## Chad A Newton

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/122985/publications.pdf

Version: 2024-02-01

26 papers 1,394 citations

16 h-index 24 g-index

26 all docs

26 docs citations

times ranked

26

1550 citing authors

#	Article	IF	Citations
1	Telomere-related lung fibrosis is diagnostically heterogeneous but uniformly progressive. European Respiratory Journal, 2016, 48, 1710-1720.	6.7	281
2	The MUC5B promoter polymorphism and telomere length in patients with chronic hypersensitivity pneumonitis: an observational cohort-control study. Lancet Respiratory Medicine, the, 2017, 5, 639-647.	10.7	206
3	Whole-Exome Sequencing in Adults With Chronic Kidney Disease. Annals of Internal Medicine, 2018, 168, 100.	3.9	154
4	Telomere length and genetic variant associations with interstitial lung disease progression and survival. European Respiratory Journal, 2019, 53, 1801641.	6.7	119
5	Telomere Length and Use of Immunosuppressive Medications in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 336-347.	5.6	99
6	Telomere length in patients with pulmonary fibrosis associated with chronic lung allograft dysfunction and post–lung transplantation survival. Journal of Heart and Lung Transplantation, 2017, 36, 845-853.	0.6	93
7	Pulmonary fibrosis in the era of stratified medicine. Thorax, 2016, 71, 1154-1160.	5.6	67
8	Somatic mutations in telomerase promoter counterbalance germline loss-of-function mutations. Journal of Clinical Investigation, 2017, 127, 982-986.	8.2	60
9	Lung function trajectory in progressive fibrosing interstitial lung disease. European Respiratory Journal, 2022, 59, 2101396.	6.7	40
10	Utility of Bronchoalveolar Lavage and Transbronchial Biopsy in Patients with Hypersensitivity Pneumonitis. Lung, 2018, 196, 617-622.	3.3	36
11	Leukocyte telomere length and mycophenolate therapy in chronic hypersensitivity pneumonitis. European Respiratory Journal, 2021, 57, 2002872.	6.7	32
12	Proteomic biomarkers of progressive fibrosing interstitial lung disease: a multicentre cohort analysis. Lancet Respiratory Medicine, the, 2022, 10, 593-602.	10.7	31
13	Family History of Pulmonary Fibrosis Predicts Worse Survival in Patients With Interstitial Lung Disease. Chest, 2021, 159, 1913-1921.	0.8	28
14	Rare and Common Variants in <i>KIF15</i> Contribute to Genetic Risk of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 56-69.	5.6	25
15	Myositis-specific antibodies identify a distinct interstitial pneumonia with autoimmune features phenotype. European Respiratory Journal, 2020, 56, 2001205.	6.7	24
16	Familial Pulmonary Fibrosis. Chest, 2021, 160, 1764-1773.	0.8	21
17	Clinical Genetics in Interstitial Lung Disease. Frontiers in Medicine, 2018, 5, 116.	2.6	19
18	The Role of Genetic Testing in Pulmonary Fibrosis. Chest, 2022, 162, 394-405.	0.8	19

#	Article	IF	CITATION
19	Role of Antigen Type in Survival in Chronic Hypersensitivity Pneumonitis. Lung, 2019, 197, 113-114.	3.3	12
20	Pleuroparenchymal fibroelastosis associated with telomerase reverse transcriptase mutations. European Respiratory Journal, 2017, 49, 1700696.	6.7	8
21	Utility of Bronchoalveolar Lavage and Transbronchial Biopsy in Patients with Interstitial Lung Disease. Lung, 2020, 198, 803-810.	3.3	6
22	Pirfenidone for Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2016, 194, 374-376.	5.6	5
23	Molecular Markers and the Promise of Precision Medicine for Interstitial Lung Disease. Clinics in Chest Medicine, 2021, 42, 357-364.	2.1	4
24	Interstitial Pneumonia with Autoimmune Features: What the Rheumatologist Needs to Know. Current Rheumatology Reports, 0, , .	4.7	3
25	Addition of antifibrotic therapy to immunosuppression in hypersensitivity pneumonitis: A case series. Respiratory Medicine Case Reports, 2021, 34, 101562.	0.4	2
26	Pulmonary fibrosis screening: quantifying the psychological impact. Thorax, 2021, 76, 532-533.	5.6	0