

James F Chmiel

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

42 papers	3,301 citations	27 h-index	45 g-index
45 ext. papers	4,132 ext. citations	8.5 avg, IF	5.15 L-index

#	Paper	IF	Citations
42	Testing the effects of combining azithromycin with inhaled tobramycin for in cystic fibrosis: a randomised, controlled clinical trial. <i>Thorax</i> , 2021 ,	7.3	4
41	Geography, generalisability, and susceptibility in clinical trials. <i>Lancet Respiratory Medicine</i> , 2021 , 9, 330-332	35.1	3
40	Safety and efficacy of lenabasum in a phase 2 randomized, placebo-controlled trial in adults with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2021 , 20, 78-85	4.1	6
39	Drug development for cystic fibrosis. <i>Pediatric Pulmonology</i> , 2021 , 56 Suppl 1, S10-S22	3.5	1
38	Somatic cell hemoglobin modulates nitrogen oxide metabolism in the human airway epithelium. <i>Scientific Reports</i> , 2021 , 11, 15498	4.9	0
37	Vitamin D for the Immune System in Cystic Fibrosis (DISC): a double-blind, multicenter, randomized, placebo-controlled clinical trial. <i>American Journal of Clinical Nutrition</i> , 2019 , 109, 544-553	7	10
36	Eradication of persistent methicillin-resistant <i>Staphylococcus aureus</i> infection in cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2019 , 18, 357-363	4.1	23
35	A phase 3 study of tezacaftor in combination with ivacaftor in children aged 6 through 11 years with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2019 , 18, 708-713	4.1	25
34	Efficacy and safety of the elxacaftor plus tezacaftor plus ivacaftor combination regimen in people with cystic fibrosis homozygous for the F508del mutation: a double-blind, randomised, phase 3 trial. <i>Lancet, The</i> , 2019 , 394, 1940-1948	40	341
33	Challenges in assessing the efficacy of systemic corticosteroids for severe wheezing episodes in preschool children. <i>Journal of Allergy and Clinical Immunology</i> , 2019 , 143, 1934-1937.e4	11.5	1
32	Airway Mucosal Host Defense Is Key to Genomic Regulation of Cystic Fibrosis Lung Disease Severity. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018 , 197, 79-93	10.2	27
31	Inflammation in cystic fibrosis: An update. <i>Pediatric Pulmonology</i> , 2018 , 53, S30-S50	3.5	106
30	Effects of endogenous sex hormones on lung function and symptom control in adolescents with asthma. <i>BMC Pulmonary Medicine</i> , 2018 , 18, 58	3.5	54
29	Azithromycin for Early <i>Pseudomonas</i> Infection in Cystic Fibrosis. The OPTIMIZE Randomized Trial. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018 , 198, 1177-1187	10.2	54
28	Preliminary comparison of normalized T1 and non-contrast perfusion MRI assessments of regional lung disease in cystic fibrosis patients. <i>Journal of Cystic Fibrosis</i> , 2017 , 16, 283-290	4.1	11
27	The Vitamin D for Enhancing the Immune System in Cystic Fibrosis (DISC) trial: Rationale and design of a multi-center, double-blind, placebo-controlled trial of high dose bolus administration of vitamin D3 during acute pulmonary exacerbation of cystic fibrosis. <i>Contemporary Clinical Trials Communications</i> , 2017 , 6, 39-45	1.8	8
26	Impact of azithromycin on the clinical and antimicrobial effectiveness of tobramycin in the treatment of cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2017 , 16, 358-366	4.1	37

25	Airway microbiota across age and disease spectrum in cystic fibrosis. <i>European Respiratory Journal</i> , 2017 , 50,	13.6	113
24	Bacterial infections in patients with primary ciliary dyskinesia: Comparison with cystic fibrosis. <i>Chronic Respiratory Disease</i> , 2017 , 14, 392-406	3	23
23	Interleukin-17 Pathophysiology and Therapeutic Intervention in Cystic Fibrosis Lung Infection and Inflammation. <i>Infection and Immunity</i> , 2016 , 84, 2410-21	3.7	29
22	Pathological Hyaluronan Matrices in Cystic Fibrosis Airways and Secretions. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2016 , 55, 576-585	5.7	16
21	Inhaled alpha1-proteinase inhibitor therapy in patients with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2016 , 15, 227-33	4.1	35
20	Neutrophils from F508del cystic fibrosis patients produce IL-17A and express IL-23 - dependent IL-17RC. <i>Clinical Immunology</i> , 2016 , 170, 53-60	9	23
19	Considerations for the Conduct of Clinical Trials with Antiinflammatory Agents in Cystic Fibrosis. A Cystic Fibrosis Foundation Workshop Report. <i>Annals of the American Thoracic Society</i> , 2015 , 12, 1398-406	4.7	28
18	Inflammation in cystic fibrosis lung disease: Pathogenesis and therapy. <i>Journal of Cystic Fibrosis</i> , 2015 , 14, 419-30	4.1	276
17	Inflammation and its genesis in cystic fibrosis. <i>Pediatric Pulmonology</i> , 2015 , 50 Suppl 40, S39-56	3.5	114
16	Use of ibuprofen to assess inflammatory biomarkers in induced sputum: Implications for clinical trials in cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2015 , 14, 720-6	4.1	43
15	Anti-PcrV antibody in cystic fibrosis: a novel approach targeting <i>Pseudomonas aeruginosa</i> airway infection. <i>Pediatric Pulmonology</i> , 2014 , 49, 650-8	3.5	73
14	Azithromycin may antagonize inhaled tobramycin when targeting <i>Pseudomonas aeruginosa</i> in cystic fibrosis. <i>Annals of the American Thoracic Society</i> , 2014 , 11, 342-50	4.7	53
13	Effect of vitamin D3 on asthma treatment failures in adults with symptomatic asthma and lower vitamin D levels: the VIDA randomized clinical trial. <i>JAMA - Journal of the American Medical Association</i> , 2014 , 311, 2083-91	27.4	179
12	Antibiotic and anti-inflammatory therapies for cystic fibrosis. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2013 , 3, a009779	5.4	40
11	Randomized trial of omalizumab (anti-IgE) for asthma in inner-city children. <i>New England Journal of Medicine</i> , 2011 , 364, 1005-15	59.2	647
10	Chronic inflammation in the cystic fibrosis lung: alterations in inter- and intracellular signaling. <i>Clinical Reviews in Allergy and Immunology</i> , 2008 , 34, 146-62	12.3	86
9	Anti-inflammatory therapies for cystic fibrosis-related lung disease. <i>Clinical Reviews in Allergy and Immunology</i> , 2008 , 35, 135-53	12.3	30
8	Sputum biomarkers of inflammation in cystic fibrosis lung disease. <i>Proceedings of the American Thoracic Society</i> , 2007 , 4, 406-17		122

7	Inflammation and anti-inflammatory therapies for cystic fibrosis. <i>Clinics in Chest Medicine</i> , 2007 , 28, 331-46	78
6	Anti-inflammatory medications for cystic fibrosis lung disease: selecting the most appropriate agent. <i>Treatments in Respiratory Medicine</i> , 2005 , 4, 255-73	27
5	Inflammatory and microbiologic markers in induced sputum after intravenous antibiotics in cystic fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2003 , 168, 1471-5	10.2 141
4	State of the art: why do the lungs of patients with cystic fibrosis become infected and why can't they clear the infection?. <i>Respiratory Research</i> , 2003 , 4, 8	7.3 154
3	The role of inflammation in the pathophysiology of CF lung disease. <i>Clinical Reviews in Allergy and Immunology</i> , 2002 , 23, 5-27	12.3 180
2	Prolonged inflammatory response to acute <i>Pseudomonas</i> challenge in interleukin-10 knockout mice. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2002 , 165, 1176-81	10.2 65
1	Murine models of CF airway infection and inflammation. <i>Methods in Molecular Medicine</i> , 2002 , 70, 495-515	7