Marlies S Wijsenbeek

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

97
papers

2,189
citations

h-index

43
g-index

114
ext. papers

9.6
avg, IF

L-index

#	Paper	IF	Citations
97	Safety and efficacy of ustekinumab or golimumab in patients with chronic sarcoidosis. <i>European Respiratory Journal</i> , 2014 , 44, 1296-307	13.6	126
96	Palliative care in interstitial lung disease: living well. Lancet Respiratory Medicine, the, 2017, 5, 968-980	35.1	115
95	Identification of Diagnostic Criteria for Chronic Hypersensitivity Pneumonitis: An International Modified Delphi Survey. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018 , 197, 1036-104	4 ^{10.2}	109
94	Effect of Recombinant Human Pentraxin 2 vs Placebo on Change in Forced Vital Capacity in Patients With Idiopathic Pulmonary Fibrosis: A Randomized Clinical Trial. <i>JAMA - Journal of the American Medical Association</i> , 2018 , 319, 2299-2307	27.4	107
93	Spectrum of Fibrotic Lung Diseases. <i>New England Journal of Medicine</i> , 2020 , 383, 958-968	59.2	102
92	European IPF Patient Charter: unmet needs and a call to action for healthcare policymakers. <i>European Respiratory Journal</i> , 2016 , 47, 597-606	13.6	71
91	A novel formulation of inhaled sodium cromoglicate (PA101) in idiopathic pulmonary fibrosis and chronic cough: a randomised, double-blind, proof-of-concept, phase 2 trial. <i>Lancet Respiratory Medicine,the</i> , 2017 , 5, 806-815	35.1	71
90	Progressive fibrosing interstitial lung diseases: current practice in diagnosis and management. <i>Current Medical Research and Opinion</i> , 2019 , 35, 2015-2024	2.5	65
89	Cough in idiopathic pulmonary fibrosis. European Respiratory Review, 2016, 25, 278-86	9.8	62
88	Recombinant human pentraxin-2 therapy in patients with idiopathic pulmonary fibrosis: safety, pharmacokinetics and exploratory efficacy. <i>European Respiratory Journal</i> , 2016 , 47, 889-97	13.6	54
87	Safety of nintedanib added to pirfenidone treatment for idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2018 , 52,	13.6	53
86	Rationale, design and objectives of two phase III, randomised, placebo-controlled studies of GLPG1690, a novel autotaxin inhibitor, in idiopathic pulmonary fibrosis (ISABELA 1 and 2). <i>BMJ Open Respiratory Research</i> , 2019 , 6, e000422	5.6	50
85	Diagnostic accuracy of a clinical diagnosis of idiopathic pulmonary fibrosis: an international case-cohort study. <i>European Respiratory Journal</i> , 2017 , 50,	13.6	50
84	Optimizing quality of life in patients with idiopathic pulmonary fibrosis. <i>Therapeutic Advances in Respiratory Disease</i> , 2017 , 11, 157-169	4.9	49
83	Increased T-helper 17.1 cells in sarcoidosis mediastinal lymph nodes. <i>European Respiratory Journal</i> , 2018 , 51,	13.6	48
82	Pharmacological Treatment of Idiopathic Pulmonary Fibrosis: Current Approaches, Unsolved Issues, and Future Perspectives. <i>BioMed Research International</i> , 2015 , 2015, 329481	3	44
81	Unfavourable effects of medically indicated oral anticoagulants on survival in idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2016 , 47, 1776-84	13.6	43

80	Treatment of Sarcoidosis. Clinics in Chest Medicine, 2015, 36, 751-67	5.3	42
79	Long-term treatment with recombinant human pentraxin 2 protein in patients with idiopathic pulmonary fibrosis: an open-label extension study. <i>Lancet Respiratory Medicine,the</i> , 2019 , 7, 657-664	35.1	42
78	Idiopathic Pulmonary Fibrosis (an Update) and Progressive Pulmonary Fibrosis in Adults: An Official ATS/ERS/JRS/ALAT Clinical Practice Guideline <i>American Journal of Respiratory and Critical Care Medicine</i> , 2022 , 205, e18-e47	10.2	38
77	Effect of pirfenidone on cough in patients with idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2017 , 50,	13.6	37
76	Consensus document for the selection of lung transplant candidates: An update from the International Society for Heart and Lung Transplantation. <i>Journal of Heart and Lung Transplantation</i> , 2021 , 40, 1349-1379	5.8	34
75	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	10.2	33
74	Idiopathic Pulmonary Fibrosis: Best Practice in Monitoring and Managing a Relentless Fibrotic Disease. <i>Respiration</i> , 2020 , 99, 73-82	3.7	30
73	The added value of comorbidities in predicting survival in idiopathic pulmonary fibrosis: a multicentre observational study. <i>European Respiratory Journal</i> , 2019 , 53,	13.6	30
72	Statin Therapy and Outcomes in Trials of Nintedanib in Idiopathic Pulmonary Fibrosis. <i>Respiration</i> , 2018 , 95, 317-326	3.7	29
71	Home Monitoring in Patients with Idiopathic Pulmonary Fibrosis. A Randomized Controlled Trial. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020 , 202, 393-401	10.2	28
70	Identifying Barriers to Idiopathic Pulmonary Fibrosis Treatment: A Survey of Patient and Physician Views. <i>Respiration</i> , 2018 , 96, 514-524	3.7	27
69	Nintedanib in Idiopathic Pulmonary Fibrosis: Practical Management Recommendations for Potential Adverse Events. <i>Respiration</i> , 2019 , 97, 173-184	3.7	27
68	No evidence found for an association between prednisone dose and FVC change in newly-treated pulmonary sarcoidosis. <i>Respiratory Medicine</i> , 2018 , 138S, S31-S37	4.6	27
67	Comprehensive Supportive Care for Patients with Fibrosing Interstitial Lung Disease. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019 , 200, 152-159	10.2	25
66	Early Experience of Pirfenidone in Daily Clinical Practice in Belgium and The Netherlands: a Retrospective Cohort Analysis. <i>Advances in Therapy</i> , 2015 , 32, 691-704	4.1	24
65	Daily home spirometry to detect early steroid treatment effects in newly treated pulmonary sarcoidosis. <i>European Respiratory Journal</i> , 2018 , 51,	13.6	24
64	Th17-lineage cells in pulmonary sarcoidosis and LEgren's syndrome: Friend or foe?. <i>Journal of Autoimmunity</i> , 2018 , 87, 82-96	15.5	22
63	What patients with pulmonary fibrosis and their partners think: a live, educative survey in the Netherlands and Germany. <i>ERJ Open Research</i> , 2017 , 3,	3.5	20

62	Acute exacerbation of idiopathic pulmonary fibrosis: international survey and call for harmonisation. <i>European Respiratory Journal</i> , 2020 , 55,	13.6	18
61	Integrating Patient Perspectives into Personalized Medicine in Idiopathic Pulmonary Fibrosis. <i>Frontiers in Medicine</i> , 2017 , 4, 226	4.9	18
60	Medical Therapy in Idiopathic Pulmonary Fibrosis. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2016 , 37, 368-77	3.9	17
59	Gaps in care of patients living with pulmonary fibrosis: a joint patient and expert statement on the results of a Europe-wide survey. <i>ERJ Open Research</i> , 2019 , 5,	3.5	17
58	Unfavourable effects of medically indicated oral anticoagulants on survival in idiopathic pulmonary fibrosis: methodological concerns. <i>European Respiratory Journal</i> , 2016 , 48, 1524-1526	13.6	16
57	Cardiovascular safety of nintedanib in subgroups by cardiovascular risk at baseline in the TOMORROW and INPULSIS[trials. <i>European Respiratory Journal</i> , 2019 , 54,	13.6	16
56	Desquamative interstitial pneumonia: a systematic review of its features and outcomes. <i>European Respiratory Review</i> , 2020 , 29,	9.8	15
55	New insights on patient-reported outcome measures in idiopathic pulmonary fibrosis: only PROMises?. <i>Current Opinion in Pulmonary Medicine</i> , 2016 , 22, 434-41	3	15
54	Patient and partner empowerment programme for idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2017 , 49,	13.6	14
53	Feasibility of a Comprehensive Home Monitoring Program for Sarcoidosis. <i>Journal of Personalized Medicine</i> , 2019 , 9,	3.6	14
52	Comprehensive Care for Patients with Sarcoidosis. Journal of Clinical Medicine, 2020, 9,	5.1	14
51	Health-related quality of life and symptoms in patients with IPF treated with nintedanib: analyses of patient-reported outcomes from the INPULSIS trials. <i>Respiratory Research</i> , 2020 , 21, 36	7.3	13
50	Cultural Differences in Palliative Care in Patients With Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2015 , 148, e56	5.3	13
49	The need for a holistic approach for SSc-ILD - achievements and ambiguity in a devastating disease. <i>Respiratory Research</i> , 2020 , 21, 197	7.3	13
48	Development and Pretesting of a Questionnaire to Assess Patient Experiences and Satisfaction with Medications (PESaM Questionnaire). <i>Patient</i> , 2017 , 10, 629-642	3.7	12
47	Enhanced Bruton's tyrosine kinase in B-cells and autoreactive IgA in patients with idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2019 , 20, 232	7.3	11
46	Nintedanib and immunomodulatory therapies in progressive fibrosing interstitial lung diseases. <i>Respiratory Research</i> , 2021 , 22, 84	7.3	11
45	Translation and validation of the KingS Brief Interstitial Lung Disease (K-BILD) questionnaire in French, Italian, Swedish, and Dutch. <i>Chronic Respiratory Disease</i> , 2017 , 14, 140-150	3	10

(2019-2020)

44	Analysis of body mass index, weight loss and progression of idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2020 , 21, 312	7.3	10
43	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	4.1	9
42	Development and feasibility of an eHealth tool for idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2018 , 51,	13.6	9
41	First patient-centred set of outcomes for pulmonary sarcoidosis: a multicentre initiative. <i>BMJ Open Respiratory Research</i> , 2019 , 6, e000394	5.6	9
40	Validity of the Patient Experiences and Satisfaction with Medications (PESaM) Questionnaire. <i>Patient</i> , 2019 , 12, 149-162	3.7	8
39	Acetylcysteine in IPF: the knockout blow?. Lancet Respiratory Medicine, the, 2016, 4, 420-1	35.1	7
38	Design of a randomized controlled trial to evaluate effectiveness of methotrexate versus prednisone as first-line treatment for pulmonary sarcoidosis: the PREDMETH study. <i>BMC Pulmonary Medicine</i> , 2020 , 20, 271	3.5	7
37	Exhaled breath analysis by use of eNose technology: a novel diagnostic tool for interstitial lung disease. <i>European Respiratory Journal</i> , 2021 , 57,	13.6	7
36	Early diagnosis of fibrotic interstitial lung disease: challenges and opportunities. <i>Lancet Respiratory Medicine,the</i> , 2021 , 9, 1065-1076	35.1	7
35	Managing Fatigue in Patients With Interstitial Lung Disease. <i>Chest</i> , 2020 , 158, 2026-2033	5.3	6
34	The impact of the new Global Lung Function Initiative reference values on trial inclusion for patients with idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2019 , 53,	13.6	5
33	Progress in the treatment of pulmonary fibrosis. Lancet Respiratory Medicine, the, 2020, 8, 424-425	35.1	5
32	Diurnal variation in forced vital capacity in patients with fibrotic interstitial lung disease using home spirometry. <i>ERJ Open Research</i> , 2020 , 6,	3.5	5
31	Patient expectations, experiences and satisfaction with nintedanib and pirfenidone in idiopathic pulmonary fibrosis: a quantitative study. <i>Respiratory Research</i> , 2020 , 21, 196	7.3	5
30	Educational aspects of rare and orphan lung diseases. Respiratory Research, 2021, 22, 92	7.3	5
29	European idiopathic pulmonary fibrosis Patient Charter: a missed opportunity. <i>European Respiratory Journal</i> , 2016 , 48, 283-4	13.6	4
28	Results of the standard set for pulmonary sarcoidosis: feasibility and multicentre outcomes. <i>ERJ Open Research</i> , 2019 , 5,	3.5	4
27	Serious adverse events in patients with idiopathic pulmonary fibrosis in the placebo arms of 6 clinical trials. <i>Respiratory Medicine</i> , 2019 , 150, 120-125	4.6	2

26	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	1.1	2
25	Evaluation of a Home Monitoring Application for Follow Up after Lung Transplantation-A Pilot Study. <i>Journal of Personalized Medicine</i> , 2020 , 10,	3.6	2
24	Gender equity in interstitial lung disease. Lancet Respiratory Medicine, the, 2020, 8, 842-843	35.1	2
23	Patient-reported outcomes and patient-reported outcome measures in interstitial lung disease: where to go from here?. <i>European Respiratory Review</i> , 2021 , 30,	9.8	2
22	Management of Idiopathic Pulmonary Fibrosis. Clinics in Chest Medicine, 2021, 42, 275-285	5.3	2
21	Research highlights from the 2018 ERS International Congress: interstitial lung diseases. <i>ERJ Open Research</i> , 2019 , 5,	3.5	2
20	Cough, an unresolved problem in interstitial lung diseases. <i>Current Opinion in Supportive and Palliative Care</i> , 2019 , 13, 143-151	2.6	2
19	Home spirometry in patients with idiopathic pulmonary fibrosis: data from the INMARK trial. <i>European Respiratory Journal</i> , 2021 , 58,	13.6	2
18	The six-minute walk test in sarcoidosis associated pulmonary hypertension: Results from an international registry <i>Respiratory Medicine</i> , 2022 , 196, 106801	4.6	2
17	Antacid Medication and Antireflux Surgery in Patients with Idiopathic Pulmonary Fibrosis: A Systematic Review and Meta-Analysis <i>Annals of the American Thoracic Society</i> , 2022 , 19, 833-844	4.7	2
16	Modelling Forced Vital Capacity in Idiopathic Pulmonary Fibrosis: Optimising Trial Design. <i>Advances in Therapy</i> , 2019 , 36, 3059-3070	4.1	1
15	COVID-19 infection in patients with sarcoidosis: susceptibility and clinical outcomes. <i>Current Opinion in Pulmonary Medicine</i> , 2021 , 27, 463-471	3	1
14	Feasibility of online home spirometry in systemic sclerosis-associated interstitial lung disease: a pilot study. <i>Rheumatology</i> , 2021 , 60, 2467-2471	3.9	1
13	Patient reported side-effects of prednisone and methotrexate in a real-world sarcoidosis population. <i>Chronic Respiratory Disease</i> , 2021 , 18, 14799731211031935	3	1
12	The Value of the Surprise Question to Predict One-Year Mortality in Idiopathic Pulmonary Fibrosis: A Prospective Cohort Study. <i>Respiration</i> , 2021 , 100, 780-785	3.7	1
11	Patient-centered Outcomes Research in Interstitial Lung Disease: An Official American Thoracic Society Research Statement. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021 , 204, e3-e2	1 ^{0.2}	1
10	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	4.1	O
9	The use of online visual analogue scales in idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2021 ,	13.6	O

LIST OF PUBLICATIONS

8	Patient Reported Experiences and Delays During the Diagnostic Pathway for Pulmonary Fibrosis: A Multinational European Survey. <i>Frontiers in Medicine</i> , 2021 , 8, 711194	4.9	O	
7	PAciFy Cough-a multicentre, double-blind, placebo-controlled, crossover trial of morphine sulphate for the treatment of pulmonary Fibrosis Cough <i>Trials</i> , 2022 , 23, 184	2.8	O	
6	Trafficking and lysosomal storage disorders 2019 , 319-332			
5	Comprehensive Care of Interstitial Lung Disease 2022 , 64-78			
4	Palliative Care in Interstitial Lung Disease. Respiratory Medicine, 2021, 189-207	0.2		
3	Challenges in the classification of fibrotic ILD: Patient case 2. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2015 , 32 Suppl 1, 13-6	1.1		
2	Non-organ-specific manifestations of sarcoidosis 2022 , 206-222			
1	Comparison of the treatment guidelines for sarcoidosis: common sense in the search for evidence <i>European Respiratory Journal</i> , 2021 ,	13.6		