

Exacerbation frequency and clinical outcomes in adult p

Thorax

66, 680-685

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Sex steroid receptors in human lung diseases. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2011, 127, 216-222.	1.2	37
2	Cystic fibrosis papers of the year 2010-2011. <i>Journal of the Royal Society of Medicine</i> , 2012, 105, 30-35.	1.1	1
3	Effect of pulmonary exacerbations on long-term lung function decline in cystic fibrosis. <i>European Respiratory Journal</i> , 2012, 40, 61-66.	3.1	201
4	The Adult Cystic Fibrosis Airway Microbiota Is Stable over Time and Infection Type, and Highly Resilient to Antibiotic Treatment of Exacerbations. <i>PLoS ONE</i> , 2012, 7, e45001.	1.1	320
5	Iron Homeostasis during Cystic Fibrosis Pulmonary Exacerbation. <i>Clinical and Translational Science</i> , 2012, 5, 368-373.	1.5	29
6	Fluoroquinolones in the treatment of bronchopulmonary disease in cystic fibrosis. <i>Therapeutic Advances in Respiratory Disease</i> , 2012, 6, 363-373.	1.0	7
7	Viral infections trigger exacerbations of cystic fibrosis in adults and children: Figure 1â€“. <i>European Respiratory Journal</i> , 2012, 40, 510-512.	3.1	67
8	I have taken my umbrella, so of course it does not rain: Figure 1. <i>Thorax</i> , 2012, 67, 88-89.	2.7	31
9	<i>Houttuynia cordata</i> inhibits lipopolysaccharide-induced rapid pulmonary fibrosis by up-regulating IFN- β and inhibiting the TGF- β 1/Smad pathway. <i>International Immunopharmacology</i> , 2012, 13, 331-340.	1.7	35
10	Inhaled antibiotics for pulmonary exacerbations in cystic fibrosis. <i>The Cochrane Library</i> , 2012, 12, CD008319.	1.5	21
11	Recent Advances in Cystic Fibrosis. <i>Clinics in Chest Medicine</i> , 2012, 33, 307-328.	0.8	20
12	Pilot study of vitamin D supplementation in adults with cystic fibrosis pulmonary exacerbation. <i>Dermato-Endocrinology</i> , 2012, 4, 191-197.	1.9	74
13	Treatment of <i>Aspergillus fumigatus</i> in Patients with Cystic Fibrosis: A Randomized, Placebo-Controlled Pilot Study. <i>PLoS ONE</i> , 2012, 7, e36077.	1.1	72
15	The Prognosis of Cystic Fibrosis - A Clinician's Perspective. , 2012, , .		1
16	Impact of exacerbations of cystic fibrosis on muscle strength. <i>Respiratory Research</i> , 2013, 14, 46.	1.4	37
17	A pulmonary exacerbation risk score among cystic fibrosis patients not receiving recommended care. <i>Pediatric Pulmonology</i> , 2013, 48, 954-961.	1.0	13
18	Cystic Fibrosis Papers of the Year 2012. <i>Paediatric Respiratory Reviews</i> , 2013, 14, 28-30.	1.2	3
19	Impact of acute antibiotic therapy on the pulmonary exacerbation endpoint in cystic fibrosis clinical trials. <i>Contemporary Clinical Trials</i> , 2013, 36, 99-105.	0.8	8

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20	Molecular analysis of changes in <i>Pseudomonas aeruginosa</i> load during treatment of a pulmonary exacerbation in cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2013, 12, 688-699.	0.3	21
21	Oral, inhaled, and intravenous antibiotic choice for treating pulmonary exacerbations in cystic fibrosis. <i>Pediatric Pulmonology</i> , 2013, 48, 666-673.	1.0	99
22	New agents to treat lung infection in cystic fibrosis: a big enough leap?. <i>Future Medicinal Chemistry</i> , 2013, 5, 117-120.	1.1	0
23	Prospective evaluation of respiratory exacerbations in children with cystic fibrosis from newborn screening to 5 years of age. <i>Thorax</i> , 2013, 68, 643-651.	2.7	83
24	Quality of life and healthcare utilisation in cystic fibrosis: a multicentre study. <i>European Respiratory Journal</i> , 2013, 41, 571-577.	3.1	45
25	Effect of residential proximity to major roadways on cystic fibrosis exacerbations. <i>International Journal of Environmental Health Research</i> , 2013, 23, 119-131.	1.3	19
26	Optimization of anti-pseudomonal antibiotics for cystic fibrosis pulmonary exacerbations: VI. Executive summary. <i>Pediatric Pulmonology</i> , 2013, 48, 525-537.	1.0	27
27	The CF-ABLE Score. <i>Chest</i> , 2013, 143, 1358-1364.	0.4	33
28	Ozone Is Associated With an Increased Risk of Respiratory Exacerbations in Patients With Cystic Fibrosis. <i>Chest</i> , 2013, 144, 1186-1192.	0.4	41
29	Peripheral Monocytes Derived from Patients with Cystic Fibrosis and Healthy Donors Secrete NGAL in Response to <i>Pseudomonas aeruginosa</i> Infection. <i>Journal of Investigative Medicine</i> , 2013, 61, 1018-1025.	0.7	19
30	Role of C-reactive protein as a biomarker for prediction of the severity of pulmonary exacerbations in patients with cystic fibrosis. <i>BMC Pulmonary Medicine</i> , 2014, 14, 150.	0.8	16
31	Gender Differences in Outcomes of Patients with Cystic Fibrosis. <i>Journal of Women's Health</i> , 2014, 23, 1012-1020.	1.5	138
32	Year in review 2013: paediatric and adult clinical studies. <i>Thorax</i> , 2014, 69, 309-311.	2.7	2
33	Vitamin D Deficiency Is Associated with Pulmonary Exacerbations in Children with Cystic Fibrosis. <i>Annals of the American Thoracic Society</i> , 2014, 11, 198-204.	1.5	63
34	Effects of puberty on cystic fibrosis related pulmonary exacerbations in women versus men. <i>Pediatric Pulmonology</i> , 2014, 49, 28-35.	1.0	41
35	Incidence and clinical impact of respiratory viruses in adults with cystic fibrosis. <i>Thorax</i> , 2014, 69, 247-253.	2.7	107
36	Aerobic fitness is associated with lower risk of hospitalization in children with cystic fibrosis. <i>Pediatric Pulmonology</i> , 2014, 49, 641-649.	1.0	67
37	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.	1.0	21

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38	Early Respiratory Infection Is Associated with Reduced Spirometry in Children with Cystic Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014, 190, 1111-1116.	2.5	142
39	Inhaled versus nebulised tobramycin: A real world comparison in adult cystic fibrosis (CF). <i>Journal of Cystic Fibrosis</i> , 2014, 13, 692-698.	0.3	54
41	Survival in Cystic Fibrosis: Trends, Clinical Factors, and Prediction Models. <i>Pediatric, Allergy, Immunology, and Pulmonology</i> , 2015, 28, 244-249.	0.3	4
42	Managing <i>Pseudomonas aeruginosa</i> respiratory infections in cystic fibrosis. <i>Current Opinion in Infectious Diseases</i> , 2015, 28, 547-556.	1.3	67
43	Respiratory tract exacerbations revisited: Ventilation, inflammation, perfusion, and structure (VIPS) monitoring to redefine treatment. <i>Pediatric Pulmonology</i> , 2015, 50, S57-65.	1.0	29
44	Intravenous antibiotics for pulmonary exacerbations in people with cystic fibrosis. <i>The Cochrane Library</i> , 2015, , CD009730.	1.5	19
45	Relationship between pulmonary exacerbations and daily physical activity in adults with cystic fibrosis. <i>BMC Pulmonary Medicine</i> , 2015, 15, 151.	0.8	27
46	Variation in lung function is associated with worse clinical outcomes in cystic fibrosis. <i>Jornal Brasileiro De Pneumologia</i> , 2015, 41, 509-515.	0.4	8
47	A statistical model to predict one-year risk of death in patients with cystic fibrosis. <i>Journal of Clinical Epidemiology</i> , 2015, 68, 1336-1345.	2.4	65
48	Mucoviscidose. <i>Revue Des Maladies Respiratoires Actualites</i> , 2015, 7, S12-S22.	0.0	0
49	Acute effects of viral respiratory tract infections on sputum bacterial density during CF pulmonary exacerbations. <i>Journal of Cystic Fibrosis</i> , 2015, 14, 482-489.	0.3	30
50	A Winogradsky-based culture system shows an association between microbial fermentation and cystic fibrosis exacerbation. <i>ISME Journal</i> , 2015, 9, 1024-1038.	4.4	59
51	Management of pulmonary exacerbations in cystic fibrosis: still an unmet medical need in clinical practice. <i>Expert Review of Respiratory Medicine</i> , 2015, 9, 183-194.	1.0	14
52	Factors associated with response to treatment of pulmonary exacerbations in cystic fibrosis patients. <i>Journal of Cystic Fibrosis</i> , 2015, 14, 755-762.	0.3	62
53	Cystic Fibrosis from Laboratory to Bedside: The Role of A20 in NF- κ B-Mediated Inflammation. <i>Medical Principles and Practice</i> , 2015, 24, 301-310.	1.1	12
54	Treatment and demographic factors affecting time to next pulmonary exacerbation in cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2015, 14, 763-769.	0.3	38
55	Rehabilitation in Patients with Chronic Respiratory Disease Other than Chronic Obstructive Pulmonary Disease: Exercise and Physical Activity Interventions in Cystic Fibrosis and Non-Cystic Fibrosis Bronchiectasis. <i>Respiration</i> , 2015, 89, 181-189.	1.2	48
56	Infectious Diseases Pharmacotherapy for Children With Cystic Fibrosis. <i>Journal of Pediatric Health Care</i> , 2015, 29, 565-578.	0.6	10

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57	Intravenous antibiotics for pulmonary exacerbations in people with cystic fibrosis. Paediatric Respiratory Reviews, 2015, 16, 246-248.	1.2	5
58	A contemporary survival analysis of individuals with cystic fibrosis: a cohort study. European Respiratory Journal, 2015, 45, 670-679.	3.1	154
59	Indicators of pulmonary exacerbation in cystic fibrosis: A Delphi survey of patients and health professionals. Journal of Cystic Fibrosis, 2015, 14, 90-96.	0.3	10
60	C-Reactive Protein in Stable Cystic Fibrosis: An Additional Indicator of Clinical Disease Activity and Risk of Future Pulmonary Exacerbations. Journal of Pulmonary & Respiratory Medicine, 2016, 6, 1000375.	0.1	19
61	Accurate reporting of adherence to inhaled therapies in adults with cystic fibrosis: methods to calculate normative adherence. Patient Preference and Adherence, 2016, 10, 887.	0.8	20
62	Aspergillus fumigatus in the cystic fibrosis lung: pros and cons of azole therapy. Infection and Drug Resistance, 2016, Volume 9, 229-238.	1.1	53
63	Recovery of baseline lung function after pulmonary exacerbation in children with primary ciliary dyskinesia. Pediatric Pulmonology, 2016, 51, 1362-1366.	1.0	31
64	Asthma attacks: should we nail our colours to the mast (cell)? European Respiratory Journal, 2016, 48, 1261-1264.	3.1	2
65	Cystic Fibrosis-Related Diabetes with strict glycaemic control is not associated with frequent intravenous antibiotics use for pulmonary infections. Diabetes Research and Clinical Practice, 2016, 116, 230-236.	1.1	6
66	IV-treated pulmonary exacerbations in the prior year: An important independent risk factor for future pulmonary exacerbation in cystic fibrosis. Journal of Cystic Fibrosis, 2016, 15, 372-379.	0.3	35
67	Healthcare resource utilization associated with ivacaftor use in patients with cystic fibrosis. Journal of Medical Economics, 2016, 19, 845-851.	1.0	13
68	Cystic fibrosis. Lancet, The, 2016, 388, 2519-2531.	6.3	1,309
69	<i>Pseudomonas aeruginosa</i> infection in cystic fibrosis: pathophysiological mechanisms and therapeutic approaches. Expert Review of Respiratory Medicine, 2016, 10, 685-697.	1.0	114
70	Constrictive Bronchiolitis in Cystic Fibrosis Adolescents with Refractory Pulmonary Decline. Annals of the American Thoracic Society, 2016, 13, 2174-2183.	1.5	8
71	Special considerations for the treatment of pulmonary exacerbations in children with cystic fibrosis. Expert Review of Respiratory Medicine, 2016, 10, 1221-1228.	1.0	5
72	Pulmonary artery enlargement and cystic fibrosis pulmonary exacerbations: a cohort study. Lancet Respiratory Medicine, the, 2016, 4, 636-645.	5.2	19
73	Advance care planning in adolescents with cystic fibrosis: A quality improvement project. Pediatric Pulmonology, 2016, 51, 1304-1310.	1.0	20
74	Ecological networking of cystic fibrosis lung infections. Npj Biofilms and Microbiomes, 2016, 2, 4.	2.9	77

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75	Impact of pulmonary exacerbations and lung function on generic health-related quality of life in patients with cystic fibrosis. <i>Health and Quality of Life Outcomes</i> , 2016, 14, 63.	1.0	37
76	A longitudinal study characterising a large adult primary ciliary dyskinesia population. <i>European Respiratory Journal</i> , 2016, 48, 441-450.	3.1	101
77	Expiratory Flow Limitation for Monitoring Cystic Fibrosis. Ready for the Starting Gun?. <i>Annals of the American Thoracic Society</i> , 2016, 13, 770-771.	1.5	0
78	Optimising treatment of CF pulmonary exacerbation: a tough nut to crack. <i>Thorax</i> , 2016, 71, 101-102.	2.7	5
79	Effect of Media Modified To Mimic Cystic Fibrosis Sputum on the Susceptibility of <i>Aspergillus fumigatus</i> , and the Frequency of Resistance at One Center. <i>Antimicrobial Agents and Chemotherapy</i> , 2016, 60, 2180-2184.	1.4	16
80	A randomised trial of hypertonic saline during hospitalisation for exacerbation of cystic fibrosis. <i>Thorax</i> , 2016, 71, 141-147.	2.7	40
81	Short-term and long-term response to pulmonary exacerbation treatment in cystic fibrosis. <i>Thorax</i> , 2016, 71, 223-229.	2.7	53
82	The diagnosis and management of respiratory viral infections in cystic fibrosis. <i>Expert Review of Respiratory Medicine</i> , 2017, 11, 221-227.	1.0	23
83	Six-Minute Walk Test Results Predict Risk of Hospitalization for Youths with Cystic Fibrosis: A 5-Year Follow-Up Study. <i>Journal of Pediatrics</i> , 2017, 182, 204-209.e1.	0.9	28
84	Early Lung Function Decline in Cystic Fibrosis. Can Registry Data Explain Divergent Phenotypes?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2017, 196, 407-409.	2.5	1
85	Standardized Treatment of Pulmonary Exacerbations (STOP) study: Physician treatment practices and outcomes for individuals with cystic fibrosis with pulmonary Exacerbations. <i>Journal of Cystic Fibrosis</i> , 2017, 16, 600-606.	0.3	76
86	A treatment evaluator tool to monitor the real-world effectiveness of inhaled aztreonam lysine in cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2017, 16, 695-701.	0.3	4
87	The treatment of the pulmonary and extrapulmonary manifestations of cystic fibrosis. <i>Presse Medicale</i> , 2017, 46, e139-e164.	0.8	12
88	Effect of Anaerobiasis or Hypoxia on <i>Pseudomonas aeruginosa</i> Inhibition of <i>Aspergillus fumigatus</i> Biofilm. <i>Archives of Microbiology</i> , 2017, 199, 881-890.	1.0	21
89	Association between glucose intolerance and bacterial colonisation in an adult population with cystic fibrosis, emergence of <i>Stenotrophomonas maltophilia</i> . <i>Journal of Cystic Fibrosis</i> , 2017, 16, 418-424.	0.3	13
90	Effect of pulmonary exacerbations treated with oral antibiotics on clinical outcomes in cystic fibrosis. <i>Thorax</i> , 2017, 72, 327-332.	2.7	58
91	Assessment of safety and efficacy of long-term treatment with combination lumacaftor and ivacaftor therapy in patients with cystic fibrosis homozygous for the F508del-CFTR mutation (PROGRESS): a phase 3, extension study. <i>Lancet Respiratory Medicine</i> , 2017, 5, 107-118.	5.2	235
92	A first step to STOP cystic fibrosis exacerbations. <i>Journal of Cystic Fibrosis</i> , 2017, 16, 529-531.	0.3	7

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93	CF-related diabetes: Containing the metabolic miscreant of cystic fibrosis. <i>Pediatric Pulmonology</i> , 2017, 52, S37-S43.	1.0	43
94	Adverse events following live-attenuated intranasal influenza vaccination of children with cystic fibrosis: Results from two influenza seasons. <i>Vaccine</i> , 2017, 35, 5019-5026.	1.7	1
95	Tezacaftor-ivacaftor in Patients with Cystic Fibrosis Homozygous for Phe508del. <i>New England Journal of Medicine</i> , 2017, 377, 2013-2023.	13.9	625
96	A prospective pilot study of home monitoring in adults with cystic fibrosis (HOME-CF): protocol for a randomised controlled trial. <i>BMC Pulmonary Medicine</i> , 2017, 17, 22.	0.8	19
97	Microbiome in the pathogenesis of cystic fibrosis and lung transplant-related disease. <i>Translational Research</i> , 2017, 179, 84-96.	2.2	29
98	Antibiotic duration and changes in FEV1 are not associated with time until next exacerbation in adult cystic fibrosis: a single center study. <i>BMC Pulmonary Medicine</i> , 2017, 17, 160.	0.8	5
99	Antibiotic perturbation of mixed-strain <i>Pseudomonas aeruginosa</i> infection in patients with cystic fibrosis. <i>BMC Pulmonary Medicine</i> , 2017, 17, 138.	0.8	11
100	Nationwide trends of hospitalizations for cystic fibrosis in the United States from 2003 to 2013. <i>Intractable and Rare Diseases Research</i> , 2017, 6, 191-198.	0.3	24
101	A smartphone application for reporting symptoms in adults with cystic fibrosis: protocol of a randomised controlled trial. <i>BMJ Open</i> , 2018, 8, e021136.	0.8	6
102	The expression of Mir1/Mir17-92 cluster in sputum samples correlates with pulmonary exacerbations in cystic fibrosis patients. <i>Journal of Cystic Fibrosis</i> , 2018, 17, 454-461.	0.3	24
103	Study design considerations for the Standardized Treatment of Pulmonary Exacerbations 2 (STOP2): A trial to compare intravenous antibiotic treatment durations in CF. <i>Contemporary Clinical Trials</i> , 2018, 64, 35-40.	0.8	42
104	Pharmacological management of cystic fibrosis related diabetes. <i>Expert Review of Clinical Pharmacology</i> , 2018, 11, 185-191.	1.3	13
105	"We can't diagnose asthma until <insert arbitrary age>". <i>Archives of Disease in Childhood</i> , 2018, 103, 729-731.	1.0	10
106	Studies of <i>Pseudomonas aeruginosa</i> Mutants Indicate Pyoverdine as the Central Factor in Inhibition of <i>Aspergillus fumigatus</i> Biofilm. <i>Journal of Bacteriology</i> , 2018, 200, .	1.0	99
107	<i>Aspergillus</i> Bronchitis in Patients with Cystic Fibrosis. <i>Mycopathologia</i> , 2018, 183, 61-69.	1.3	65
108	Multiple reaction monitoring mass spectrometry to identify novel plasma protein biomarkers of treatment response in cystic fibrosis pulmonary exacerbations. <i>Journal of Cystic Fibrosis</i> , 2018, 17, 333-340.	0.3	9
109	Daily spirometry in an acute exacerbation of adult cystic fibrosis patients. <i>Chronic Respiratory Disease</i> , 2018, 15, 258-264.	1.0	4
110	Fungal Pathogens in CF Airways: Leave or Treat?. <i>Mycopathologia</i> , 2018, 183, 119-137.	1.3	32

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111	Niche partitioning of a pathogenic microbiome driven by chemical gradients. <i>Science Advances</i> , 2018, 4, eaau1908.	4.7	40
112	ASSOCIAÇÃO DO ESTADO NUTRICIONAL COM FUNÇÃO PULMONAR E MORBIDADE EM CRIANÇAS E ADOLESCENTES COM FIBROSE CÍSTICA: COORTE DE 36 MESES. <i>Revista Paulista De Pediatria</i> , 2018, 36, 31-38.	0.4	8
113	Unmet needs in cystic fibrosis: the next steps in improving outcomes. <i>Expert Review of Respiratory Medicine</i> , 2018, 12, 585-593.	1.0	17
114	Course of Illness after Viral Infection in Indian Children with Cystic Fibrosis. <i>Journal of Tropical Pediatrics</i> , 2019, 65, 176-182.	0.7	6
115	Adult patients'™ experiences of symptom management during pulmonary exacerbations in cystic fibrosis: A thematic synthesis of qualitative research. <i>Chronic Illness</i> , 2019, 15, 245-263.	0.6	5
116	Long-Term Consequences of Childhood Respiratory Disease. , 2019, , 247-256.e4.		0
117	Lumacaftor/Ivacaftor reduces pulmonary exacerbations in patients irrespective of initial changes in FEV1. <i>Journal of Cystic Fibrosis</i> , 2019, 18, 94-101.	0.3	36
118	“Thrust out of normality” How adults living with cystic fibrosis experience pulmonary exacerbations: A qualitative study. <i>Journal of Clinical Nursing</i> , 2019, 28, 190-200.	1.4	10
119	Elexacaftor“Tezacaftor“Ivacaftor for Cystic Fibrosis with a Single Phe508del Allele. <i>New England Journal of Medicine</i> , 2019, 381, 1809-1819.	13.9	1,231
120	Intermicrobial interaction: <i>Aspergillus fumigatus</i> siderophores protect against competition by <i>Pseudomonas aeruginosa</i> . <i>PLoS ONE</i> , 2019, 14, e0216085.	1.1	53
121	The Role of HMGB1, a Nuclear Damage-Associated Molecular Pattern Molecule, in the Pathogenesis of Lung Diseases. <i>Antioxidants and Redox Signaling</i> , 2019, 31, 954-993.	2.5	50
122	Lung transplant referral for individuals with cystic fibrosis: Cystic Fibrosis Foundation consensus guidelines. <i>Journal of Cystic Fibrosis</i> , 2019, 18, 321-333.	0.3	154
123	Oral Azithromycin Use and the Recovery of Lung Function from Pulmonary Exacerbations Treated with Intravenous Tobramycin or Colistimethate in Adults with Cystic Fibrosis. <i>Annals of the American Thoracic Society</i> , 2019, 16, 853-860.	1.5	12
124	New insights into exogenous surfactant as a carrier of pulmonary therapeutics. <i>Biochemical Pharmacology</i> , 2019, 164, 64-73.	2.0	30
125	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.	2.2	27
126	Adult Care in Cystic Fibrosis. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2019, 40, 857-868.	0.8	8
127	Caregiver Burden Due to Pulmonary Exacerbations in Patients with Cystic Fibrosis. <i>Journal of Pediatrics</i> , 2019, 215, 164-171.e2.	0.9	16
128	Cystic Fibrosis: Emerging Understanding and Therapies. <i>Annual Review of Medicine</i> , 2019, 70, 197-210.	5.0	39

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129	Application of multiple event analysis as an alternative approach to studying pulmonary exacerbations as an outcome measure. <i>Journal of Cystic Fibrosis</i> , 2020, 19, 114-118.	0.3	7
130	A smartphone application for reporting symptoms in adults with cystic fibrosis improves the detection of exacerbations: Results of a randomised controlled trial. <i>Journal of Cystic Fibrosis</i> , 2020, 19, 271-276.	0.3	23
131	Real-Life Safety and Effectiveness of Lumacaftor/ivacaftor in Patients with Cystic Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 201, 188-197.	2.5	95
132	Lumacaftor/ivacaftor reduces exacerbations in adults homozygous for Phe508del mutation with severe lung disease. <i>Journal of Cystic Fibrosis</i> , 2020, 19, 415-420.	0.3	15
133	Extracellular DNA in sputum is associated with pulmonary function and hospitalization in patients with cystic fibrosis. <i>Respiratory Medicine</i> , 2020, 172, 106144.	1.3	15
134	Impact of Socioeconomic Position on Access to the U.S. Lung Transplant Waiting List in a Matched Cystic Fibrosis Cohort. <i>Annals of the American Thoracic Society</i> , 2020, 17, 1384-1392.	1.5	19
135	Timing it right: the challenge of recipient selection for lung transplantation. <i>Annals of Translational Medicine</i> , 2020, 8, 408-408.	0.7	4
136	CFTR modulator therapies – Effect on life expectancy in people with cystic fibrosis. <i>Paediatric Respiratory Reviews</i> , 2022, 42, 3-8.	1.2	26
137	Optimizing Exogenous Surfactant as a Pulmonary Delivery Vehicle for Chicken Cathelicidin-2. <i>Scientific Reports</i> , 2020, 10, 9392.	1.6	5
138	Review of Potential Pseudomonas Weaponry, Relevant to the Pseudomonas/Aspergillus Interplay, for the Mycology Community. <i>Journal of Fungi (Basel, Switzerland)</i> , 2020, 6, 81.	1.5	32
139	Procalcitonin Predicts the Severity of Cystic Fibrosis Pulmonary Exacerbations and Readmissions in Adult Patients: A Prospective Cohort Study. <i>Journal of Investigative Medicine</i> , 2020, 68, 856-863.	0.7	5
140	Complete versus Limited Endoscopic Sinus Surgery for Chronic Rhinosinusitis in Adults with Cystic Fibrosis. <i>Otolaryngology - Head and Neck Surgery</i> , 2020, 162, 572-580.	1.1	6
141	Enhancing care for individuals with advanced cystic fibrosis lung disease. <i>Pediatric Pulmonology</i> , 2021, 56, S69-S78.	1.0	0
142	Impulse oscillometry and spirometry measurements relative to personal best values at the time of acute exacerbations of cystic fibrosis in adults. <i>Clinical Physiology and Functional Imaging</i> , 2021, 41, 76-84.	0.5	2
143	Lung Clearance Index to Track Acute Respiratory Events in School-Age Children with Cystic Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021, 203, 977-986.	2.5	34
144	Acute hyperglycaemia in cystic fibrosis pulmonary exacerbations. <i>Endocrinology, Diabetes and Metabolism</i> , 2021, 4, e00208.	1.0	4
145	Under nonlimiting iron conditions pyocyanin is a major antifungal molecule, and differences between prototypic <i>Pseudomonas aeruginosa</i> strains. <i>Medical Mycology</i> , 2021, 59, 453-464.	0.3	16
146	Antibiotics in Adult Cystic Fibrosis Patients: A Review of Population Pharmacokinetic Analyses. <i>Clinical Pharmacokinetics</i> , 2021, 60, 447-470.	1.6	5

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147	Reflections on the Importance of Cost of Illness Analysis in Rare Diseases: A Proposal. <i>International Journal of Environmental Research and Public Health</i> , 2021, 18, 1101.	1.2	8
148	Long-term safety and efficacy of tezacaftor-ivacaftor in individuals with cystic fibrosis aged 12 years or older who are homozygous or heterozygous for Phe508del CFTR (EXTEND): an open-label extension study. <i>Lancet Respiratory Medicine</i> , 2021, 9, 733-746.	5.2	33
149	Cost-effectiveness of implementing routine hearing screening using a tablet audiometer for pediatric cystic fibrosis patients receiving high-dose IV aminoglycosides. <i>Journal of Managed Care & Specialty Pharmacy</i> , 2021, 27, 157-165.	0.5	0
150	Early Interleukin-22 and Neutrophil Proteins Are Correlated to Future Lung Damage in Children With Cystic Fibrosis. <i>Frontiers in Pediatrics</i> , 2021, 9, 640184.	0.9	4
151	A pilot study of cystic fibrosis exacerbation response phenotypes reveals contrasting serum and sputum iron trends. <i>Scientific Reports</i> , 2021, 11, 4897.	1.6	3
152	SPLUNC1: a novel marker of cystic fibrosis exacerbations. <i>European Respiratory Journal</i> , 2021, 58, 2000507.	3.1	20
153	Sex differences in treatment patterns in cystic fibrosis pulmonary exacerbations. <i>Journal of Cystic Fibrosis</i> , 2021, 20, 920-925.	0.3	10
154	Clinical impact of levofloxacin inhalation solution in cystic fibrosis patients in a real-world setting. <i>Journal of Cystic Fibrosis</i> , 2021, 20, 1035-1039.	0.3	7
155	Consensus document for the selection of lung transplant candidates: An update from the International Society for Heart and Lung Transplantation. <i>Journal of Heart and Lung Transplantation</i> , 2021, 40, 1349-1379.	0.3	293
156	Disease burden in people with cystic fibrosis heterozygous for F508del and a minimal function mutation. <i>Journal of Cystic Fibrosis</i> , 2022, 21, 96-103.	0.3	2
157	Variation in treatment preferences of pulmonary exacerbations among Australian and New Zealand cystic fibrosis physicians. <i>BMJ Open Respiratory Research</i> , 2021, 8, e000956.	1.2	4
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