546	Mediastinal lymph node enlargement in idiopathic pulmonary fibrosis: relationships with disease progression and pulmonary function trends. <i>BMC Pulmonary Medicine</i> , 2020, 20, 249.	0.8	7
1547	CA 19-9 serum levels in patients with end-stage idiopathic pulmonary fibrosis (IPF) and other interstitial lung diseases (ILDs): Correlation with functional decline. <i>Chronic Respiratory Disease</i> , 2020, 17, 147997312095842.	1.0	9
1548	Tocilizumab in systemic sclerosis: a randomised, double-blind, placebo-controlled, phase 3 trial. <i>Lancet Respiratory Medicine</i> , the, 2020, 8, 963-974.	5.2	348
1549	Antifibrotic treatment improves clinical outcomes in patients with idiopathic pulmonary fibrosis: a propensity score matching analysis. <i>Scientific Reports</i> , 2020, 10, 15620.	1.6	38

#	ARTICLE	IF	CITATIONS
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1551	Neuraxial anesthesia and nintedanib use: a word of caution. <i>Regional Anesthesia and Pain Medicine</i> , 2020, 45, 482.3-483.	1.1	0
1552	Real-World Data on Bleeding Risk and Anticoagulation in Patients with IPF Treated with Antifibrotics. <i>Drug Safety</i> , 2020, 43, 953-955.	1.4	1
1553	Identification of a unique temporal signature in blood and BAL associated with IPF progression. <i>Scientific Reports</i> , 2020, 10, 12049.	1.6	10
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1555	The Efficacy and Safety of First-Line Chemotherapy in Patients With Non-small Cell Lung Cancer and Interstitial Lung Disease: A Systematic Review and Meta-Analysis. <i>Frontiers in Oncology</i> , 2020, 10, 1636.	1.3	8
1556	Translational pharmacology of an inhaled small molecule $\hat{=}$ 26 integrin inhibitor for idiopathic pulmonary fibrosis. <i>Nature Communications</i> , 2020, 11, 4659.	5.8	65
1557	Changes in management of idiopathic pulmonary fibrosis: impact on disease severity and mortality. <i>European Clinical Respiratory Journal</i> , 2020, 7, 1807682.	0.7	7
1558	Cell Therapy for Idiopathic Pulmonary Fibrosis: Rationale and Progress to Date. <i>BioDrugs</i> , 2020, 34, 543-556.	2.2	8
1559	Radiation-induced lung toxicity $\hat{=}$ cellular and molecular mechanisms of pathogenesis, management, and literature review. <i>Radiation Oncology</i> , 2020, 15, 214.	1.2	103
1560	Lung complications of Sjogren syndrome. <i>European Respiratory Review</i> , 2020, 29, 200021.	3.0	31
1561	Nintedanib in Bronchiolitis Obliterans Syndrome After Allogeneic Hematopoietic Stem Cell Transplantation. <i>Chest</i> , 2020, 158, e89-e91.	0.4	10
1563	Transbronchial cryobiopsy increases diagnostic confidence in interstitial lung disease: a prospective multicentre trial. <i>European Respiratory Journal</i> , 2020, 56, 1901520.	3.1	41
1564	Gender equity in interstitial lung disease. <i>Lancet Respiratory Medicine</i> , 2020, 8, 842-843.	5.2	6
1565	Impact of Lung Biopsy on Lung Function in Idiopathic Pulmonary Fibrosis. <i>Respiration</i> , 2020, 99, 1101-1108.	1.2	3
1566	Infections and systemic sclerosis: an emerging challenge. <i>Revista Colombiana De ReumatologÃa (English Edition)</i> , 2020, 27, 62-84.	0.1	1
1567	The effect of statin therapy on disease-related outcomes in idiopathic pulmonary fibrosis: A systematic review and meta-analysis. <i>Respiratory Medicine and Research</i> , 2021, 80, 100792.	0.4	1
1568	Galectin-3 levels are elevated following nintedanib treatment. <i>Therapeutic Advances in Chronic Disease</i> , 2020, 11, 204062232096841.	1.1	5

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1570	ILDGDB: a manually curated database of genomics, transcriptomics, proteomics and drug information for interstitial lung diseases. <i>BMC Pulmonary Medicine</i> , 2020, 20, 323.	0.8	3
1571	Supportive care of patients with fibrosing interstitial lung disease: answering a great clinical need. <i>Breathe</i> , 2020, 16, 200066.	0.6	0
1572	Nanoapproaches to Modifying Epigenetics of Epithelial Mesenchymal Transition for Treatment of Pulmonary Fibrosis. <i>Frontiers in Pharmacology</i> , 2020, 11, 607689.	1.6	28
1573	Pharmacokinetic evaluation of two pirfenidone formulations in patients with idiopathic pulmonary fibrosis and chronic hypersensitivity pneumonitis. <i>Heliyon</i> , 2020, 6, e05279.	1.4	8
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1575	Toward Realizing the Full Potential of Registries in Interstitial Lung Disease. <i>Annals of the American Thoracic Society</i> , 2020, 17, 1534-1535.	1.5	0
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1577	Peritubular Capillary Rarefaction: An Underappreciated Regulator of CKD Progression. <i>International Journal of Molecular Sciences</i> , 2020, 21, 8255.	1.8	33
1578	Inhaled nintedanib is well-tolerated and delivers key pharmacokinetic parameters required to treat bleomycin-induced pulmonary fibrosis. <i>Pulmonary Pharmacology and Therapeutics</i> , 2020, 63, 101938.	1.1	20
1579	Human pluripotent stem cell-derived lung organoids: Potential applications in development and disease modeling. <i>Wiley Interdisciplinary Reviews: Developmental Biology</i> , 2021, 10, e399.	5.9	32
1581	Assessment and Management of Occupational Hypersensitivity Pneumonitis. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2020, 8, 3295-3309.	2.0	7
1582	Cardiovascular and Pulmonary Challenges After Treatment of Childhood Cancer. <i>Pediatric Clinics of North America</i> , 2020, 67, 1155-1170.	0.9	7
1583	Serial CT analysis in idiopathic pulmonary fibrosis: comparison of visual features that determine patient outcome. <i>Thorax</i> , 2020, 75, 648-654.	2.7	26
1584	Tannic acid alleviates experimental pulmonary fibrosis in mice by inhibiting inflammatory response and fibrotic process. <i>Inflammopharmacology</i> , 2020, 28, 1301-1314.	1.9	10
1585	Blockade of Pan-Fibroblast Growth Factor Receptors Mediates Bidirectional Effects in Lung Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2020, 63, 317-326.	1.4	8
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#	ARTICLE	IF	CITATIONS
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1590	A multicentre retrospective observational study on Polish experience of pirfenidone therapy in patients with idiopathic pulmonary fibrosis: the PolExPIR study. <i>BMC Pulmonary Medicine</i> , 2020, 20, 122.	0.8	9
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1592	Smoking-associated interstitial lung disease: update and review. <i>Expert Review of Respiratory Medicine</i> , 2020, 14, 825-834.	1.0	23
1593	The world is not enough – the value of increasing registry data in idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2020, 21, 105.	1.4	3
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1596	Respiratory surveillance in mineral dust-exposed workers. <i>Breathe</i> , 2020, 16, 190632.	0.6	12
1597	A modified scar model with controlled tension on secondary wound healing in mice. <i>Burns and Trauma</i> , 2020, 8, tkaa013.	2.3	4
1598	Heart failure with preserved ejection fraction diagnosis and treatment: An updated review of the evidence. <i>Progress in Cardiovascular Diseases</i> , 2020, 63, 570-584.	1.6	53
1599	Idiopathic pulmonary fibrosis: airway volume measurement identifies progressive disease on computed tomography scans. <i>ERJ Open Research</i> , 2020, 6, 00290-2019.	1.1	8
1600	Update in Interstitial Lung Disease 2019. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 202, 500-507.	2.5	17
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1603	A new therapy for systemic sclerosis-associated interstitial lung disease. <i>Respiratory Investigation</i> , 2020, 58, 227-229.	0.9	0
1604	Meaningful survival benefit for single lung transplantation in idiopathic pulmonary fibrosis patients over 65 years of age. <i>European Respiratory Journal</i> , 2020, 56, 1902413.	3.1	7
1605	TRIM72 promotes alveolar epithelial cell membrane repair and ameliorates lung fibrosis. <i>Respiratory Research</i> , 2020, 21, 132.	1.4	13

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1609	Idiopathic pulmonary fibrosis: do scientists focus on publishing rather than on clinical relevance?. <i>European Respiratory Journal</i> , 2020, 55, 2000811.	3.1	0
1610	Multicenter phase 2 trial of nintedanib in advanced nonpancreatic neuroendocrine tumors. <i>Cancer</i> , 2020, 126, 3689-3697.	2.0	11
1611	Deep learning in interstitial lung disease—how long until daily practice. <i>European Radiology</i> , 2020, 30, 6285-6292.	2.3	37
1612	Shedding light on developmental drugs for idiopathic pulmonary fibrosis. <i>Expert Opinion on Investigational Drugs</i> , 2020, 29, 797-808.	1.9	8
1613	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
1614	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
1615	Interstitial lung diseases in children. <i>Presse Medicale</i> , 2020, 49, 103909.	0.8	26
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1617	Discovery of a Potent and Selective Covalent Inhibitor and Activity-Based Probe for the Deubiquitylating Enzyme UCHL1, with Antifibrotic Activity. <i>Journal of the American Chemical Society</i> , 2020, 142, 12020-12026.	6.6	51
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1621	Radiation-Induced Lung Fibrosis: Preclinical Animal Models and Therapeutic Strategies. <i>Cancers</i> , 2020, 12, 1561.	1.7	56
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1626	Contemporary approaches in the use of extracorporeal membrane oxygenation to support patients waiting for lung transplantation. <i>Annals of Cardiothoracic Surgery</i> , 2020, 9, 29-41.	0.6	9
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1628	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
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1630	Hope in Patients With Progressive Fibrosis Interstitial Lung Disease (PF-ILD). <i>Clinical Pulmonary Medicine</i> , 2020, 27, 38-38.	0.3	0
1631	The Prophylactic Use of Macrolide Antibiotics to Prevent Acute Exacerbations in Bronchiectasis. <i>Clinical Pulmonary Medicine</i> , 2020, 27, 37-38.	0.3	0
1632	NADPH Oxidase Inhibition in Fibrotic Pathologies. <i>Antioxidants and Redox Signaling</i> , 2020, 33, 455-479.	2.5	20
1633	FLT1: a potential therapeutic target in sepsis-associated ARDS?. <i>Lancet Respiratory Medicine</i> , 2020, 8, 219-220.	5.2	1
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1635	Pantoea agglomerans chronic exposure induces epithelial-mesenchymal transition in human lung epithelial cells and mice lungs. <i>Ecotoxicology and Environmental Safety</i> , 2020, 194, 110416.	2.9	5
1636	Emerging biomarkers in chronic lung allograft dysfunction. <i>Expert Review of Molecular Diagnostics</i> , 2020, 20, 467-475.	1.5	6
1637	Disease progression across the spectrum of idiopathic pulmonary fibrosis: A multicentre study. <i>Respirology</i> , 2020, 25, 1144-1151.	1.3	6
1638	Patientsâ€™ and Healthcare Professionalsâ€™ Experiences of Idiopathic Pulmonary Fibrosis Treatment with the Pirfenidone 801Âµg Tablet Formulation: A Multinational Survey. <i>Pulmonary Therapy</i> , 2020, 6, 93-105.	1.1	1
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1641	Effects of Nintedanib in an Animal Model of Liver Fibrosis. <i>BioMed Research International</i> , 2020, 2020, 1-9.	0.9	19
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1646	Transcriptomic profiling reveals disease-specific characteristics of epithelial cells in idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2020, 21, 165.	1.4	11
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1650	Genetic Variation in CCL18 Gene Influences CCL18 Expression and Correlates with Survival in Idiopathic Pulmonary Fibrosis – Part B. <i>Journal of Clinical Medicine</i> , 2020, 9, 1993.	1.0	10
1651	Early referral to palliative care in IPF – pitfalls and opportunities in clinical trials. <i>Respiratory Research</i> , 2020, 21, 174.	1.4	2
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1654	Novel Therapeutic Approaches for Pulmonary Manifestations of Systemic Sclerosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 202, 878-880.	2.5	0
1655	Genetic Variation in CCL18 Gene Influences CCL18 Expression and Correlates with Survival in Idiopathic Pulmonary Fibrosis: Part A. <i>Journal of Clinical Medicine</i> , 2020, 9, 1940.	1.0	18
1656	Progressive fibrosing interstitial lung disease: we know it behaves badly, but what does that mean?. <i>European Respiratory Journal</i> , 2020, 55, 2000894.	3.1	1
1657	An aberrantly sustained emergency granulopoiesis response accelerates postchemotherapy relapse in MLL1-rearranged acute myeloid leukemia in mice. <i>Journal of Biological Chemistry</i> , 2020, 295, 9663-9675.	1.6	2
1658	The myofibroblast at a glance. <i>Journal of Cell Science</i> , 2020, 133, .	1.2	167
1659	Synthesis, structural properties, enzyme inhibition and molecular docking studies of (Z)-N'-(1-allyl-2-oxoindolin-3-ylidene) methanesulfonylhydrazide and (Z)-N'-(1-allyl-2-oxoindolin-3-ylidene)-3-nitrobenzenesulfonylhydrazide. <i>Journal of Molecular Structure</i> , 2020, 1221, 128880.	1.8	6
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1664	Nintedanib inhibits intrahepatic cholangiocarcinoma aggressiveness via suppression of cytokines extracted from activated cancer-associated fibroblasts. <i>British Journal of Cancer</i> , 2020, 122, 986-994.	2.9	44
1665	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
1666	Therapeutic Options for the Treatment of Interstitial Lung Disease Related to Connective Tissue Diseases. A Narrative Review. <i>Journal of Clinical Medicine</i> , 2020, 9, 407.	1.0	40
1667	Impact of Depression on Patients With Idiopathic Pulmonary Fibrosis. <i>Frontiers in Medicine</i> , 2020, 7, 29.	1.2	18
1668	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
1669	Antibody-based therapies for idiopathic pulmonary fibrosis. <i>Expert Opinion on Biological Therapy</i> , 2020, 20, 779-786.	1.4	13
1670	Deterioration of high-resolution computed tomography findings predicts disease progression after initial decline in forced vital capacity in idiopathic pulmonary fibrosis patients treated with pirfenidone. <i>Respiratory Investigation</i> , 2020, 58, 185-189.	0.9	5
1671	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
1672	Left ventricular dysfunction in an idiopathic pulmonary fibrosis patient on nintedanib. <i>Respirology Case Reports</i> , 2020, 8, e00533.	0.3	2
1673	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
1674	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
1675	Acute exacerbation of idiopathic pulmonary fibrosis: international survey and call for harmonisation. <i>European Respiratory Journal</i> , 2020, 55, 1901760.	3.1	61
1676	Patient gender bias on the diagnosis of idiopathic pulmonary fibrosis. <i>Thorax</i> , 2020, 75, 407-412.	2.7	30
1677	Acute Exacerbation of Interstitial Lung Disease in Adult Patients With Idiopathic Inflammatory Myopathies: A Retrospective Case-Control Study. <i>Frontiers in Medicine</i> , 2020, 7, 12.	1.2	12
1678	Small airways pathology in idiopathic pulmonary fibrosis: a retrospective cohort study. <i>Lancet Respiratory Medicine</i> , 2020, 8, 573-584.	5.2	70

#	ARTICLE	IF	CITATIONS
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1680	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
1681	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
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1708	Nonspecific Interstitial Pneumonia. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2020, 41, 184-201.	0.8	7
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1751	Keloid disorder: Fibroblast differentiation and gene expression profile in fibrotic skin diseases. <i>Experimental Dermatology</i> , 2021, 30, 132-145.	1.4	59
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1760	Tolerability and safety of nintedanib in elderly patients with idiopathic pulmonary fibrosis. <i>Respiratory Investigation</i> , 2021, 59, 99-105.	0.9	13
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1762	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
1763	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
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1798	Mouse Models of Lung Fibrosis. <i>Methods in Molecular Biology</i> , 2021, 2299, 291-321.	0.4	8
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1804	Association of proangiogenic and profibrotic serum markers with lung function and quality of life in sarcoidosis. <i>PLoS ONE</i> , 2021, 16, e0247197.	1.1	3
1805	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
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1808	Chronic Lung Allograft Dysfunction: Review of CT and Pathologic Findings. <i>Radiology: Cardiothoracic Imaging</i> , 2021, 3, e200314.	0.9	15
1809	Pulmonary rehabilitation for interstitial lung disease. <i>The Cochrane Library</i> , 2021, 2021, CD006322.	1.5	67
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1822	Blood Transcriptomics Predicts Progression of Pulmonary Fibrosis and Associated Natural Killer Cells. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021, 204, 197-208.	2.5	27
1823	Real-world retrospective observational study exploring the effectiveness and safety of antifibrotics in idiopathic pulmonary fibrosis. <i>BMJ Open Respiratory Research</i> , 2021, 8, e000782.	1.2	22
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1826	Nintedanib: A Review in Fibrotic Interstitial Lung Diseases. <i>Drugs</i> , 2021, 81, 575-586.	4.9	51
1827	Pharmacological Interventions for Pulmonary Involvement in Rheumatic Diseases. <i>Pharmaceuticals</i> , 2021, 14, 251.	1.7	2
1828	Gene therapy strategies for idiopathic pulmonary fibrosis: recent advances, current challenges, and future directions. <i>Molecular Therapy - Methods and Clinical Development</i> , 2021, 20, 483-496.	1.8	21
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1832	Membrane particles from mesenchymal stromal cells reduce the expression of fibrotic markers on pulmonary cells. <i>PLoS ONE</i> , 2021, 16, e0248415.	1.1	1
1833	A mixed-methods pilot study of handheld fan for breathlessness in interstitial lung disease. <i>Scientific Reports</i> , 2021, 11, 6874.	1.6	8
1834	Pharmacological treatment of idiopathic pulmonary fibrosis and fibrosing interstitial lung diseases: current trends and future directions. <i>Precision and Future Medicine</i> , 2021, 5, 31-40.	0.5	2
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1836	Interstitial lung disease. <i>Canadian Journal of Respiratory, Critical Care, and Sleep Medicine</i> , 2021, 5, 93-96.	0.2	0
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1840	Nintedanib and immunomodulatory therapies in progressive fibrosing interstitial lung diseases. <i>Respiratory Research</i> , 2021, 22, 84.	1.4	33
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1842	The Protective Effects of IL-31RA Deficiency During Bleomycin-Induced Pulmonary Fibrosis. <i>Frontiers in Immunology</i> , 2021, 12, 645717.	2.2	13
1843	A Phase 2 Randomized, Double-Blind, Placebo-Controlled Study Evaluating Nintedanib Versus Placebo as Prophylaxis Against Radiation Pneumonitis in Patients With Unresectable NSCLC Undergoing Chemoradiation Therapy. <i>Journal of Thoracic Oncology</i> , 2021, 16, e19-e20.	0.5	10
1844	The Effect of Nintedanib on T-Cell Activation, Subsets and Functions. <i>Drug Design, Development and Therapy</i> , 2021, Volume 15, 997-1011.	2.0	11
1845	Disparate Interferon Signaling and Shared Aberrant Basaloid Cells in Single-Cell Profiling of Idiopathic Pulmonary Fibrosis and Systemic Sclerosis-Associated Interstitial Lung Disease. <i>Frontiers in Immunology</i> , 2021, 12, 595811.	2.2	54
1846	Simple method for detecting idiopathic interstitial pneumonias by measuring vertical lung length on chest X-ray. <i>Scientific Reports</i> , 2021, 11, 7669.	1.6	2
1847	Wound dehiscence with nintedanib after cardiac surgery: A cautionary tale. <i>JTCVS Techniques</i> , 2021, 6, 99-101.	0.2	6

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1849	Remote Assessment of Lung Disease and Impact on Physical and Mental Health (RALPMH): Protocol for a Prospective Observational Study. <i>JMIR Research Protocols</i> , 2021, 10, e28873.	0.5	10
1850	Diverse Pathological Findings of Interstitial Lung Disease in a Patient with Dyskeratosis Congenita. <i>Internal Medicine</i> , 2021, 60, 1257-1263.	0.3	3
1851	The transition from normal lung anatomy to minimal and established fibrosis in idiopathic pulmonary fibrosis (IPF). <i>EBioMedicine</i> , 2021, 66, 103325.	2.7	16
1852	Protective Effects of Extracellular Matrix-Derived Hydrogels in Idiopathic Pulmonary Fibrosis. <i>Tissue Engineering - Part B: Reviews</i> , 2022, 28, 517-530.	2.5	5
1853	Antifibrotic drugs for pulmonary sarcoidosis: A treatment in search of an indication. <i>Respiratory Medicine</i> , 2021, 180, 106371.	1.3	5
1854	Inhibition of the Proliferation of Human Lung Fibroblasts by Prostacyclin Receptor Agonists is Linked to a Sustained cAMP Signal in the Nucleus. <i>Frontiers in Pharmacology</i> , 2021, 12, 669227.	1.6	16
1855	Fibroblasts as immune regulators in infection, inflammation and cancer. <i>Nature Reviews Immunology</i> , 2021, 21, 704-717.	10.6	229
1856	Current and future treatment for idiopathic pulmonary fibrosis. <i>Journal of the Korean Medical Association</i> , 2021, 64, 256-263.	0.1	1
1857	Extracellular matrix proteins produced by stromal cells in idiopathic pulmonary fibrosis and lung adenocarcinoma. <i>PLoS ONE</i> , 2021, 16, e0250109.	1.1	15
1858	<i>H. sinensis</i> mycelium inhibits epithelial-mesenchymal transition by inactivating the midkine pathway in pulmonary fibrosis. <i>Frontiers of Medicine</i> , 2021, 15, 313-329.	1.5	3
1859	Impact of bronchoalveolar lavage lymphocytosis on the effects of anti-inflammatory therapy in idiopathic non-specific interstitial pneumonia, idiopathic pleuroparenchymal fibroelastosis, and unclassifiable idiopathic interstitial pneumonia. <i>Respiratory Research</i> , 2021, 22, 115.	1.4	5
1860	Precision medicine in idiopathic pulmonary fibrosis therapy: From translational research to patient-centered care. <i>Current Opinion in Pharmacology</i> , 2021, 57, 71-80.	1.7	7
1861	Pulmonary fibrosis secondary to COVID-19: a narrative review. <i>Expert Review of Respiratory Medicine</i> , 2021, 15, 791-803.	1.0	64
1862	Post-COVID-19 pneumonia lung fibrosis: a worrisome sequelae in surviving patients. <i>Egyptian Journal of Radiology and Nuclear Medicine</i> , 2021, 52, .	0.3	60
1863	Efficacy and safety of nintedanib in patients with idiopathic pulmonary fibrosis who are elderly or have comorbidities. <i>Respiratory Research</i> , 2021, 22, 125.	1.4	22
1864	Regulatory Immune Cells in Idiopathic Pulmonary Fibrosis: Friends or Foes?. <i>Frontiers in Immunology</i> , 2021, 12, 663203.	2.2	33
1865	Prognostic classification in acute exacerbation of idiopathic pulmonary fibrosis: a multicentre retrospective cohort study. <i>Scientific Reports</i> , 2021, 11, 9120.	1.6	9

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1867	Isolation and characterisation of lymphatic endothelial cells from lung tissues affected by lymphangioleiomyomatosis. <i>Scientific Reports</i> , 2021, 11, 8406.	1.6	5
1868	Inhibitory effects of somatostatin analogue in bleomycin-induced pulmonary fibrosis. <i>Experimental Lung Research</i> , 2021, 47, 280-288.	0.5	1
1871	The Role of Macrophages in the Development of Acute and Chronic Inflammatory Lung Diseases. <i>Cells</i> , 2021, 10, 897.	1.8	97
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1873	Exposure-efficacy analyses of nintedanib in patients with chronic fibrosing interstitial lung disease. <i>Respiratory Medicine</i> , 2021, 180, 106369.	1.3	9
1874	Treatment of chronic fibrosing interstitial lung diseases. <i>Journal of the Korean Medical Association</i> , 2021, 64, 277-285.	0.1	1
1875	Current advances in the treatment of autoimmune-associated interstitial lung diseases. <i>Journal of the Korean Medical Association</i> , 2021, 64, 264-276.	0.1	0
1876	Endobronchial autologous bone marrowâ€mesenchymal stromal cells in idiopathic pulmonary fibrosis: a phase I trial. <i>ERJ Open Research</i> , 2021, 7, 00773-2020.	1.1	10
1877	A role for cardiopulmonary exercise testing in detecting physiological changes underlying health status in Idiopathic pulmonary fibrosis: a feasibility study. <i>BMC Pulmonary Medicine</i> , 2021, 21, 147.	0.8	1
1878	Comparative outcomes in patients receiving pirfenidone or nintedanib for idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2021, 22, 135.	1.4	18
1879	Usual interstitial pneumonia progressing to nonspecific interstitial pneumonia-like pattern on high-resolution CT with histologic confirmation. <i>Radiology Case Reports</i> , 2021, 16, 1019-1022.	0.2	1
1880	Aberrant B Cell Receptor Signaling in Na ⁺ -ve B Cells from Patients with Idiopathic Pulmonary Fibrosis. <i>Cells</i> , 2021, 10, 1321.	1.8	12
1882	Chronic Lung Allograft Dysfunction: Evolving Concepts and Therapies. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2021, 42, 392-410.	0.8	4
1883	Deep Learning Approach for Auto-Detecting Idiopathic Pulmonary Fibrosis Prediction. , 2021, , .		0
1884	Antibody-mediated depletion of CCR10+ EphA3+ cells ameliorates fibrosis in IPF. <i>JCI Insight</i> , 2021, 6, .	2.3	9
1885	Antifibrotic Drugs for COVID-19: From Orphan Drugs to Blockbusters?. <i>Current Respiratory Medicine Reviews</i> , 2021, 17, 8-12.	0.1	2
1886	Nurse-Led Palliative Care Clinical Trial Improves Knowledge and Preparedness in Caregivers of Patients with Idiopathic Pulmonary Fibrosis. <i>Annals of the American Thoracic Society</i> , 2021, 18, 1811-1821.	1.5	25

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1888	Pirfenidone in patients with non-IPF progressive fibrotic interstitial lung diseases: expert guidance is urgently needed. <i>Lancet Respiratory Medicine</i> , 2021, 9, 437-438.	5.2	5
1889	Safety and tolerability of combination therapy with pirfenidone and nintedanib for idiopathic pulmonary fibrosis: A multicenter retrospective observational study in Japan. <i>Respiratory Investigation</i> , 2021, 59, 819-826.	0.9	10
1890	An updated safety review of the drug treatments for idiopathic pulmonary fibrosis. <i>Expert Opinion on Drug Safety</i> , 2021, 20, 1035-1048.	1.0	6
1891	A study design for statistical learning technique to predict radiological progression with an application of idiopathic pulmonary fibrosis using chest CT images. <i>Contemporary Clinical Trials</i> , 2021, 104, 106333.	0.8	3
1892	Baseline plasma KL-6 level predicts adverse outcomes in patients with idiopathic pulmonary fibrosis receiving nintedanib: a retrospective real-world cohort study. <i>BMC Pulmonary Medicine</i> , 2021, 21, 165.	0.8	9
1893	Recent advances in the diagnosis and management of interstitial pneumonia with autoimmune features: the perspective of rheumatologists. <i>Korean Journal of Internal Medicine</i> , 2021, 36, 515-526.	0.7	3
1894	Azithromycin for the Treatment of Chronic Cough in Idiopathic Pulmonary Fibrosis: A Randomized Controlled Crossover Trial. <i>Annals of the American Thoracic Society</i> , 2021, 18, 2018-2026.	1.5	19
1895	Patient-reported outcomes and patient-reported outcome measures in interstitial lung disease: where to go from here?. <i>European Respiratory Review</i> , 2021, 30, 210026.	3.0	17
1896	TGF β 1 induces resistance of human lung myofibroblasts to cell death via downregulation of TRPA1 channels. <i>British Journal of Pharmacology</i> , 2021, 178, 2948-2962.	2.7	8
1897	Antifibrotic Therapies and Progressive Fibrosing Interstitial Lung Disease (PF-ILD): Building on INBUILD. <i>Journal of Clinical Medicine</i> , 2021, 10, 2285.	1.0	18
1898	Micellar Hyaluronidase and Spiperone as a Potential Treatment for Pulmonary Fibrosis. <i>International Journal of Molecular Sciences</i> , 2021, 22, 5599.	1.8	6
1899	PerFECT 2.0: A Web-Based Platform Designed to Facilitate and Support the Diagnosis of Patients with Idiopathic Pulmonary Fibrosis in Italy. <i>Pulmonary Therapy</i> , 2021, 7, 267-279.	1.1	0
1900	Posttranslational regulation of PGC α modulates fibrotic repair. <i>FASEB Journal</i> , 2021, 35, e21675.	0.2	6
1901	Longitudinal Changes in Clinical Features, Management, and Outcomes of Idiopathic Pulmonary Fibrosis. A Nationwide Cohort Study. <i>Annals of the American Thoracic Society</i> , 2021, 18, 780-787.	1.5	14
1902	Putting the spotlight on macrophage-derived cathepsin in the pathophysiology of obliterative bronchiolitis. <i>European Respiratory Journal</i> , 2021, 57, 2004607.	3.1	1
1903	COVID-19 and pulmonary fibrosis: A potential role for lung epithelial cells and fibroblasts. <i>Immunological Reviews</i> , 2021, 302, 228-240.	2.8	126
1904	Increased monocyte count and red cell distribution width as prognostic biomarkers in patients with Idiopathic Pulmonary Fibrosis. <i>Respiratory Research</i> , 2021, 22, 140.	1.4	37

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1906	Use of Nintedanib and Pirfenidone in Non-Idiopathic Pulmonary Fibrosis Lung Disease. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021, 204, 92-94.	2.5	2
1907	A systematic review of the incidence, risk factors and prognosis of acute exacerbation of systemic autoimmune disease-associated interstitial lung disease. <i>BMC Pulmonary Medicine</i> , 2021, 21, 150.	0.8	11
1908	Treatment of Idiopathic Pulmonary Fibrosis. <i>Cureus</i> , 2021, 13, e15360.	0.2	4
1909	Ground glass and fibrotic change in children with surfactant protein C dysfunction mutations. <i>Pediatric Pulmonology</i> , 2021, 56, 2223-2231.	1.0	10
1910	Travel Distance to Subspecialty Clinic and Outcomes in Patients with Fibrotic Interstitial Lung Disease. <i>Annals of the American Thoracic Society</i> , 2022, 19, 20-27.	1.5	16
1911	Identification of Potential Pathogenic Super-Enhancers-Driven Genes in Pulmonary Fibrosis. <i>Frontiers in Genetics</i> , 2021, 12, 644143.	1.1	6
1912	Antifibrotic Role of Nintedanib in Tracheal Stenosis After a Tracheal Wound. <i>Laryngoscope</i> , 2021, 131, E2496-E2505.	1.1	11
1913	The safety of nintedanib for the treatment of interstitial lung disease: A systematic review and meta-analysis of randomized controlled trials. <i>PLoS ONE</i> , 2021, 16, e0251636.	1.1	13
1914	Awake or intubated surgery in diagnosis for interstitial lung diseases? A prospective study. <i>ERJ Open Research</i> , 2021, 7, 00630-2020.	1.1	12
1915	Anti-angiogenic Agents: A Review on Vascular Endothelial Growth Factor Receptor-2 (VEGFR-2) Inhibitors. <i>Current Medicinal Chemistry</i> , 2021, 28, 2540-2564.	1.2	27
1916	Molecular pathways in idiopathic pulmonary fibrosis pathogenesis: Transcending barriers to optimally targeted pharmacotherapies. <i>EBioMedicine</i> , 2021, 67, 103373.	2.7	1
1917	Expectations about treatment of idiopathic pulmonary fibrosis: Comparative survey of patients, carers and physicians (the RESPIR French survey). <i>Respiratory Medicine and Research</i> , 2021, 79, 100811.	0.4	1
1918	Esomeprazole attenuates inflammatory and fibrotic response in lung cells through the MAPK/Nrf2/HO1 pathway. <i>Journal of Inflammation</i> , 2021, 18, 17.	1.5	9
1919	Emerging drugs for the treatment of idiopathic pulmonary fibrosis: 2020 phase II clinical trials. <i>Expert Opinion on Emerging Drugs</i> , 2021, 26, 93-101.	1.0	6
1920	Biomarkers in Progressive Fibrosing Interstitial Lung Disease: Optimizing Diagnosis, Prognosis, and Treatment Response. <i>Frontiers in Medicine</i> , 2021, 8, 680997.	1.2	31
1921	Looking Ahead. <i>Clinics in Chest Medicine</i> , 2021, 42, 375-384.	0.8	2
1922	IL-18 binding protein can be a prognostic biomarker for idiopathic pulmonary fibrosis. <i>PLoS ONE</i> , 2021, 16, e0252594.	1.1	11

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1924	Automated Digital Quantification of Pulmonary Fibrosis in Human Histopathology Specimens. <i>Frontiers in Medicine</i> , 2021, 8, 607720.	1.2	13
1925	Cellular Senescence in Lung Fibrosis. <i>International Journal of Molecular Sciences</i> , 2021, 22, 7012.	1.8	33
1926	Genetically increased circulating FUT3 level leads to reduced risk of Idiopathic Pulmonary Fibrosis: a Mendelian Randomisation Study. <i>European Respiratory Journal</i> , 2021, , 2003979.	3.1	9
1927	Rheumatological evaluation of patients with interstitial lung disease. <i>Scandinavian Journal of Rheumatology</i> , 2022, 51, 34-41.	0.6	6
1928	Efficacy and safety of nintedanib in Japanese patients with early-stage idiopathic pulmonary fibrosis: a study protocol for an observational study. <i>BMJ Open</i> , 2021, 11, e047249.	0.8	1
1929	Detection and Early Referral of Patients With Interstitial Lung Abnormalities. <i>Chest</i> , 2022, 161, 470-482.	0.4	26
1930	Protocol for long-term effect of pulmonary rehabilitation under nintedanib in idiopathic pulmonary fibrosis. <i>ERJ Open Research</i> , 2021, 7, 00321-2021.	1.1	3
1931	Treatment of Persistent Cough in Subjects with Idiopathic Pulmonary Fibrosis (IPF) with Gefapixant, a P2X3 Antagonist, in a Randomized, Placebo-Controlled Clinical Trial. <i>Pulmonary Therapy</i> , 2021, 7, 471-486.	1.1	25
1932	Minimal clinically important difference in idiopathic pulmonary fibrosis. <i>Breathe</i> , 2021, 17, 200345.	0.6	0
1933	An update on targeted therapies in systemic sclerosis based on a systematic review from the last 3–years. <i>Arthritis Research and Therapy</i> , 2021, 23, 155.	1.6	42
1934	Interstitial Lung Disease in 2020. <i>Clinics in Chest Medicine</i> , 2021, 42, 229-239.	0.8	8
1935	Importance of chest HRCT in the diagnostic evaluation of fibrosing interstitial lung diseases. <i>Jornal Brasileiro De Pneumologia</i> , 2021, 47, e20200096.	0.4	11
1936	Resistance Training for Rehabilitation in Patients with Idiopathic Pulmonary Fibrosis. <i>Translational Medicine and Exercise Prescription</i> , 0, , 62-69.	0.0	0
1937	Antifibrotics Modify B-Cell–induced Fibroblast Migration and Activation in Patients with Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2021, 64, 722-733.	1.4	26
1938	VEGFR1-tyrosine kinase signaling in pulmonary fibrosis. <i>Inflammation and Regeneration</i> , 2021, 41, 16.	1.5	14
1939	A bird–s eye view of fibroblast heterogeneity: A pan–disease, pan–cancer perspective. <i>Immunological Reviews</i> , 2021, 302, 299-320.	2.8	23
1940	Clinical Trials for Idiopathic Pulmonary Fibrosis and the Role of Health Systems. <i>Clinics in Chest Medicine</i> , 2021, 42, 287-294.	0.8	3

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1942	Management of Idiopathic Pulmonary Fibrosis. <i>Clinics in Chest Medicine</i> , 2021, 42, 275-285.	0.8	7
1945	New Insights into Profibrotic Myofibroblast Formation in Systemic Sclerosis: When the Vascular Wall Becomes the Enemy. <i>Life</i> , 2021, 11, 610.	1.1	14
1946	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
1947	Differentiation of Idiopathic Pulmonary Fibrosis from Connective Tissue Disease-Related Interstitial Lung Disease Using Quantitative Imaging. <i>Journal of Clinical Medicine</i> , 2021, 10, 2663.	1.0	11
1948	Diagnosis and Management of Fibrotic Interstitial Lung Diseases. <i>Clinics in Chest Medicine</i> , 2021, 42, 321-335.	0.8	7
1949	Idiopathic pulmonary fibrosis: Disease mechanisms and drug development. , 2021, 222, 107798.		216
1950	Molecular Markers and the Promise of Precision Medicine for Interstitial Lung Disease. <i>Clinics in Chest Medicine</i> , 2021, 42, 357-364.	0.8	4
1951	Molecular Pathogenesis of Pulmonary Fibrosis, with Focus on Pathways Related to TGF- β 2 and the Ubiquitin-Proteasome Pathway. <i>International Journal of Molecular Sciences</i> , 2021, 22, 6107.	1.8	55
1952	Inhaled treprostinil and forced vital capacity in patients with interstitial lung disease and associated pulmonary hypertension: a post-hoc analysis of the INCREASE study. <i>Lancet Respiratory Medicine</i> , 2021, 9, 1266-1274.	5.2	62
1953	Disease Behaviour During the Peri-Diagnostic Period in Patients with Suspected Interstitial Lung Disease: The STARLINER Study. <i>Advances in Therapy</i> , 2021, 38, 4040-4056.	1.3	6
1954	Salvianolic acid B inhalation solution enhances antifibrotic and anticoagulant effects in a rat model of pulmonary fibrosis. <i>Biomedicine and Pharmacotherapy</i> , 2021, 138, 111475.	2.5	18
1955	Efficacy of nintedanib in the treatment of interstitial lung disease associated with systemic sclerosis. <i>Pulmonologiya</i> , 2021, 31, 391-396.	0.2	0
1956	Antifibrotic drugs in lung transplantation and chronic lung allograft dysfunction: a review. <i>European Respiratory Review</i> , 2021, 30, 210050.	3.0	8
1957	Idiopathic pulmonary fibrosis complications: what a radiologist should know. <i>Journal of Radiological Review</i> , 2021, 8, .	0.1	1
1958	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
1959	Morbidity and mortality reduction associated with polysomnography testing in idiopathic pulmonary fibrosis: a population-based cohort study. <i>BMC Pulmonary Medicine</i> , 2021, 21, 185.	0.8	2
1960	Clinical Features and Outcomes of Combined Pulmonary Fibrosis and Emphysema After Lung Transplantation. <i>Chest</i> , 2021, 160, 1743-1750.	0.4	12

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1962	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
1963	Unclassifiable, or simply unclassified interstitial lung disease?. <i>Current Opinion in Pulmonary Medicine</i> , 2021, 27, 405-413.	1.2	5
1964	Clinical Significance of Interstitial Lung Disease and Its Acute Exacerbation in Microscopic Polyangiitis. <i>Chest</i> , 2021, 159, 2334-2345.	0.4	18
1965	The Role of miRNAs in Extracellular Matrix Repair and Chronic Fibrotic Lung Diseases. <i>Cells</i> , 2021, 10, 1706.	1.8	13
1966	<i>in vivo</i> assessment of pulmonary fibrosis and edema in rodents using the backscatter coefficient and envelope statistics. <i>Journal of the Acoustical Society of America</i> , 2021, 150, 183-192.	0.5	5
1967	Validation and minimum important difference of the UCSD Shortness of Breath Questionnaire in fibrotic interstitial lung disease. <i>Respiratory Research</i> , 2021, 22, 202.	1.4	5
1968	Kindlin-2 Acts as a Key Mediator of Lung Fibroblast Activation and Pulmonary Fibrosis Progression. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2021, 65, 54-69.	1.4	8
1969	Serum Biomarkers in Differential Diagnosis of Idiopathic Pulmonary Fibrosis and Connective Tissue Disease-Associated Interstitial Lung Disease. <i>Journal of Clinical Medicine</i> , 2021, 10, 3167.	1.0	10
1970	Prognostic significance of serum cytokines during acute exacerbation of idiopathic interstitial pneumonias treated with thrombomodulin. <i>BMJ Open Respiratory Research</i> , 2021, 8, e000889.	1.2	2
1971	Potential for inhibition of checkpoint kinases 1/2 in pulmonary fibrosis and secondary pulmonary hypertension. <i>Thorax</i> , 2022, 77, 247-258.	2.7	11
1972	Broad Adoption of Antifibrotics in Idiopathic Pulmonary Fibrosis: Still a Long Way to Go. <i>Annals of the American Thoracic Society</i> , 2021, 18, 1115-1116.	1.5	2
1973	Early Intervention of Pulmonary Rehabilitation for Fibrotic Interstitial Lung Disease Is a Favorable Factor for Short-Term Improvement in Health-Related Quality of Life. <i>Journal of Clinical Medicine</i> , 2021, 10, 3153.	1.0	9
1974	Turnover of type I and III collagen predicts progression of idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2021, 22, 205.	1.4	20
1975	Repurposing Nintedanib for pathological cardiac remodeling and dysfunction. <i>Pharmacological Research</i> , 2021, 169, 105605.	3.1	10
1976	Outcomes of patients with advanced idiopathic pulmonary fibrosis treated with nintedanib or pirfenidone in a real-world multicentre cohort. <i>Respirology</i> , 2021, 26, 982-988.	1.3	13
1977	New insights into the Hippo/YAP pathway in idiopathic pulmonary fibrosis. <i>Pharmacological Research</i> , 2021, 169, 105635.	3.1	18
1978	Effects of Exercise Training on Cardiopulmonary Function and Quality of Life in Elderly Patients with Pulmonary Fibrosis: A Meta-Analysis. <i>International Journal of Environmental Research and Public Health</i> , 2021, 18, 7643.	1.2	9

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1981	Nintedanib for treatment of progressive interstitial lung disease after peripheral blood stem cell transplantation in a patient with recurrent acute lymphoblastic leukemia. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29211.	0.8	2
1982	NOX4 regulates macrophage apoptosis resistance to induce fibrotic progression. <i>Journal of Biological Chemistry</i> , 2021, 297, 100810.	1.6	12
1983	Harnessing the Role of HDAC6 in Idiopathic Pulmonary Fibrosis: Design, Synthesis, Structural Analysis, and Biological Evaluation of Potent Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 9960-9988.	2.9	26
1984	Efficacy of early antifibrotic treatment for idiopathic pulmonary fibrosis. <i>BMC Pulmonary Medicine</i> , 2021, 21, 218.	0.8	9
1985	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
1986	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
1987	Management and support of patients with fibrosing interstitial lung diseases. <i>Nurse Practitioner</i> , 2021, 46, 39-44.	0.2	2
1988	Outcomes for hospitalized patients with idiopathic pulmonary fibrosis treated with antifibrotic medications. <i>BMC Pulmonary Medicine</i> , 2021, 21, 239.	0.8	6
1989	Extracellular matrix and Hippo signaling as therapeutic targets of antifibrotic compounds for uterine fibroids. <i>Clinical and Translational Medicine</i> , 2021, 11, e475.	1.7	27
1990	Recognition of Connective Tissue Disease-Related Interstitial Pneumonia Based on Histological Scoreâ€‘A Validation Study of an Online Diagnostic Decision Support Tool. <i>Diagnostics</i> , 2021, 11, 1359.	1.3	0
1991	50-gene risk profiles in peripheral blood predict COVID-19 outcomes: A retrospective, multicenter cohort study. <i>EBioMedicine</i> , 2021, 69, 103439.	2.7	20
1992	Consensus document for the selection of lung transplant candidates: An update from the International Society for Heart and Lung Transplantation. <i>Journal of Heart and Lung Transplantation</i> , 2021, 40, 1349-1379.	0.3	293
1993	Efficacy and safety of nintedanib for pulmonary fibrosis in severe pneumonia induced by COVID-19: An interventional study. <i>International Journal of Infectious Diseases</i> , 2021, 108, 454-460.	1.5	65
1994	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
1996	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
1997	Elucidation of prognostic factors and the effect of anti-fibrotic therapy on waitlist mortality in lung transplant candidates with idiopathic interstitial pneumonias. <i>Respiratory Investigation</i> , 2021, 59, 428-435.	0.9	5
1998	Impact of Antifibrotic Therapy on Mortality and Acute Exacerbation in Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2021, 160, 1751-1763.	0.4	88

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2000	The Paradigm of Targeting an Oncogenic Tyrosine Kinase: Lesson from BCR-ABL. , 0, , .		3
2001	Patient-centered Outcomes Research in Interstitial Lung Disease: An Official American Thoracic Society Research Statement. American Journal of Respiratory and Critical Care Medicine, 2021, 204, e3-e23.	2.5	41
2002	Pterostilbene alleviates pulmonary fibrosis by regulating ASIC2. Chinese Medicine, 2021, 16, 66.	1.6	6
2003	Characteristics and evaluation of acute exacerbations in chronic interstitial lung diseases. Respiratory Medicine, 2021, 183, 106400.	1.3	6
2004	An update on interstitial lung disease. British Journal of Hospital Medicine (London, England: 2005), 2021, 82, 1-14.	0.2	2
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2006	The Development of 3-substituted Indolin-2-one Derivatives as Kinase Inhibitors for Cancer Therapy. Current Medicinal Chemistry, 2022, 29, 1891-1919.	1.2	7
2007	The histologic diagnosis of usual interstitial pneumonia of idiopathic pulmonary fibrosis. Where we are and where we need to go. Modern Pathology, 2022, 35, 8-14.	2.9	13
2008	Lung transplantation for interstitial lung disease. European Respiratory Review, 2021, 30, 210017.	3.0	36
2009	Fibrometabolismâ€”An emerging therapeutic frontier in pulmonary fibrosis. Science Signaling, 2021, 14, .	1.6	31
2010	Human bronchial epithelial cellâ€”derived extracellular vesicle therapy for pulmonary fibrosis via inhibition of TGFâ€”WNT crosstalk. Journal of Extracellular Vesicles, 2021, 10, e12124.	5.5	74
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2012	Survival after inpatient or outpatient pulmonary rehabilitation in patients with fibrotic interstitial lung disease: a multicentre retrospective cohort study. Thorax, 2022, 77, 589-595.	2.7	21
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2018	Cellular Senescence in Idiopathic Pulmonary Fibrosis. <i>Current Molecular Biology Reports</i> , 2021, 7, 31-40.	0.8	29
2019	Enzymatic cross-linking of collagens in organ fibrosis – resolution and assessment. <i>Expert Review of Molecular Diagnostics</i> , 2021, 21, 1049-1064.	1.5	20
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2022	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
2023	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
2024	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
2025	Clinical Impact of Surgical Lung Biopsy for Interstitial Lung Disease in a Reference Center. <i>Annals of Thoracic Surgery</i> , 2021, , .	0.7	0
2026	Inspiratory muscle training in interstitial lung disease: a systematic scoping review. <i>Jornal Brasileiro De Pneumologia</i> , 2021, 47, e20210089.	0.4	6
2027	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
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2030	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
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2036	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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2038	CT of Post-Acute Lung Complications of COVID-19. <i>Radiology</i> , 2021, 301, E383-E395.	3.6	115
2039	Fibrotic Idiopathic Interstitial Lung Disease: The Molecular and Cellular Key Players. <i>International Journal of Molecular Sciences</i> , 2021, 22, 8952.	1.8	27
2040	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
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2043	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
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2047	Dose Optimization of Sotorasib: Is the US Food and Drug Administration Sending a Message?. <i>Journal of Clinical Oncology</i> , 2021, 39, 3423-3426.	0.8	25
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2049	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
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2051	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
2052	<scp>SHP2</scp>: The protein tyrosine phosphatase involved in chronic pulmonary inflammation and fibrosis. <i>IUBMB Life</i> , 2022, 74, 131-142.	1.5	6
2053	Nintedanib in progressive interstitial lung diseases: data from the whole INBUILD trial. <i>European Respiratory Journal</i> , 2022, 59, 2004538.	3.1	47
2054	Special considerations for pulmonary rehabilitation in conditions other than COPD. , 2021, , 145-164.		3

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2056	Progressive Fibrosing Interstitial Lung Diseases: A Current Perspective. <i>Biomedicines</i> , 2021, 9, 1237.	1.4	10
2057	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
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2062	GED-0507 attenuates lung fibrosis by counteracting myofibroblast transdifferentiation in vivo and in vitro. <i>PLoS ONE</i> , 2021, 16, e0257281.	1.1	5
2063	Therapeutic targets in lung tissue remodelling and fibrosis. , 2021, 225, 107839.		98
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2070	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
2071	The Aggregate Index of Systemic Inflammation (AIS): A Novel Prognostic Biomarker in Idiopathic Pulmonary Fibrosis. <i>Journal of Clinical Medicine</i> , 2021, 10, 4134.	1.0	32
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2073	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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2077	Treatment of fibrotic interstitial lung disease: current approaches and future directions. <i>Lancet</i> , The, 2021, 398, 1450-1460.	6.3	47
2078	Improved Radiolytic Stability of a ⁶⁸ Ga-labelled Collagelin Analogue for the Imaging of Fibrosis. <i>Pharmaceuticals</i> , 2021, 14, 990.	1.7	3
2079	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
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2081	Ten Commandments for Randomized Trials of Pharmacological Therapy for COPD and Other Lung Diseases. <i>COPD: Journal of Chronic Obstructive Pulmonary Disease</i> , 2021, 18, 1-8.	0.7	1
2082	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
2083	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
2084	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
2085	Hyperpolarized ¹²⁹ Xe MRI and Spectroscopy of Gas-Exchange Abnormalities in Nonspecific Interstitial Pneumonia. <i>Radiology</i> , 2021, 301, 211-220.	3.6	11
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2087	Beyond TGFβ1 - novel treatment strategies targeting lung fibrosis. <i>International Journal of Biochemistry and Cell Biology</i> , 2021, 141, 106090.	1.2	2
2088	Defining and predicting progression in non-IPF interstitial lung disease. <i>Respiratory Medicine</i> , 2021, 189, 106626.	1.3	5
2089	Molecular Signatures of Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2021, 65, 430-441.	1.4	23
2090	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
2091	Moving beyond usual interstitial pneumonia to define progressive fibrotic interstitial lung disease. <i>Lancet Respiratory Medicine</i> , the, 2021, 9, 1087-1089.	5.2	0
2092	Interstitial Lung Abnormalities: State of the Art. <i>Radiology</i> , 2021, 301, 19-34.	3.6	63

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2095	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
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2097	Peptide DR8 analogs alleviate pulmonary fibrosis via suppressing TGF- β 21 mediated epithelial-mesenchymal transition and ERK1/2 pathway in vivo and in vitro. <i>European Journal of Pharmaceutical Sciences</i> , 2021, 167, 106009.	1.9	11
2098	The phosphodiesterase 4 inhibitor AA6216 suppresses activity of fibrosis-specific macrophages. <i>Biochemistry and Biophysics Reports</i> , 2021, 28, 101118.	0.7	1
2099	Scleroderma Associated Interstitial Lung Disease. , 2022, , 319-325.		0
2100	Rheumatoid Arthritis Interstitial Lung Disease. , 2022, , 307-318.		0
2101	Interstitial Pneumonia With Autoimmune Features. , 2022, , 298-306.		1
2103	A Comprehensive Guide to Lung Transplantation for the Recipient With Pulmonary Fibrosis. , 2022, , 661-675.		0
2104	Other Idiopathic Interstitial Pneumonias and Unclassifiable Interstitial Lung Disease. , 2022, , 257-274.		0
2105	Comprehensive Care of Interstitial Lung Disease. , 2022, , 64-78.		0
2106	Idiopathic Pulmonary Fibrosis-Treatment and Management. , 2022, , 218-233.		0
2107	The Role of the Innate Immune System in Interstitial Lung Disease. , 2022, , 135-143.		0
2108	Palliative Care in Interstitial Lung Disease. <i>Respiratory Medicine</i> , 2021, , 189-207.	0.1	0
2109	Patient-Centredness and Patient-Reported Measures (PRMs) in Palliation of Lung Disease. <i>Respiratory Medicine</i> , 2021, , 43-75.	0.1	1
2110	An Introduction to Advanced Lung Disease. <i>Respiratory Medicine</i> , 2021, , 11-25.	0.1	0
2111	<i>Hirsutella sinensis</i> mycelium regulates autophagy of alveolar macrophages via TLR4/NF- β B signaling pathway. <i>International Journal of Medical Sciences</i> , 2021, 18, 1810-1823.	1.1	7

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2113	Phase three clinical trials in idiopathic pulmonary fibrosis. <i>Expert Opinion on Orphan Drugs</i> , 2021, 9, 1-11.	0.5	2
2114	Chronic Fibrosing Interstitial Lung Disease with Progressive Phenotype. <i>Nauchno-Prakticheskaya Revmatologiya</i> , 2021, 58, 631-636.	0.2	3
2115	<i>Pulmonary and Critical Care Medicine.</i> , 2021, , 325-338.		0
2116	Post-COVID lung fibrosis: The tsunami that will follow the earthquake. <i>Lung India</i> , 2021, 38, 41.	0.3	69
2117	Immune dysregulation as a driver of idiopathic pulmonary fibrosis. <i>Journal of Clinical Investigation</i> , 2021, 131, .	3.9	114
2118	Molecular Imaging of Fibrosis. , 2021, , 1447-1468.		0
2119	S2K Guideline for Diagnosis of Idiopathic Pulmonary Fibrosis. <i>Respiration</i> , 2021, 100, 238-271.	1.2	19
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2121	The Promise (and Pitfalls) of Administrative Data for Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2021, 159, 9-10.	0.4	0
2122	Spontaneous pneumothorax during nintedanib therapy in patients with systemic sclerosis-associated interstitial lung disease. <i>Respirology Case Reports</i> , 2021, 9, e00716.	0.3	3
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2124	Post-COVID-19 pulmonary fibrosis: A case series and review of literature. <i>Journal of Family Medicine and Primary Care</i> , 2021, 10, 2028.	0.3	11
2125	Combined Pulmonary Fibrosis and Emphysema (CPFE) Clinical Features and Management. <i>International Journal of COPD</i> , 2021, Volume 16, 167-177.	0.9	16
2126	Pulmonary Fibrosis Progression Prediction Using Image Processing and Machine Learning. <i>Advances in Science, Technology and Innovation</i> , 2021, , 159-177.	0.2	1
2127	Pretreatment of aged mice with retinoic acid supports alveolar regeneration via upregulation of reciprocal PDGFA signalling. <i>Thorax</i> , 2021, 76, 456-467.	2.7	19
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2129	Genetic Risk Factors for Idiopathic Pulmonary Fibrosis: Insights into Immunopathogenesis. <i>Journal of Inflammation Research</i> , 2020, Volume 13, 1305-1318.	1.6	29

#	ARTICLE	IF	CITATIONS
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2133	Risk Factors and Biomarkers of RA-ILD. <i>Respiratory Medicine</i> , 2018, , 59-72.	0.1	3
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2137	Cardiovascular implications of idiopathic pulmonary fibrosis: A way forward together?. <i>American Heart Journal</i> , 2020, 226, 69-74.	1.2	7
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2144	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
2145	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
2150	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
2151	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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2157	Soluble Thy-1 reverses lung fibrosis via its integrin-binding motif. JCI Insight, 2019, 4, .	2.3	20
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2164	miR-323a-3p regulates lung fibrosis by targeting multiple profibrotic pathways. JCI Insight, 2016, 1, e90301.	2.3	37
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2167	Fra-2 α -expressing macrophages promote lung fibrosis. Journal of Clinical Investigation, 2019, 129, 3293-3309.	3.9	67
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2172	The therapy of idiopathic pulmonary fibrosis: what is next?. <i>European Respiratory Review</i> , 2019, 28, 190021.	3.0	157
2173	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
2174	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
2176	Pharmacological management. , 0, , 196-217.		1
2177	Progression of fibrosing interstitial lung disease. <i>Respiratory Research</i> , 2020, 21, 32.	1.4	94
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2386	The evaluation of disease severity/staging for prognosis. , 0, , 97-105.		1
2387	Acute exacerbations. , 0, , 143-150.		1
2388	Symptom management: dyspnoea and cough. , 0, , 218-229.		0
2389	Key ongoing issues in trial design. , 0, , 253-259.		0
2390	Perspectives for the future. , 0, , 260-274.		1
2393	IPF: definition, severity and impact of pulmonary exacerbations. , 0, , 58-65.		0
2394	Coexistent COPD and ILD. , 0, , 109-120.		1
2395	Patterns of cardiopulmonary response to exercise in fibrotic ILD. , 0, , 128-145.		0
2396	Ways to improve the diagnosis and treatment of interstitial lung disease associated with systemic sclerosis in the Siberian Federal District (materials of the advisory board of rheumatologists and) Tj ETQq0 0 0 rgBTQ Overlock10 Tf 50 1		0
2399	Differentiating combined pulmonary fibrosis and emphysema from pure emphysema: utility of late gadolinium-enhanced MRI. <i>European Radiology Experimental</i> , 2020, 4, 61.	1.7	3
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#	ARTICLE	IF	CITATIONS
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#	ARTICLE	IF	CITATIONS
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