

Alex R Horsley

List of Publications by Year in descending order

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Version: 2024-02-01

85
papers

4,062
citations

201674

27
h-index

128289

60
g-index

96
all docs

96
docs citations

96
times ranked

6045
citing authors

#	ARTICLE	IF	CITATIONS
1	Consensus statement for inert gas washout measurement using multiple- and single- breath tests. <i>European Respiratory Journal</i> , 2013, 41, 507-522.	6.7	631
2	Physical, cognitive, and mental health impacts of COVID-19 after hospitalisation (PHOSP-COVID): a UK multicentre, prospective cohort study. <i>Lancet Respiratory Medicine</i> , 2021, 9, 1275-1287.	10.7	394
3	Safety and efficacy of inhaled nebulised interferon beta-1a (SNG001) for treatment of SARS-CoV-2 infection: a randomised, double-blind, placebo-controlled, phase 2 trial. <i>Lancet Respiratory Medicine</i> , 2021, 9, 196-206.	10.7	370
4	Robust SARS-CoV-2-specific T cell immunity is maintained at 6 months following primary infection. <i>Nature Immunology</i> , 2021, 22, 620-626.	14.5	320
5	VX-659â€“Tezacaftorâ€“Ivacaftor in Patients with Cystic Fibrosis and One or Two Phe508del Alleles. <i>New England Journal of Medicine</i> , 2018, 379, 1599-1611.	27.0	280
6	Lung clearance index is a sensitive, repeatable and practical measure of airways disease in adults with cystic fibrosis. <i>Thorax</i> , 2007, 63, 135-140.	5.6	193
7	Efficacy and Complications of Small-Bore, Wire-Guided Chest Drains. <i>Chest</i> , 2006, 130, 1857-1863.	0.8	135
8	Lung clearance index in the assessment of airways disease. <i>Respiratory Medicine</i> , 2009, 103, 793-799.	2.9	128
9	Changes in physiological, functional and structural markers of cystic fibrosis lung disease with treatment of a pulmonary exacerbation. <i>Thorax</i> , 2013, 68, 532-539.	5.6	121
10	Ventilation heterogeneity in children with well controlled asthma with normal spirometry indicates residual airways disease. <i>Thorax</i> , 2008, 64, 33-37.	5.6	97
11	Preschool Multiple-Breath Washout Testing. An Official American Thoracic Society Technical Statement. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018, 197, e1-e19.	5.6	92
12	Effects of Ivacaftor in Patients With Cystic Fibrosis Who Carry the G551D Mutation and Have Severe Lung Disease. <i>Chest</i> , 2014, 146, 152-158.	0.8	85
13	Detection of early subclinical lung disease in children with cystic fibrosis by lung ventilation imaging with hyperpolarised gas MRI. <i>Thorax</i> , 2017, 72, 760-762.	5.6	70
14	Lung Clearance Index Is a Repeatable and Sensitive Indicator of Radiological Changes in Bronchiectasis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014, 189, 586-592.	5.6	67
15	Lung clearance index in cystic fibrosis subjects treated for pulmonary exacerbations. <i>European Respiratory Journal</i> , 2015, 46, 1055-1064.	6.7	61
16	Serological surveillance of SARS-CoV-2: Six-month trends and antibody response in a cohort of public health workers. <i>Journal of Infection</i> , 2021, 82, 162-169.	3.3	61
17	Longitudinal Assessment of Children with Mild Cystic Fibrosis Using Hyperpolarized Gas Lung Magnetic Resonance Imaging and Lung Clearance Index. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018, 197, 397-400.	5.6	54
18	Effects of cystic fibrosis lung disease on gas mixing indices derived from alveolar slope analysis. <i>Respiratory Physiology and Neurobiology</i> , 2008, 162, 197-203.	1.6	50

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19	Putting lung function and physiology into perspective: cystic fibrosis in adults. <i>Respirology</i> , 2015, 20, 33-45.	2.3	43
20	Lung clearance index in adults with non-cystic fibrosis bronchiectasis. <i>Respiratory Research</i> , 2014, 15, 59.	3.6	39
21	Antibiotic treatment for <i>Burkholderia cepacia</i> complex in people with cystic fibrosis experiencing a pulmonary exacerbation. <i>The Cochrane Library</i> , 2016, , CD009529.	2.8	39
22	Can Early <i>Burkholderia cepacia</i> Complex Infection in Cystic Fibrosis be Eradicated with Antibiotic Therapy?. <i>Frontiers in Cellular and Infection Microbiology</i> , 2011, 1, 18.	3.9	38
23	Sweat chloride is not a useful marker of clinical response to Ivacaftor. <i>Thorax</i> , 2014, 69, 586-587.	5.6	35
24	Patterns of regional lung physiology in cystic fibrosis using ventilation magnetic resonance imaging and multiple-breath washout. <i>European Respiratory Journal</i> , 2018, 52, 1800821.	6.7	35
25	Itraconazole and inhaled fluticasone causing hypothalamic-pituitary-adrenal axis suppression in adults with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2013, 12, 399-402.	0.7	33
26	Antibiotic treatment for <i>Burkholderia cepacia</i> complex in people with cystic fibrosis experiencing a pulmonary exacerbation. , 2012, 10, CD009529.		30
27	Feasibility and challenges of using multiple breath washout in COPD. <i>International Journal of COPD</i> , 2018, Volume 13, 2113-2119.	2.3	30
28	Evaluation of the Impact of Alveolar Nitrogen Excretion on Indices Derived from Multiple Breath Nitrogen Washout. <i>PLoS ONE</i> , 2013, 8, e73335.	2.5	29
29	Aspergillosis and the role of mucins in cystic fibrosis. <i>Pediatric Pulmonology</i> , 2017, 52, 548-555.	2.0	28
30	The assessment of short- and long-term changes in lung function in cystic fibrosis using ¹²⁹ Xe MRI. <i>European Respiratory Journal</i> , 2020, 56, 2000441.	6.7	25
31	What is the importance of classifying <i>Aspergillus</i> disease in cystic fibrosis patients?. <i>Expert Review of Respiratory Medicine</i> , 2014, 8, 389-392.	2.5	21
32	Simultaneous sulfur hexafluoride and nitrogen multiple-breath washout (MBW) to examine inherent differences in MBW outcomes. <i>ERJ Open Research</i> , 2019, 5, 00234-2018.	2.6	20
33	Antibiotic treatment for <i>Burkholderia cepacia</i> complex in people with cystic fibrosis experiencing a pulmonary exacerbation. <i>The Cochrane Library</i> , 2020, 4, CD009529.	2.8	20
34	Enhanced Photoacoustic Gas Analyser Response Time and Impact on Accuracy at Fast Ventilation Rates during Multiple Breath Washout. <i>PLoS ONE</i> , 2014, 9, e98487.	2.5	20
35	Reassessment of the importance of mucins in determining sputum properties in cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2014, 13, 260-266.	0.7	18
36	Ventilation heterogeneity and the benefits and challenges of multiple breath washout testing in patients with cystic fibrosis. <i>Paediatric Respiratory Reviews</i> , 2015, 16, 15-18.	1.8	18

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37	Monitoring early stage lung disease in cystic fibrosis. <i>Current Opinion in Pulmonary Medicine</i> , 2020, 26, 671-678.	2.6	16
38	Joint patient and clinician priority setting to identify 10 key research questions regarding the long-term sequelae of COVID-19. <i>Thorax</i> , 2022, 77, 717-720.	5.6	16
39	Handheld computers in medicine: the way forward. <i>Postgraduate Medical Journal</i> , 2005, 81, 481-482.	1.8	15
40	Closed circuit rebreathing to achieve inert gas wash-in for multiple breath wash-out. <i>ERJ Open Research</i> , 2016, 2, 00042-2015.	2.6	14
41	Spectral graph theory efficiently characterizes ventilation heterogeneity in lung airway networks. <i>Journal of the Royal Society Interface</i> , 2020, 17, 20200253.	3.4	12
42	Longitudinal assessment of lung clearance index to monitor disease progression in children and adults with cystic fibrosis. <i>Thorax</i> , 2022, 77, 357-363.	5.6	11
43	An immunocytochemical assay to detect human CFTR expression following gene transfer. <i>Molecular and Cellular Probes</i> , 2009, 23, 272-280.	2.1	10
44	Using social media to improve communication with people with cystic fibrosis. <i>ERJ Open Research</i> , 2016, 2, 00015-2016.	2.6	10
45	Assessing arthritis in the context of cystic fibrosis. <i>Pediatric Pulmonology</i> , 2019, 54, 770-777.	2.0	10
46	The effect of acute maximal exercise on the regional distribution of ventilation using ventilation MRI in CF. <i>Journal of Cystic Fibrosis</i> , 2020, 20, 625-631.	0.7	10
47	Accurate lung volume measurements in vitro using a novel inert gas washout method suitable for infants. <i>Pediatric Pulmonology</i> , 2016, 51, 491-497.	2.0	9
48	Lung clearance index in detection of post-transplant bronchiolitis obliterans syndrome. <i>ERJ Open Research</i> , 2019, 5, 00164-2019.	2.6	9
49	Empire-CF study: A phase 2 clinical trial of leukotriene A4 hydrolase inhibitor acebilustat in adult subjects with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2021, 20, 1026-1034.	0.7	9
50	Modelling structural determinants of ventilation heterogeneity: A perturbative approach. <i>PLoS ONE</i> , 2018, 13, e0208049.	2.5	8
51	Implications of fatherhood in cystic fibrosis. <i>Paediatric Respiratory Reviews</i> , 2019, 31, 18-20.	1.8	7
52	A multimodal approach to detect and monitor early lung disease in cystic fibrosis. <i>Expert Review of Respiratory Medicine</i> , 2021, 15, 761-772.	2.5	7
53	Supine posture changes lung volumes and increases ventilation heterogeneity in cystic fibrosis. <i>PLoS ONE</i> , 2017, 12, e0188275.	2.5	7
54	Large scale clinical trials: lessons from the COVID-19 pandemic. <i>BMJ Open Respiratory Research</i> , 2022, 9, e001226.	3.0	7

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55	Burkholderia latens infection in cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2011, 10, 291-292.	0.7	6
56	Gout and hyperuricaemia in adults with cystic fibrosis. <i>Journal of the Royal Society of Medicine</i> , 2011, 104, 36-39.	2.0	6
57	A novel method for infant multiple breath washout: First report in clinical practice. <i>Pediatric Pulmonology</i> , 2019, 54, 1284-1290.	2.0	6
58	Composition of airway bacterial community correlates with chest HRCT in adults with bronchiectasis. <i>Respirology</i> , 2020, 25, 64-70.	2.3	6
59	Surface-tension-driven evolution of a viscoplastic liquid coating the interior of a cylindrical tube. <i>Journal of Fluid Mechanics</i> , 2022, 944, .	3.4	6
60	Commentaries on Viewpoint: Using the same cut-off for sulfur hexafluoride and nitrogen multiple-breath washout may not be appropriate. <i>Journal of Applied Physiology</i> , 2015, 119, 1513-1514.	2.5	5
61	Orkambi in patients with severe disease – Bumps in the road to CFTR modulation. <i>Journal of Cystic Fibrosis</i> , 2017, 16, 311-312.	0.7	4
62	Lung clearance index in healthy volunteers, measured using a novel portable system with a closed circuit wash-in. <i>PLoS ONE</i> , 2020, 15, e0229300.	2.5	4
63	Impact of airway <i>Exophiala</i> spp. on children with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2021, 20, 702-707.	0.7	4
64	Discordance between clinical, physiological, and radiological measures in cystic fibrosis. <i>Respirology Case Reports</i> , 2014, 2, 129-131.	0.6	3
65	Better late than never: correcting the error in the Exhalyzer nitrogen washout system. <i>Journal of Applied Physiology</i> , 2021, 131, 1286-1287.	2.5	3
66	Reply: Lung Clearance Index in Primary Ciliary Dyskinesia and Bronchiectasis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014, 189, 1148-1149.	5.6	2
67	ERS International Congress 2020: highlights from the Paediatric Assembly. <i>ERJ Open Research</i> , 2021, 7, 00893-2020.	2.6	2
68	112 – Musculoskeletal symptoms represent a significant burden for adults with cystic fibrosis. <i>Rheumatology</i> , 2018, 57, .	1.9	1
69	Comment on Comparison of lung clearance index determined by washout of N2 and SF6 in infants and preschool children with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2019, 18, e26-e27.	0.7	1
70	ERS International Congress, Madrid, 2019: highlights from the Paediatric Assembly. <i>ERJ Open Research</i> , 2020, 6, 00063-2020.	2.6	1
71	Chapter 1 Genetics and pathophysiology. , 2010, , .		1
72	The potential of closed circuit lung clearance index (LCI) to provide longitudinal clinical utility in cystic fibrosis (CF). , 2019, , .		1

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73	HPV-associated complications post lung transplantation: why prevention is better than a cure. Thorax, 2022, 77, 539-539.	5.6	1
74	A very breathless woman. BMJ: British Medical Journal, 2011, 343, d5935-d5935.	2.3	0
75	Author's response: heterogeneity of change in LCI in patients with cystic fibrosis following antibiotic treatment. Thorax, 2014, 69, 184.2-184.	5.6	0
76	Challenge for a new era—the importance of ensuring accuracy of genotype in cystic fibrosis registries. Journal of Cystic Fibrosis, 2016, 15, e50-e51.	0.7	0
77	Effect of intermittent inspiratory leaks on measurement of lung clearance index using nitrogen and sulfur hexafluoride. ERJ Open Research, 2018, 4, 00132-2018.	2.6	0
78	Chapter 5 Management of respiratory exacerbations. , 2010, , .		0
79	Title is missing!. , 2020, 15, e0229300.		0
80	Title is missing!. , 2020, 15, e0229300.		0
81	Title is missing!. , 2020, 15, e0229300.		0
82	Title is missing!. , 2020, 15, e0229300.		0
83	Title is missing!. , 2020, 15, e0229300.		0
84	Title is missing!. , 2020, 15, e0229300.		0
85	Model-based Bayesian inference of the ventilation distribution in patients with Cystic Fibrosis from multiple breath washout, with comparison to ventilation MRI. Respiratory Physiology and Neurobiology, 2022, 302, 103919.	1.6	0