Luca Richeldi

List of Publications by Citations

248 papers	25,164 citations	59 h-index	157 g-index
320 ext. papers	32,896 ext. citations	9.8 avg, IF	6.82 L-index

#	Paper	IF	Citations
248	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	10.2	4665
247	Efficacy and safety of nintedanib in idiopathic pulmonary fibrosis. *New England Journal of Medicine*, **2014**, 370, 2071-82	59.2	2337
246	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, 733-48	10.2	2176
245	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	10.2	1426
244	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, e3-19	10.2	1122
243	Efficacy of a tyrosine kinase inhibitor in idiopathic pulmonary fibrosis. *New England Journal of Medicine*, **2011**, 365, 1079-87	59.2	728
242	Acute Exacerbation of Idiopathic Pulmonary Fibrosis. An International Working Group Report. *American Journal of Respiratory and Critical Care Medicine*, **2016**, 194, 265-75	10.2	653
241	Idiopathic pulmonary fibrosis. *Lancet, The*, **2017**, 389, 1941-1952	40	617
240	Nintedanib in Progressive Fibrosing Interstitial Lung Diseases. *New England Journal of Medicine*, **2019**, 381, 1718-1727	59.2	585
239	An official European Respiratory Society/American Thoracic Society research statement: interstitial pneumonia with autoimmune features. *European Respiratory Journal*, **2015**, 46, 976-87	13.6	541
238	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	20.5	471
237	Diagnostic criteria for idiopathic pulmonary fibrosis: a Fleischner Society White Paper. *Lancet Respiratory Medicine,the*, **2018**, 6, 138-153	35.1	452
236	Idiopathic pulmonary fibrosis. *Nature Reviews Disease Primers*, **2017**, 3, 17074	51.1	395
235	The Role of Chest Imaging in Patient Management During the COVID-19 Pandemic: A Multinational Consensus Statement From the Fleischner Society. *Chest*, **2020**, 158, 106-116	5.3	384
234	Use in routine clinical practice of two commercial blood tests for diagnosis of infection with Mycobacterium tuberculosis: a prospective study. *Lancet, The*, **2006**, 367, 1328-34	40	382
233	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	11.6	336
232	Treatment of idiopathic pulmonary fibrosis with ambrisentan: a parallel, randomized trial. *Annals of Internal Medicine*, **2013**, 158, 641-9	8	327

231	Nintedanib in patients with idiopathic pulmonary fibrosis: Combined evidence from the TOMORROW and INPULSIS( ) trials. *Respiratory Medicine*, **2016**, 113, 74-9	4.6	245
230	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	11.6	237
229	An update on the diagnosis of tuberculosis infection. *American Journal of Respiratory and Critical Care Medicine*, **2006**, 174, 736-42	10.2	229
228	Routine hospital use of a new commercial whole blood interferon-gamma assay for the diagnosis of tuberculosis infection. *American Journal of Respiratory and Critical Care Medicine*, **2005**, 172, 631-5	10.2	207
227	Idiopathic pulmonary fibrosis: pathogenesis and management. *Respiratory Research*, **2018**, 19, 32	7.3	174
226	Efficacy of Nintedanib in Idiopathic Pulmonary Fibrosis across Prespecified Subgroups in INPULSIS. *American Journal of Respiratory and Critical Care Medicine*, **2016**, 193, 178-85	10.2	169
225	Prevalence and prognosis of unclassifiable interstitial lung disease. *European Respiratory Journal*, **2013**, 42, 750-7	13.6	164
224	Nintedanib in patients with progressive fibrosing interstitial lung diseases-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	35.1	154
223	Suspected acute exacerbation of idiopathic pulmonary fibrosis as an outcome measure in clinical trials. *Respiratory Research*, **2013**, 14, 73	7.3	136
222	Nintedanib plus Sildenafil in Patients with Idiopathic Pulmonary Fibrosis. *New England Journal of Medicine*, **2018**, 379, 1722-1731	59.2	135
221	Nintedanib in patients with idiopathic pulmonary fibrosis and preserved lung volume. *Thorax*, **2017**, 72, 340-346	7.3	130
220	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	10.2	128
219	Performance of tests for latent tuberculosis in different groups of immunocompromised patients. *Chest*, **2009**, 136, 198-204	5.3	119
218	Relative versus absolute change in forced vital capacity in idiopathic pulmonary fibrosis. *Thorax*, **2012**, 67, 407-11	7.3	117
217	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	10.2	109
216	Effect of Nintedanib in Subgroups of Idiopathic Pulmonary Fibrosis by Diagnostic Criteria. *American Journal of Respiratory and Critical Care Medicine*, **2017**, 195, 78-85	10.2	108
215	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	5.6	107
214	Effect of Recombinant Human Pentraxin 2 vs Placebo on Change in Forced Vital Capacity in Patients With Idiopathic Pulmonary Fibrosis: A Randomized Clinical Trial. *JAMA - Journal of the American Medical Association*, **2018**, 319, 2299-2307	27.4	107

213	Performance of commercial blood tests for the diagnosis of latent tuberculosis infection in children and adolescents. *Pediatrics*, **2009**, 123, e419-24	7.4	106
212	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	10.2	105
211	T cell-based tracking of multidrug resistant tuberculosis infection after brief exposure. *American Journal of Respiratory and Critical Care Medicine*, **2004**, 170, 288-95	10.2	104
210	Bronchoalveolar lavage enzyme-linked immunospot for a rapid diagnosis of tuberculosis: a Tuberculosis Network European Trialsgroup study. *American Journal of Respiratory and Critical Care Medicine*, **2009**, 180, 666-73	10.2	103
209	Interaction of genetic and exposure factors in the prevalence of berylliosis. *American Journal of Industrial Medicine*, **1997**, 32, 337-40	2.7	103
208	Combination therapy: the future of management for idiopathic pulmonary fibrosis?. *Lancet Respiratory Medicine,the*, **2014**, 2, 933-942	35.1	97
207	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, 483-96	35.1	95
206	Corticosteroids for idiopathic pulmonary fibrosis. *The Cochrane Library*, **2003**, CD002880	5.2	95
205	Idiopathic pulmonary fibrosis: Diagnosis, epidemiology and natural history. *Respirology*, **2016**, 21, 427-37	3.6	90
204	A multicentre evaluation of the accuracy and performance of IP-10 for the diagnosis of infection with M. tuberculosis. *Tuberculosis*, **2011**, 91, 260-7	2.6	86
203	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	35.1	81
202	Safety, tolerability and appropriate use of nintedanib in idiopathic pulmonary fibrosis. *Respiratory Research*, **2015**, 16, 116	7.3	79
201	Idiopathic pulmonary fibrosis: lung function is a clinically meaningful endpoint for phase III trials. *American Journal of Respiratory and Critical Care Medicine*, **2012**, 186, 712-5	10.2	78
200	Interstitial lung abnormalities detected incidentally on CT: a Position Paper from the Fleischner Society. *Lancet Respiratory Medicine,the*, **2020**, 8, 726-737	35.1	77
199	Progressive fibrosing interstitial lung disease: clinical uncertainties, consensus recommendations, and research priorities. *Lancet Respiratory Medicine,the*, **2020**, 8, 925-934	35.1	77
198	Time for a change: is idiopathic pulmonary fibrosis still idiopathic and only fibrotic?. *Lancet Respiratory Medicine,the*, **2018**, 6, 154-160	35.1	76
197	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, 428-36	7.3	75
196	Acute exacerbations in the INPULSIS trials of nintedanib in idiopathic pulmonary fibrosis. *European Respiratory Journal*, **2017**, 49,	13.6	74

195	Design of the INPULSIS trials: two phase 3 trials of nintedanib in patients with idiopathic pulmonary fibrosis. *Respiratory Medicine*, **2014**, 108, 1023-30	4.6	69
194	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	5.6	66
193	Idiopathic pulmonary fibrosis: CT and risk of death. *Radiology*, **2014**, 273, 570-9	20.5	66
192	Non-steroid agents for idiopathic pulmonary fibrosis. *The Cochrane Library*, **2010**, CD003134	5.2	66
191	Prevalence and clinical significance of circulating autoantibodies in idiopathic pulmonary fibrosis. *Respiratory Medicine*, **2013**, 107, 249-55	4.6	64
190	Idiopathic pulmonary fibrosis: Recent advances on pharmacological therapy. *Pharmacology & Therapeutics*, **2015**, 152, 18-27	13.9	61
189	Autophagy inhibition-mediated epithelial-mesenchymal transition augments local myofibroblast differentiation in pulmonary fibrosis. *Cell Death and Disease*, **2019**, 10, 591	9.8	59
188	Treatment of idiopathic pulmonary fibrosis: a network meta-analysis. *BMC Medicine*, **2016**, 14, 18	11.4	54
187	Optimising experimental research in respiratory diseases: an ERS statement. *European Respiratory Journal*, **2018**, 51,	13.6	53
186	The diagnosis of idiopathic pulmonary fibrosis: current and future approaches. *Lancet Respiratory Medicine,the*, **2017**, 5, 61-71	35.1	52
185	The revised ATS/ERS/JRS/ALAT diagnostic criteria for idiopathic pulmonary fibrosis (IPF)--practical implications. *Respiratory Research*, **2013**, 14 Suppl 1, S2	7.3	51
184	Diagnostic accuracy of a clinical diagnosis of idiopathic pulmonary fibrosis: an international case-cohort study. *European Respiratory Journal*, **2017**, 50,	13.6	50
183	Pharmacological management of progressive-fibrosing interstitial lung diseases: a review of the current evidence. *European Respiratory Review*, **2018**, 27,	9.8	50
182	Paracrine signalling during ZEB1-mediated epithelial-mesenchymal transition augments local myofibroblast differentiation in lung fibrosis. *Cell Death and Differentiation*, **2019**, 26, 943-957	12.7	49
181	Neglected evidence in idiopathic pulmonary fibrosis and the importance of early diagnosis and treatment. *European Respiratory Review*, **2014**, 23, 106-10	9.8	49
180	Nanoscale dysregulation of collagen structure-function disrupts mechano-homeostasis and mediates pulmonary fibrosis. *ELife*, **2018**, 7,	8.9	48
179	Precision Medicine: The New Frontier in Idiopathic Pulmonary Fibrosis. *American Journal of Respiratory and Critical Care Medicine*, **2016**, 193, 1213-8	10.2	47
178	Antacid therapy in idiopathic pulmonary fibrosis: more questions than answers?. *Lancet Respiratory Medicine,the*, **2017**, 5, 591-598	35.1	47

177	Three-dimensional characterization of fibroblast foci in idiopathic pulmonary fibrosis. *JCI Insight*, **2016**, 1,	9.9	47
176	Nintedanib in the treatment of idiopathic pulmonary fibrosis. *Therapeutic Advances in Respiratory Disease*, **2015**, 9, 121-9	4.9	46
175	Immunomodulatory agents for idiopathic pulmonary fibrosis. *Cochrane Database of Systematic Reviews*, **2003**, CD003134		44
174	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	35.1	42
173	SAR156597 in idiopathic pulmonary fibrosis: a phase 2 placebo-controlled study (DRI11772). *European Respiratory Journal*, **2018**, 52,	13.6	41
172	Lung cancer in scleroderma: results from an Italian rheumatologic center and review of the literature. *Autoimmunity Reviews*, **2013**, 12, 374-9	13.6	40
171	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	7.3	38
170	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	10.2	38
169	Current approaches to the management of idiopathic pulmonary fibrosis. *Respiratory Medicine*, **2017**, 129, 24-30	4.6	37
168	Nintedanib for the treatment of idiopathic pulmonary fibrosis. *Expert Opinion on Pharmacotherapy*, **2018**, 19, 167-175	4	36
167	The histone deacetylase inhibitor, romidepsin, as a potential treatment for pulmonary fibrosis. *Oncotarget*, **2017**, 8, 48737-48754	3.3	36
166	A new era in idiopathic pulmonary fibrosis: considerations for future clinical trials. *European Respiratory Journal*, **2015**, 46, 243-9	13.6	35
165	Idiopathic pulmonary fibrosis: diagnostic pitfalls and therapeutic challenges. *Multidisciplinary Respiratory Medicine*, **2012**, 7, 42	3	34
164	Diagnostic Likelihood Thresholds That Define a Working Diagnosis of Idiopathic Pulmonary Fibrosis. *American Journal of Respiratory and Critical Care Medicine*, **2019**, 200, 1146-1153	10.2	33
163	Fibulin-1 predicts disease progression in patients with idiopathic pulmonary fibrosis. *Chest*, **2014**, 146, 1055-1063	5.3	32
162	Sarcoidosis: challenging diagnostic aspects of an old disease. *American Journal of Medicine*, **2012**, 125, 118-25	2.4	32
161	Post-COVID lung fibrosis: The tsunami that will follow the earthquake. *Lung India*, **2021**, 38, S41-S47	1.1	30
160	Efficacy and safety of nintedanib in patients with advanced idiopathic pulmonary fibrosis. *BMC Pulmonary Medicine*, **2020**, 20, 3	3.5	29

159	Statin Therapy and Outcomes in Trials of Nintedanib in Idiopathic Pulmonary Fibrosis. *Respiration*, **2018**, 95, 317-326	3.7	29
158	Idiopathic pulmonary fibrosis in BRIC countries: the cases of Brazil, Russia, India, and China. *BMC Medicine*, **2015**, 13, 237	11.4	27
157	The big clinical trials in idiopathic pulmonary fibrosis. *Current Opinion in Pulmonary Medicine*, **2012**, 18, 428-32	3	27
156	Idiopathic pulmonary fibrosis: recent trials and current drug therapy. *Respiration*, **2013**, 86, 353-63	3.7	26
155	Idiopathic Pulmonary Fibrosis: Prospective, Case-Controlled Study of Natural History and Circulating Biomarkers. *Chest*, **2018**, 154, 1359-1370	5.3	25
154	X-ray Micro-Computed Tomography for Nondestructive Three-Dimensional (3D) X-ray Histology. *American Journal of Pathology*, **2019**, 189, 1608-1620	5.8	24
153	The characterisation of interstitial lung disease multidisciplinary team meetings: a global study. *ERJ Open Research*, **2019**, 5,	3.5	24
152	Improved pulmonary function following pirfenidone treatment in a patient with progressive interstitial lung disease associated with systemic sclerosis. *Lung India*, **2015**, 32, 50-2	1.1	24
151	Treating heart failure with preserved ejection fraction: learning from pulmonary fibrosis. *European Journal of Heart Failure*, **2018**, 20, 1385-1391	12.3	24
150	Cross-disciplinary collaboration in connective tissue disease-related lung disease. *Seminars in Respiratory and Critical Care Medicine*, **2014**, 35, 159-65	3.9	23
149	The reply. *American Journal of Medicine*, **2013**, 126, e19	2.4	23
148	Pirfenidone in idiopathic pulmonary fibrosis: the CAPACITY program. *Expert Review of Respiratory Medicine*, **2011**, 5, 473-81	3.8	23
147	"Velcro-type" crackles predict specific radiologic features of fibrotic interstitial lung disease. *BMC Pulmonary Medicine*, **2018**, 18, 103	3.5	22
146	Treatments for idiopathic pulmonary fibrosis. *New England Journal of Medicine*, **2014**, 371, 783	59.2	22
145	Levels of circulating endothelial cells are low in idiopathic pulmonary fibrosis and are further reduced by anti-fibrotic treatments. *BMC Medicine*, **2015**, 13, 277	11.4	21
144	Lung function outcomes in the INPULSIS trials of nintedanib in idiopathic pulmonary fibrosis. *Respiratory Medicine*, **2019**, 146, 42-48	4.6	20
143	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	10.2	20
142	Stability or improvement in forced vital capacity with nintedanib in patients with idiopathic pulmonary fibrosis. *European Respiratory Journal*, **2018**, 52,	13.6	20

141	Emerging drugs for idiopathic pulmonary fibrosis. *Expert Opinion on Emerging Drugs*, **2011**, 16, 341-62	3.7	19
140	The complex interrelationships between chronic lung and liver disease: a review. *Journal of Viral Hepatitis*, **2010**, 17, 381-90	3.4	19
139	A Systematically Derived Exposure Assessment Instrument for Chronic Hypersensitivity Pneumonitis. *Chest*, **2020**, 157, 1506-1512	5.3	18
138	Acute myocardial infarction other cardiovascular events in community-acquired pneumonia. *ERJ Open Research*, **2015**, 1,	3.5	18
137	Structured reporting for fibrosing lung disease: a model shared by radiologist and pulmonologist. *Radiologia Medica*, **2018**, 123, 245-253	6.5	18
136	Idiopathic Pulmonary Fibrosis: Molecular Endotypes of Fibrosis Stratifying Existing and Emerging Therapies. *Respiration*, **2017**, 93, 379-395	3.7	17
135	Differing severities of acute exacerbations of idiopathic pulmonary fibrosis (IPF): insights from the INPULSIS trials. *Respiratory Research*, **2019**, 20, 71	7.3	16
134	Novel drug targets for idiopathic pulmonary fibrosis. *Expert Review of Respiratory Medicine*, **2016**, 10, 393-405	3.8	16
133	Assessing the treatment effect from multiple trials in idiopathic pulmonary fibrosis. *European Respiratory Review*, **2012**, 21, 147-51	9.8	16
132	Existing and emerging biomarkers for disease progression in idiopathic pulmonary fibrosis. *Expert Review of Respiratory Medicine*, **2019**, 13, 39-51	3.8	16
131	Lung ultrasonography for early management of patients with respiratory symptoms during COVID-19 pandemic. *Journal of Ultrasound*, **2020**, 23, 449-456	3.4	15
130	The safety of new drug treatments for idiopathic pulmonary fibrosis. *Expert Opinion on Drug Safety*, **2016**, 15, 1483-1489	4.1	15
129	Pamrevlumab for the treatment of idiopathic pulmonary fibrosis. *Expert Opinion on Investigational Drugs*, **2020**, 29, 771-777	5.9	15
128	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	20.5	15
127	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. *Chest*, **2021**, 159, 1107-1125	5.3	15
126	Safety and tolerability of nintedanib for the treatment of idiopathic pulmonary fibrosis in routine UK clinical practice. *ERJ Open Research*, **2018**, 4,	3.5	15
125	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	5.6	13
124	Aortic pulse wave velocity measurement in systemic sclerosis patients. *Reumatismo*, **2012**, 64, 360-7	1.1	13

123	Recommendations on treatment for IPF. *Respiratory Research*, **2013**, 14 Suppl 1, S6	7.3	13
122	Exploring the immune response against Mycobacterium tuberculosis for a better diagnosis of the infection. *Archivum Immunologiae Et Therapiae Experimentalis*, **2009**, 57, 425-33	4	13
121	Effects of nintedanib in patients with idiopathic pulmonary fibrosis by GAP stage. *ERJ Open Research*, **2019**, 5,	3.5	12
120	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	4.6	12
119	Environmental triggers and susceptibility factors in idiopathic granulomatous diseases. *Seminars in Respiratory and Critical Care Medicine*, **2008**, 29, 610-9	3.9	12
118	No relevant pharmacokinetic drug-drug interaction between nintedanib and pirfenidone. *European Respiratory Journal*, **2019**, 53,	13.6	12
117	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	4.1	12
116	The role of biomarkers in low respiratory tract infections. *European Journal of Internal Medicine*, **2012**, 23, 429-35	3.9	11
115	Role of the quantiferon-TB test in ruling out pleural tuberculosis: a multi-centre study. *International Journal of Immunopathology and Pharmacology*, **2011**, 24, 159-65	3	11
114	Treatment strategies for asthma: reshaping the concept of asthma management. *Allergy, Asthma and Clinical Immunology*, **2020**, 16, 75	3.2	11
113	Nintedanib and immunomodulatory therapies in progressive fibrosing interstitial lung diseases. *Respiratory Research*, **2021**, 22, 84	7.3	11
112	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	10.2	10
111	Validation of multidisciplinary diagnosis in IPF. *Lancet Respiratory Medicine,the*, **2018**, 6, 88-89	35.1	10
110	Investigational drugs for idiopathic pulmonary fibrosis. *Expert Opinion on Investigational Drugs*, **2017**, 26, 1019-1031	5.9	10
109	Using ELISpot technology to improve the diagnosis of tuberculosis infection: from the bench to the T-SPOT.TB assay. *Expert Review of Respiratory Medicine*, **2008**, 2, 253-60	3.8	10
108	New treatment directions for IPF: current status of ongoing and upcoming clinical trials. *Expert Review of Respiratory Medicine*, **2017**, 11, 533-548	3.8	9
107	Agreement between chest ultrasonography and chest X-ray in patients who have undergone thoracic surgery: preliminary results. *Multidisciplinary Respiratory Medicine*, **2019**, 14, 9	3	9
106	A quantitative proteomic approach to identify significantly altered protein networks in the serum of patients with lymphangioleiomyomatosis (LAM). *PLoS ONE*, **2014**, 9, e105365	3.7	9

105	Viruses and acute exacerbations of idiopathic pulmonary fibrosis: rest in peace?. *American Journal of Respiratory and Critical Care Medicine*, **2011**, 183, 1583-4	10.2	9
104	What if we made stratified medicine work for patients?. *Lancet Respiratory Medicine,the*, **2016**, 4, 8-10	35.1	9
103	Time for Prevention of Idiopathic Pulmonary Fibrosis Exacerbation. *Annals of the American Thoracic Society*, **2015**, 12 Suppl 2, S181-5	4.7	9
102	Current and Future Idiopathic Pulmonary Fibrosis Therapy. *American Journal of the Medical Sciences* , **2019**, 357, 370-373	2.2	8
101	Clinical trials of investigational agents for IPF: a review of a Cochrane report. *Respiratory Research*, **2013**, 14 Suppl 1, S4	7.3	8
100	Safety of Nintedanib in Patients with Idiopathic Pulmonary Fibrosis: Global Pharmacovigilance Data. *Advances in Therapy*, **2020**, 37, 4209-4219	4.1	8
99	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	4.5	8
98	Subclinical Interstitial Lung Abnormalities: Lumping and Splitting Revisited. *American Journal of Respiratory and Critical Care Medicine*, **2019**, 200, 121-123	10.2	7
97	Management of idiopathic pulmonary fibrosis. *Clinics in Chest Medicine*, **2012**, 33, 85-94	5.3	7
96	A culture-proven case of community-acquired legionella pneumonia apparently classified as nosocomial: diagnostic and public health implications. *Case Reports in Medicine*, **2013**, 2013, 303712	0.7	7
95	Alemtuzumab-induced lung injury in multiple sclerosis: Learning from adversity in three patients. *Multiple Sclerosis and Related Disorders*, **2020**, 37, 101450	4	7
94	Update in Pulmonary Fibrosis 2018. *American Journal of Respiratory and Critical Care Medicine*, **2019** , 200, 292-300	10.2	6
93	Paracrine SPARC signaling dysregulates alveolar epithelial barrier integrity and function in lung fibrosis. *Cell Death Discovery*, **2020**, 6, 54	6.9	6
92	Antibody-based therapies for idiopathic pulmonary fibrosis. *Expert Opinion on Biological Therapy*, **2020**, 20, 779-786	5.4	6
91	Individualizing duration of antibiotic therapy in community-acquired pneumonia. *Pulmonary Pharmacology and Therapeutics*, **2017**, 45, 191-201	3.5	6
90	Acute exacerbations of chronic obstructive pulmonary disease: are antibiotics needed?. *American Journal of Respiratory and Critical Care Medicine*, **2010**, 181, 102-3	10.2	6
89	Effect of baseline FVC on lung function decline with nintedanib in patients with IPF **2015**,		6
88	Multidrug-resistant tuberculosis outbreak in an Italian prison: tolerance of pyrazinamide plus levofloxacin prophylaxis and serial interferon gamma release assays. *New Microbes and New Infections*, **2016**, 12, 45-51	4.1	6

87	Quantitative analysis of lung sounds for monitoring idiopathic pulmonary fibrosis: a prospective pilot study. *European Respiratory Journal*, **2019**, 53,	13.6	6
86	Fibrotic Hypersensitivity Pneumonitis: Diagnosis and Management. *Lung*, **2020**, 198, 429-440	2.9	5
85	Ultrasonography of the Mediastinum: Techniques, Current Practice, and Future Directions. *Respiratory Care*, **2018**, 63, 1421-1438	2.1	5
84	Genetic testing in diffuse parenchymal lung disease. *Orphanet Journal of Rare Diseases*, **2012**, 7, 79	4.2	5
83	Tuberculosis infection in foreign-born children: a screening survey based on skin and blood testing. *International Journal of Tuberculosis and Lung Disease*, **2011**, 15, 1182-4, i	2.1	5
82	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,	3.5	5
81	Current Diagnosis and Management of Hypersensitivity Pneumonitis. *Tuberculosis and Respiratory Diseases*, **2020**, 83, 122-131	3.2	5
80	Mesenchymal Stromal Cell Secretome for Post-COVID-19 Pulmonary Fibrosis: A New Therapy to Treat the Long-Term Lung Sequelae?. *Cells*, **2021**, 10,	7.9	5
79	Residual respiratory impairment after COVID-19 pneumonia. *BMC Pulmonary Medicine*, **2021**, 21, 241	3.5	5
78	Chronic beryllium disease: a model for pulmonary sarcoidosis?. *Acta Biomedica*, **2005**, 76 Suppl 2, 11-4	3.2	5
77	Do all patients with idiopathic pulmonary fibrosis warrant a trial of therapeutic intervention? A pro-con perspective. *Respirology*, **2015**, 20, 389-94	3.6	4
76	Opportunities to diagnose fibrotic lung diseases in routine care: A primary care cohort study. *Respirology*, **2020**, 25, 1274-1282	3.6	4
75	Do Randomized Clinical Trials Always Provide Certain Results? The Case of Tralokinumab in Idiopathic Pulmonary Fibrosis. *American Journal of Respiratory and Critical Care Medicine*, **2018**, 197, 9-10	10.2	4
74	Are newly launched pharmacotherapies efficacious in treating idiopathic pulmonary fibrosis? Or is there still more work to be done?. *Expert Opinion on Pharmacotherapy*, **2017**, 18, 1583-1594	4	4
73	To BAL or not to BAL: is this a problem in diagnosing IPF?. *American Journal of Respiratory and Critical Care Medicine*, **2009**, 180, 379-80; author reply 380	10.2	4
72	Twenty-five years of Respirology: Advances in idiopathic pulmonary fibrosis. *Respirology*, **2020**, 25, 20-22	3.6	4
71	Detection and Early Referral of Patients With Interstitial Lung Abnormalities: An Expert Survey Initiative. *Chest*, **2021**,	5.3	4
70	Nintedanib in progressive interstitial lung diseases: data from the whole INBUILD trial. *European Respiratory Journal*, **2021**,	13.6	4

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