

# Moira L Aitken

## List of Publications by Year in descending order

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Version: 2024-02-01

27  
papers

541  
citations

687363

13  
h-index

677142

22  
g-index

27  
all docs

27  
docs citations

27  
times ranked

748  
citing authors

#	ARTICLE	IF	CITATIONS
1	Pregnancy in cystic fibrosis: Review of the literature and expert recommendations. <i>Journal of Cystic Fibrosis</i> , 2022, 21, 387-395.	0.7	28
2	Family building and parenting considerations for people with cystic fibrosis. <i>Pediatric Pulmonology</i> , 2022, 57, .	2.0	9
3	Optimizing sexual and reproductive health across the lifespan in people with cystic fibrosis. <i>Pediatric Pulmonology</i> , 2022, 57, .	2.0	8
4	Low body mass index as a barrier to lung transplant in cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2022, 21, 475-481.	0.7	8
5	Hemoptysis and the Risk for Lung Transplant or Death without Transplant in Individuals with Cystic Fibrosis in the United States. <i>Annals of the American Thoracic Society</i> , 2022, 19, 1986-1992.	3.2	2
6	Glycated Albumin Triggers an Inflammatory Response in the Human Airway Epithelium and Causes an Increase in Ciliary Beat Frequency. <i>Frontiers in Physiology</i> , 2021, 12, 653177.	2.8	4
7	Strong toll-like receptor responses in cystic fibrosis patients are associated with higher lung function. <i>Journal of Cystic Fibrosis</i> , 2020, 19, 608-613.	0.7	6
8	Underweight Patients With Cystic Fibrosis Have Acceptable Survival Following Lung Transplantation. <i>Chest</i> , 2020, 157, 898-906.	0.8	35
9	Impaired counterregulatory responses to hypoglycaemia following oral glucose in adults with cystic fibrosis. <i>Diabetologia</i> , 2020, 63, 1055-1065.	6.3	13
10	The expanding phenotype of <i>CFTR</i> -related disorders: Hemizygous loss-of-function variants in three patients with primary ciliary dyskinesia. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2019, 7, e911.	1.2	31
11	Addressing lung transplant with adults with cystic fibrosis: A qualitative analysis of patients' perspectives and experiences. <i>Journal of Cystic Fibrosis</i> , 2019, 18, 416-419.	0.7	13
12	Transcriptional and functional diversity of human macrophage repolarization. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 1536-1548.	2.9	49
13	Exploring Opportunities for Primary Outpatient Palliative Care for Adults with Cystic Fibrosis: A Mixed-Methods Study of Patients' Needs. <i>Journal of Palliative Medicine</i> , 2018, 21, 513-521.	1.1	16
14	Changing Rates of Chronic <i>Pseudomonas aeruginosa</i> Infections in Cystic Fibrosis: A Population-Based Cohort Study. <i>Clinical Infectious Diseases</i> , 2018, 67, 1089-1095.	5.8	47
15	Common clinical features of CF (respiratory disease and exocrine pancreatic insufficiency). <i>Presse Medicale</i> , 2017, 46, e109-e124.	1.9	10
16	Deaths Related to Bronchial Arterial Embolization in Patients With Cystic Fibrosis. <i>Chest</i> , 2016, 150, e93-e98.	0.8	15
17	Change in <i>Pseudomonas aeruginosa</i> prevalence in cystic fibrosis adults over time. <i>BMC Pulmonary Medicine</i> , 2016, 16, 176.	2.0	37
18	Gallium Compounds Exhibit Potential as New Therapeutic Agents against <i>Mycobacterium abscessus</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 4826-4834.	3.2	43

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19	Prevalence of Symptoms of Depression and Anxiety in Adults With Cystic Fibrosis Based on the PHQ-9 and GAD-7 Screening Questionnaires. <i>Psychosomatics</i> , 2015, 56, 345-353.	2.5	42
20	Response. <i>Chest</i> , 2013, 143, 272.	0.8	0
21	Risk of Post-Lung Transplant Renal Dysfunction in Adults With Cystic Fibrosis. <i>Chest</i> , 2012, 142, 185-191.	0.8	33
22	Analysis of Sequential Aliquots of Hypertonic Saline Solution-Induced Sputum From Clinically Stable Patients With Cystic Fibrosis. <i>Chest</i> , 2003, 123, 792-799.	0.8	34
23	Pseudomonal pericarditis complicating cystic fibrosis. , 1999, 27, 62-64.		10
24	Sensation of smell does not determine nutritional status in patients with cystic fibrosis. , 1997, 24, 52-56.		9
25	Characterization of a Marker for Tracheal Basal Cells. <i>Experimental Lung Research</i> , 1995, 21, 1-16.	1.2	7
26	Disposition of drugs in cystic fibrosis. V. In vivo CYP2C9 activity as probed by (S)-warfarin is not enhanced in cystic fibrosis. <i>Clinical Pharmacology and Therapeutics</i> , 1993, 54, 323-328.	4.7	20
27	Enrichment of Subpopulations of Respiratory Epithelial Cells Using Flow Cytometry. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 1991, 4, 174-178.	2.9	12