

Ayodeji Adegunsoye

List of Publications by Year in descending order

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Version: 2024-02-01

68
papers

2,218
citations

201575

27
h-index

243529

44
g-index

71
all docs

71
docs citations

71
times ranked

2248
citing authors

#	ARTICLE	IF	CITATIONS
1	Characterisation of patients with interstitial pneumonia with autoimmune features. <i>European Respiratory Journal</i> , 2016, 47, 1767-1775.	3.1	219
2	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
3	Telomere length and genetic variant associations with interstitial lung disease progression and survival. <i>European Respiratory Journal</i> , 2019, 53, 1801641.	3.1	119
4	CT Features of the Usual Interstitial Pneumonia Pattern: Differentiating Connective Tissue Disease-Associated Interstitial Lung Disease From Idiopathic Pulmonary Fibrosis. <i>American Journal of Roentgenology</i> , 2018, 210, 307-313.	1.0	98
5	Computed Tomography Honeycombing Identifies a Progressive Fibrotic Phenotype with Increased Mortality across Diverse Interstitial Lung Diseases. <i>Annals of the American Thoracic Society</i> , 2019, 16, 580-588.	1.5	93
6	Azathioprine response in patients with fibrotic connective tissue disease-associated interstitial lung disease. <i>Respiratory Medicine</i> , 2016, 121, 117-122.	1.3	85
7	Outcomes of immunosuppressive therapy in chronic hypersensitivity pneumonitis. <i>ERJ Open Research</i> , 2017, 3, 00016-2017.	1.1	84
8	Rare Protein-Altering Telomere-related Gene Variants in Patients with Chronic Hypersensitivity Pneumonitis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 1154-1163.	2.5	81
9	Comprehensive Care of the Lung Transplant Patient. <i>Chest</i> , 2017, 152, 150-164.	0.4	74
10	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
11	Treatment of fibrotic interstitial lung disease: current approaches and future directions. <i>Lancet</i> , The, 2021, 398, 1450-1460.	6.3	47
12	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
13	Association of Black Race with Outcomes in COVID-19 Disease: A Retrospective Cohort Study. <i>Annals of the American Thoracic Society</i> , 2020, 17, 1336-1339.	1.5	44
14	Predictors of survival in coexistent hypersensitivity pneumonitis with autoimmune features. <i>Respiratory Medicine</i> , 2016, 114, 53-60.	1.3	42
15	CT findings associated with survival in chronic hypersensitivity pneumonitis. <i>European Radiology</i> , 2017, 27, 5127-5135.	2.3	41
16	Phenotypic Clusters Predict Outcomes in a Longitudinal Interstitial Lung Disease Cohort. <i>Chest</i> , 2018, 153, 349-360.	0.4	40
17	Lung function trajectory in progressive fibrosing interstitial lung disease. <i>European Respiratory Journal</i> , 2022, 59, 2101396.	3.1	40
18	Therapeutic Approach to Adult Fibrotic Lung Diseases. <i>Chest</i> , 2016, 150, 1371-1386.	0.4	39

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19	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
20	Integrating Genomics Into Management of Fibrotic Interstitial Lung Disease. <i>Chest</i> , 2019, 155, 1026-1040.	0.4	37
21	Prognosticating Outcomes in Interstitial Lung Disease by Mediastinal Lymph Node Assessment. An Observational Cohort Study with Independent Validation. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 199, 747-759.	2.5	36
22	Diagnostic test interpretation and referral delay in patients with interstitial lung disease. <i>Respiratory Research</i> , 2019, 20, 253.	1.4	35
23	Imaging of Hypersensitivity Pneumonitis. <i>Radiologic Clinics of North America</i> , 2016, 54, 1033-1046.	0.9	34
24	Leukocyte telomere length and mycophenolate therapy in chronic hypersensitivity pneumonitis. <i>European Respiratory Journal</i> , 2021, 57, 2002872.	3.1	32
25	Circulating Plasma Biomarkers of Survival in Antifibrotic-Treated Patients With Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2020, 158, 1526-1534.	0.4	31
26	Underreporting of Interstitial Lung Abnormalities on Lung Cancer Screening Computed Tomography. <i>Annals of the American Thoracic Society</i> , 2018, 15, 764-766.	1.5	30
27	Circulating Plasma Biomarkers of Progressive Interstitial Lung Disease. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 201, 250-253.	2.5	30
28	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
29	Interstitial Pneumonia With Autoimmune Features: Value of Histopathology. <i>Archives of Pathology and Laboratory Medicine</i> , 2017, 141, 960-969.	1.2	29
30	Etiopathogenetic Mechanisms of Pulmonary Hypertension in Sleep-Related Breathing Disorders. <i>Pulmonary Medicine</i> , 2012, 2012, 1-10.	0.5	27
31	Inflammatory Response Mechanisms Exacerbating Hypoxemia in Coexistent Pulmonary Fibrosis and Sleep Apnea. <i>Mediators of Inflammation</i> , 2015, 2015, 1-13.	1.4	27
32	Blood Transcriptomics Predicts Progression of Pulmonary Fibrosis and Associated Natural Killer Cells. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021, 204, 197-208.	2.5	27
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	Myositis-specific antibodies identify a distinct interstitial pneumonia with autoimmune features phenotype. <i>European Respiratory Journal</i> , 2020, 56, 2001205.	3.1	24
35	ICOS protects against mortality from acute lung injury through activation of IL-5+ ILC2s. <i>Mucosal Immunology</i> , 2018, 11, 61-70.	2.7	23
36	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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37	CPAP Adherence, Mortality, and Progression-Free Survival in Interstitial Lung Disease and OSA. <i>Chest</i> , 2020, 158, 1701-1712.	0.4	19
38	Integration and Application of Clinical Practice Guidelines for the Diagnosis of Idiopathic Pulmonary Fibrosis and Fibrotic Hypersensitivity Pneumonitis. <i>Chest</i> , 2022, 162, 614-629.	0.4	19
39	Characteristics and Prevalence of Domestic and Occupational Inhalational Exposures Across Interstitial Lung Diseases. <i>Chest</i> , 2021, 160, 209-218.	0.4	18
40	Echocardiographic Evaluation of Calcific Aortic Stenosis in the Older Adult. <i>Echocardiography</i> , 2011, 28, 117-129.	0.3	16
41	Automated CT Analysis of Major Forms of Interstitial Lung Disease. <i>Journal of Clinical Medicine</i> , 2020, 9, 3776.	1.0	14
42	De Novo Development of Bronchiectasis in Patients With Hematologic Malignancy. <i>Chest</i> , 2017, 152, 683-685.	0.4	12
43	Inhalational exposures in patients with fibrotic interstitial lung disease: Presentation, pulmonary function and survival in the <sc>Canadian Registry</sc> for <sc>Pulmonary Fibrosis</sc>. <i>Respirology</i> , 2022, 27, 635-644.	1.3	12
44	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
45	Autoimmune Hypothyroidism As a Predictor of Mortality in Chronic Hypersensitivity Pneumonitis. <i>Frontiers in Medicine</i> , 2017, 4, 170.	1.2	10
46	Vessel-related structures predict UIP pathology in those with a non-IPF pattern on CT. <i>European Radiology</i> , 2021, 31, 7295-7302.	2.3	10
47	A series of <sc>COVID</sc>-19 autopsies with clinical and pathologic comparisons to both seasonal and pandemic influenza. <i>Journal of Pathology: Clinical Research</i> , 2021, 7, 459-470.	1.3	9
48	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
49	Hospitalizations in patients with idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2021, 22, 257.	1.4	8
50	High-titer rheumatoid factor seropositivity predicts mediastinal lymphadenopathy and mortality in rheumatoid arthritis-related interstitial lung disease. <i>Scientific Reports</i> , 2021, 11, 22821.	1.6	8
51	Rituximab for interstitial pneumonia with autoimmune features at two medical centres. <i>Rheumatology Advances in Practice</i> , 2021, 5, ii1-ii9.	0.3	7
52	A 20-year-old woman with rapidly progressive dyspnea and diffuse pulmonary infiltrates. <i>Respiratory Medicine Case Reports</i> , 2012, 5, 34-36.	0.2	6
53	The role of echocardiography in the evaluation and management of aortic stenosis in the older adult. <i>International Journal of Cardiology</i> , 2012, 155, 39-48.	0.8	6
54	Diagnostic Classification of Interstitial Lung Disease in Clinical Practice. <i>Clinics in Chest Medicine</i> , 2021, 42, 251-261.	0.8	6

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55	Interstitial Lung Disease in Firefighters: An Emerging Occupational Hazard. <i>Frontiers in Medicine</i> , 2022, 9, 864658.	1.2	6
56	Novel Therapeutic Strategies for Reducing Right Heart Failure Associated Mortality in Fibrotic Lung Diseases. <i>BioMed Research International</i> , 2015, 2015, 1-16.	0.9	5
57	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
58	Racial Disparities in Pulmonary Fibrosis and the Impact on the Black Population. <i>Archivos De Bronconeumologia</i> , 2022, 58, 590-592.	0.4	5
59	The Progression to Interstitial Lung Disease. <i>Chest</i> , 2021, 160, 400-402.	0.4	4
60	Inhaled Nitric Oxide for Fibrotic Interstitial Lung Disease: A Step Forward. <i>Annals of the American Thoracic Society</i> , 2022, 19, 536-538.	1.5	4
61	Diagnostic Delay in Idiopathic Pulmonary Fibrosis: Where the Rubber Meets the Road. <i>Annals of the American Thoracic Society</i> , 2019, 16, 310-312.	1.5	2
62	MUC5B promoter variant: genomic fingerprint for early identification of undiagnosed pulmonary fibrosis. <i>Thorax</i> , 2019, 74, 1111-1112.	2.7	2
63	Anti-RNA binding protein positivity in idiopathic interstitial pneumonia. <i>Respiratory Medicine</i> , 2019, 146, 23-27.	1.3	1
64	Molecular pathways in idiopathic pulmonary fibrosis pathogenesis: Transcending barriers to optimally targeted pharmacotherapies. <i>EBioMedicine</i> , 2021, 67, 103373.	2.7	1
65	Risk, Race, and Structural Racism. <i>Annals of the American Thoracic Society</i> , 2021, 18, 1289-1290.	1.5	1
66	Reply to Lescoat et al. and to Khamis et al.. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 199, 1041-1042.	2.5	0
67	Anticoagulation and Pulmonary Fibrosis. <i>Chest</i> , 2021, 159, 1321-1323.	0.4	0
68	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