

Paul W Noble

List of Publications by Year in descending order

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Version: 2024-02-01

86
papers

23,532
citations

31976

53
h-index

62596

80
g-index

94
all docs

94
docs citations

94
times ranked

18875
citing authors

#	ARTICLE	IF	CITATIONS
1	Efficacy and Safety of Nintedanib in Idiopathic Pulmonary Fibrosis. <i>New England Journal of Medicine</i> , 2014, 370, 2071-2082.	27.0	3,351
2	A Phase 3 Trial of Pirfenidone in Patients with Idiopathic Pulmonary Fibrosis. <i>New England Journal of Medicine</i> , 2014, 370, 2083-2092.	27.0	2,959
3	Pirfenidone in patients with idiopathic pulmonary fibrosis (CAPACITY): two randomised trials. <i>Lancet</i> , The, 2011, 377, 1760-1769.	13.7	1,711
4	Type 2 alveolar cells are stem cells in adult lung. <i>Journal of Clinical Investigation</i> , 2013, 123, 3025-3036.	8.2	1,352
5	Regulation of lung injury and repair by Toll-like receptors and hyaluronan. <i>Nature Medicine</i> , 2005, 11, 1173-1179.	30.7	1,291
6	Efficacy of a Tyrosine Kinase Inhibitor in Idiopathic Pulmonary Fibrosis. <i>New England Journal of Medicine</i> , 2011, 365, 1079-1087.	27.0	930
7	Multiple stromal populations contribute to pulmonary fibrosis without evidence for epithelial to mesenchymal transition. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, E1475-83.	7.1	849
8	Hyaluronan as an Immune Regulator in Human Diseases. <i>Physiological Reviews</i> , 2011, 91, 221-264.	28.8	848
9	Hyaluronan in Tissue Injury and Repair. <i>Annual Review of Cell and Developmental Biology</i> , 2007, 23, 435-461.	9.4	727
10	Resolution of Lung Inflammation by CD44. <i>Science</i> , 2002, 296, 155-158.	12.6	611
11	The Clinical Course of Patients with Idiopathic Pulmonary Fibrosis. <i>Annals of Internal Medicine</i> , 2005, 142, 963.	3.9	530
12	Hyaluronan and its catabolic products in tissue injury and repair. <i>Matrix Biology</i> , 2002, 21, 25-29.	3.6	491
13	Pulmonary fibrosis: patterns and perpetrators. <i>Journal of Clinical Investigation</i> , 2012, 122, 2756-2762.	8.2	429
14	Single-Cell Deconvolution of Fibroblast Heterogeneity in Mouse Pulmonary Fibrosis. <i>Cell Reports</i> , 2018, 22, 3625-3640.	6.4	392
15	Forced Vital Capacity in Patients with Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2011, 184, 1382-1389.	5.6	390
16	Six-Minute-Walk Test in Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2011, 183, 1231-1237.	5.6	369
17	Pirfenidone for idiopathic pulmonary fibrosis: analysis of pooled data from three multinational phase 3 trials. <i>European Respiratory Journal</i> , 2016, 47, 243-253.	6.7	349
18	Idiopathic pulmonary fibrosis. <i>Orphanet Journal of Rare Diseases</i> , 2008, 3, 8.	2.7	332

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19	Severe lung fibrosis requires an invasive fibroblast phenotype regulated by hyaluronan and CD44. <i>Journal of Experimental Medicine</i> , 2011, 208, 1459-1471.	8.5	322
20	Hyaluronan Fragments Induce Nitric-oxide Synthase in Murine Macrophages through a Nuclear Factor κ B-dependent Mechanism. <i>Journal of Biological Chemistry</i> , 1997, 272, 8013-8018.	3.4	264
21	Effect of pirfenidone on mortality: pooled analyses and meta-analyses of clinical trials in idiopathic pulmonary fibrosis. <i>Lancet Respiratory Medicine</i> , 2017, 5, 33-41.	10.7	240
22	Hyaluronan and TLR4 promote surfactant-protein-C-positive alveolar progenitor cell renewal and prevent severe pulmonary fibrosis in mice. <i>Nature Medicine</i> , 2016, 22, 1285-1293.	30.7	211
23	Alveolar Epithelial Type II Cells as Drivers of Lung Fibrosis in Idiopathic Pulmonary Fibrosis. <i>International Journal of Molecular Sciences</i> , 2020, 21, 2269.	4.1	202
24	Efficacy of simtuzumab versus placebo in patients with idiopathic pulmonary fibrosis: a randomised, double-blind, controlled, phase 2 trial. <i>Lancet Respiratory Medicine</i> , 2017, 5, 22-32.	10.7	200
25	Flow Cytometric Analysis of Myeloid Cells in Human Blood, Bronchoalveolar Lavage, and Lung Tissues. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2016, 54, 13-24.	2.9	191
26	Interleukin-11 is a therapeutic target in idiopathic pulmonary fibrosis. <i>Science Translational Medicine</i> , 2019, 11, .	12.4	189
27	6-minute walk distance is an independent predictor of mortality in patients with idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2014, 43, 1421-1429.	6.7	180
28	Airway Epithelial Progenitors Are Region Specific and Show Differential Responses to Bleomycin-Induced Lung Injury. <i>Stem Cells</i> , 2012, 30, 1948-1960.	3.2	171
29	Hyaluronan as a therapeutic target in human diseases. <i>Advanced Drug Delivery Reviews</i> , 2016, 97, 186-203.	13.7	167
30	Hyaluronan Fragments Synergize with Interferon- γ to Induce the C-X-C Chemokines Mig and Interferon-inducible Protein-10 in Mouse Macrophages. <i>Journal of Biological Chemistry</i> , 1998, 273, 35088-35094.	3.4	161
31	Effect of continued treatment with pirfenidone following clinically meaningful declines in forced vital capacity: analysis of data from three phase 3 trials in patients with idiopathic pulmonary fibrosis. <i>Thorax</i> , 2016, 71, 429-435.	5.6	151
32	Time for a change: is idiopathic pulmonary fibrosis still idiopathic and only fibrotic?. <i>Lancet Respiratory Medicine</i> , 2018, 6, 154-160.	10.7	137
33	Cellular Mechanisms of Tissue Fibrosis. 7. New insights into the cellular mechanisms of pulmonary fibrosis. <i>American Journal of Physiology - Cell Physiology</i> , 2014, 306, C987-C996.	4.6	133
34	Blocking follistatin-like 1 attenuates bleomycin-induced pulmonary fibrosis in mice. <i>Journal of Experimental Medicine</i> , 2015, 212, 235-252.	8.5	130
35	The role of Toll-like receptors in non-infectious lung injury. <i>Cell Research</i> , 2006, 16, 693-701.	12.0	129
36	Pre-existing traits associated with Covid-19 illness severity. <i>PLoS ONE</i> , 2020, 15, e0236240.	2.5	129

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37	CD44 Is a Negative Regulator of Acute Pulmonary Inflammation and Lipopolysaccharide-TLR Signaling in Mouse Macrophages. <i>Journal of Immunology</i> , 2007, 178, 2469-2475.	0.8	127
38	CD44 Deficiency Leads to Enhanced Neutrophil Migration and Lung Injury in Escherichia coli Pneumonia in Mice. <i>American Journal of Pathology</i> , 2002, 161, 2219-2228.	3.8	119
39	An Open-Label Study of the Long-Term Safety of Pirfenidone in Patients with Idiopathic Pulmonary Fibrosis (RECAP). <i>Respiration</i> , 2017, 94, 408-415.	2.6	116
40	Single-Cell Reconstruction of Human Basal Cell Diversity in Normal and Idiopathic Pulmonary Fibrosis Lungs. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 202, 1540-1550.	5.6	107
41	Transcription factor TBX4 regulates myofibroblast accumulation and lung fibrosis. <i>Journal of Clinical Investigation</i> , 2016, 126, 3063-3079.	8.2	101
42	Safety of pirfenidone in patients with idiopathic pulmonary fibrosis: integrated analysis of cumulative data from 5 clinical trials. <i>BMJ Open Respiratory Research</i> , 2016, 3, e000105.	3.0	96
43	Matrix Regulation of Lung Injury, Inflammation, and Repair: The Role of Innate Immunity. <i>Proceedings of the American Thoracic Society</i> , 2006, 3, 401-404.	3.5	93
44	Idiopathic pulmonary fibrosis: new insights into pathogenesis. <i>Clinics in Chest Medicine</i> , 2004, 25, 749-758.	2.1	92
45	Hyaluronan synthase 2-mediated hyaluronan production mediates Notch1 activation and liver fibrosis. <i>Science Translational Medicine</i> , 2019, 11, .	12.4	91
46	Role of hyaluronan and hyaluronan-binding proteins in human asthma. <i>Journal of Allergy and Clinical Immunology</i> , 2011, 128, 403-411.e3.	2.9	89
47	Validation of test performance characteristics and minimal clinically important difference of the 6-minute walk test in patients with idiopathic pulmonary fibrosis. <i>Respiratory Medicine</i> , 2015, 109, 914-922.	2.9	85
48	Back to the Future. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2005, 33, 113-120.	2.9	83
49	Hyaluronan synthase 2 regulates fibroblast senescence in pulmonary fibrosis. <i>Matrix Biology</i> , 2016, 55, 35-48.	3.6	72
50	Lung Transplantation for Covid-19-Related Respiratory Failure in the United States. <i>New England Journal of Medicine</i> , 2022, 386, 1187-1188.	27.0	72
51	Targeting of TAM Receptors Ameliorates Fibrotic Mechanisms in Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018, 197, 1443-1456.	5.6	66
52	PD-L1 on invasive fibroblasts drives fibrosis in a humanized model of idiopathic pulmonary fibrosis. <i>JCI Insight</i> , 2019, 4, .	5.0	64
53	Regulation of plasminogen activator inhibitor-1 and urokinase by hyaluronan fragments in mouse macrophages. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2000, 279, L707-L715.	2.9	63
54	Idiopathic Pulmonary Fibrosis: Natural History and Prognosis. <i>Clinics in Chest Medicine</i> , 2006, 27, 11-16.	2.1	56

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55	Regulation of Non-infectious Lung Injury, Inflammation, and Repair by the Extracellular Matrix Glycosaminoglycan Hyaluronan. <i>Anatomical Record</i> , 2010, 293, 982-985.	1.4	54
56	Pirfenidone in patients with idiopathic pulmonary fibrosis and more advanced lung function impairment. <i>Respiratory Medicine</i> , 2019, 153, 44-51.	2.9	54
57	Syndecan-1 promotes lung fibrosis by regulating epithelial reprogramming through extracellular vesicles. <i>JCI Insight</i> , 2019, 4, .	5.0	50
58	MicroRNA-29c Prevents Pulmonary Fibrosis by Regulating Epithelial Cell Renewal and Apoptosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2017, 57, 721-732.	2.9	46
59	Categorization of lung mesenchymal cells in development and fibrosis. <i>IScience</i> , 2021, 24, 102551.	4.1	46
60	CD44 Deficiency Is Associated with Increased Bacterial Clearance but Enhanced Lung Inflammation During Gram-Negative Pneumonia. <i>American Journal of Pathology</i> , 2010, 177, 2483-2494.	3.8	43
61	Pirfenidone Initiates a New Era in the Treatment of Idiopathic Pulmonary Fibrosis. <i>Annual Review of Medicine</i> , 2016, 67, 487-495.	12.2	37
62	miR-323a-3p regulates lung fibrosis by targeting multiple profibrotic pathways. <i>JCI Insight</i> , 2016, 1, e90301.	5.0	37
63	The ZIP8/SIRT1 axis regulates alveolar progenitor cell renewal in aging and idiopathic pulmonary fibrosis. <i>Journal of Clinical Investigation</i> , 2022, 132, .	8.2	37
64	Methylation-mediated BMPER expression in fibroblast activation in vitro and lung fibrosis in mice in vivo. <i>Scientific Reports</i> , 2015, 5, 14910.	3.3	35
65	Sensitivity Analyses of the Change in FVC in a Phase 3 Trial of Pirfenidone for Idiopathic Pulmonary Fibrosis. <i>Chest</i> , 2015, 148, 196-201.	0.8	35
66	Dose modification and dose intensity during treatment with pirfenidone: analysis of pooled data from three multinational phase III trials. <i>BMJ Open Respiratory Research</i> , 2018, 5, e000323.	3.0	35
67	Effect of pirfenidone in patients with more advanced idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2019, 20, 55.	3.6	33
68	Idiopathic pulmonary fibrosis. New insights into classification and pathogenesis usher in a new era therapeutic approaches. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2003, 29, S27-31.	2.9	31
69	Risk factors for disease progression in idiopathic pulmonary fibrosis. <i>Thorax</i> , 2020, 75, 78-80.	5.6	22
70	Mitogen-activated Protein Kinase-activated Protein Kinase 2 Inhibition Attenuates Fibroblast Invasion and Severe Lung Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2019, 60, 41-48.	2.9	18
71	Cardiovascular Risks, Bleeding Risks, and Clinical Events from 3 Phase III Trials of Pirfenidone in Patients with Idiopathic Pulmonary Fibrosis. <i>Advances in Therapy</i> , 2019, 36, 2910-2926.	2.9	18
72	CC-90001, a c-Jun N-terminal kinase (JNK) inhibitor, in patients with pulmonary fibrosis: design of a phase 2, randomised, placebo-controlled trial. <i>BMJ Open Respiratory Research</i> , 2022, 9, e001060.	3.0	17

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73	Pirfenidone Treatment in Individuals with Idiopathic Pulmonary Fibrosis: Impact of Timing of Treatment Initiation. <i>Annals of the American Thoracic Society</i> , 2019, 16, 927-930.	3.2	16
74	Mesenchymal growth hormone receptor deficiency leads to failure of alveolar progenitor cell function and severe pulmonary fibrosis. <i>Science Advances</i> , 2021, 7, .	10.3	10
75	Abnormal respiratory progenitors in fibrotic lung injury. <i>Stem Cell Research and Therapy</i> , 2022, 13, 64.	5.5	10
76	Antibody-mediated depletion of CCR10+ EphA3+ cells ameliorates fibrosis in IPF. <i>JCI Insight</i> , 2021, 6, .	5.0	9
77	Quantitative Image Analysis at Chronic Lung Allograft Dysfunction Onset Predicts Mortality. <i>Transplantation</i> , 2022, 106, 1253-1261.	1.0	6
78	Disruption of respiratory epithelial basement membrane in COVID-19 patients. <i>Molecular Biomedicine</i> , 2021, 2, 8.	4.4	4
79	Case Study Review. <i>Chest</i> , 2005, 128, 540S-546S.	0.8	3
80	The allograft injury marker CXCL9 determines prognosis of anti- α HLA antibodies after lung transplantation. <i>American Journal of Transplantation</i> , 2021, , .	4.7	2
81	Stem Cells and Progenitor Cells in Interstitial Lung Disease. , 2022, , 158-168.		2
82	Take a deep breath: pulmonary research inspires. <i>Journal of Clinical Investigation</i> , 2012, 122, 2722-2723.	8.2	0
83	Pre-existing traits associated with Covid-19 illness severity. , 2020, 15, e0236240.		0
84	Pre-existing traits associated with Covid-19 illness severity. , 2020, 15, e0236240.		0
85	Pre-existing traits associated with Covid-19 illness severity. , 2020, 15, e0236240.		0
86	Pre-existing traits associated with Covid-19 illness severity. , 2020, 15, e0236240.		0