

# Mary E Streck

## List of Publications by Year in descending order

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Version: 2024-02-01

65  
papers

5,005  
citations

147566

31  
h-index

118652

62  
g-index

68  
all docs

68  
docs citations

68  
times ranked

4122  
citing authors

#	ARTICLE	IF	CITATIONS
1	Detection and Early Referral of Patients With Interstitial Lung Abnormalities. <i>Chest</i> , 2022, 161, 470-482.	0.4	26
2	Hypersensitivity pneumonitis: Current concepts in pathogenesis, diagnosis, and treatment. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2022, 77, 442-453.	2.7	28
3	Lung function trajectory in progressive fibrosing interstitial lung disease. <i>European Respiratory Journal</i> , 2022, 59, 2101396.	3.1	40
4	Interstitial Lung Disease in Firefighters: An Emerging Occupational Hazard. <i>Frontiers in Medicine</i> , 2022, 9, 864658.	1.2	6
5	Occupational Exposures in Rheumatoid Arthritis-Related Airway Disease: A Missing Link?. <i>Annals of the American Thoracic Society</i> , 2022, , .	1.5	0
6	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.	2.5	780
7	Inhalational exposures in patients with fibrotic interstitial lung disease: Presentation, pulmonary function and survival in the <scp>Canadian Registry</scp> for <scp>Pulmonary Fibrosis</scp>. <i>Respirology</i> , 2022, 27, 635-644.	1.3	12
8	COVIDOSE: A Phase II Clinical Trial of Low-Dose Tocilizumab in the Treatment of Noncritical COVID-19 Pneumonia. <i>Clinical Pharmacology and Therapeutics</i> , 2021, 109, 688-696.	2.3	42
9	Leukocyte telomere length and mycophenolate therapy in chronic hypersensitivity pneumonitis. <i>European Respiratory Journal</i> , 2021, 57, 2002872.	3.1	32
10	Differentiation of Idiopathic Pulmonary Fibrosis from Connective Tissue Disease-Related Interstitial Lung Disease Using Quantitative Imaging. <i>Journal of Clinical Medicine</i> , 2021, 10, 2663.	1.0	11
11	Characteristics and Prevalence of Domestic and Occupational Inhalational Exposures Across Interstitial Lung Diseases. <i>Chest</i> , 2021, 160, 209-218.	0.4	18
12	Association of antinuclear antibody seropositivity with inhaled environmental exposures in patients with interstitial lung disease. <i>ERJ Open Research</i> , 2021, 7, 00254-2021.	1.1	0
13	Hospitalizations in patients with idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2021, 22, 257.	1.4	8
14	Rituximab for interstitial pneumonia with autoimmune features at two medical centres. <i>Rheumatology Advances in Practice</i> , 2021, 5, ii1-ii9.	0.3	7
15	Recent Advances in Severe Asthma. <i>Chest</i> , 2020, 157, 516-528.	0.4	96
16	Genome-Wide Association Study of Susceptibility to Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 201, 564-574.	2.5	208
17	Myositis-specific antibodies identify a distinct interstitial pneumonia with autoimmune features phenotype. <i>European Respiratory Journal</i> , 2020, 56, 2001205.	3.1	24
18	Telomere biology disorder prevalence and phenotypes in adults with familial hematologic and/or pulmonary presentations. <i>Blood Advances</i> , 2020, 4, 4873-4886.	2.5	23

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19	Automated CT Analysis of Major Forms of Interstitial Lung Disease. <i>Journal of Clinical Medicine</i> , 2020, 9, 3776.	1.0	14
20	Gender equity in interstitial lung disease. <i>Lancet Respiratory Medicine</i> , 2020, 8, 842-843.	5.2	6
21	Recognition and Management of Myositis-Associated Rapidly Progressive Interstitial Lung Disease. <i>Chest</i> , 2020, 158, 252-263.	0.4	40
22	Gender in idiopathic pulmonary fibrosis diagnosis: time to address unconscious bias. <i>Thorax</i> , 2020, 75, 365-366.	2.7	4
23	Circulating Plasma Biomarkers of Survival in Antifibrotic-Treated Patients With Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2020, 158, 1526-1534.	0.4	31
24	CPAP Adherence, Mortality, and Progression-Free Survival in Interstitial Lung Disease and OSA. <i>Chest</i> , 2020, 158, 1701-1712.	0.4	19
25	Prevalence and Clinical Significance of Antineutrophil Cytoplasmic Antibodies in North American Patients With Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2019, 156, 715-723.	0.4	45
26	Diagnostic test interpretation and referral delay in patients with interstitial lung disease. <i>Respiratory Research</i> , 2019, 20, 253.	1.4	35
27	Antifibrotic therapy for idiopathic pulmonary fibrosis: time to treat. <i>Respiratory Research</i> , 2019, 20, 205.	1.4	166
28	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.	2.5	90
29	T cell Co-Stimulatory molecules ICOS and CD28 stratify idiopathic pulmonary fibrosis survival. <i>Respiratory Medicine: X</i> , 2019, 1, 100002.	1.4	7
30	Anti-RNA binding protein positivity in idiopathic interstitial pneumonia. <i>Respiratory Medicine</i> , 2019, 146, 23-27.	1.3	1
31	Telomere length and genetic variant associations with interstitial lung disease progression and survival. <i>European Respiratory Journal</i> , 2019, 53, 1801641.	3.1	119
32	African-American race and mortality in interstitial lung disease: a multicentre propensity-matched analysis. <i>European Respiratory Journal</i> , 2018, 51, 1800255.	3.1	37
33	CT-Pathologic Correlation of Major Types of Pulmonary Fibrosis: Insights for Revisions to Current Guidelines. <i>American Journal of Roentgenology</i> , 2018, 210, 1034-1041.	1.0	26
34	Phenotypic Clusters Predict Outcomes in a Longitudinal Interstitial Lung Disease Cohort. <i>Chest</i> , 2018, 153, 349-360.	0.4	40
35	Asthma Outcomes and Management During Pregnancy. <i>Chest</i> , 2018, 153, 515-527.	0.4	91
36	N-acetylcysteine exposure is associated with improved survival in anti-nuclear antibody seropositive patients with usual interstitial pneumonia. <i>BMC Pulmonary Medicine</i> , 2018, 18, 30.	0.8	5

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37	Systemic sclerosis-associated interstitial lung disease: Role of the oesophagus in outcomes. <i>Respirology</i> , 2018, 23, 885-886.	1.3	3
38	Allergic and Noninvasive Infectious Pulmonary Aspergillosis Syndromes. <i>Clinics in Chest Medicine</i> , 2017, 38, 521-534.	0.8	33
39	CT Findings, Radiologic-Pathologic Correlation, and Imaging Predictors of Survival for Patients With Interstitial Pneumonia With Autoimmune Features. <i>American Journal of Roentgenology</i> , 2017, 208, 1229-1236.	1.0	47
40	De Novo Development of Bronchiectasis in Patients With Hematologic Malignancy. <i>Chest</i> , 2017, 152, 683-685.	0.4	12
41	Outcomes of immunosuppressive therapy in chronic hypersensitivity pneumonitis. <i>ERJ Open Research</i> , 2017, 3, 00016-2017.	1.1	84
42	Myositis-associated Interstitial Lung Disease: Predictors of Failure of Conventional Treatment and Response to Tacrolimus in a US Cohort. <i>Journal of Rheumatology</i> , 2017, 44, 1612-1618.	1.0	40
43	Interstitial pneumonia with autoimmune features: Clinical, radiologic, and histological characteristics and outcome in a series of 57 patients. <i>Respiratory Medicine</i> , 2017, 123, 56-62.	1.3	119
44	Comprehensive Care of the Lung Transplant Patient. <i>Chest</i> , 2017, 152, 150-164.	0.4	74
45	Autoimmune Hypothyroidism As a Predictor of Mortality in Chronic Hypersensitivity Pneumonitis. <i>Frontiers in Medicine</i> , 2017, 4, 170.	1.2	10
46	Skewed Lung CCR4 to CCR6 CD4+ T Cell Ratio in Idiopathic Pulmonary Fibrosis Is Associated with Pulmonary Function. <i>Frontiers in Immunology</i> , 2016, 7, 516.	2.2	29
47	A call for uniformity in implementing the IPAF (interstitial pneumonia with autoimmune features) criteria. <i>European Respiratory Journal</i> , 2016, 48, 1813-1814.	3.1	8
48	The Diagnosis and Treatment of Antisynthetase Syndrome. <i>Clinical Pulmonary Medicine</i> , 2016, 23, 218-226.	0.3	151
49	Predictors of survival in coexistent hypersensitivity pneumonitis with autoimmune features. <i>Respiratory Medicine</i> , 2016, 114, 53-60.	1.3	42
50	Characterisation of patients with interstitial pneumonia with autoimmune features. <i>European Respiratory Journal</i> , 2016, 47, 1767-1775.	3.1	219
51	Therapeutic Approach to Adult Fibrotic Lung Diseases. <i>Chest</i> , 2016, 150, 1371-1386.	0.4	39
52	Azathioprine response in patients with fibrotic connective tissue disease-associated interstitial lung disease. <i>Respiratory Medicine</i> , 2016, 121, 117-122.	1.3	85
53	Response. <i>Chest</i> , 2016, 149, 600-601.	0.4	0
54	Treatment of Idiopathic Pulmonary Fibrosis. <i>Annals of the American Thoracic Society</i> , 2016, 13, 115-117.	1.5	19

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55	Thyroid Disease Is Prevalent and Predicts Survival in Patients With Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2015, 148, 692-700.	0.4	50
56	A functional genomic model for predicting prognosis in idiopathic pulmonary fibrosis. <i>BMC Pulmonary Medicine</i> , 2015, 15, 147.	0.8	42
57	An official European Respiratory Society/American Thoracic Society research statement: interstitial pneumonia with autoimmune features. <i>European Respiratory Journal</i> , 2015, 46, 976-987.	3.1	803
58	<i>TOLLIP</i> , <i>MUC5B</i> , and the Response to <i>N</i> -Acetylcysteine among Individuals with Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2015, 192, 1475-1482.	2.5	257
59	Diagnosis and Treatment of Connective Tissue Disease-Associated Interstitial Lung Disease. <i>Chest</i> , 2013, 143, 814-824.	0.4	257
60	Barriers to Optimal Palliative Care of Lung Transplant Candidates. <i>Chest</i> , 2013, 143, 736-743.	0.4	60
61	Bronchiectasis in a Diverse US Population. <i>Chest</i> , 2012, 142, 159-167.	0.4	88
62	Improving Asthma Management. <i>Chest</i> , 2012, 142, 1085-1087.	0.4	3
63	Autoimmune-Featured Interstitial Lung Disease. <i>Chest</i> , 2011, 140, 1292-1299.	0.4	236
64	Difficult Asthma. <i>Proceedings of the American Thoracic Society</i> , 2006, 3, 116-123.	3.5	30
65	Consensus guidelines for asthma therapy. <i>Annals of Allergy, Asthma and Immunology</i> , 2001, 86, 40-44.	0.5	18