Luca Richeldi

List of Publications by Year
in descending order

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312 papers	39,367 citations	13827 67 h-index	2883 190 g-index
320 all docs	320 docs citations	320 times ranked	22064 citing authors

#	Article	IF	Citations
1	An Official ATS/ERS/JRS/ALAT Statement: Idiopathic Pulmonary Fibrosis: Evidence-based Guidelines for Diagnosis and Management. American Journal of Respiratory and Critical Care Medicine, 2011, 183, 788-824.	2.5	6,033
2	Efficacy and Safety of Nintedanib in Idiopathic Pulmonary Fibrosis. New England Journal of Medicine, 2014, 370, 2071-2082.	13.9	3,351
3	An Official American Thoracic Society/European Respiratory Society Statement: Update of the International Multidisciplinary Classification of the Idiopathic Interstitial Pneumonias. American Journal of Respiratory and Critical Care Medicine, 2013, 188, 733-748.	2.5	3,134
4	Diagnosis of Idiopathic Pulmonary Fibrosis. An Official ATS/ERS/JRS/ALAT Clinical Practice Guideline. American Journal of Respiratory and Critical Care Medicine, 2018, 198, e44-e68.	2.5	2,678
5	An Official ATS/ERS/JRS/ALAT Clinical Practice Guideline: Treatment of Idiopathic Pulmonary Fibrosis. An Update of the 2011 Clinical Practice Guideline. American Journal of Respiratory and Critical Care Medicine, 2015, 192, e3-e19.	2.5	1,521
6	Nintedanib in Progressive Fibrosing Interstitial Lung Diseases. New England Journal of Medicine, 2019, 381, 1718-1727.	13.9	1,338
7	Idiopathic pulmonary fibrosis. Lancet, The, 2017, 389, 1941-1952.	6.3	1,199
8	Acute Exacerbation of Idiopathic Pulmonary Fibrosis. An International Working Group Report. American Journal of Respiratory and Critical Care Medicine, 2016, 194, 265-275.	2.5	1,006
9	Efficacy of a Tyrosine Kinase Inhibitor in Idiopathic Pulmonary Fibrosis. New England Journal of Medicine, 2011, 365, 1079-1087.	13.9	930
10	The Role of Chest Imaging in Patient Management During the COVID-19 Pandemic. Chest, 2020, 158, 106-116.	0.4	832
11	An official European Respiratory Society/American Thoracic Society research statement: interstitial pneumonia with autoimmune features. European Respiratory Journal, 2015, 46, 976-987.	3.1	803
12	Idiopathic pulmonary fibrosis. Nature Reviews Disease Primers, 2017, 3, 17074.	18.1	786
13	Idiopathic Pulmonary Fibrosis (an Update) and Progressive Pulmonary Fibrosis in Adults: An Official ATS/ERS/JRS/ALAT Clinical Practice Guideline. American Journal of Respiratory and Critical Care Medicine, 2022, 205, e18-e47.	2.5	780
14	Diagnostic criteria for idiopathic pulmonary fibrosis: a Fleischner Society White Paper. Lancet Respiratory Medicine,the, 2018, 6, 138-153.	5.2	739
15	The Role of Chest Imaging in Patient Management during the COVID-19 Pandemic: A Multinational Consensus Statement from the Fleischner Society. Radiology, 2020, 296, 172-180.	3.6	721
16	Official American Thoracic Society/Infectious Diseases Society of America/Centers for Disease Control and Prevention Clinical Practice Guidelines: Diagnosis of Tuberculosis in Adults and Children. Clinical Infectious Diseases, 2017, 64, e1-e33.	2.9	501
17	Official American Thoracic Society/Infectious Diseases Society of America/Centers for Disease Control and Prevention Clinical Practice Guidelines: Diagnosis of Tuberculosis in Adults and Children. Clinical Infectious Diseases, 2017, 64, 111-115.	2.9	492
18	Use in routine clinical practice of two commercial blood tests for diagnosis of infection with Mycobacterium tuberculosis: a prospective study. Lancet, The, 2006, 367, 1328-1334.	6.3	468

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19	Treatment of Idiopathic Pulmonary Fibrosis With Ambrisentan. Annals of Internal Medicine, 2013, 158, 641.	2.0	437
20	Idiopathic pulmonary fibrosis: pathogenesis and management. Respiratory Research, 2018, 19, 32.	1.4	339
21	Nintedanib in patients with idiopathic pulmonary fibrosis: Combined evidence from the TOMORROW and INPULSISÂ® trials. Respiratory Medicine, 2016, 113, 74-79.	1.3	335
22	Nintedanib in patients with progressive fibrosing interstitial lung diseasesâ€"subgroup analyses by interstitial lung disease diagnosis in the INBUILD trial: a randomised, double-blind, placebo-controlled, parallel-group trial. Lancet Respiratory Medicine,the, 2020, 8, 453-460.	5.2	331
23	An Update on the Diagnosis of Tuberculosis Infection. American Journal of Respiratory and Critical Care Medicine, 2006, 174, 736-742.	2.5	287
24	Interstitial lung abnormalities detected incidentally on CT: a Position Paper from the Fleischner Society. Lancet Respiratory Medicine,the, 2020, 8, 726-737.	5.2	279
25	Routine Hospital Use of a New Commercial Whole Blood Interferon-Î³ Assay for the Diagnosis of Tuberculosis Infection. American Journal of Respiratory and Critical Care Medicine, 2005, 172, 631-635.	2.5	240
26	Prevalence and prognosis of unclassifiable interstitial lung disease. European Respiratory Journal, 2013, 42, 750-757.	3.1	238
27	Efficacy of Nintedanib in Idiopathic Pulmonary Fibrosis across Prespecified Subgroups in INPULSIS. American Journal of Respiratory and Critical Care Medicine, 2016, 193, 178-185.	2.5	209
28	Nintedanib plus Sildenafil in Patients with Idiopathic Pulmonary Fibrosis. New England Journal of Medicine, 2018, 379, 1722-1731.	13.9	207
29	Progressive fibrosing interstitial lung disease: clinical uncertainties, consensus recommendations, and research priorities. Lancet Respiratory Medicine,the, 2020, 8, 925-934.	5.2	198
30	Nintedanib with Add-on Pirfenidone in Idiopathic Pulmonary Fibrosis. Results of the INJOURNEY Trial. American Journal of Respiratory and Critical Care Medicine, 2018, 197, 356-363.	2.5	193
31	Nintedanib in patients with idiopathic pulmonary fibrosis and preserved lung volume. Thorax, 2017, 72, 340-346.	2.7	191
32	Identification of Diagnostic Criteria for Chronic Hypersensitivity Pneumonitis. An International Modified Delphi Survey. American Journal of Respiratory and Critical Care Medicine, 2018, 197, 1036-1044.	2.5	174
33	Suspected acute exacerbation of idiopathic pulmonary fibrosis as an outcome measure in clinical trials. Respiratory Research, 2013, 14, 73.	1.4	173
34	Effect of Recombinant Human Pentraxin 2 vs Placebo on Change in Forced Vital Capacity in Patients With Idiopathic Pulmonary Fibrosis. JAMA - Journal of the American Medical Association, 2018, 319, 2299.	3.8	170
35	A Standardized Diagnostic Ontology for Fibrotic Interstitial Lung Disease. An International Working Group Perspective. American Journal of Respiratory and Critical Care Medicine, 2017, 196, 1249-1254.	2.5	166
36	Pamrevlumab, an anti-connective tissue growth factor therapy, for idiopathic pulmonary fibrosis (PRAISE): a phase 2, randomised, double-blind, placebo-controlled trial. Lancet Respiratory Medicine,the, 2020, 8, 25-33.	5.2	165

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37	Relative versus absolute change in forced vital capacity in idiopathic pulmonary fibrosis. Thorax, 2012, 67, 407-411.	2.7	160
38	Design of the PF-ILD trial: a double-blind, randomised, placebo-controlled phase III trial of nintedanib in patients with progressive fibrosing interstitial lung disease. BMJ Open Respiratory Research, 2017, 4, e000212.	1.2	151
39	CT staging and monitoring of fibrotic interstitial lung diseases in clinical practice and treatment trials: a Position Paper from the Fleischner society. Lancet Respiratory Medicine,the, 2015, 3, 483-496.	5.2	149
40	Effect of Nintedanib in Subgroups of Idiopathic Pulmonary Fibrosis by Diagnostic Criteria. American Journal of Respiratory and Critical Care Medicine, 2017, 195, 78-85.	2.5	147
41	Performance of Tests for Latent Tuberculosis in Different Groups of Immunocompromised Patients. Chest, 2009, 136, 198-204.	0.4	137
42	Idiopathic pulmonary fibrosis: Diagnosis, epidemiology and natural history. Respirology, 2016, 21, 427-437.	1.3	137
43	Time for a change: is idiopathic pulmonary fibrosis still idiopathic and only fibrotic?. Lancet Respiratory Medicine,the, 2018, 6, 154-160.	5.2	137
44	Corticosteroids for idiopathic pulmonary fibrosis. The Cochrane Library, 2003, , CD002880.	1.5	133
45	Performance of Commercial Blood Tests for the Diagnosis of Latent Tuberculosis Infection in Children and Adolescents. Pediatrics, 2009, 123, e419-e424.	1.0	132
46	T Cellâ€“Based Tracking of Multidrug Resistant Tuberculosis Infection after Brief Exposure. American Journal of Respiratory and Critical Care Medicine, 2004, 170, 288-295.	2.5	131
47	Interaction of genetic and exposure factors in the prevalence of berylliosis. , 1997, 32, 337-340.		128
48	Combination therapy: the future of management for idiopathic pulmonary fibrosis?. Lancet Respiratory Medicine,the, 2014, 2, 933-942.	5.2	128
49	Bronchoalveolar Lavage Enzyme-linked Immunospot for a Rapid Diagnosis of Tuberculosis. American Journal of Respiratory and Critical Care Medicine, 2009, 180, 666-673.	2.5	125
50	Safety and survival data in patients with idiopathic pulmonary fibrosis treated with nintedanib: pooled data from six clinical trials. BMJ Open Respiratory Research, 2019, 6, e000397.	1.2	121
51	Safety, tolerability and appropriate use of nintedanib in idiopathic pulmonary fibrosis. Respiratory Research, 2015, 16, 116.	1.4	114
52	A multicentre evaluation of the accuracy and performance of IP-10 for the diagnosis of infection with M. tuberculosis. Tuberculosis, 2011, 91, 260-267.	0.8	113
53	Acute exacerbations in the INPULSIS trials of nintedanib in idiopathic pulmonary fibrosis. European Respiratory Journal, 2017, 49, 1601339.	3.1	109
54	Autophagy inhibition-mediated epithelialâ€“mesenchymal transition augments local myofibroblast differentiation in pulmonary fibrosis. Cell Death and Disease, 2019, 10, 591.	2.7	107

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55	Paracrine signalling during ZEB1-mediated epithelialâ€“mesenchymal transition augments local myofibroblast differentiation in lung fibrosis. Cell Death and Differentiation, 2019, 26, 943-957.	5.0	104
56	Non-steroid agents for idiopathic pulmonary fibrosis. The Cochrane Library, 2010, , CD003134.	1.5	103
57	Computed Tomographic Biomarkers in Idiopathic Pulmonary Fibrosis. The Future of Quantitative Analysis. American Journal of Respiratory and Critical Care Medicine, 2019, 199, 12-21.	2.5	102
58	Connective tissue disease related interstitial lung diseases and idiopathic pulmonary fibrosis: provisional core sets of domains and instruments for use in clinical trials. Thorax, 2014, 69, 436-444.	2.7	100
59	Nanoscale dysregulation of collagen structure-function disrupts mechano-homeostasis and mediates pulmonary fibrosis. ELife, 2018, 7, .	2.8	99
60	Optimising experimental research in respiratory diseases: an ERS statement. European Respiratory Journal, 2018, 51, 1702133.	3.1	98
61	Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2012, 186, 712-715.	2.5	92
62	Idiopathic Pulmonary Fibrosis: CT and Risk of Death. Radiology, 2014, 273, 570-579.	3.6	85
63	Prevalence and clinical significance of circulating autoantibodies in idiopathic pulmonary fibrosis. Respiratory Medicine, 2013, 107, 249-255.	1.3	84
64	Design of the INPULSISâ„¢ trials: Two phase 3 trials of nintedanib in patients with idiopathic pulmonary fibrosis. Respiratory Medicine, 2014, 108, 1023-1030.	1.3	82
65	Treatment of idiopathic pulmonary fibrosis: a network meta-analysis. BMC Medicine, 2016, 14, 18.	2.3	79
66	The diagnosis of idiopathic pulmonary fibrosis: current and future approaches. Lancet Respiratory Medicine,the, 2017, 5, 61-71.	5.2	79
67	Trial of a Preferential Phosphodiesterase 4B Inhibitor for Idiopathic Pulmonary Fibrosis. New England Journal of Medicine, 2022, 386, 2178-2187.	13.9	77
68	Diagnostic accuracy of a clinical diagnosis of idiopathic pulmonary fibrosis: an international caseâ€“cohort study. European Respiratory Journal, 2017, 50, 1700936.	3.1	75
69	Neglected evidence in idiopathic pulmonary fibrosis and the importance of early diagnosis and treatment. European Respiratory Review, 2014, 23, 106-110.	3.0	74
70	Idiopathic pulmonary fibrosis: Recent advances on pharmacological therapy. , 2015, 152, 18-27.		74
71	Pharmacological management of progressive-fibrosing interstitial lung diseases: a review of the current evidence. European Respiratory Review, 2018, 27, 180074.	3.0	73
72	Long-term treatment with recombinant human pentraxin 2 protein in patients with idiopathic pulmonary fibrosis: an open-label extension study. Lancet Respiratory Medicine,the, 2019, 7, 657-664.	5.2	73

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73	Three-dimensional characterization of fibroblast foci in idiopathic pulmonary fibrosis. JCI Insight, 2016, 1, .	2.3	73
74	The revised ATS/ERS/JRS/ALAT diagnostic criteria for idiopathic pulmonary fibrosis (IPF) - practical implications. Respiratory Research, 2013, 14, S2.	1.4	72
75	Hot of the breath: Mortality as a primary end-point in IPF treatment trials: the best is the enemy of the good. Thorax, 2012, 67, 938-940.	2.7	71
76	Antacid therapy in idiopathic pulmonary fibrosis: more questions than answers?. Lancet Respiratory Medicine,the, 2017, 5, 591-598.	5.2	71
77	Post-COVID lung fibrosis: The tsunami that will follow the earthquake. Lung India, 2021, 38, 41.	0.3	69
78	Immunomodulatory agents for idiopathic pulmonary fibrosis. , 2003, , CD003134.		62
79	Efficacy and safety of nintedanib in patients with advanced idiopathic pulmonary fibrosis. BMC Pulmonary Medicine, 2020, 20, 3.	0.8	61
80	Diagnostic Likelihood Thresholds That Define a Working Diagnosis of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 1146-1153.	2.5	60
81	Precision Medicine: The New Frontier in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2016, 193, 1213-1218.	2.5	59
82	Long-term treatment of patients with idiopathic pulmonary fibrosis with nintedanib: results from the TOMORROW trial and its open-label extension. Thorax, 2018, 73, 581-583.	2.7	59
83	SAR156597 in idiopathic pulmonary fibrosis: a phase 2 placebo-controlled study (DRI11772). European Respiratory Journal, 2018, 52, 1801130.	3.1	59
84	Nintedanib in the treatment of idiopathic pulmonary fibrosis. Therapeutic Advances in Respiratory Disease, 2015, 9, 121-129.	1.0	57
85	X-ray Micro-Computed Tomography for Nondestructive Three-Dimensional (3D) X-ray Histology. American Journal of Pathology, 2019, 189, 1608-1620.	1.9	57
86	Treatment strategies for asthma: reshaping the concept of asthma management. Allergy, Asthma and Clinical Immunology, 2020, 16, 75.	0.9	55
87	Utility of a Molecular Classifier as a Complement to High-Resolution Computed Tomography to Identify Usual Interstitial Pneumonia. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 211-220.	2.5	55
88	Nintedanib for the treatment of idiopathic pulmonary fibrosis. Expert Opinion on Pharmacotherapy, 2018, 19, 167-175.	0.9	53
89	Estimation of the Prevalence of Progressive Fibrosing Interstitial Lung Diseases: Systematic Literature Review and Data from a Physician Survey. Advances in Therapy, 2021, 38, 854-867.	1.3	53
90	Chest CT Diagnosis and Clinical Management of Drug-related Pneumonitis in Patients Receiving Molecular Targeting Agents and Immune Checkpoint Inhibitors: A Position Paper from the Fleischner Society. Radiology, 2021, 298, 550-566.	3.6	53

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91	Chest CT Diagnosis and Clinical Management of Drug-Related Pneumonitis in Patients Receiving Molecular Targeting Agents and Immune Checkpoint Inhibitors. Chest, 2021, 159, 1107-1125.	0.4	53
92	Current approaches to the management of idiopathic pulmonary fibrosis. Respiratory Medicine, 2017, 129, 24-30.	1.3	52
93	Lung cancer in scleroderma: Results from an Italian rheumatologic center and review of the literature. Autoimmunity Reviews, 2013, 12, 374-379.	2.5	50
94	The characterisation of interstitial lungÂdisease multidisciplinary team meetings:ÂaÂglobal study. ERJ Open Research, 2019, 5, 00209-2018.	1.1	49
95	A new era in idiopathic pulmonary fibrosis: considerations for future clinical trials. European Respiratory Journal, 2015, 46, 243-249.	3.1	48
96	The histone deacetylase inhibitor, romidepsin, as a potential treatment for pulmonary fibrosis. Oncotarget, 2017, 8, 48737-48754.	0.8	48
97	Pirfenidone. Nature Reviews Drug Discovery, 2011, 10, 489-490.	21.5	47
98	Nintedanib in progressive interstitial lung diseases: data from the whole INBUILD trial. European Respiratory Journal, 2022, 59, 2004538.	3.1	47
99	â€œVelcro-typeâ€•crackles predict specific radiologic features of fibrotic interstitial lung disease. BMC Pulmonary Medicine, 2018, 18, 103.	0.8	45
100	The 2018 Diagnosis of Idiopathic Pulmonary Fibrosis Guidelines: Surgical Lung Biopsy for Radiological Pattern of Probable Usual Interstitial Pneumonia Is Not Mandatory. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 1089-1092.	2.5	45
101	Idiopathic Pulmonary Fibrosis. Chest, 2018, 154, 1359-1370.	0.4	44
102	Sarcoidosis: Challenging Diagnostic Aspects of an Old Disease. American Journal of Medicine, 2012, 125, 118-125.	0.6	43
103	Idiopathic pulmonary fibrosis: diagnostic pitfalls and therapeutic challenges. Multidisciplinary Respiratory Medicine, 2012, 7, 42.	0.6	42
104	Fibulin-1 Predicts Disease Progression in Patients With Idiopathic Pulmonary Fibrosis. Chest, 2014, 146, 1055-1063.	0.4	42
105	Statin Therapy and Outcomes in Trials of Nintedanib in Idiopathic Pulmonary Fibrosis. Respiration, 2018, 95, 317-326.	1.2	42
106	Pamrevlumab for the treatment of idiopathic pulmonary fibrosis. Expert Opinion on Investigational Drugs, 2020, 29, 771-777.	1.9	40
107	The big clinical trials in idiopathic pulmonary fibrosis. Current Opinion in Pulmonary Medicine, 2012, 18, 428-432.	1.2	38
108	Treating heart failure with preserved ejection fraction: learning from pulmonary fibrosis. European Journal of Heart Failure, 2018, 20, 1385-1391.	2.9	38

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109	Idiopathic pulmonary fibrosis in BRIC countries: the cases of Brazil, Russia, India, and China. BMC Medicine, 2015, 13, 237.	2.3	34
110	Structured reporting for fibrosing lung disease: a model shared by radiologist and pulmonologist. Radiologia Medica, 2018, 123, 245-253.	4.7	34
111	Lung function outcomes in the INPULSISÂ® trials of nintedanib in idiopathic pulmonary fibrosis. Respiratory Medicine, 2019, 146, 42-48.	1.3	34
112	A Systematically Derived Exposure Assessment Instrument for Chronic Hypersensitivity Pneumonitis. Chest, 2020, 157, 1506-1512.	0.4	33
113	Nintedanib and immunomodulatory therapies in progressive fibrosing interstitial lung diseases. Respiratory Research, 2021, 22, 84.	1.4	33
114	Rising to the Challenge of COVID-19: Advice for Pulmonary and Critical Care and an Agenda for Research. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 1019-1022.	2.5	32
115	Treatments for Idiopathic Pulmonary Fibrosis. New England Journal of Medicine, 2014, 371, 781-784.	13.9	31
116	Improved pulmonary function following pirfenidone treatment in a patient with progressive interstitial lung disease associated with systemic sclerosis. Lung India, 2015, 32, 50.	0.3	31
117	Pseudohypoxic HIF pathway activation dysregulates collagen structure-function in human lung fibrosis. ELife, 2022, 11, .	2.8	31
118	Differing severities of acute exacerbations of idiopathic pulmonary fibrosis (IPF): insights from the INPULSISÂ® trials. Respiratory Research, 2019, 20, 71.	1.4	30
119	Stability or improvement in forced vital capacity with nintedanib in patients with idiopathic pulmonary fibrosis. European Respiratory Journal, 2018, 52, 1702593.	3.1	29
120	Lung ultrasonography for early management of patients with respiratory symptoms during COVID-19 pandemic. Journal of Ultrasound, 2020, 23, 449-456.	0.7	29
121	Pirfenidone in idiopathic pulmonary fibrosis: the CAPACITY program. Expert Review of Respiratory Medicine, 2011, 5, 473-481.	1.0	28
122	Cross-Disciplinary Collaboration in Connective Tissue Disease-Related Lung Disease. Seminars in Respiratory and Critical Care Medicine, 2014, 35, 159-165.	0.8	28
123	Novel drug targets for idiopathic pulmonary fibrosis. Expert Review of Respiratory Medicine, 2016, 10, 393-405.	1.0	27
124	Idiopathic Pulmonary Fibrosis: Recent Trials and Current Drug Therapy. Respiration, 2013, 86, 353-363.	1.2	26
125	Acute myocardial infarction versus other cardiovascular events in community-acquired pneumonia. ERJ Open Research, 2015, 1, 00020-2015.	1.1	26
126	Detection and Early Referral of Patients With Interstitial Lung Abnormalities. Chest, 2022, 161, 470-482.	0.4	26

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127	A global registry for idiopathic pulmonary fibrosis: the time is now. European Respiratory Journal, 2014, 44, 273-276.	3.1	25
128	Existing and emerging biomarkers for disease progression in idiopathic pulmonary fibrosis. Expert Review of Respiratory Medicine, 2019, 13, 39-51.	1.0	25
129	The complex interrelationships between chronic lung and liver disease: a review. Journal of Viral Hepatitis, 2010, 17, 381-390.	1.0	24
130	The Reply. American Journal of Medicine, 2013, 126, e19.	0.6	24
131	Safety and tolerability of nintedanib for the treatment of idiopathic pulmonary fibrosis in routine UK clinical practice. ERJ Open Research, 2018, 4, 00049-2018.	1.1	24
132	Bidirectional epithelialâ€“mesenchymal crosstalk provides self-sustaining profibrotic signals in pulmonary fibrosis. Journal of Biological Chemistry, 2021, 297, 101096.	1.6	24
133	Levels of circulating endothelial cells are low in idiopathic pulmonary fibrosis and are further reduced by anti-fibrotic treatments. BMC Medicine, 2015, 13, 277.	2.3	23
134	Diagnosing idiopathic pulmonary fibrosis in 2018: bridging recommendations made by experts serving different societies. European Respiratory Journal, 2018, 52, 1801485.	3.1	23
135	Update in Pulmonary Fibrosis 2018. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 292-300.	2.5	23
136	Fibrotic Hypersensitivity Pneumonitis: Diagnosis and Management. Lung, 2020, 198, 429-440.	1.4	23
137	Paracrine SPARC signaling dysregulates alveolar epithelial barrier integrity and function in lung fibrosis. Cell Death Discovery, 2020, 6, 54.	2.0	23
138	Residual respiratory impairment after COVID-19 pneumonia. BMC Pulmonary Medicine, 2021, 21, 241.	0.8	23
139	Assessing the treatment effect from multiple trials in idiopathic pulmonary fibrosis. European Respiratory Review, 2012, 21, 147-151.	3.0	22
140	Mindfulness-based stress reduction in patients with interstitial lung diseases: a pilot, single-centre observational study on safety and efficacy. BMJ Open Respiratory Research, 2015, 2, e000065.	1.2	22
141	Idiopathic Pulmonary Fibrosis: Molecular Endotypes of Fibrosis Stratifying Existing and Emerging Therapies. Respiration, 2017, 93, 379-395.	1.2	22
142	Outcomes following decline in forced vital capacity in patients with idiopathic pulmonary fibrosis: Results from the INPULSIS and INPULSIS-ON trials of nintedanib. Respiratory Medicine, 2019, 156, 20-25.	1.3	22
143	No relevant pharmacokinetic drugâ€“drug interaction between nintedanib and pirfenidone. European Respiratory Journal, 2019, 53, 1801060.	3.1	22
144	Emerging drugs for idiopathic pulmonary fibrosis. Expert Opinion on Emerging Drugs, 2011, 16, 341-362.	1.0	21

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145	The safety of new drug treatments for idiopathic pulmonary fibrosis. Expert Opinion on Drug Safety, 2016, 15, 1483-1489.	1.0	21
146	Effects of nintedanib in patients with idiopathic pulmonary fibrosis by GAP stage. ERJ Open Research, 2019, 5, 00127-2018.	1.1	21
147	Safety of Nintedanib in Patients with Idiopathic Pulmonary Fibrosis: Global Pharmacovigilance Data. Advances in Therapy, 2020, 37, 4209-4219.	1.3	21
148	Telemedicine-enabled Accelerated Discharge of Patients Hospitalized with COVID-19 to Isolation in Repurposed Hotel Rooms. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 508-510.	2.5	21
149	Current Diagnosis and Management of Hypersensitivity Pneumonitis. Tuberculosis and Respiratory Diseases, 2020, 83, 122.	0.7	20
150	Environmental Triggers and Susceptibility Factors in Idiopathic Granulomatous Diseases. Seminars in Respiratory and Critical Care Medicine, 2008, 29, 610-619.	0.8	18
151	Investigational drugs for idiopathic pulmonary fibrosis. Expert Opinion on Investigational Drugs, 2017, 26, 1019-1031.	1.9	18
152	Mesenchymal Stromal Cell Secretome for Post-COVID-19 Pulmonary Fibrosis: A New Therapy to Treat the Long-Term Lung Sequelae?. Cells, 2021, 10, 1203.	1.8	18
153	Aortic pulse wave velocity measurement in systemic sclerosis patients. Reumatismo, 2012, 64, 360-7.	0.4	17
154	Alpha-1 antitrypsin deficiency as a common treatable mechanism in chronic respiratory disorders and for conditions different from pulmonary emphysema? A commentary on the new European Respiratory Society statement. Multidisciplinary Respiratory Medicine, 2018, 13, 39.	0.6	17
155	CC-90001, a c-Jun N-terminal kinase (JNK) inhibitor, in patients with pulmonary fibrosis: design of a phase 2, randomised, placebo-controlled trial. BMJ Open Respiratory Research, 2022, 9, e001060.	1.2	17
156	Time for Prevention of Idiopathic Pulmonary Fibrosis Exacerbation. Annals of the American Thoracic Society, 2015, 12, S181-S185.	1.5	17
157	Long-term evaluation of the safety and efficacy of recombinant human pentraxin-2 (rhPTX-2) in patients with idiopathic pulmonary fibrosis (IPF): an open-label extension study. Respiratory Research, 2022, 23, .	1.4	17
158	Exploring the immune response against Mycobacterium tuberculosis for a better diagnosis of the infection. Archivum Immunologiae Et Therapiae Experimentalis, 2009, 57, 425-433.	1.0	16
159	Recommendations on treatment for IPF. Respiratory Research, 2013, 14, S6.	1.4	16
160	COVID-19 Vaccine in Patients with Exacerbation of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 219-221.	2.5	16
161	Time taken from primary care referral to a specialist centre diagnosis of idiopathic pulmonary fibrosis: an opportunity to improve patient outcomes?. ERJ Open Research, 2020, 6, 00120-2020.	1.1	15
162	Multidisciplinary Evaluation of Interstitial Lung Diseases: New Opportunities Linked to Rheumatologist Involvement. Diagnostics, 2020, 10, 664.	1.3	15

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163	Integrating Clinical Probability into the Diagnostic Approach to Idiopathic Pulmonary Fibrosis: An International Working Group Perspective. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 247-259.	2.5	15
164	The role of biomarkers in low respiratory tract infections. European Journal of Internal Medicine, 2012, 23, 429-435.	1.0	14
165	A Quantitative Proteomic Approach to Identify Significantly Altered Protein Networks in the Serum of Patients with Lymphangioleiomyomatosis (LAM). PLoS ONE, 2014, 9, e105365.	1.1	14
166	Agreement between chest ultrasonography and chest X-ray in patients who have undergone thoracic surgery: preliminary results. Multidisciplinary Respiratory Medicine, 2019, 14, 9.	0.6	14
167	Reticulation Is a Risk Factor of Progressive Subpleural Nonfibrotic Interstitial Lung Abnormalities. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 178-185.	2.5	14
168	Role of the Quantiferon-TB Test in Ruling Out Pleural Tuberculosis: A Multi-Centre Study. International Journal of Immunopathology and Pharmacology, 2011, 24, 159-165.	1.0	13
169	New treatment directions for IPF: current status of ongoing and upcoming clinical trials. Expert Review of Respiratory Medicine, 2017, 11, 533-548.	1.0	13
170	Individualizing duration of antibiotic therapy in community-acquired pneumonia. Pulmonary Pharmacology and Therapeutics, 2017, 45, 191-201.	1.1	13
171	Validation of multidisciplinary diagnosis in IPF. Lancet Respiratory Medicine,the, 2018, 6, 88-89.	5.2	13
172	Subclinical Interstitial Lung Abnormalities: Lumping and Splitting Revisited. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 121-123.	2.5	13
173	Current and Future Idiopathic Pulmonary Fibrosis Therapy. American Journal of the Medical Sciences, 2019, 357, 370-373.	0.4	13
174	Antibody-based therapies for idiopathic pulmonary fibrosis. Expert Opinion on Biological Therapy, 2020, 20, 779-786.	1.4	13
175	Using ELISpot technology to improve the diagnosis of tuberculosis infection: from the bench to the T-SPOT.TBassay. Expert Review of Respiratory Medicine, 2008, 2, 253-260.	1.0	12
176	What if we made stratified medicine work for patients?. Lancet Respiratory Medicine,the, 2016, 4, 8-10.	5.2	12
177	Molecular Testing in EBUS-TBNA Specimens of Lung Adenocarcinoma: A Study of Concordance Between Cell Block Method and Liquid-Based Cytology in Appraising Sample Cellularity and EGFR Mutations. Molecular Diagnosis and Therapy, 2018, 22, 723-728.	1.6	12
178	Quantitative analysis of lung sounds for monitoring idiopathic pulmonary fibrosis:Âa prospective pilot study. European Respiratory Journal, 2019, 53, 1802093.	3.1	12
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