## Nazia Chaudhuri

List of Publications by Year in descending order

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279487 214527 2,477 96 23 47 citations h-index g-index papers 100 100 100 3233 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	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. Lancet Respiratory Medicine,the, 2020, 8, 453-460.	5.2	331
2	Drug-Induced Interstitial Lung Disease: A Systematic Review. Journal of Clinical Medicine, 2018, 7, 356.	1.0	215
3	Idiopathic Pulmonary Fibrosis (IPF): An Overview. Journal of Clinical Medicine, 2018, 7, 201.	1.0	215
4	Outcome of Hospitalization for COVID-19 in Patients with Interstitial Lung Disease. An International Multicenter Study. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 1656-1665.	2.5	171
5	The therapy of idiopathic pulmonary fibrosis: what is next?. European Respiratory Review, 2019, 28, 190021.	3.0	157
6	Real World Experiences: Pirfenidone and Nintedanib are Effective and Well Tolerated Treatments for Idiopathic Pulmonary Fibrosis. Journal of Clinical Medicine, 2016, 5, 78.	1.0	139
7	Association of Cardiovascular Disease With Respiratory Disease. Journal of the American College of Cardiology, 2019, 73, 2166-2177.	1.2	104
8	Persistent self-reported changes in hearing and tinnitus in post-hospitalisation COVID-19 cases. International Journal of Audiology, 2020, 59, 889-890.	0.9	73
9	Toll-like receptors and chronic lung disease. Clinical Science, 2005, 109, 125-133.	1.8	66
10	Biomarkers of extracellular matrix turnover in patients with idiopathic pulmonary fibrosis given nintedanib (INMARK study): a randomised, placebo-controlled study. Lancet Respiratory Medicine, the, 2019, 7, 771-779.	5.2	65
11	Acute exacerbation of idiopathic pulmonary fibrosis: international survey and call for harmonisation. European Respiratory Journal, 2020, 55, 1901760.	3.1	61
12	Summary of the British Thoracic Society guideline for diagnostic flexible bronchoscopy in adults. Thorax, 2013, 68, 786-787.	2.7	60
13	Real world experiences: Pirfenidone is well tolerated in patients with idiopathic pulmonary fibrosis. Respiratory Medicine, 2014, 108, 224-226.	1.3	60
14	Nintedanib in the management of idiopathic pulmonary fibrosis: clinical trial evidence and real-world experience. Therapeutic Advances in Respiratory Disease, 2018, 12, 175346661880061.	1.0	59
15	Treatment of fibrotic interstitial lung disease: current approaches and future directions. Lancet, The, 2021, 398, 1450-1460.	6.3	47
16	Reducing the Toll of Inflammatory Lung Disease. Chest, 2007, 131, 1550-1556.	0.4	46
17	IPF Care: A Support Program for Patients with Idiopathic Pulmonary Fibrosis Treated with Pirfenidone in Europe. Advances in Therapy, 2015, 32, 87-107.	1.3	44
18	A Review of the Multidisciplinary Diagnosis of Interstitial Lung Diseases: A Retrospective Analysis in a Single UK Specialist Centre. Journal of Clinical Medicine, 2016, 5, 66.	1.0	44

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19	Nintedanib in Idiopathic Pulmonary Fibrosis: Practical Management Recommendations for Potential Adverse Events. Respiration, 2019, 97, 173-184.	1.2	39
20	The burden of progressive fibrotic interstitial lung disease across the UK. European Respiratory Journal, 2021, 58, 2100221.	3.1	39
21	Early clinical experiences with nintedanib in three UK tertiary interstitial lung disease centres. Clinical and Translational Medicine, 2017, 6, 41.	1.7	32
22	Home spirometry in patients with idiopathic pulmonary fibrosis: data from the INMARK trial. European Respiratory Journal, 2021, 58, 2001518.	3.1	30
23	Understanding the burden of interstitial lung disease post-COVID-19: the UK Interstitial Lung Disease-Long COVID Study (UKILD-Long COVID). BMJ Open Respiratory Research, 2021, 8, e001049.	1.2	28
24	Breath biomarkers in idiopathic pulmonary fibrosis: a systematic review. Respiratory Research, 2019, 20, 7.	1.4	25
25	Diesel Exhaust Particle Exposure In Vitro Alters Monocyte Differentiation and Function. PLoS ONE, 2012, 7, e51107.	1.1	24
26	Basic science of the innate immune system and the lung. Paediatric Respiratory Reviews, 2008, 9, 236-242.	1.2	23
27	Pulmonary Sequelae at 4 Months After COVID-19 Infection: A Single-Centre Experience of a COVID Follow-Up Service. Advances in Therapy, 2021, 38, 4505-4519.	1.3	23
28	Idiopathic pulmonary fibrosis: a holistic approach to disease management in the antifibrotic age. Journal of Thoracic Disease, 2017, 9, 4700-4707.	0.6	22
29	No relevant pharmacokinetic drug–drug interaction between nintedanib and pirfenidone. European Respiratory Journal, 2019, 53, 1801060.	3.1	22
30	Substrate for the MyocardialÂInflammation–Heart Failure Hypothesis Identified Using NovelÂUSPIOÂMethodology. JACC: Cardiovascular Imaging, 2021, 14, 365-376.	2.3	20
31	Bagpipe lung; a new type of interstitial lung disease?. Thorax, 2017, 72, 380-382.	2.7	17
32	Idiopathic pulmonary fibrosis in the UK: analysis of the British Thoracic Society electronic registry between 2013 and 2019. ERJ Open Research, 2021, 7, 00187-2020.	1.1	17
33	Sarcoidosis in the UK: insights from British Thoracic Society registry data. BMJ Open Respiratory Research, 2019, 6, e000357.	1.2	16
34	Real-World Study Analysing Progression and Survival of Patients with Idiopathic Pulmonary Fibrosis with Preserved Lung Function on Antifibrotic Treatment. Advances in Therapy, 2021, 38, 268-277.	1.3	13
35	Conventional oxygen therapy versus CPAP as a ceiling of care in ward-based patients with COVID-19: a multi-centre cohort evaluation EClinicalMedicine, 2021, 40, 101122.	3.2	13
36	Drug induced interstitial lung disease: a systematic review. , 2018, , .		12

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37	Managing Idiopathic Pulmonary Fibrosis: Which Drug for Which Patient?. Drugs and Aging, 2017, 34, 647-653.	1.3	10
38	Myocardial involvement in eosinophilic granulomatosis with polyangiitis evaluated with cardiopulmonary magnetic resonance. International Journal of Cardiovascular Imaging, 2021, 37, 1371-1381.	0.7	10
39	Management of Acute Exacerbation of Idiopathic Pulmonary Fibrosis in Specialised and Non-specialised ILD Centres Around the World. Frontiers in Medicine, 2021, 8, 699644.	1.2	8
40	Transbronchial Lung Cryobiopsy in Idiopathic Pulmonary Fibrosis: A State of the Art Review. Advances in Therapy, 2019, 36, 2193-2204.	1.3	7
41	S91â€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. Thorax, 2015, 70, A52.1-A52.	2.7	6
42	The extended utility of antifibrotic therapy in progressive fibrosing interstitial lung disease. Expert Review of Respiratory Medicine, 2020, 14, 1001-1008.	1.0	6
43	Current Treatments in the Management of Idiopathic Pulmonary Fibrosis: Pirfenidone and Nintedanib. Clinical Medicine Insights Therapeutics, 2017, 9, 1179559X1771912.	0.4	5
44	Physiological predictors of Hypoxic Challenge Testing (HCT) outcomes in Interstitial Lung Disease (ILD). Respiratory Medicine, 2018, 135, 51-56.	1.3	5
45	A global perspective on acute exacerbation of idiopathic pulmonary fibrosis (AE-IPF): results from an international survey. , 2018, , .		5
46	Real world experience of response to pirfenidone in patients with idiopathic pulmonary fibrosis: a two centre retrospective study. Sarcoidosis Vasculitis and Diffuse Lung Diseases, 2020, 37, 218-224.	0.2	5
47	How consistently do physicians diagnose and manage drug-induced interstitial lung disease? Two surveys of European ILD specialist physicians. ERJ Open Research, 2020, 6, 00286-2019.	1.1	4
48	P28â€Real World MDT Diagnosis of Idiopathic Pulmonary Fibrosis. Thorax, 2015, 70, A90.1-A90.	2.7	3
49	Feather bedding as a cause of hypersensitivity pneumonitis. QJM - Monthly Journal of the Association of Physicians, 2017, 110, hcx010.	0.2	3
50	Pulmonary Therapy Podcastâ€"COVID-19: Research and Real-World Experiences from the Editorial Board. Pulmonary Therapy, 2021, 7, 1-7.	1.1	2
51	P168â€Patient symptoms following discharge from hospital after COVID-19 Pneumonia. , 2021, , .		2
52	Adherence to home spirometry among patients with IPF: results from the INMARK trial. , 2019, , .		2
53	Interstitial lung disease associated with ANCA positivity: a retrospective analysis. , 2019, , .		2
54	Does nintedanib have the same effect on FVC decline in patients with progressive fibrosing ILDs treated with DMARDs or glucocorticoids?. , 2020, , .		2

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55	Idiopathic pulmonary fibrosis: a clinical update. British Journal of General Practice, 2018, 68, 249-250.	0.7	1
56	Treatment of severe idiopathic pulmonary fibrosisâ€"is sildenafil the next (in)stage?. Journal of Thoracic Disease, 2019, 11, 339-340.	0.6	1
57	Beware Weakening the Ivory Tower of MDT Diagnosis in Interstitial Lung Disease. Journal of Clinical Medicine, 2019, 8, 1964.	1.0	1
58	Introducing a new formulation of pirfenidone to reduce tablet burden for the IPF patient: Is it tolerable? Is it easy to take? What do our patients think?. , 2018, , .		1
59	Oxygen enhanced MRI biomarkers of lung function in interstitial lung disease. , 2020, , .		1
60	Rapidly non-ipf progressive fibrosing interstitial lung disease: a phenotype with an ipf-like behavior. Sarcoidosis Vasculitis and Diffuse Lung Diseases, 2020, 37, 231-233.	0.2	1
61	IPF Care, a support program for patients with idiopathic pulmonary fibrosis in the UK. , 2015, , .		1
62	First insights from the BTS idiopathic pulmonary fibrosis (IPF) registry. , 2016, , .		1
63	Hyperpolarised 129-xenon diffusion-weighted MRI in interstitial lung disease. , 2019, , .		1
64	Longitudinal change in hyperpolarised 129-xenon MR spectroscopy in interstitial lung disease., 2019,,.		1
65	Diesel Exhaust Particles Alter Monocyte Differentiation In Vitro. , 2010, , .		0
66	S13â€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. Thorax, 2013, 68, A10.1-A10.	2.7	0
67	P280 Extended Clinical Experience With Pirfenidone During A Named Patient Programme For Idiopathic Pulmonary Fibrosis (ipf): Interim Results. Thorax, 2014, 69, A196-A196.	2.7	0
68	M263 A Quarter Of Ipf Patients Not Eligible For Pirfenidone Treatment Due To The Nice Criteria Significantly Decline Over Time. Thorax, 2014, 69, A218-A218.	2.7	0
69	M264 Health And Economic Impact Of Prescribing Pirfenidone. Thorax, 2014, 69, A218-A219.	2.7	0
70	P6â€Early Clinical Experience With Nintedanib – a two centre review. Thorax, 2015, 70, A77.2-A77.	2.7	0
71	P34â€Sarcoidosis and co-existent Aspergillus lung disease. Thorax, 2015, 70, A92.3-A93.	2.7	0
72	Nintedanib for treating idiopathic pulmonary fibrosis. British Journal of Health Care Management, 2016, 22, 250-251.	0.1	0

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73	P151â€The role of bronchoalveolar lavage and its quality in the diagnosis of interstitial lung disease., 2017,,.		0
74	M24â€Do antifibrotics impact on lung transplantation outcomes in idiopathic pulmonary fibrosis?. , 2017, , .		0
75	P157â€Can baseline physiological tests help predict the outcome of hypoxic challenge testing (hct) in interstitial lung disease (ild)?. , 2017, , .		0
76	P148â€ldiopathic pulmonary fibrosis: "lost in the system―in the north west of england?., 2017,,.		0
77	Prevalence of incidental interstitial lung disease in the Manchester lung cancer screening pilot. Lung Cancer, 2019, 127, S23-S24.	0.9	0
78	AB0593â€INTERSTITIAL LUNG DISEASE ASSOCIATED WITH ANCA POSITIVITY: A RETROSPECTIVE ANALYSIS. , 20	)19, ,	0
79	Comment from the Editor of the Special Issue: "Lung Disease on COPD, Asthma, Bronchiectasis, Lung Cancer Screening, IPF― Journal of Clinical Medicine, 2019, 8, 2060.	1.0	0
80	Reply to Althuwaybi et al.: Hospitalization Outcomes for COVID-19 in Patients with Interstitial Lung Disease: A Potential Role for Aerodigestive Pathophysiology?. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 522-524.	2.5	0
81	Sarcoidosis in the UK: Insights from the BTS interstitial lung disease registry. , 2016, , .		0
82	Clinical experience with nintedanib for IPF in 3 UK tertiary referral centres., 2016,,.		0
83	The impact of a network based approach on lung function and symptom duration at diagnosis in idiopathic pulmonary fibrosis. , $2018,  ,  .$		0
84	What can we learn from Idiopathic Pulmonary Fibrosis Registries?., 2018,,.		0
85	Antifibrotic choice in idiopathic pulmonary fibrosis. , 2018, , .		0
86	Breath Biomarkers in Idiopathic Pulmonary Fibrosis: A Systematic Review., 2018,,.		0
87	Predictive variables to obtain a bronchoalveolar lavage of adequate quality in patients with interstitial lung diseases. , 2019, , .		0
88	Association of demographic, laboratory and clinical parameters with HRCT-chest findings in patients with sarcoidosis. , 2019, , .		0
89	Safety and tolerability of immunosuppression in non-IPF ILD: clinical experience from a tertiary ILD centre. , 2019, , .		0
90	P56â€What happens to patients with idiopathic pulmonary fibrosis who are not eligible for antifibrotic treatment due to current NICE guidelines. , 2019, , .		0

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91	S125â€Quantitative CT and hyperpolarised 129-xenon diffusion-weighted MRI in interstitial lung disease. , 2019, , .		O
92	Pulmonary involvement in antiphospholipid syndrome., 2019,, 124-139.		O
93	Exhaled volatile organic compounds in idiopathic pulmonary fibrosis and disease progression. , 2020, , .		O
94	Continued nintedanib treatment in patients with progressive fibrosing ILDs: interim analysis of INBUILD-ON. , 2021, , .		0
95	The impact of Covid-19 on hospital length of stay and resources: an experience from a tertiary respiratory centre in the UK. , 2021, , .		0
96	Admission clinical parameters in predicting in-hospital mortality in COVID-19., 2021, , .		0