Bruno Crestani

List of Publications by Year
in descending order

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Version: 2024-02-01

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	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
2	EULAR/ERA-EDTA recommendations for the management of ANCA-associated vasculitis. Annals of the Rheumatic Diseases, 2016, 75, 1583-1594.	0.9	940
3	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
4	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
5	MUC5B Promoter Variant and Rheumatoid Arthritis with Interstitial Lung Disease. New England Journal of Medicine, 2018, 379, 2209-2219.	27.0	326
6	The European IPF registry (eurIPFreg): baseline characteristics and survival of patients with idiopathic pulmonary fibrosis. Respiratory Research, 2018, 19, 141.	3.6	199
7	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
8	Ectopic respiratory epithelial cell differentiation in bronchiolised distal airspaces in idiopathic pulmonary fibrosis. Thorax, 2011, 66, 651-657.	5.6	159
9	Physiology of the lung in idiopathic pulmonary fibrosis. European Respiratory Review, 2018, 27, 170062.	7.1	159
10	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
11	Shared genetic predisposition in rheumatoid arthritis-interstitial lung disease and familial pulmonary fibrosis. European Respiratory Journal, 2017, 49, 1602314.	6.7	154
12	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
13	Prevalence and characteristics ofTERTandTERCmutations in suspected genetic pulmonary fibrosis. European Respiratory Journal, 2016, 48, 1721-1731.	6.7	136
14	HeterozygousRTEL1mutations are associated with familial pulmonary fibrosis. European Respiratory Journal, 2015, 46, 474-485.	6.7	135
15	Combination therapy: the future of management for idiopathic pulmonary fibrosis?. Lancet Respiratory Medicine,the, 2014, 2, 933-942.	10.7	128
16	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
17	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
18	GermlineSFTPA1mutation in familial idiopathic interstitial pneumonia and lung cancer. Human Molecular Genetics, 2016, 25, 1457-1467.	2.9	119

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19	Safety, tolerability and appropriate use of nintedanib in idiopathic pulmonary fibrosis. Respiratory Research, 2015, 16, 116.	3.6	114
20	Methotrexate and rheumatoid arthritis associated interstitial lung disease. European Respiratory Journal, 2021, 57, 2000337.	6.7	114
21	Severe Pulmonary Fibrosis as the First Manifestation of Interferonopathy (TMEM173 Mutation). Chest, 2016, 150, e65-e71.	0.8	112
22	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
23	Idiopathic Pulmonary Fibrosis: From Epithelial Injury to Biomarkers - Insights from the Bench Side. Respiration, 2013, 86, 441-452.	2.6	108
24	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
25	Dendritic Cells Accumulate in Human Fibrotic Interstitial Lung Disease. American Journal of Respiratory and Critical Care Medicine, 2007, 176, 1007-1014.	5.6	97
26	Idiopathic pulmonary fibrosis: An update. Annals of Medicine, 2015, 47, 15-27.	3.8	97
27	Inhibition of HSP27 blocks fibrosis development and EMT features by promoting Snail degradation. FASEB Journal, 2013, 27, 1549-1560.	0.5	95
28	Modulation of bleomycin-induced lung fibrosis by serotonin receptor antagonists in mice. European Respiratory Journal, 2008, 32, 426-436.	6.7	92
29	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
30	European Respiratory Society statement on long COVID follow-up. European Respiratory Journal, 2022, 60, 2102174.	6.7	81
31	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
32	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
33	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
34	Obstructive sleep apnoea and related comorbidities in incident idiopathic pulmonary fibrosis. European Respiratory Journal, 2017, 49, 1601934.	6.7	72
35	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
36	Patients with IPF and lung cancer: diagnosis and management. Lancet Respiratory Medicine,the, 2018, 6, 86-88.	10.7	67

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37	[18F]FDG PET/CT predicts progression-free survival in patients with idiopathic pulmonary fibrosis. Respiratory Research, 2017, 18, 74.	3.6	66
38	Quantitative analysis of ciliary beating in primary ciliary dyskinesia: a pilot study. Orphanet Journal of Rare Diseases, 2012, 7, 78.	2.7	62
39	Diagnosis and management of idiopathic pulmonary fibrosis: French practical guidelines. European Respiratory Review, 2014, 23, 193-214.	7.1	62
40	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
41	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
42	The impaired proteases and anti-proteases balance in Idiopathic Pulmonary Fibrosis. Matrix Biology, 2018, 68-69, 382-403.	3.6	56
43	Management of suspected monogenic lung fibrosis in a specialised centre. European Respiratory Review, 2017, 26, 160122.	7.1	54
44	Heterogeneity of lung disease associated with NK2 homeobox 1 mutations. Respiratory Medicine, 2017, 129, 16-23.	2.9	54
45	Pulmonary phenotypes associated with genetic variation in telomere-related genes. Current Opinion in Pulmonary Medicine, 2018, 24, 269-280.	2.6	54
46	Hepatocyte Growth Factor and Lung Fibrosis. Proceedings of the American Thoracic Society, 2012, 9, 158-163.	3.5	52
47	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
48	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
49	Antifibrotic role of vascular endothelial growth factor in pulmonary fibrosis. JCI Insight, 2017, 2, .	5.0	51
50	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
51	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
52	TRIM33 prevents pulmonary fibrosis by impairing TGF-Î²1 signalling. European Respiratory Journal, 2020, 55, 1901346.	6.7	45
53	Critical Evaluation of Sinonasal Disease in 64 Adults with Primary Ciliary Dyskinesia. Journal of Clinical Medicine, 2019, 8, 619.	2.4	44
54	Increased uptake of 111In-octreotide in idiopathic pulmonary fibrosis. Journal of Nuclear Medicine, 2006, 47, 1281-7.	5.0	44

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55	Usual interstitial pneumonia in ANCA-associated vasculitis: A poor prognostic factor. Journal of Autoimmunity, 2020, 106, 102338.	6.5	43
56	Anti-acid therapy in idiopathic pulmonary fibrosis: insights from the INPULSISÂ® trials. Respiratory Research, 2018, 19, 167.	3.6	42
57	The genetics of interstitial lung diseases. European Respiratory Review, 2019, 28, 190053.	7.1	41
58	Severe chronic bronchiolitis as the presenting feature of primary SjÃ¶grenâ€™s syndrome. Respiratory Medicine, 2011, 105, 130-136.	2.9	40
59	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
60	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
61	Safety and efficacy of pirfenidone in patients carrying telomerase complex mutation. European Respiratory Journal, 2018, 51, 1701875.	6.7	34
62	Detection of Alveolar Fibrocytes in Idiopathic Pulmonary Fibrosis and Systemic Sclerosis. PLoS ONE, 2013, 8, e53736.	2.5	33
63	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
64	Pharmacological management of IPF. Respirology, 2016, 21, 615-625.	2.3	32
65	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
66	Lung CT Densitometry in Idiopathic Pulmonary Fibrosis for the Prediction of Natural Course, Severity, and Mortality. Chest, 2019, 155, 972-981.	0.8	32
67	Psychometric properties and minimal important differences of SF-36 in Idiopathic Pulmonary Fibrosis. Respiratory Research, 2019, 20, 47.	3.6	31
68	Chaotic activation of developmental signalling pathways drives idiopathic pulmonary fibrosis. European Respiratory Review, 2020, 29, 190140.	7.1	31
69	Recent advances in rheumatoid arthritis-associated interstitial lung disease. Current Opinion in Pulmonary Medicine, 2020, 26, 477-486.	2.6	31
70	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
71	NADPH oxidase DUOX1 sustains TGF-Î²1 signalling and promotes lung fibrosis. European Respiratory Journal, 2021, 57, 1901949.	6.7	30
72	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

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73	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
74	Antineutrophil Cytoplasmic Antibodyâ€“Associated Lung Fibrosis. Seminars in Respiratory and Critical Care Medicine, 2018, 39, 465-470.	2.1	27
75	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
76	Pharmacological Targeting of Protease-Activated Receptor 2 Affords Protection from Bleomycin-Induced Pulmonary Fibrosis. Molecular Medicine, 2015, 21, 576-583.	4.4	24
77	Pilot experience of multidisciplinary team discussion dedicated to inherited pulmonary fibrosis. Orphanet Journal of Rare Diseases, 2019, 14, 280.	2.7	24
78	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
79	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
80	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
81	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
82	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
83	Familial forms of nonspecific interstitial pneumonia/idiopathic pulmonary fibrosis. Current Opinion in Pulmonary Medicine, 2012, 18, 455-461.	2.6	19
84	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
85	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
86	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
87	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
88	Licence to kill senescent cells in idiopathic pulmonary fibrosis?. European Respiratory Journal, 2017, 50, 1701360.	6.7	16
89	Lung function in Birt-Hogg-DubÃ© syndrome: a retrospective analysis of 96 patients. Orphanet Journal of Rare Diseases, 2020, 15, 120.	2.7	15
90	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

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91	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
92	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
93	Octreotide treatment of idiopathic pulmonary fibrosis: a proof-of-concept study. European Respiratory Journal, 2012, 39, 772-775.	6.7	14
94	Eight novel variants in the SLC34A2 gene in pulmonary alveolar microlithiasis. European Respiratory Journal, 2020, 55, 1900806.	6.7	14
95	Basophils and IgE contribute to mixed connective tissue disease development. Journal of Allergy and Clinical Immunology, 2021, 147, 1478-1489.e11.	2.9	14
96	First heterozygous NOP10 mutation in familial pulmonary fibrosis. European Respiratory Journal, 2020, 55, 1902465.	6.7	13
97	Identification of periplakin as a major regulator of lung injury and repair in mice. JCI Insight, 2018, 3, .	5.0	13
98	Pneumocystosis revealing immunodeficiency secondary to TERC mutation. European Respiratory Journal, 2017, 50, 1701443.	6.7	12
99	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
100	Observer agreement and clinical significance of chest CT reporting in patients suspected of COVID-19. European Radiology, 2021, 31, 1081-1089.	4.5	11
101	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
102	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
103	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
104	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
105	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
106	Genetic testing in interstitial lung disease: An international survey. Respirology, 0, , .	2.3	10
107	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
108	Immune checkpoint blockade for patients with lung cancer and idiopathic pulmonary fibrosis. European Journal of Cancer, 2021, 145, 179-182.	2.8	9

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109	Impact of genetic factors on fibrosing interstitial lung diseases. Incidence and clinical presentation in adults. Presse Medicale, 2020, 49, 104024.	1.9	9
110	Fibroblasts: the missing link between fibrotic lung diseases of different etiologies?. Respiratory Research, 2013, 14, 81.	3.6	8
111	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
112	NKX2.1 (TTF1) germline mutation associated with pulmonary fibrosis and lung cancer. ERJ Open Research, 2021, 7, 00356-2021.	2.6	8
113	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
114	Myelodysplastic syndromes and idiopathic pulmonary fibrosis: a dangerous liaison. Respiratory Research, 2019, 20, 182.	3.6	7
115	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
116	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
117	Human airway trypsinâ€like protease exerts potent, antifibrotic action in pulmonary fibrosis. FASEB Journal, 2018, 32, 1250-1264.	0.5	6
118	Rheumatological evaluation of patients with interstitial lung disease. Scandinavian Journal of Rheumatology, 2022, 51, 34-41.	1.1	6
119	Interstitial lung diseases associated with mutations of poly(A)â€specific ribonuclease: A multicentre retrospective study. Respirology, 2022, 27, 226-235.	2.3	6
120	A 70-Year-Old Woman With Acute Chest Pain and a Paracardiac Mass. Chest, 2013, 143, 866-869.	0.8	5
121	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
122	Towards a global initiative for fibrosis treatment (GIFT). ERJ Open Research, 2017, 3, 00106-2017.	2.6	5
123	Lung Fibroblasts from Idiopathic Pulmonary Fibrosis Patients Harbor Short and Unstable Telomeres Leading to Chromosomal Instability. Biomedicines, 2022, 10, 310.	3.2	5
124	Detection of anti-periplakin auto-antibodies during idiopathic pulmonary fibrosis. Clinica Chimica Acta, 2014, 433, 242.	1.1	4
125	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
126	Gastro-oesophageal reflux and idiopathic pulmonary fibrosis: in search of evidence. European Respiratory Journal, 2016, 48, 623-625.	6.7	3

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127	Familial pulmonary fibrosis: a world without frontiers. Jornal Brasileiro De Pneumologia, 2019, 45, e20190303.	0.7	3
128	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
129	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
130	Giant hiatal hernia: beware of the supine ICU chest X-ray!. BMJ Case Reports, 2017, 2017, bcr-2017-219668.	0.5	2
131	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
132	European IPF Patient Charter: an SOS to the world. European Respiratory Journal, 2016, 47, 403-405.	6.7	1
133	Alveolar epithelial TET2 is not involved in the development of bleomycinâ€induced pulmonary fibrosis. FASEB Journal, 2021, 35, e21599.	0.5	1
134	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
135	Molecular biomarkers in idiopathic pulmonary fibrosis and disease severity. Sarcoidosis Vasculitis and Diffuse Lung Diseases, 2013, 30 Suppl 1, 27-32.	0.2	1
136	Targeting the nasty nestin to shoot lung fibrosis. European Respiratory Journal, 2022, 59, 2103146.	6.7	1
137	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
138	The FLORA study: presenting a novel IPF trial design. Lancet Respiratory Medicine,the, 2018, 6, 572-573.	10.7	0
139	The Genetics of Interstitial Lung Diseases. , 2022, , 96-113.		0
140	Pneumopathies interstitielles. Revue Des Maladies Respiratoires Actualites, 2018, 10, S21-S23.	0.0	0
141	Telomere syndrome and the lung. , 2019, , 391-403.		0
142	Title is missing!. , 2020, 15, e0243961.		0
143	Title is missing!. , 2020, 15, e0243961.		0
144	Title is missing!. , 2020, 15, e0243961.		0

#	Article	IF	Citations
145	Title is missing!. , 2020, 15, e0243961.		0
146	Fibrotic-Like CT Alterations in COVID-19: Distinct Patterns of Temporal Evolution. , 2022, , .		0
147	Higher basal tryptase, asthma and loss of consciousness in anaphylaxis are associated with biphasic reactions. Clinical and Translational Allergy, 2022, 12, .	3.2	0