

David N O'dwyer

List of Publications by Year in descending order

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Version: 2024-02-01

33
papers

1,479
citations

471371

17
h-index

414303

32
g-index

34
all docs

34
docs citations

34
times ranked

2370
citing authors

#	ARTICLE	IF	CITATIONS
1	The Lung Microbiome, Immunity, and the Pathogenesis of Chronic Lung Disease. <i>Journal of Immunology</i> , 2016, 196, 4839-4847.	0.4	291
2	Lung Microbiota Contribute to Pulmonary Inflammation and Disease Progression in Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 199, 1127-1138.	2.5	205
3	The Toll-like Receptor 3 L412F Polymorphism and Disease Progression in Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2013, 188, 1442-1450.	2.5	149
4	The role of periostin in lung fibrosis and airway remodeling. <i>Cellular and Molecular Life Sciences</i> , 2017, 74, 4305-4314.	2.4	99
5	Methods in Lung Microbiome Research. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2020, 62, 283-299.	1.4	94
6	Rheumatoid Arthritis (RA) associated interstitial lung disease (ILD). <i>European Journal of Internal Medicine</i> , 2013, 24, 597-603.	1.0	93
7	Influences of innate immunity, autophagy, and fibroblast activation in the pathogenesis of lung fibrosis. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2016, 311, L590-L601.	1.3	74
8	The peripheral blood proteome signature of idiopathic pulmonary fibrosis is distinct from normal and is associated with novel immunological processes. <i>Scientific Reports</i> , 2017, 7, 46560.	1.6	51
9	Six-SOMAmer Index Relating to Immune, Protease and Angiogenic Functions Predicts Progression in IPF. <i>PLoS ONE</i> , 2016, 11, e0159878.	1.1	43
10	Lung Dysbiosis, Inflammation, and Injury in Hematopoietic Cell Transplantation. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018, 198, 1312-1321.	2.5	42
11	A Germline Mutation in the C2 Domain of PLCÎ³2 Associated with Gain-of-Function Expands the Phenotype for PLCG2-Related Diseases. <i>Journal of Clinical Immunology</i> , 2020, 40, 267-276.	2.0	31
12	First-Onset Herpesviral Infection and Lung Injury in Allogeneic Hematopoietic Cell Transplantation. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 63-74.	2.5	30
13	Therapeutic Targeting of the Respiratory Microbiome. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2022, 206, 535-544.	2.5	24
14	Targeting defective Toll-like receptor-3 function and idiopathic pulmonary fibrosis. <i>Expert Opinion on Therapeutic Targets</i> , 2015, 19, 507-514.	1.5	23
15	Loss of CCR2 signaling alters leukocyte recruitment and exacerbates Î³-herpesvirus-induced pneumonitis and fibrosis following bone marrow transplantation. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2016, 311, L611-L627.	1.3	22
16	Pulmonary immunity and extracellular matrix interactions. <i>Matrix Biology</i> , 2018, 73, 122-134.	1.5	21
17	Radiographic Honeycombing and Altered Lung Microbiota in Patients with Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 1544-1547.	2.5	20
18	The evolving role of the lung microbiome in pulmonary fibrosis. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2020, 319, L675-L682.	1.3	18

#	ARTICLE	IF	CITATIONS
19	Animal Models of Pulmonary Fibrosis. <i>Methods in Molecular Biology</i> , 2018, 1809, 363-378.	0.4	17
20	Proteomics: Clinical and research applications in respiratory diseases. <i>Respirology</i> , 2018, 23, 993-1003.	1.3	15
21	Candidate Role for Toll-like Receptor 3 L412F Polymorphism and Infection in Acute Exacerbation of Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2022, 205, 550-562.	2.5	12
22	Macrophage migration inhibitory factor enhances <i>Pseudomonas aeruginosa</i> biofilm formation, potentially contributing to cystic fibrosis pathogenesis. <i>FASEB Journal</i> , 2017, 31, 5102-5110.	0.2	10
23	Identification of a unique temporal signature in blood and BAL associated with IPF progression. <i>Scientific Reports</i> , 2020, 10, 12049.	1.6	10
24	Fibrotic lung disease inhibits immune responses to staphylococcal pneumonia via impaired neutrophil and macrophage function. <i>JCI Insight</i> , 2022, 7, .	2.3	9
25	Toll-like receptors, environmental caging, and lung dysbiosis. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2021, 321, L404-L415.	1.3	8
26	Diffuse pulmonary meningotheliomatosis. <i>Clinical Imaging</i> , 2021, 70, 111-113.	0.8	6
27	Pretransplant Antifibrotic Therapy Is Associated with Resolution of Primary Graft Dysfunction. <i>Annals of the American Thoracic Society</i> , 2022, 19, 335-338.	1.5	4
28	The Lung Microbiome in Health, Hypersensitivity Pneumonitis, and Idiopathic Pulmonary Fibrosis: A Heavy Bacterial Burden to Bear. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021, 203, 281-283.	2.5	3
29	Host-Microbial Interactions: Idiopathic Pulmonary Fibrosis in Technicolor. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2017, 195, 1554-1556.	2.5	2
30	Ironing Out the Roles of Macrophages in Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 127-129.	2.5	2
31	Concurrent Reductions in Spirometry Predict Mortality and Bronchiolitis Obliterans in Chronic Graft-versus-Host Disease. <i>Annals of the American Thoracic Society</i> , 2021, 18, 720-723.	1.5	1
32	Hypothyroidism Is Associated with Increased Mortality in Interstitial Pneumonia with Autoimmune Features. <i>Annals of the American Thoracic Society</i> , 2022, 19, 1772-1776.	1.5	1
33	Toll-Interacting Protein and Altered Lung Microbiota in Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2022, , .	2.5	0