Shane Shapera

List of Publications by Year in descending order

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623734 610901 34 645 14 24 citations g-index h-index papers 34 34 34 805 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Association of BMI and Change in Weight With Mortality in Patients With Fibrotic Interstitial Lung Disease. Chest, 2022, 161, 1320-1329.	0.8	25
2	Effect of continued antifibrotic therapy after forced vital capacity decline in patients with idiopathic pulmonary fibrosis; a real world multicenter cohort study. Respiratory Medicine, 2022, 191, 106722.	2.9	3
3	Prevalence and characteristics of progressive fibrosing interstitial lung disease in a prospective registry. European Respiratory Journal, 2022, 60, 2102571.	6.7	57
4	Correlation of respiratory oscillometry with CT image analysis in a prospective cohort of idiopathic pulmonary fibrosis. BMJ Open Respiratory Research, 2022, 9, e001163.	3.0	7
5	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> . Respirology, 2022, 27, 635-644.	2.3	12
6	Minimum important difference of the EQ-5D-5L and EQ-VAS in fibrotic interstitial lung disease. Thorax, 2021, 76, 37-43.	5.6	28
7	Validation and minimum important difference of the UCSD Shortness of Breath Questionnaire in fibrotic interstitial lung disease. Respiratory Research, 2021, 22, 202.	3.6	5
8	Feasibility and Outcomes of a Standardized Management Protocol for Acute Exacerbation of Interstitial Lung Disease. Lung, 2021, 199, 379-387.	3.3	4
9	Utility of anti-neutrophil cytoplasmic antibody screening in idiopathic interstitial lung disease. Sarcoidosis Vasculitis and Diffuse Lung Diseases, 2021, 38, e2021015.	0.2	0
10	Real-world patterns of pirfenidone use and safety in patients with idiopathic pulmonary fibrosis in Canada: Data from INSPIRATION PLUS. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2020, 4, 25-30.	0.5	0
11	Development of the Functional Assessment of Cancer Therapy–Immune Checkpoint Modulator (FACTâ€ŀCM): A toxicity subscale to measure quality of life in patients with cancer who are treated with ICMs. Cancer, 2020, 126, 1550-1558.	4.1	26
12	Idiopathic Pulmonary Fibrosis: A Review of Disease, Pharmacological, and Nonpharmacological Strategies With a Focus on Symptoms, Function, and Health-Related Quality of Life. Journal of Pain and Symptom Management, 2020, 59, 1362-1378.	1.2	20
13	EFFICACY AND SAFETY OF NINTEDANIB IN US/CANADIAN PATIENTS WITH PROGRESSIVE FIBROSING INTERSTITIAL LUNG DISEASES: FURTHER ANALYSES OF THE INBUILD TRIAL. Chest, 2020, 158, A2613-A2615.	0.8	0
14	Long-term monitoring of patients with fibrotic interstitial lung disease: A Canadian Thoracic Society Position Statement. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2020, 4, 147-155.	0.5	3
15	Optimizing care for patients with interstitial lung disease during the COVID-19 pandemic. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2020, 4, 226-228.	0.5	2
16	Management of idiopathic pulmonary fibrosis: A survey of Canadian respirologists. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2020, 4, 165-173.	0.5	1
17	A cluster-based analysis evaluating the impact of comorbidities in fibrotic interstitial lung disease. Respiratory Research, 2020, 21, 322.	3.6	18
18	Costs of Workplace Productivity Loss in Patients with Connective Tissue Disease–associated Interstitial Lung Disease. Annals of the American Thoracic Society, 2020, 17, 1077-1084.	3.2	5

#	Article	IF	CITATIONS
19	Practical Considerations for the Diagnosis and Treatment of Fibrotic Interstitial Lung Disease During the Coronavirus Disease 2019 Pandemic. Chest, 2020, 158, 1069-1078.	0.8	57
20	Costs of Workplace Productivity Loss in Patients With Fibrotic Interstitial Lung Disease. Chest, 2019, 156, 887-895.	0.8	14
21	Screening for Myositis Antibodies in Idiopathic Interstitial Lung Disease. Lung, 2019, 197, 277-284.	3.3	17
22	Procedure volume and mortality after surgical lung biopsy in interstitial lungÂdisease. European Respiratory Journal, 2019, 53, 1801164.	6.7	54
23	Diagnostic criteria for idiopathic pulmonary fibrosis. Lancet Respiratory Medicine, the, 2018, 6, e6.	10.7	1
24	Treatment of Gastroesophageal Reflux in Patients With Idiopathic Pulmonary Fibrosis. Chest, 2018, 153, 1405-1415.	0.8	44
25	Impact of Pretreatment Interstitial Lung Disease on Radiation Pneumonitis and Survival in Patients Treated With Lung Stereotactic Body Radiation Therapy (SBRT). Clinical Lung Cancer, 2018, 19, e219-e226.	2.6	57
26	Comprehensive management of fibrotic interstitial lung diseases: A Canadian Thoracic Society position statement. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2018, 2, 234-243.	0.5	8
27	Transbronchial lung cryobiopsy for ILD: Ready or not, here it comes?. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2018, 2, 257-258.	0.5	0
28	Multi-dimensional scores to predict mortality in patients with idiopathic pulmonary fibrosis undergoing lung transplantation assessment. Respiratory Medicine, 2017, 125, 65-71.	2.9	13
29	Evaluation of patients with fibrotic interstitial lung disease: A Canadian Thoracic Society position statement. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2017, 1, 133-141.	0.5	15
30	The Canadian Registry for Pulmonary Fibrosis: Design and Rationale of a National Pulmonary Fibrosis Registry. Canadian Respiratory Journal, 2016, 2016, 1-7.	1.6	45
31	Accuracy and Reliability of Internet Resources for Information on Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2016, 194, 218-225.	5.6	98
32	Diagnostic Disparity of Previous and Revised American Thoracic Society Guidelines for Idiopathic Pulmonary Fibrosis. Canadian Respiratory Journal, 2015, 22, 86-90.	1.6	3
33	Acute Exacerbation of Idiopathic Pulmonary Fibrosis. Canadian Respiratory Journal, 2015, 22, 308-308.	1.6	1
34	What to Do with All of These Lung Nodules?. Canadian Respiratory Journal, 2014, 21, e52-e54.	1.6	2