Bruno Crestani

List of Publications by Year
in descending order

147	11,444	41344	30922
papers	citations	49	102
		h-index	g-index

159	159	159	10523
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Rheumatological evaluation of patients with interstitial lung disease. Scandinavian Journal of Rheumatology, 2022, 51, 34-41.	1.1	6
2	The Genetics of Interstitial Lung Diseases. , 2022, , 96-113.		0
3	Determinants of survival after lung transplantation in telomerase-related gene mutation carriers: A retrospective cohort. American Journal of Transplantation, 2022, 22, 1236-1244.	4.7	11
4	Lung Fibroblasts from Idiopathic Pulmonary Fibrosis Patients Harbor Short and Unstable Telomeres Leading to Chromosomal Instability. Biomedicines, 2022, 10, 310.	3.2	5
5	Interstitial lung diseases associated with mutations of poly(A)â€specific ribonuclease: A multicentre retrospective study. Respirology, 2022, 27, 226-235.	2.3	6
6	European Respiratory Society statement on long COVID follow-up. European Respiratory Journal, 2022, 60, 2102174.	6.7	81
7	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.	5.6	780
8	Fibrotic-Like CT Alterations in COVID-19: Distinct Patterns of Temporal Evolution. , 2022, , .		0
9	Targeting the nasty nestin to shoot lung fibrosis. European Respiratory Journal, 2022, 59, 2103146.	6.7	1
10	FGF19 is Downregulated in Idiopathic Pulmonary Fibrosis and Inhibits Lung Fibrosis in Mice. American Journal of Respiratory Cell and Molecular Biology, 2022, , .	2.9	10
11	Higher basal tryptase, asthma and loss of consciousness in anaphylaxis are associated with biphasic reactions. Clinical and Translational Allergy, 2022, 12, .	3.2	0
12	Methotrexate and rheumatoid arthritis associated interstitial lung disease. European Respiratory Journal, 2021, 57, 2000337.	6.7	114
13	NADPH oxidase DUOX1 sustains TGF-Î²1 signalling and promotes lung fibrosis. European Respiratory Journal, 2021, 57, 1901949.	6.7	30
14	Safety and efficacy of pirfenidone and nintedanib in patients with idiopathic pulmonary fibrosis and carrying a telomere-related gene mutation. European Respiratory Journal, 2021, 57, 2003198.	6.7	36
15	Basophils and IgE contribute to mixed connective tissue disease development. Journal of Allergy and Clinical Immunology, 2021, 147, 1478-1489.e11.	2.9	14
16	Observer agreement and clinical significance of chest CT reporting in patients suspected of COVID-19. European Radiology, 2021, 31, 1081-1089.	4.5	11
17	Immune checkpoint blockade for patients with lung cancer and idiopathic pulmonary fibrosis. European Journal of Cancer, 2021, 145, 179-182.	2.8	9
18	Alveolar epithelial TET2 is not involved in the development of bleomycinâ€induced pulmonary fibrosis. FASEB Journal, 2021, 35, e21599.	0.5	1

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19	OP0099â€…EPIDEMIOLOGY AND MORTALITY OF RA-ASSOCIATED INTERSTITIAL LUNG DISEASE: DATA FROM A FRENCH ADMINISTRATIVE HEALTHCARE DATABASE. Annals of the Rheumatic Diseases, 2021, 80, 54.2-55.	0.9	2
20	Response to letter entitled: Re: Immune checkpoint blockade for patients with lung cancer and idiopathic pulmonary fibrosis. European Journal of Cancer, 2021, 151, 252-253.	2.8	1
21	NKX2.1 (TTF1) germline mutation associated with pulmonary fibrosis and lung cancer. ERJ Open Research, 2021, 7, 00356-2021.	2.6	8
22	Blood fibrocytes are associated with severity and prognosis in COVID-19 pneumonia. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2021, 321, L847-L858.	2.9	11
23	Usual interstitial pneumonia in ANCA-associated vasculitis: A poor prognostic factor. Journal of Autoimmunity, 2020, 106, 102338.	6.5	43
24	Efficacy and safety of rituximab in patients with chronic hypersensitivity pneumonitis (cHP): A retrospective, multicentric, observational study. Respiratory Medicine, 2020, 172, 106146.	2.9	28
25	Chaotic activation of developmental signalling pathways drives idiopathic pulmonary fibrosis. European Respiratory Review, 2020, 29, 190140.	7.1	31
26	Clinical and Functional Characteristics of Patients with Unclassifiable Interstitial Lung Disease (uILD): Long-Term Follow-Up Data from European IPF Registry (eurIPFreg). Journal of Clinical Medicine, 2020, 9, 2499.	2.4	17
27	Recent advances in rheumatoid arthritis-associated interstitial lung disease. Current Opinion in Pulmonary Medicine, 2020, 26, 477-486.	2.6	31
28	Functional assessment and phenotypic heterogeneity of SFTPA1 and SFTPA2 mutations in interstitial lung diseases and lung cancer. European Respiratory Journal, 2020, 56, 2002806.	6.7	23
29	Eight novel variants in the SLC34A2 gene in pulmonary alveolar microlithiasis. European Respiratory Journal, 2020, 55, 1900806.	6.7	14
30	Lung function in Birt-Hogg-DubÃ© syndrome: a retrospective analysis of 96 patients. Orphanet Journal of Rare Diseases, 2020, 15, 120.	2.7	15
31	TRIM33 prevents pulmonary fibrosis by impairing TGF-Î²1 signalling. European Respiratory Journal, 2020, 55, 1901346.	6.7	45
32	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.	10.7	331
33	First heterozygous NOP10 mutation in familial pulmonary fibrosis. European Respiratory Journal, 2020, 55, 1902465.	6.7	13
34	Impact of genetic factors on fibrosing interstitial lung diseases. Incidence and clinical presentation in adults. Presse Medicale, 2020, 49, 104024.	1.9	9
35	Glucocorticoids with low-dose anti-IL1 anakinra rescue in severe non-ICU COVID-19 infection: A cohort study. PLoS ONE, 2020, 15, e0243961.	2.5	15
36	Title is missing!. , 2020, 15, e0243961.		0

#	Article	IF	Citations
37	Title is missing!. , 2020, 15, e0243961.		0
38	Title is missing!. , 2020, 15, e0243961.		0
39	Title is missing!. , 2020, 15, e0243961.		0
40	Myelodysplastic syndromes and idiopathic pulmonary fibrosis: a dangerous liaison. Respiratory Research, 2019, 20, 182.	3.6	7
41	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.	2.9	22
42	Follow-Up and Management of Chronic Rhinosinusitis in Adults with Primary Ciliary Dyskinesia: Review and Experience of Our Reference Centers. Journal of Clinical Medicine, 2019, 8, 1495.	2.4	12
43	Calcium-solubilizing sodium thiosulfate failed to improve pulmonary alveolar microlithiasis: Evaluation of calcium content with CT scan. Respiratory Medicine and Research, 2019, 75, 10-12.	0.6	7
44	The Ariane-IPF ERS Clinical Research Collaboration: seeking collaboration through launch of a federation of European registries on idiopathic pulmonary fibrosis. European Respiratory Journal, 2019, 53, 1900539.	6.7	8
45	Critical Evaluation of Sinonasal Disease in 64 Adults with Primary Ciliary Dyskinesia. Journal of Clinical Medicine, 2019, 8, 619.	2.4	44
46	Resequencing Study Confirms That Host Defense and Cell Senescence Gene Variants Contribute to the Risk of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 199-208.	5.6	90
47	Psychometric properties and minimal important differences of SF-36 in Idiopathic Pulmonary Fibrosis. Respiratory Research, 2019, 20, 47.	3.6	31
48	Lung CT Densitometry in Idiopathic Pulmonary Fibrosis for the Prediction of Natural Course, Severity, and Mortality. Chest, 2019, 155, 972-981.	0.8	32
49	The Long Noncoding RNA DNM3OS Is a Reservoir of FibromiRs with Major Functions in Lung Fibroblast Response to TGF-Î² and Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 184-198.	5.6	78
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.	3.0	121
51	Dysregulated balance of lung macrophage populations in idiopathic pulmonary fibrosis revealed by single-cell RNA seq: an unstable â€œmÃ©nage-Ã-troisâ€• European Respiratory Journal, 2019, 54, 1901229.	6.7	7
52	The genetics of interstitial lung diseases. European Respiratory Review, 2019, 28, 190053.	7.1	41
53	Pilot experience of multidisciplinary team discussion dedicated to inherited pulmonary fibrosis. Orphanet Journal of Rare Diseases, 2019, 14, 280.	2.7	24
54	Long-term safety and tolerability of nintedanib in patients with idiopathic pulmonary fibrosis: results from the open-label extension study, INPULSIS-ON. Lancet Respiratory Medicine,the, 2019, 7, 60-68.	10.7	160

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55	Combined pulmonary fibrosis and emphysema in systemic sclerosis: A syndrome associated with heavy morbidity and mortality. Seminars in Arthritis and Rheumatism, 2019, 49, 98-104.	3.4	33
56	Interleukin-8 release by endothelial colony-forming cells isolated from idiopathic pulmonary fibrosis patients might contribute to their pathogenicity. Angiogenesis, 2019, 22, 325-339.	7.2	23
57	Regulator of telomere length 1 (RTEL1) mutations are associated with heterogeneous pulmonary and extra-pulmonary phenotypes. European Respiratory Journal, 2019, 53, 1800508.	6.7	45
58	Familial pulmonary fibrosis: a world without frontiers. Jornal Brasileiro De Pneumologia, 2019, 45, e20190303.	0.7	3
59	Telomere syndrome and the lung. , 2019, , 391-403.		0
60	Pulmonary phenotypes associated with genetic variation in telomere-related genes. Current Opinion in Pulmonary Medicine, 2018, 24, 269-280.	2.6	54
61	The impaired proteases and anti-proteases balance in Idiopathic Pulmonary Fibrosis. Matrix Biology, 2018, 68-69, 382-403.	3.6	56
62	Safety and efficacy of pirfenidone in patients carrying telomerase complex mutation. European Respiratory Journal, 2018, 51, 1701875.	6.7	34
63	Physiology of the lung in idiopathic pulmonary fibrosis. European Respiratory Review, 2018, 27, 170062.	7.1	159
64	Is chronic exposure to air pollutants a risk factor for the development of idiopathic pulmonary fibrosis?. European Respiratory Journal, 2018, 51, 1702663.	6.7	10
65	Anti-parietal cell autoimmunity is associated with an accelerated decline of lung function in IPF patients. Respiratory Medicine, 2018, 135, 15-21.	2.9	10
66	Patients with IPF and lung cancer: diagnosis and management. Lancet Respiratory Medicine,the, 2018, 6, 86-88.	10.7	67
67	Human airway trypsinâ€like protease exerts potent, antifibrotic action in pulmonary fibrosis. FASEB Journal, 2018, 32, 1250-1264.	0.5	6
68	Antineutrophil Cytoplasmic Antibodyâ€“Associated Lung Fibrosis. Seminars in Respiratory and Critical Care Medicine, 2018, 39, 465-470.	2.1	27
69	MUC5B Promoter Variant and Rheumatoid Arthritis with Interstitial Lung Disease. New England Journal of Medicine, 2018, 379, 2209-2219.	27.0	326
70	Anti-acid therapy in idiopathic pulmonary fibrosis: insights from the INPULSISÂ® trials. Respiratory Research, 2018, 19, 167.	3.6	42
71	The FLORA study: presenting a novel IPF trial design. Lancet Respiratory Medicine,the, 2018, 6, 572-573.	10.7	0
72	The European IPF registry (eurIPFreg): baseline characteristics and survival of patients with idiopathic pulmonary fibrosis. Respiratory Research, 2018, 19, 141.	3.6	199

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73	Identification of periplakin as a major regulator of lung injury and repair in mice. JCI Insight, 2018, 3, .	5.0	13
74	Pneumopathies interstitielles. Revue Des Maladies Respiratoires Actualites, 2018, 10, S21-S23.	0.0	0
75	CCAAT/enhancer binding protein delta (C/EBPÎ´) deficiency does not affect bleomycin-induced pulmonary fibrosis. Journal of Clinical and Translational Research, 2018, 3, 358-365.	0.3	3
76	Human airway trypsin-like protease, a serine protease involved in respiratory diseases. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2017, 312, L657-L668.	2.9	32
77	Shared genetic predisposition in rheumatoid arthritis-interstitial lung disease and familial pulmonary fibrosis. European Respiratory Journal, 2017, 49, 1602314.	6.7	154
78	Management of suspected monogenic lung fibrosis in a specialised centre. European Respiratory Review, 2017, 26, 160122.	7.1	54
79	Heterogeneity of lung disease associated with NK2 homeobox 1 mutations. Respiratory Medicine, 2017, 129, 16-23.	2.9	54
80	The Genetic Diagnosis of Interstitial Lung Disease: A Need for an International Consensus. American Journal of Respiratory and Critical Care Medicine, 2017, 195, 1538-1539.	5.6	7
81	Obstructive sleep apnoea and related comorbidities in incident idiopathic pulmonary fibrosis. European Respiratory Journal, 2017, 49, 1601934.	6.7	72
82	French practical guidelines for the diagnosis and management of idiopathic pulmonary fibrosisÂâ€“Â2017 update. Full-length version. Revue Des Maladies Respiratoires, 2017, 34, 900-968.	1.7	51
83	French practical guidelines for the diagnosis and management of idiopathic pulmonary fibrosisÂâ€“Â2017 update. Short-length version. Revue Des Maladies Respiratoires, 2017, 34, 852-899.	1.7	2
84	FGF9 prevents pleural fibrosis induced by intrapleural adenovirus injection in mice. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2017, 313, L781-L795.	2.9	18
85	Licence to kill senescent cells in idiopathic pulmonary fibrosis?. European Respiratory Journal, 2017, 50, 1701360.	6.7	16
86	Pneumocystosis revealing immunodeficiency secondary to TERC mutation. European Respiratory Journal, 2017, 50, 1701443.	6.7	12
87	Towards a global initiative for fibrosis treatment (GIFT). ERJ Open Research, 2017, 3, 00106-2017.	2.6	5
88	Validation of the EULAR/ERA-EDTA recommendations for the management of ANCA-associated vasculitis by disease content experts. RMD Open, 2017, 3, e000449.	3.8	23
89	[18F]FDG PET/CT predicts progression-free survival in patients with idiopathic pulmonary fibrosis. Respiratory Research, 2017, 18, 74.	3.6	66
90	Giant hiatal hernia: beware of the supine ICU chest X-ray!. BMJ Case Reports, 2017, 2017, bcr-2017-219668.	0.5	2

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91	Antifibrotic role of vascular endothelial growth factor in pulmonary fibrosis. JCI Insight, 2017, 2, .	5.0	51
92	Transcriptome of Cultured Lung Fibroblasts in Idiopathic Pulmonary Fibrosis: Meta-Analysis of Publically Available Microarray Datasets Reveals Repression of Inflammation and Immunity Pathways. International Journal of Molecular Sciences, 2016, 17, 2091.	4.1	28
93	Serum Amyloid P Contained in Alveolar Fluid From Patients With Acute Respiratory Distress Syndrome Mediates the Inhibition of Monocyte Differentiation into Fibrocyte. Critical Care Medicine, 2016, 44, e563-e573.	0.9	5
94	EULAR/ERA-EDTA recommendations for the management of ANCA-associated vasculitis. Annals of the Rheumatic Diseases, 2016, 75, 1583-1594.	0.9	940
95	Prevalence and characteristics ofTERTandTERCmutations in suspected genetic pulmonary fibrosis. European Respiratory Journal, 2016, 48, 1721-1731.	6.7	136
96	New targets in idiopathic pulmonary fibrosis: from inflammation and immunity to remodeling and repair. Expert Opinion on Orphan Drugs, 2016, 4, 511-520.	0.8	4
97	FGF9 and FGF18 in idiopathic pulmonary fibrosis promote survival and migration and inhibit myofibroblast differentiation of human lung fibroblasts in vitro. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2016, 310, L615-L629.	2.9	75
98	Pharmacological management of IPF. Respirology, 2016, 21, 615-625.	2.3	32
99	Gastro-oesophageal reflux and idiopathic pulmonary fibrosis: in search of evidence. European Respiratory Journal, 2016, 48, 623-625.	6.7	3
100	Severe Pulmonary Fibrosis as the First Manifestation of Interferonopathy (TMEM173 Mutation). Chest, 2016, 150, e65-e71.	0.8	112
101	THU0338â€…Cystic Lung Disease in SjÃ¶gren's Syndrome: An Observational Study. Annals of the Rheumatic Diseases, 2016, 75, 309.2-309.	0.9	0
102	GermlineSFTPA1mutation in familial idiopathic interstitial pneumonia and lung cancer. Human Molecular Genetics, 2016, 25, 1457-1467.	2.9	119
103	European IPF Patient Charter: an SOS to the world. European Respiratory Journal, 2016, 47, 403-405.	6.7	1
104	Increased volume of conducting airways in idiopathic pulmonary fibrosis is independent of disease severity: a volumetric capnography study. Journal of Breath Research, 2016, 10, 016005.	3.0	19
105	Membrane-anchored Serine Protease Matriptase Is a Trigger of Pulmonary Fibrogenesis. American Journal of Respiratory and Critical Care Medicine, 2016, 193, 847-860.	5.6	47
106	Safety, tolerability and appropriate use of nintedanib in idiopathic pulmonary fibrosis. Respiratory Research, 2015, 16, 116.	3.6	114
107	Pharmacological Targeting of Protease-Activated Receptor 2 Affords Protection from Bleomycin-Induced Pulmonary Fibrosis. Molecular Medicine, 2015, 21, 576-583.	4.4	24
108	Idiopathic pulmonary fibrosis: An update. Annals of Medicine, 2015, 47, 15-27.	3.8	97

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109	Proteaseâ€activated receptor (<scp>PAR</scp>)â€2 is required for <scp>PAR</scp>â€1 signalling in pulmonary fibrosis. Journal of Cellular and Molecular Medicine, 2015, 19, 1346-1356.	3.6	21
110	Antifibrotic Role of Î±B-Crystallin Inhibition in Pleural and Subpleural Fibrosis. American Journal of Respiratory Cell and Molecular Biology, 2015, 52, 244-252.	2.9	19
111	Severe hematologic complications after lung transplantation in patients with telomerase complex mutations. Journal of Heart and Lung Transplantation, 2015, 34, 538-546.	0.6	109
112	HeterozygousRTEL1mutations are associated with familial pulmonary fibrosis. European Respiratory Journal, 2015, 46, 474-485.	6.7	135
113	Missing data in IPF trials: do not let methodological issues undermine a major therapeutic breakthrough. European Respiratory Journal, 2015, 46, 607-614.	6.7	9
114	Increased expression of protease nexin-1 in fibroblasts during idiopathic pulmonary fibrosis regulates thrombin activity and fibronectin expression. Laboratory Investigation, 2014, 94, 1237-1246.	3.7	24
115	Forkhead Box F1 represses cell growth and inhibits COL1 and ARPC2 expression in lung fibroblasts in vitro. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2014, 307, L838-L847.	2.9	30
116	Targeting the Hedgehogâ€“Glioma-Associated Oncogene Homolog Pathway Inhibits Bleomycin-Induced Lung Fibrosis in Mice. American Journal of Respiratory Cell and Molecular Biology, 2014, 51, 11-25.	2.9	76
117	Diagnosis and management of idiopathic pulmonary fibrosis: French practical guidelines. European Respiratory Review, 2014, 23, 193-214.	7.1	62
118	Detection of anti-periplakin auto-antibodies during idiopathic pulmonary fibrosis. Clinica Chimica Acta, 2014, 433, 242.	1.1	4
119	Combination therapy: the future of management for idiopathic pulmonary fibrosis?. Lancet Respiratory Medicine,the, 2014, 2, 933-942.	10.7	128
120	The small heatâ€shock protein Î±<scp>B</scp>â€crystallin is essential for the nuclear localization of Smad4: impact on pulmonary fibrosis. Journal of Pathology, 2014, 232, 458-472.	4.5	52
121	Fibroblasts: the missing link between fibrotic lung diseases of different etiologies?. Respiratory Research, 2013, 14, 81.	3.6	8
122	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.	5.6	3,134
123	Idiopathic Pulmonary Fibrosis: From Epithelial Injury to Biomarkers - Insights from the Bench Side. Respiration, 2013, 86, 441-452.	2.6	108
124	Inhibition of HSP27 blocks fibrosis development and EMT features by promoting Snail degradation. FASEB Journal, 2013, 27, 1549-1560.	0.5	95
125	A 70-Year-Old Woman With Acute Chest Pain and a Paracardiac Mass. Chest, 2013, 143, 866-869.	0.8	5
126	Detection of Alveolar Fibrocytes in Idiopathic Pulmonary Fibrosis and Systemic Sclerosis. PLoS ONE, 2013, 8, e53736.	2.5	33

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127	The MUC5B Variant Is Associated with Idiopathic Pulmonary Fibrosis but Not with Systemic Sclerosis Interstitial Lung Disease in the European Caucasian Population. PLoS ONE, 2013, 8, e70621.	2.5	142
128	Molecular biomarkers in idiopathic pulmonary fibrosis and disease severity. Sarcoidosis Vasculitis and Diffuse Lung Diseases, 2013, 30 Suppl 1, 27-32.	0.2	1
129	Familial forms of nonspecific interstitial pneumonia/idiopathic pulmonary fibrosis. Current Opinion in Pulmonary Medicine, 2012, 18, 455-461.	2.6	19
130	Hepatocyte Growth Factor and Lung Fibrosis. Proceedings of the American Thoracic Society, 2012, 9, 158-163.	3.5	52
131	Octreotide treatment of idiopathic pulmonary fibrosis: a proof-of-concept study. European Respiratory Journal, 2012, 39, 772-775.	6.7	14
132	Alveolar fibrocyte percentage is an independent predictor of poor outcome in patients with acute lung injury*. Critical Care Medicine, 2012, 40, 21-28.	0.9	37
133	The Hedgehog System Machinery Controls Transforming Growth Factor-Î²â€“Dependent Myofibroblastic Differentiation in Humans. American Journal of Pathology, 2012, 181, 2126-2137.	3.8	119
134	Quantitative analysis of ciliary beating in primary ciliary dyskinesia: a pilot study. Orphanet Journal of Rare Diseases, 2012, 7, 78.	2.7	62
135	Severe chronic bronchiolitis as the presenting feature of primary SjÃ¶grenâ€™s syndrome. Respiratory Medicine, 2011, 105, 130-136.	2.9	40
136	Identification of Periplakin as a New Target for Autoreactivity in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2011, 183, 759-766.	5.6	102
137	Ectopic respiratory epithelial cell differentiation in bronchiolised distal airspaces in idiopathic pulmonary fibrosis. Thorax, 2011, 66, 651-657.	5.6	159
138	Imbalance in the Proâ€“Hepatocyte Growth Factor Activation System in Bleomycin-Induced Lung Fibrosis in Mice. American Journal of Respiratory Cell and Molecular Biology, 2010, 42, 286-293.	2.9	14
139	Prevalence of Telomere Shortening in Familial and Sporadic Pulmonary Fibrosis Is Increased in Men. American Journal of Respiratory and Critical Care Medicine, 2009, 179, 1073-1073.	5.6	14
140	Modulation of bleomycin-induced lung fibrosis by serotonin receptor antagonists in mice. European Respiratory Journal, 2008, 32, 426-436.	6.7	92
141	Regulation of hepatocyte growth factor secretion by fibroblasts in patients with acute lung injury. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2008, 294, L334-L343.	2.9	61
142	Dendritic Cells Accumulate in Human Fibrotic Interstitial Lung Disease. American Journal of Respiratory and Critical Care Medicine, 2007, 176, 1007-1014.	5.6	97
143	Defect of Pro-Hepatocyte Growth Factor Activation by Fibroblasts in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2006, 174, 58-66.	5.6	57
144	Cutting Edge: Nonproliferating Mature Immune Cells Form a Novel Type of Organized Lymphoid Structure in Idiopathic Pulmonary Fibrosis. Journal of Immunology, 2006, 176, 5735-5739.	0.8	157

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145	Increased uptake of 111In-octreotide in idiopathic pulmonary fibrosis. Journal of Nuclear Medicine, 2006, 47, 1281-7.	5.0	44
146	Defect of Hepatocyte Growth Factor Secretion by Fibroblasts in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2003, 168, 1156-1161.	5.6	71
147	Genetic testing in interstitial lung disease: An international survey. Respirology, 0, , .	2.3	10