

Nazia Chaudhuri

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.

61
papers

1,316
citations

19
h-index

35
g-index

99
ext. papers

1,924
ext. citations

7
avg, IF

4.71
L-index

#	Paper	IF	Citations
61	Nintedanib in patients with progressive fibrosing interstitial lung diseases-subgroup analyses by interstitial lung disease diagnosis in the INBUILD trial: a randomised, double-blind, placebo-controlled, parallel-group trial. <i>Lancet Respiratory Medicine</i> , 2020 , 8, 453-460	35.1	154
60	Drug-Induced Interstitial Lung Disease: A Systematic Review. <i>Journal of Clinical Medicine</i> , 2018 , 7,	5.1	117
59	Idiopathic Pulmonary Fibrosis (IPF): An Overview. <i>Journal of Clinical Medicine</i> , 2018 , 7,	5.1	111
58	Real World Experiences: Pirfenidone and Nintedanib are Effective and Well Tolerated Treatments for Idiopathic Pulmonary Fibrosis. <i>Journal of Clinical Medicine</i> , 2016 , 5,	5.1	102
57	Outcome of Hospitalization for COVID-19 in Patients with Interstitial Lung Disease. An International Multicenter Study. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020 , 202, 1656-1665	10.2	72
56	The therapy of idiopathic pulmonary fibrosis: what is next?. <i>European Respiratory Review</i> , 2019 , 28,	9.8	68
55	Association of Cardiovascular Disease With Respiratory Disease. <i>Journal of the American College of Cardiology</i> , 2019 , 73, 2166-2177	15.1	60
54	Toll-like receptors and chronic lung disease. <i>Clinical Science</i> , 2005 , 109, 125-33	6.5	59
53	Real world experiences: pirfenidone is well tolerated in patients with idiopathic pulmonary fibrosis. <i>Respiratory Medicine</i> , 2014 , 108, 224-6	4.6	52
52	Reducing the toll of inflammatory lung disease. <i>Chest</i> , 2007 , 131, 1550-6	5.3	39
51	Persistent self-reported changes in hearing and tinnitus in post-hospitalisation COVID-19 cases. <i>International Journal of Audiology</i> , 2020 , 59, 889-890	2.6	38
50	Summary of the British Thoracic Society guideline for diagnostic flexible bronchoscopy in adults. <i>Thorax</i> , 2013 , 68, 786-7	7.3	37
49	Biomarkers of extracellular matrix turnover in patients with idiopathic pulmonary fibrosis given nintedanib (INMARK study): a randomised, placebo-controlled study. <i>Lancet Respiratory Medicine</i> , 2019 , 7, 771-779	35.1	34
48	IPF Care: a support program for patients with idiopathic pulmonary fibrosis treated with pirfenidone in Europe. <i>Advances in Therapy</i> , 2015 , 32, 87-107	4.1	28
47	Nintedanib in Idiopathic Pulmonary Fibrosis: Practical Management Recommendations for Potential Adverse Events. <i>Respiration</i> , 2019 , 97, 173-184	3.7	27
46	Nintedanib in the management of idiopathic pulmonary fibrosis: clinical trial evidence and real-world experience. <i>Therapeutic Advances in Respiratory Disease</i> , 2018 , 12, 1753466618800618	4.9	27
45	A Review of the Multidisciplinary Diagnosis of Interstitial Lung Diseases: A Retrospective Analysis in a Single UK Specialist Centre. <i>Journal of Clinical Medicine</i> , 2016 , 5,	5.1	26

44	Early clinical experiences with nintedanib in three UK tertiary interstitial lung disease centres. <i>Clinical and Translational Medicine</i> , 2017 , 6, 41	5.7	23
43	Basic science of the innate immune system and the lung. <i>Paediatric Respiratory Reviews</i> , 2008 , 9, 236-42	4.8	22
42	Acute exacerbation of idiopathic pulmonary fibrosis: international survey and call for harmonisation. <i>European Respiratory Journal</i> , 2020 , 55,	13.6	18
41	Diesel exhaust particle exposure in vitro alters monocyte differentiation and function. <i>PLoS ONE</i> , 2012 , 7, e51107	3.7	18
40	Idiopathic pulmonary fibrosis: a holistic approach to disease management in the antifibrotic age. <i>Journal of Thoracic Disease</i> , 2017 , 9, 4700-4707	2.6	17
39	Bagpipe lung; a new type of interstitial lung disease?. <i>Thorax</i> , 2017 , 72, 380-382	7.3	14
38	Sarcoidosis in the UK: insights from British Thoracic Society registry data. <i>BMJ Open Respiratory Research</i> , 2019 , 6, e000357	5.6	13
37	Breath biomarkers in idiopathic pulmonary fibrosis: a systematic review. <i>Respiratory Research</i> , 2019 , 20, 7	7.3	13
36	Substrate for the Myocardial Inflammation-Heart Failure Hypothesis Identified Using Novel USPIO Methodology. <i>JACC: Cardiovascular Imaging</i> , 2021 , 14, 365-376	8.4	13
35	No relevant pharmacokinetic drug-drug interaction between nintedanib and pirfenidone. <i>European Respiratory Journal</i> , 2019 , 53,	13.6	12
34	Managing Idiopathic Pulmonary Fibrosis: Which Drug for Which Patient?. <i>Drugs and Aging</i> , 2017 , 34, 647-653	4.7	9
33	Treatment of fibrotic interstitial lung disease: current approaches and future directions. <i>Lancet, The</i> , 2021 , 398, 1450-1460	40	7
32	Understanding the burden of interstitial lung disease post-COVID-19: the UK Interstitial Lung Disease-Long COVID Study (UKILD-Long COVID). <i>BMJ Open Respiratory Research</i> , 2021 , 8,	5.6	6
31	Drug induced interstitial lung disease: a systematic review 2018 ,		5
30	Idiopathic pulmonary fibrosis in the UK: analysis of the British Thoracic Society electronic registry between 2013 and 2019. <i>ERJ Open Research</i> , 2021 , 7,	3.5	5
29	The burden of progressive fibrotic interstitial lung disease across the UK. <i>European Respiratory Journal</i> , 2021 , 58,	13.6	5
28	Myocardial involvement in eosinophilic granulomatosis with polyangiitis evaluated with cardiopulmonary magnetic resonance. <i>International Journal of Cardiovascular Imaging</i> , 2021 , 37, 1371-1381	2.5	5
27	Conventional oxygen therapy versus CPAP as a ceiling of care in ward-based patients with COVID-19: a multi-centre cohort evaluation. <i>EClinicalMedicine</i> , 2021 , 40, 101122	11.3	5

26	Physiological predictors of Hypoxic Challenge Testing (HCT) outcomes in Interstitial Lung Disease (ILD). <i>Respiratory Medicine</i> , 2018 , 135, 51-56	4.6	4
25	Transbronchial Lung Cryobiopsy in Idiopathic Pulmonary Fibrosis: A State of the Art Review. <i>Advances in Therapy</i> , 2019 , 36, 2193-2204	4.1	4
24	S91 A randomised, double-blind, placebo-controlled crossover study to assess the efficacy of a single dose of 100 mg of VRP700 by inhalation in reducing the frequency and severity of cough in adult patients with Idiopathic Pulmonary Fibrosis. <i>Thorax</i> , 2015 , 70, A52.1-A52	7.3	4
23	A global perspective on acute exacerbation of idiopathic pulmonary fibrosis (AE-IPF): results from an international survey 2018 ,		4
22	How consistently do physicians diagnose and manage drug-induced interstitial lung disease? Two surveys of European ILD specialist physicians. <i>ERJ Open Research</i> , 2020 , 6,	3.5	4
21	Feather bedding as a cause of hypersensitivity pneumonitis. <i>QJM - Monthly Journal of the Association of Physicians</i> , 2017 , 110, 233-234	2.7	3
20	P28 Real World MDT Diagnosis of Idiopathic Pulmonary Fibrosis. <i>Thorax</i> , 2015 , 70, A90.1-A90	7.3	3
19	Outcome of hospitalisation for COVID-19 in patients with Interstitial Lung Disease: An international multicentre study.		3
18	Pulmonary Sequelae at 4 Months After COVID-19 Infection: A Single-Centre Experience of a COVID Follow-Up Service. <i>Advances in Therapy</i> , 2021 , 38, 4505-4519	4.1	3
17	Management of Acute Exacerbation of Idiopathic Pulmonary Fibrosis in Specialised and Non-specialised ILD Centres Around the World. <i>Frontiers in Medicine</i> , 2021 , 8, 699644	4.9	3
16	Current Treatments in the Management of Idiopathic Pulmonary Fibrosis: Pirfenidone and Nintedanib. <i>Clinical Medicine Insights Therapeutics</i> , 2017 , 9, 1179559X1771912	0	2
15	Real world experience of response to pirfenidone in patients with idiopathic pulmonary fibrosis: a two centre retrospective study. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2020 , 37, 218-224	1.1	2
14	Real-World Study Analysing Progression and Survival of Patients with Idiopathic Pulmonary Fibrosis with Preserved Lung Function on Antifibrotic Treatment. <i>Advances in Therapy</i> , 2021 , 38, 268-277	4.1	2
13	Home spirometry in patients with idiopathic pulmonary fibrosis: data from the INMARK trial. <i>European Respiratory Journal</i> , 2021 , 58,	13.6	2
12	The extended utility of antifibrotic therapy in progressive fibrosing interstitial lung disease. <i>Expert Review of Respiratory Medicine</i> , 2020 , 14, 1001-1008	3.8	1
11	Rapidly non-ipf progressive fibrosing interstitial lung disease: a phenotype with an ipf-like behavior. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2020 , 37, 231-233	1.1	1
10	Nintedanib for treating idiopathic pulmonary fibrosis. <i>British Journal of Health Care Management</i> , 2016 , 22, 250-251	0.4	
9	Idiopathic pulmonary fibrosis: a clinical update. <i>British Journal of General Practice</i> , 2018 , 68, 249-250	1.6	

8	P280 Extended Clinical Experience With Pirfenidone During A Named Patient Programme For Idiopathic Pulmonary Fibrosis (ipf): Interim Results. <i>Thorax</i> , 2014 , 69, A196-A196	7.3
7	M263 A Quarter Of Ipf Patients Not Eligible For Pirfenidone Treatment Due To The Nice Criteria Significantly Decline Over Time. <i>Thorax</i> , 2014 , 69, A218-A218	7.3
6	M264 Health And Economic Impact Of Prescribing Pirfenidone. <i>Thorax</i> , 2014 , 69, A218-A219	7.3
5	P6 Early Clinical Experience With Nintedanib In two centre review. <i>Thorax</i> , 2015 , 70, A77.2-A77	7.3
4	P34 Sarcoidosis and co-existent Aspergillus lung disease. <i>Thorax</i> , 2015 , 70, A92.3-A93	7.3
3	S13 Sole use of forced vital capacity as per national institute of health and care excellence criteria disadvantage 2 in 5 people with idiopathic pulmonary fibrosis. <i>Thorax</i> , 2013 , 68, A10.1-A10	7.3
2	Pulmonary involvement in antiphospholipid syndrome 2019 , 124-139	
1	Reply to Althwaybi : Hospitalization Outcomes for COVID-19 in Patients with Interstitial Lung Disease: A Potential Role for Aerodigestive Pathophysiology?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021 , 203, 522-524	10.2