

Elisabeth Bendstrup

List of Publications by Year in descending order

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Version: 2024-02-01

152
papers

4,463
citations

147786

31
h-index

123420

61
g-index

160
all docs

160
docs citations

160
times ranked

4299
citing authors

#	ARTICLE	IF	CITATIONS
1	Diagnosis of Hypersensitivity Pneumonitis in Adults: An Official ATS/JRS/ALAT Clinical Practice Guideline. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 202, e36-e69.	5.6	508
2	Multicentre evaluation of multidisciplinary team meeting agreement on diagnosis in diffuse parenchymal lung disease: a case-cohort study. <i>Lancet Respiratory Medicine</i> , 2016, 4, 557-565.	10.7	337
3	A population-based cohort study of rheumatoid arthritis-associated interstitial lung disease: comorbidity and mortality. <i>Annals of the Rheumatic Diseases</i> , 2017, 76, 1700-1706.	0.9	263
4	Palliative care in interstitial lung disease: living well. <i>Lancet Respiratory Medicine</i> , 2017, 5, 968-980.	10.7	185
5	Outcome of Hospitalization for COVID-19 in Patients with Interstitial Lung Disease. An International Multicenter Study. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 202, 1656-1665.	5.6	171
6	Global incidence and prevalence of idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2021, 22, 197.	3.6	170
7	Progressive fibrosing interstitial lung diseases: current practice in diagnosis and management. <i>Current Medical Research and Opinion</i> , 2019, 35, 2015-2024.	1.9	148
8	A cohort study of interstitial lung diseases in central Denmark. <i>Respiratory Medicine</i> , 2014, 108, 793-799.	2.9	128
9	How does comorbidity influence survival in idiopathic pulmonary fibrosis?. <i>Respiratory Medicine</i> , 2014, 108, 647-653.	2.9	125
10	Safety and tolerability of acetylcysteine and pirfenidone combination therapy in idiopathic pulmonary fibrosis: a randomised, double-blind, placebo-controlled, phase 2 trial. <i>Lancet Respiratory Medicine</i> , 2016, 4, 445-453.	10.7	108
11	Managing the supportive care needs of those affected by COVID-19. <i>European Respiratory Journal</i> , 2020, 55, 2000815.	6.7	95
12	Resequencing Study Confirms That Host Defense and Cell Senescence Gene Variants Contribute to the Risk of Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 199-208.	5.6	90
13	Updated guidance on the management of COVID-19: from an American Thoracic Society/European Respiratory Society coordinated International Task Force (29 July 2020). <i>European Respiratory Review</i> , 2020, 29, 200287.	7.1	82
14	Risk factors for diagnostic delay in idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2019, 20, 103.	3.6	78
15	Diagnostic accuracy of a clinical diagnosis of idiopathic pulmonary fibrosis: an international case-cohort study. <i>European Respiratory Journal</i> , 2017, 50, 1700936.	6.7	75
16	Acute exacerbation of idiopathic pulmonary fibrosis: international survey and call for harmonisation. <i>European Respiratory Journal</i> , 2020, 55, 1901760.	6.7	61
17	Diagnostic Likelihood Thresholds That Define a Working Diagnosis of Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 1146-1153.	5.6	60
18	Autoimmune pulmonary alveolar proteinosis: Treatment options in year 2013. <i>Respirology</i> , 2013, 18, 82-91.	2.3	58

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19	Unclassifiable interstitial lung diseases: Clinical characteristics and survival. <i>Respirology</i> , 2017, 22, 494-500.	2.3	58
20	Idiopathic Pulmonary Fibrosis: Best Practice in Monitoring and Managing a Relentless Fibrotic Disease. <i>Respiration</i> , 2020, 99, 73-82.	2.6	58
21	Preferred Place of Care and Death in Terminally Ill Patients with Lung and Heart Disease Compared to Cancer Patients. <i>Journal of Palliative Medicine</i> , 2017, 20, 1217-1224.	1.1	54
22	Mindfulness-based cognitive therapy in COPD: a cluster randomised controlled trial. <i>European Respiratory Journal</i> , 2018, 51, 1702082.	6.7	52
23	Interstitial Lung Disease in Rheumatoid Arthritis Remains a Challenge for Clinicians. <i>Journal of Clinical Medicine</i> , 2019, 8, 2038.	2.4	51
24	Opioids: an unexplored option for treatment of dyspnea in IPF. <i>European Clinical Respiratory Journal</i> , 2016, 3, 30629.	1.5	48
25	Rheumatoid Arthritis-Associated Interstitial Lung Disease: Clinical Characteristics and Predictors of Mortality. <i>Respiration</i> , 2019, 98, 455-460.	2.6	47
26	Pulmonary alveolar microlithiasis: two case reports and review of the literature. <i>European Respiratory Review</i> , 2012, 21, 249-256.	7.1	45
27	Introduction of cryobiopsies in the diagnostics of interstitial lung diseases – experiences in a referral center. <i>European Clinical Respiratory Journal</i> , 2017, 4, 1274099.	1.5	44
28	Statin Therapy and Outcomes in Trials of Nintedanib in Idiopathic Pulmonary Fibrosis. <i>Respiration</i> , 2018, 95, 317-326.	2.6	42
29	Nintedanib in Idiopathic Pulmonary Fibrosis: Practical Management Recommendations for Potential Adverse Events. <i>Respiration</i> , 2019, 97, 173-184.	2.6	39
30	Validation of the IPF-specific version of St. George's Respiratory Questionnaire. <i>Respiratory Research</i> , 2019, 20, 199.	3.6	37
31	A systematic review of occupational exposure to coal dust and the risk of interstitial lung diseases. <i>European Clinical Respiratory Journal</i> , 2017, 4, 1264711.	1.5	35
32	The need for a holistic approach for SSC-ILD – achievements and ambiguity in a devastating disease. <i>Respiratory Research</i> , 2020, 21, 197.	3.6	33
33	Desquamative interstitial pneumonia: a systematic review of its features and outcomes. <i>European Respiratory Review</i> , 2020, 29, 190181.	7.1	32
34	Acute exacerbation of interstitial lung disease associated with rheumatic disease. <i>Nature Reviews Rheumatology</i> , 2022, 18, 85-96.	8.0	28
35	Treatment of idiopathic pulmonary fibrosis: a position paper from a Nordic expert group. <i>Journal of Internal Medicine</i> , 2017, 281, 149-166.	6.0	27
36	Tele-Rehabilitation Program in Idiopathic Pulmonary Fibrosis – A Single-Center Randomized Trial. <i>International Journal of Environmental Research and Public Health</i> , 2021, 18, 10016.	2.6	27

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37	Pulmonary manifestations of pyoderma gangrenosum: 2 cases and a review of the literature. <i>Respiratory Medicine</i> , 2015, 109, 443-450.	2.9	26
38	Pirfenidone treatment in idiopathic pulmonary fibrosis: nationwide Danish results. <i>European Clinical Respiratory Journal</i> , 2016, 3, 32608.	1.5	26
39	Birtâ€“Hoggâ€“DubÃ© syndrome: a case report and a review of the literature. <i>European Clinical Respiratory Journal</i> , 2017, 4, 1292378.	1.5	24
40	Increased mortality among patients with rheumatoid arthritis and COPD: A population-based study. <i>Respiratory Medicine</i> , 2018, 140, 101-107.	2.9	22
41	Validation of the Kingâ€™s Brief Interstitial Lung Disease questionnaire in Idiopathic Pulmonary Fibrosis. <i>BMC Pulmonary Medicine</i> , 2019, 19, 255.	2.0	22
42	Palliation of chronic breathlessness with morphine in patients with fibrotic interstitial lung disease â€“ a randomised placebo-controlled trial. <i>Respiratory Research</i> , 2020, 21, 195.	3.6	22
43	The DIAMORFOSIS (DIagnosis and Management Of lung cancer and FibrOSIS) survey: international survey and call for consensus. <i>ERJ Open Research</i> , 2021, 7, 00529-2020.	2.6	22
44	Efficacy and safety of nintedanib in patients with idiopathic pulmonary fibrosis who are elderly or have comorbidities. <i>Respiratory Research</i> , 2021, 22, 125.	3.6	22
45	SLC34A2 Gene Mutation May Explain Comorbidity of Pulmonary Alveolar Microlithiasis and Aortic Valve Sclerosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2012, 185, 464-464.	5.6	21
46	Rituximab-induced interstitial lung disease: five case reports. <i>European Clinical Respiratory Journal</i> , 2015, 2, 27178.	1.5	21
47	Responsiveness and minimal clinically important difference of SGRQ-I and K-BILD in idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2020, 21, 91.	3.6	21
48	Risk of hypersensitivity pneumonitis and interstitial lung diseases among pigeon breeders. <i>European Respiratory Journal</i> , 2016, 48, 818-825.	6.7	20
49	Turnover of type I and III collagen predicts progression of idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2021, 22, 205.	3.6	20
50	Voriconazole Concentrations in Plasma and Epithelial Lining Fluid after Inhalation and Oral Treatment. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2017, 121, 430-434.	2.5	19
51	Screening tools for evaluation of depression in Chronic Obstructive Pulmonary Disease (COPD). A systematic review. <i>European Clinical Respiratory Journal</i> , 2017, 4, 1332931.	1.5	19
52	Clinical characteristics and outcome in patients with antisynthetase syndrome associated interstitial lung disease: a retrospective cohort study. <i>European Clinical Respiratory Journal</i> , 2019, 6, 1583516.	1.5	19
53	Effect of a New Tele-Rehabilitation Program versus Standard Rehabilitation in Patients with Chronic Obstructive Pulmonary Disease. <i>Journal of Clinical Medicine</i> , 2022, 11, 11.	2.4	19
54	Patient-reported outcomes and patient-reported outcome measures in interstitial lung disease: where to go from here?. <i>European Respiratory Review</i> , 2021, 30, 210026.	7.1	17

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55	Immunoglobulin G4-related pleuritis – A case report. <i>Respiratory Medicine Case Reports</i> , 2016, 19, 18-20.	0.4	16
56	Pirfenidone Treatment in Individuals with Idiopathic Pulmonary Fibrosis: Impact of Timing of Treatment Initiation. <i>Annals of the American Thoracic Society</i> , 2019, 16, 927-930.	3.2	16
57	Childhood pneumothorax in Birt-Hogg-Åub© syndrome: A cohort study and review of the literature. <i>Molecular Genetics & Genomic Medicine</i> , 2018, 6, 332-338.	1.2	15
58	Design of a Study Assessing Disease Behaviour During the Peri-Diagnostic Period in Patients with Interstitial Lung Disease: The STARLINER Study. <i>Advances in Therapy</i> , 2019, 36, 232-243.	2.9	15
59	Augmented reality glasses as a new tele-rehabilitation tool for home use: patients'™ perception and expectations. <i>Disability and Rehabilitation: Assistive Technology</i> , 2022, 17, 480-486.	2.2	15
60	Patient reported outcome measures (PROMs) in sarcoidosis. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2017, 34, 2-17.	0.2	15
61	Eight novel variants in the <i>SLC34A2</i> gene in pulmonary alveolar microlithiasis. <i>European Respiratory Journal</i> , 2020, 55, 1900806.	6.7	14
62	Interstitial Lung Disease in Connective Tissue Diseases: Survival Patterns in a Population-Based Cohort. <i>Journal of Clinical Medicine</i> , 2021, 10, 4830.	2.4	14
63	Cultural Differences in Palliative Care in Patients With Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2015, 148, e56.	0.8	13
64	NT-proBNP ≤ 95 ng/l can exclude pulmonary hypertension on echocardiography at diagnostic workup in patients with interstitial lung disease. <i>European Clinical Respiratory Journal</i> , 2016, 3, 32027.	1.5	13
65	Continuous Rituximab treatment for recurrent diffuse alveolar hemorrhage in a patient with systemic lupus erythematosus and antiphospholipid syndrome. <i>Respiratory Medicine Case Reports</i> , 2017, 22, 263-265.	0.4	13
66	Comorbidity and mortality among patients with interstitial lung diseases: A population-based study. <i>Respirology</i> , 2018, 23, 606-612.	2.3	13
67	Danish respiratory society position paper: palliative care in patients with chronic progressive non-malignant lung diseases. <i>European Clinical Respiratory Journal</i> , 2018, 5, 1530029.	1.5	13
68	Tele-delivered mindfulness-based cognitive therapy in chronic obstructive pulmonary disease: A mixed-methods feasibility study. <i>Journal of Telemedicine and Telecare</i> , 2019, 25, 468-475.	2.7	13
69	Outcomes of patients with advanced idiopathic pulmonary fibrosis treated with nintedanib or pirfenidone in a real-world multicentre cohort. <i>Respirology</i> , 2021, 26, 982-988.	2.3	13
70	Clusters of comorbidities in idiopathic pulmonary fibrosis. <i>Respiratory Medicine</i> , 2021, 185, 106490.	2.9	13
71	PD-L1 Expression in Patients with Idiopathic Pulmonary Fibrosis. <i>Journal of Clinical Medicine</i> , 2021, 10, 5562.	2.4	13
72	Lung ultrasound has limited diagnostic value in rare cystic lung diseases: a cross-sectional study. <i>European Clinical Respiratory Journal</i> , 2017, 4, 1330111.	1.5	12

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73	COPD: an overlooked cause of excess mortality in patients with rheumatoid arthritis. <i>Lancet Respiratory Medicine</i> , 2018, 6, 326-327.	10.7	12
74	Advance care planning for patients with lung, heart and cancer diseases and their relatives. <i>International Journal of Palliative Nursing</i> , 2019, 25, 112-127.	0.5	12
75	Attitude and Barriers in Palliative Care and Advance Care Planning in Nonmalignant Chronic Lung Disease: Results From a Danish National Survey. <i>Journal of Palliative Care</i> , 2020, 35, 232-235.	1.0	12
76	Integration of cryobiopsies for interstitial lung disease diagnosis is a valid and safe diagnostic strategy—experiences based on 250 biopsy procedures. <i>Journal of Thoracic Disease</i> , 2021, 13, 1455-1465.	1.4	12
77	Pulmonary alveolar microlithiasis: no longer in the stone age. <i>ERJ Open Research</i> , 2020, 6, 00289-2020.	2.6	12
78	Review of IPF diagnosis and management recommendations in Europe. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2013, 30, 249-61.	0.2	12
79	Fatigue Is a Major Symptom at COVID-19 Hospitalization Follow-Up. <i>Journal of Clinical Medicine</i> , 2022, 11, 2411.	2.4	12
80	Cryobiopsies are diagnostic in Pleuroparenchymal and Airway-centered Fibroelastosis. <i>Respiratory Research</i> , 2018, 19, 135.	3.6	11
81	Report Standardization in Transbronchial Lung Cryobiopsy. <i>Archives of Pathology and Laboratory Medicine</i> , 2019, 143, 416-417.	2.5	10
82	Identifying unmet needs in SSc-ILD by semi-qualitative in-depth interviews. <i>Rheumatology</i> , 2021, 60, 5601-5609.	1.9	10
83	The case of methotrexate and the lung: Dr Jekyll and Mr Hyde. <i>European Respiratory Journal</i> , 2021, 57, 2100079.	6.7	10
84	Lung Ultrasound to Phenotype Chronic Lung Allograft Dysfunction in Lung Transplant Recipients. A Prospective Observational Study. <i>Journal of Clinical Medicine</i> , 2021, 10, 1078.	2.4	10
85	Organisation of diagnosis and treatment of idiopathic pulmonary fibrosis and other interstitial lung diseases in the Nordic countries. <i>European Clinical Respiratory Journal</i> , 2015, 2, 28348.	1.5	9
86	Hypersensitivity pneumonitis among wind musicians — an overlooked disease?. <i>European Clinical Respiratory Journal</i> , 2017, 4, 1351268.	1.5	9
87	Granulomatous-lymphocytic interstitial lung disease and recurrent sinopulmonary infections in a patient with Good's syndrome. <i>BMJ Case Reports</i> , 2015, 2015, bcr2014205635.	0.5	9
88	Advance care planning and longer survival in the terminally ill: a randomised controlled trial unexpected finding. <i>BMJ Supportive and Palliative Care</i> , 2020, 10, 221-222.	1.6	8
89	Screening Tools for Depression and Anxiety in Patients with Chronic Obstructive Pulmonary Disease — A Systematic Review. <i>COPD: Journal of Chronic Obstructive Pulmonary Disease</i> , 2021, 18, 683-689.	1.6	8
90	Management of Acute Exacerbation of Idiopathic Pulmonary Fibrosis in Specialised and Non-specialised ILD Centres Around the World. <i>Frontiers in Medicine</i> , 2021, 8, 699644.	2.6	8

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91	A pilot study on the use of the super dimension navigation system for optimal cryobiopsy location in interstitial lung disease diagnostics. <i>Pulmonology</i> , 2023, 29, 119-123.	2.1	8
92	Changes in management of idiopathic pulmonary fibrosis: impact on disease severity and mortality. <i>European Clinical Respiratory Journal</i> , 2020, 7, 1807682.	1.5	7
93	High turnover of types <scp>III</scp> and <scp>VI</scp> collagen in progressive idiopathic pulmonary fibrosis. <i>Respirology</i> , 2021, 26, 582-589.	2.3	7
94	An RCT of acute health effects in COPD-patients after passive vape exposure from e-cigarettes. <i>European Clinical Respiratory Journal</i> , 2021, 8, 1861580.	1.5	7
95	Eosinophil alveolitis in two patients with idiopathic pulmonary fibrosis. <i>Respiratory Medicine Case Reports</i> , 2016, 19, 61-64.	0.4	6
96	Fatigue in idiopathic pulmonary fibrosis measured by the Fatigue Assessment Scale during antifibrotic treatment. <i>European Clinical Respiratory Journal</i> , 2021, 8, 1853658.	1.5	6
97	Disease Behaviour During the Peri-Diagnostic Period in Patients with Suspected Interstitial Lung Disease: The STARLINER Study. <i>Advances in Therapy</i> , 2021, 38, 4040-4056.	2.9	6
98	Lung Ultrasound â€œ A Novel Diagnostic Tool To Phenotype Chronic Lung Allograft Dysfunction?. <i>Ultrasound International Open</i> , 2017, 03, E117-E119.	0.6	5
99	Remarkable benefits of intravenous immunoglobulin (IVIG) in a patient with polymyositis-associated acute interstitial lung disease. <i>European Clinical Respiratory Journal</i> , 2020, 7, 1840706.	1.5	5
100	A global perspective on acute exacerbation of idiopathic pulmonary fibrosis (AE-IPF): results from an international survey. , 2018, , .		5
101	Longitudinal serological assessment of type VI collagen turnover is related to progression in a real-world cohort of idiopathic pulmonary fibrosis. <i>BMC Pulmonary Medicine</i> , 2021, 21, 382.	2.0	5
102	Comorbidities in unclassifiable interstitial lung disease. <i>Respiratory Research</i> , 2022, 23, 59.	3.6	5
103	Increasing awareness of corticosteroid hypersensitivity reactions is important. <i>Respirology Case Reports</i> , 2013, 1, 43-45.	0.6	4
104	Tracheal collapse diagnosed by multidetector computed tomography: evaluation of different image analysis methods. <i>European Clinical Respiratory Journal</i> , 2017, 4, 1407624.	1.5	4
105	Autoimmune pulmonary alveolar proteinosis in an adolescent successfully treated with inhaled rhGM-CSF (molgramostim). <i>Respiratory Medicine Case Reports</i> , 2018, 23, 167-169.	0.4	4
106	Economic consequences of idiopathic pulmonary fibrosis in Denmark. <i>ERJ Open Research</i> , 2018, 4, 00045-2017.	2.6	4
107	Familial idiopathic pulmonary fibrosis in a young female. <i>Respiratory Medicine Case Reports</i> , 2018, 24, 1-4.	0.4	4
108	Surgical Lung Biopsy and Cryobiopsy in Fibrosing Interstitial Lung Diseases: One Swallow Does Not Make a Summer. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 939-940.	5.6	4

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109	Tracheal collapsibility in adults is dynamic over time. <i>Respiratory Medicine</i> , 2019, 146, 124-128.	2.9	4
110	Interstitial lung diseases: quo vadis?. <i>Lancet Respiratory Medicine</i> , 2021, 9, 1084-1087.	10.7	4
111	Emphysema mimicking interstitial lung disease: Two case reports. <i>Respiratory Medicine Case Reports</i> , 2015, 15, 24-26.	0.4	3
112	The effect of the walk-bike on quality of life and exercise capacity in patients with idiopathic pulmonary fibrosis: a feasibility study. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2020, 37, 192-202.	0.2	3
113	Impaired phosphate transport in SLC34A2 variants in patients with pulmonary alveolar microlithiasis. <i>Human Genomics</i> , 2022, 16, 13.	2.9	3
114	No effect of pirfenidone treatment in fulminant bleomycin-induced pneumonitis. <i>Respiratory Medicine Case Reports</i> , 2014, 12, 47-49.	0.4	2
115	Pulmonary hemorrhage following anabolic agent abuse: Two cases. <i>Respiratory Medicine Case Reports</i> , 2016, 18, 45-47.	0.4	2
116	Early referral to palliative care in IPF – pitfalls and opportunities in clinical trials. <i>Respiratory Research</i> , 2020, 21, 174.	3.6	2
117	Comparison of Palliative Care Models in Idiopathic Pulmonary Fibrosis. <i>Applied Sciences (Switzerland)</i> , 2021, 11, 9028.	2.5	2
118	Suggestions for improving clinical utility of future guidelines for diagnosis and management of idiopathic pulmonary fibrosis: results of a Delphi survey. <i>European Respiratory Journal</i> , 2021, 57, 2004219.	6.7	2
119	Validation of a derived version of the IPF-specific Saint George’s Respiratory Questionnaire. <i>Respiratory Research</i> , 2021, 22, 259.	3.6	2
120	Erectile dysfunction is a common problem in interstitial lung diseases. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2017, 34, 356-364.	0.2	2
121	Challenges in the classification of fibrotic ILD. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2015, 32 Suppl 1, 4-9.	0.2	2
122	Careful Planning Reduces Cryobiopsy Complications. <i>Annals of the American Thoracic Society</i> , 2017, 14, 1229.	3.2	2
123	Acute Exacerbations in Patients with Progressive Fibrosing Interstitial Lung Diseases: Data from the INBUILD Trial. , 2022, , .		2
124	Real-World Data on Bleeding Risk and Anticoagulation in Patients with IPF Treated with Antifibrotics. <i>Drug Safety</i> , 2020, 43, 953-955.	3.2	1
125	International multidisciplinary team discussions on the diagnosis of idiopathic non-specific interstitial pneumonia and the development of connective tissue disease. <i>European Clinical Respiratory Journal</i> , 2021, 8, 1933878.	1.5	1
126	Decline in Forced Vital Capacity as a Surrogate for Mortality in Patients with Fibrosing Interstitial Lung Diseases. , 2021, , .		1

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127	Efficacy and safety of nintedanib in the elderly patient with IPF. , 2019, , .		1
128	New-onset asthma in a bilateral lung transplant patient. BMJ Case Reports, 2019, 12, e231654.	0.5	1
129	Pulmonary Alveolar Microlithiasis - A Review.. Yale Journal of Biology and Medicine, 2021, 94, 637-644.	0.2	1
130	Intravenous immunoglobulin treatment stabilizing a patient with Anti-PL7 antisynthetase syndrome with interstitial lung disease and eosinophilic inflammation. Respiratory Medicine Case Reports, 2022, , 101686.	0.4	1
131	0191â€¦Are indoor concentrations of airborne mould spores in non-industrial environments sufficiently high to cause hypersensitivity pneumonitis?. Occupational and Environmental Medicine, 2014, 71, A85.2-A85.	2.8	0
132	O40-1â€¦Risk of hypersensitivity pneumonitis and other interstitial lung diseases among pigeon breeders. , 2016, , .		0
133	FRI0174â€¦A population based cohort study of rheumatoid arthritis-associated interstitial lung disease: comorbidity and mortality. , 2017, , .		0
134	O435â€¦Pigeon breeding and the risk of interstitial lung disease, does number of pigeons matter?. , 2017, , .		0
135	Differences in the Approach to Acute Exacerbation of Idiopathic Pulmonary Fibrosis (AE IPF) Between Expert Centres and General Pulmonologists: Results from an International Survey. , 2019, , .		0
136	Pulmonary alveolar proteinosis - a crazy presentation of dyspnea. European Clinical Respiratory Journal, 2019, 6, 1552065.	1.5	0
137	Efficacy and Safety of Nintedanib in Patients with Idiopathic Pulmonary Fibrosis and Multiple Comorbidities. , 2020, , .		0
138	Inhaled GM-CSF (Molgramostim) Therapy Reduces the Need for Whole Lung Lavage in Patients with Autoimmune Pulmonary Alveolar Proteinosis - Long-Term Results from a Randomized, Double-Blind Trial (IMPALA). , 2020, , .		0
139	A Pilot Study on the Utility of SuperDimension to Guide Cryobiopsies in ILDs. , 2021, , .		0
140	Fibrin Formation Measured in Serum Is Related to Disease Severity in IPF Patients. , 2021, , .		0
141	Estimated Global Prevalence of Idiopathic Pulmonary Fibrosis. , 2021, , .		0
142	Health Related Quality of Life and Symptoms Are Related to Mortality in Idiopathic Pulmonary Fibrosis. , 2021, , .		0
143	Ein globaler Blick auf akute Exazerbationen der idiopathischen Lungenfibrose (AE-IPF): Ergebnisse einer internationalen Umfrage. , 2019, 73, .		0
144	P-101â€¦Occupational dust exposures and CT findings of interstitial lung disease and chronic obstructive pulmonary disease. , 2021, , .		0

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145	Efficacy and safety of nintedanib in the elderly patient with IPF*. , 2020, 74, .		0
146	Design and rationale of ProSar, the first Danish sarcoidosis registry.. Sarcoidosis Vasculitis and Diffuse Lung Diseases, 2022, 38, e2021044.	0.2	0
147	Characteristics of Danish Patients with Pulmonary Sarcoidosis. , 2022, , .		0
148	Efficacy and Safety of Nintedanib in Elderly Patients with Progressive Fibrosing Interstitial Lung Diseases (ILDs). , 2022, , .		0
149	A Neo-Epitope Biomarker of Type VI Collagen Formation Reflects Patient-Reported Outcomes. , 2022, , .		0
150	Lung Cancer in Patients with Idiopathic Pulmonary Fibrosis: A Retrospective Multicenter Study in Europe. , 2022, , .		0
151	Effect of a Telerehabilitation program in sarcoidosis.. Sarcoidosis Vasculitis and Diffuse Lung Diseases, 2022, 39, e2022003.	0.2	0
152	Decline in forced vital capacity as a surrogate for mortality in patients with fibrosing interstitial lung diseases*. Pneumologie, 2022, , .	0.1	0