Alan R Smyth

List of Publications by Year
in descending order

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190 papers	8,821 citations	61984 43 h-index	49909 87 g-index
206 all docs	206 docs citations	206 times ranked	9117 citing authors

#	Article	IF	Citations
1	Daily Bread: Womenâ€™s Self-Help Microfinance and the Social Meanings of Money. Sociological Research Online, 2023, 28, 442-461.	1.1	0
2	Parenchymal lung abnormalities following hospitalisation for COVID-19 and viral pneumonitis: a systematic review and meta-analysis. Thorax, 2023, 78, 191-201.	5.6	49
3	Novel detection of specific bacterial quorum sensing molecules in saliva: Potential non-invasive biomarkers for pulmonary Pseudomonas aeruginosa in cystic fibrosis. Journal of Cystic Fibrosis, 2022, 21, 626-629.	0.7	6
4	Exploring the challenges of accessing medication for patients with cystic fibrosis. Thorax, 2022, 77, 295-297.	5.6	1
5	Cystic Fibrosis Therapies. , 2022, , 179-187.		0
6	Preferred health outcome states following treatment for pulmonary exacerbations of cystic fibrosis. Journal of Cystic Fibrosis, 2022, 21, 581-587.	0.7	4
7	Perspectives of patients, family members, health professionals and the public on the impact of COVID-19 on mental health. Journal of Mental Health, 2022, 31, 524-533.	1.9	2
8	Porphyromonas pasteri and Prevotella nanceiensis in the sputum microbiota are associated with increased decline in lung function in individuals with cystic fibrosis. Journal of Medical Microbiology, 2022, 71, .	1.8	9
9	Intestinal function and transit associate with gut microbiota dysbiosis in cystic fibrosis. Journal of Cystic Fibrosis, 2022, 21, 506-513.	0.7	16
10	Magnetic resonance imaging of the gastrointestinal tract shows reduced small bowel motility and altered chyme in cystic fibrosis compared to controls. Journal of Cystic Fibrosis, 2022, 21, 502-505.	0.7	12
11	Industry influence in healthcare harms patients: myth or maxim?. Breathe, 2022, 18, 220010.	1.3	3
12	Wheeze in the time of COVID-19: overcoming obstacles to an unusual diagnosis. Thorax, 2022, 77, 1050-1053.	5.6	0
13	Postprandial changes in gastrointestinal function and transit in cystic fibrosis assessed by Magnetic Resonance Imaging. Journal of Cystic Fibrosis, 2021, 20, 591-597.	0.7	29
14	Antimicrobial resistance: Concerns of healthcare providers and people with CF. Journal of Cystic Fibrosis, 2021, 20, 407-412.	0.7	13
15	Core Outcome Measures for Trials in People With Coronavirus Disease 2019: Respiratory Failure, Multiorgan Failure, Shortness of Breath, and Recovery. Critical Care Medicine, 2021, 49, 503-516.	0.9	41
16	An ex vivo cystic fibrosis model recapitulates key clinical aspects of chronic Staphylococcus aureus infection. Microbiology (United Kingdom), 2021, 167, .	1.8	12
17	Systematic review and meta-analysis of anakinra, sarilumab, siltuximab and tocilizumab for COVID-19. Thorax, 2021, 76, 907-919.	5.6	90
18	Time to get serious about the detection and monitoring of early lung disease in cystic fibrosis. Thorax, 2021, 76, 1255-1265.	5.6	24

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19	The measurement properties of tests and tools used in cystic fibrosis studies: a systematic review. European Respiratory Review, 2021, 30, 200354.	7.1	12
20	Professor Pangloss and the Pangenome: Does Staphylococcus aureus Have the Best of All Possible Worlds?. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 1055-1057.	5.6	0
21	Timing of pancreatic enzyme replacement therapy (PERT) in cystic fibrosis. The Cochrane Library, 2021, 2021, CD013488.	2.8	4
22	Telehealth after the pandemic: Will the inverse care law apply? (Commentary). Journal of Cystic Fibrosis, 2021, 20, 47-48.	0.7	7
23	A systematic cochrane review of the timing of pancreatic enzyme replacement therapy (PERT) in cystic fibrosis. Paediatric Respiratory Reviews, 2021, 40, 44-45.	1.8	0
24	2-Alkyl-4-quinolone quorum sensing molecules are biomarkers for culture-independent Pseudomonas aeruginosa burden in adults with cystic fibrosis. Journal of Medical Microbiology, 2021, 70, .	1.8	5
25	Novel method to select meaningful outcomes for evaluation in clinical trials. BMJ Open Respiratory Research, 2021, 8, e000877.	3.0	4
26	Intravenous or oral antibiotic treatment in adults and children with cystic fibrosis and Pseudomonas aeruginosa infection: the TORPEDO-CF RCT. Health Technology Assessment, 2021, 25, 1-128.	2.8	6
27	Antibiotic strategies for eradicating Pseudomonas aeruginosa in people with cystic fibrosis. The Cochrane Library, 2020, 2020, CD004197.	2.8	79
28	Do current clinical trials in cystic fibrosis match the priorities of patients and clinicans? A systematic review. Journal of Cystic Fibrosis, 2020, 19, 26-33.	0.7	9
29	Can exercise replace airway clearance techniques in cystic fibrosis? A survey of patients and healthcare professionals. Journal of Cystic Fibrosis, 2020, 19, e19-e24.	0.7	17
30	Infection prevention and control in cystic fibrosis: One size fits all The argument against. Paediatric Respiratory Reviews, 2020, 36, 94-96.	1.8	4
31	Using digital technology for home monitoring, adherence and self-management in cystic fibrosis: a state-of-the-art review. Thorax, 2020, 75, 72-77.	5.6	35
32	Characterising burden of treatment in cystic fibrosis to identify priority areas for clinical trials. Journal of Cystic Fibrosis, 2020, 19, 499-502.	0.7	60
33	Treatment of pulmonary exacerbations in cystic fibrosis. Current Opinion in Pulmonary Medicine, 2020, 26, 679-684.	2.6	11
34	International Survey to Establish Prioritized Outcomes for Trials in People With Coronavirus Disease 2019. Critical Care Medicine, 2020, 48, 1612-1621.	0.9	12
35	Intravenous versus oral antibiotics for eradication of Pseudomonas aeruginosa in cystic fibrosis (TORPEDO-CF): a randomised controlled trial. Lancet Respiratory Medicine,the, 2020, 8, 975-986.	10.7	38
36	Antibiotic adjuvant therapy for pulmonary infection in cystic fibrosis. The Cochrane Library, 2020, 2020, CD008037.	2.8	2

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37	Core Outcomes Set for Trials in People With Coronavirus Disease 2019. Critical Care Medicine, 2020, 48, 1622-1635.	0.9	47
38	Outcomes and endpoints reported in studies of pulmonary exacerbations in people with cystic fibrosis: A systematic review. Journal of Cystic Fibrosis, 2020, 19, 858-867.	0.7	13
39	Response to Journal Club: Cluster Randomized Trial Evaluating Impact of a Community-based Microfinance Scheme on Childhood Nutritional Status: Evidence-based Medicine Viewpoint. Indian Pediatrics, 2020, 57, 688-690.	0.4	0
40	A systematic Cochrane Review of antibiotic adjuvant therapy for pulmonary infection in cystic fibrosis. Paediatric Respiratory Reviews, 2020, 36, 109-111.	1.8	0
41	What effective ways of motivation, support and technologies help people with cystic fibrosis improve and sustain adherence to treatment?. BMJ Open Respiratory Research, 2020, 7, e000601.	3.0	17
42	How can we relieve gastrointestinal symptoms in people with cystic fibrosis? An international qualitative survey. BMJ Open Respiratory Research, 2020, 7, e000614.	3.0	22
43	Rojiroti microfinance and child nutrition: a cluster randomised trial. Archives of Disease in Childhood, 2020, 105, 229-235.	1.9	13
44	Smoking ban in cars protects children, but is vaping â€˜The Elephant in the Carâ€™?. Thorax, 2020, 75, 297-297.	5.6	2
45	Preprint servers: a â€˜rush to publishâ€™ or â€˜just in time deliveryâ€™ for science?. Thorax, 2020, 75, 532-533.	5.6	17
46	A randomised controlled trial of rosuvastatin for the prevention of aminoglycoside-induced kidney toxicity in children with cystic fibrosis. Scientific Reports, 2020, 10, 1796.	3.3	4
47	Development and Reporting of Prediction Models: Guidance for Authors From Editors of Respiratory, Sleep, and Critical Care Journals. Critical Care Medicine, 2020, 48, 623-633.	0.9	188
48	Prophylactic anti-staphylococcal antibiotics for cystic fibrosis. The Cochrane Library, 2020, 2020, CD001912.	2.8	7
49	Response to Journal Club: Cluster Randomized Trial Evaluating Impact of a Community-based Microfinance Scheme on Childhood Nutritional Status: Evidence-based Medicine Viewpoint. Indian Pediatrics, 2020, 57, 688-689.	0.4	0
50	Adapting the James Lind Alliance priority setting process to better support patient participation: an example from cystic fibrosis. Research Involvement and Engagement, 2019, 5, 24.	2.9	24
51	Is microfinance associated with changes in womenâ€™s well-being and childrenâ€™s nutrition? A systematic review and meta-analysis. BMJ Open, 2019, 9, e023658.	1.9	16
52	Patient engagement to prioritise CF research: Inclusive or selective?. Journal of Cystic Fibrosis, 2019, 18, 307-308.	0.7	2
53	Climate change and lung health: presidential failure, professional responsibility. Thorax, 2019, 74, 627-628.	5.6	1
54	Reconciling Antimicrobial Susceptibility Testing and Clinical Response in Antimicrobial Treatment of Chronic Cystic Fibrosis Lung Infections. Clinical Infectious Diseases, 2019, 69, 1812-1816.	5.8	62

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55	Infection prevention and control in cystic fibrosis: a systematic review of interventions. Expert Review of Respiratory Medicine, 2019, 13, 425-434.	2.5	14
56	Once-daily versus multiple-daily dosing with intravenous aminoglycosides for cystic fibrosis. The Cochrane Library, 2019, 2019, CD002009.	2.8	5
57	Treatments for preventing recurrence of infection with Pseudomonas aeruginosa in people with cystic fibrosis. The Cochrane Library, 2019, 12, CD012300.	2.8	8
58	Control of Confounding and Reporting of Results in Causal Inference Studies. Guidance for Authors from Editors of Respiratory, Sleep, and Critical Care Journals. Annals of the American Thoracic Society, 2019, 16, 22-28.	3.2	458
59	The prevalence, clinical status and genotype of cystic fibrosis patients living in Cuba using national registry data. Journal of Cystic Fibrosis, 2019, 18, 522-524.	0.7	2
60	Gaps in the evidence for treatment decisions in cystic fibrosis: a systematic review. Thorax, 2019, 74, 229-236.	5.6	12
61	Clinical significance of Pseudomonas aeruginosa 2-alkyl-4-quinolone quorum-sensing signal molecules for long-term outcomes in adults with cystic fibrosis. Journal of Medical Microbiology, 2019, 68, 1823-1828.	1.8	6
62	ECFS best practice guidelines: the 2018 revision. Journal of Cystic Fibrosis, 2018, 17, 153-178.	0.7	521
63	Do guidelines for treating chest disease in children use Cochrane Reviews effectively? A systematic review. Thorax, 2018, 73, 670-673.	5.6	3
64	Early Respiratory Bacterial Detection and Antistaphylococcal Antibiotic Prophylaxis in Young Children with Cystic Fibrosis. Annals of the American Thoracic Society, 2018, 15, 42-48.	3.2	24
65	The top 10 research priorities in cystic fibrosis developed by a partnership between people with CF and healthcare providers. Thorax, 2018, 73, 388-390.	5.6	181
66	Staphylococcus aureus in cystic fibrosis. Current Opinion in Pulmonary Medicine, 2018, 24, 586-591.	2.6	14
67	Interventions for the eradication of meticillin-resistant Staphylococcus aureus (MRSA) in people with cystic fibrosis. The Cochrane Library, 2018, 7, CD009650.	2.8	17
68	Climate change and lung health: the challenge for a new president. Thorax, 2017, 72, 295-296.	5.6	5
69	Prophylactic anti-staphylococcal antibiotics for cystic fibrosis. The Cochrane Library, 2017, 4, CD001912.	2.8	33
70	Once-daily versus multiple-daily dosing with intravenous aminoglycosides for cystic fibrosis. The Cochrane Library, 2017, 3, CD002009.	2.8	34
71	The patient voice in research â€" Supporting actor or starring role?. Journal of Cystic Fibrosis, 2017, 16, 313-314.	0.7	4
72	Weight gain during acute treatment of an initial pulmonary exacerbation is associated with a longer interval to the next exacerbation in adults with cystic fibrosis. ERJ Open Research, 2017, 3, 00057-2017.	2.6	1

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73	Diagnostic and prognostic significance of systemic alkyl quinolones for P. aeruginosa in cystic fibrosis: A longitudinal study. Journal of Cystic Fibrosis, 2017, 16, 230-238.	0.7	36
74	Bronchoscopy-guided antimicrobial therapy for cystic fibrosis. The Cochrane Library, 2016, , CD009530.	2.8	17
75	Addressing resistance to antibiotics in systematic reviews of antibiotic interventions. Journal of Antimicrobial Chemotherapy, 2016, 71, 2367-2369.	3.0	45
76	Thoraxprotocol review: working with trialists to improve trial quality. Thorax, 2016, 71, 491-492.	5.6	0
77	First year of the thoracic triumvirate. Thorax, 2016, 71, 579-580.	5.6	0
78	Perception of first respiratory infection with Pseudomonas aeruginosa by people with cystic fibrosis and those close to them: an online qualitative study. BMJ Open, 2016, 6, e012303.	1.9	19
79	US Cystic Fibrosis Foundation and European Cystic Fibrosis Society consensus recommendations for the management of non-tuberculous mycobacteria in individuals with cystic fibrosis: executive summary. Thorax, 2016, 71, 88-90.	5.6	274
80	Glutamine supplementation in cystic fibrosis: A randomized placebo-controlled trial. Pediatric Pulmonology, 2016, 51, 253-257.	2.0	5
81	Treatment of pulmonary exacerbations in cystic fibrosis â€“ could do better?. Paediatric Respiratory Reviews, 2016, 20, 6-7.	1.8	6
82	US Cystic Fibrosis Foundation and European Cystic Fibrosis Society consensus recommendations for the management of non-tuberculous mycobacteria in individuals with cystic fibrosis. Thorax, 2016, 71, i1-i22.	5.6	348
83	A phase 3, multi-center, multinational, randomized, double-blind, placebo-controlled study to evaluate the efficacy and safety of levofloxacin inhalation solution (APT-1026) in stable cystic fibrosis patients. Journal of Cystic Fibrosis, 2016, 15, 495-502.	0.7	59
84	Interventions for the eradication of meticillin-resistant Staphylococcus aureus (MRSA) in people with cystic fibrosis. The Cochrane Library, 2015, , CD009650.	2.8	35
85	Embracing social media: TableÂ1. Thorax, 2015, 70, 1112-1112.	5.6	4
86	Editorial â€¢ Cochrane review: antibiotic strategies for eradicating Pseudomonas aeruginosa in people with cystic fibrosis. Journal of Evidence-Based Medicine, 2015, 8, 168-169.	2.4	1
87	Exhaled breath hydrogen cyanide as a marker of earlyPseudomonas aeruginosainfection in children with cystic fibrosis. ERJ Open Research, 2015, 1, 00044-2015.	2.6	40
88	Pseudomonas aeruginosaquorum sensing molecules correlate with clinical status in cystic fibrosis. European Respiratory Journal, 2015, 46, 1046-1054.	6.7	95
89	The first thoracic triumvirate. Thorax, 2015, 70, 917-917.	5.6	0
90	Optimising respiratory health in children with cystic fibrosis. Paediatrics and Child Health (United) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50	0.4	1

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91	A phase 3, open-label, randomized trial to evaluate the safety and efficacy of levofloxacin inhalation solution (APT-1026) versus tobramycin inhalation solution in stable cystic fibrosis patients. Journal of Cystic Fibrosis, 2015, 14, 507-514.	0.7	62
92	Cystic fibrosis microbiology: Advances in antimicrobial therapy. Journal of Cystic Fibrosis, 2015, 14, 551-560.	0.7	83
93	Testing children of HIV-positive parents: a multidisciplinary approach. Archives of Disease in Childhood, 2014, 99, 789-790.	1.9	0
94	Once-daily versus multiple-daily dosing with intravenous aminoglycosides for cystic fibrosis. , 2014, , CD002009.		8
95	Antibiotic strategies for eradicating Pseudomonas aeruginosa in people with cystic fibrosis. , 2014, , CD004197.		120
96	Evidence into practice: How do we get past the roadblocks?. Paediatric Respiratory Reviews, 2014, 15, 45-46.	1.8	0
97	Prescribing practices for intravenous aminoglycosides in UK Cystic Fibrosis clinics: A questionnaire survey. Journal of Cystic Fibrosis, 2014, 13, 424-427.	0.7	12
98	Feasibility and pilot study of the effects of microfinance on mortality and nutrition in children under five amongst the very poor in India: study protocol for a cluster randomized controlled trial. Trials, 2014, 15, 298.	1.6	2
99	Risk-proportionate clinical trial monitoring: an example approach from a non-commercial trials unit. Trials, 2014, 15, 127.	1.6	14
100	Rate of improvement of CF life expectancy exceeds that of general populationâ€"Observational death registration study. Journal of Cystic Fibrosis, 2014, 13, 410-415.	0.7	66
101	Standards of Care for Cystic Fibrosis ten years later. Journal of Cystic Fibrosis, 2014, 13, S1-S2.	0.7	17
102	European Cystic Fibrosis Society Standards of Care: Best Practice guidelines. Journal of Cystic Fibrosis, 2014, 13, S23-S42.	0.7	438
103	Prophylactic anti-staphylococcal antibiotics for cystic fibrosis. , 2014, , CD001912.		13
104	Bronchoscopy-guided antimicrobial therapy for cystic fibrosis. , 2013, , CD009530.		13
105	20 Years of Cochrane Glancing backwards â€“ Moving ahead: a tale of two Cochrane Review Groups. Paediatric Respiratory Reviews, 2013, 14, 165-167.	1.8	1
106	Measures of body habitus are associated with lung function in adults with cystic fibrosis: A population-based study. Journal of Cystic Fibrosis, 2013, 12, 284-289.	0.7	8
107	Inhaled antibiotics for pulmonary exacerbations in cystic fibrosis. Paediatric Respiratory Reviews, 2013, 14, 27-28.	1.8	1
108	From pipeline to patient: new developments in cystic fibrosis therapeutics. Expert Opinion on Pharmacotherapy, 2013, 14, 323-329.	1.8	3

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109	Interventions for the eradication of methicillin-resistantStaphylococcus aureus(MRSA) in people with cystic fibrosis. , 2013, , CD009650.		10
110	New agents to treat lung infection in cystic fibrosis: a big enough leap?. Future Medicinal Chemistry, 2013, 5, 117-120.	2.3	0
111	Finding and filling the gaps in the evidence with high quality clinical trials â€“ the experience of one Cochrane Review Group. Journal of Evidence-Based Medicine, 2013, 6, 229-231.	2.4	1
112	Antibiotic adjuvant therapy for pulmonary infection in cystic fibrosis. The Cochrane Library, 2013, , CD008037.	2.8	18
113	Compliance with mandatory reporting of clinical trial results on ClinicalTrials.gov: cross sectional study. BMJ: British Medical Journal, 2012, 344, d7373-d7373.	2.3	235
114	Fluoroquinolones in the treatment of bronchopulmonary disease in cystic fibrosis. Therapeutic Advances in Respiratory Disease, 2012, 6, 363-373.	2.6	7
115	Birth Cohorts in Childhood Asthma: Lessons and Limitations. American Journal of Respiratory and Critical Care Medicine, 2012, 185, 238-239.	5.6	3
116	Novel approaches to the treatment ofPseudomonas aeruginosainfections in cystic fibrosis. European Respiratory Journal, 2012, 40, 1014-1023.	6.7	100
117	Are Measures of Body Habitus Associated With Mortality in Cystic Fibrosis?. Chest, 2012, 142, 712-717.	0.8	35
118	Prophylactic anti-staphylococcal antibiotics for cystic fibrosis. , 2012, 12, CD001912.		18
119	Current dilemmas in antimicrobial therapy in cystic fibrosis. Expert Review of Respiratory Medicine, 2012, 6, 407-422.	2.5	14
120	Delayed publication of clinical trials in cystic fibrosis. Journal of Cystic Fibrosis, 2012, 11, 14-17.	0.7	8
121	Results of antibiotic susceptibility testing do not influence clinical outcome in children with cystic fibrosis. Journal of Cystic Fibrosis, 2012, 11, 288-292.	0.7	113
122	A clinical approach to a wheezy infant. Paediatrics and Child Health (United Kingdom), 2012, 22, 307-309.	0.4	0
123	Once-daily versus multiple-daily dosing with intravenous aminoglycosides for cystic fibrosis. , 2012, , CD002009.		19
124	Twice vs three times daily antibiotics in the treatment of pulmonary exacerbations of cystic fibrosis. Journal of Cystic Fibrosis, 2011, 10, 383.	0.7	1
125	Optimizing respiratory health in children with cystic fibrosis. Paediatrics and Child Health (United) Tj ETQq1 1 0.784314 rgBT /Overlo	0.4	0
126	Therapeutic approaches to chronic cystic fibrosis respiratory infections with available, emerging aerosolized antibiotics. Respiratory Medicine, 2011, 105, S2-S8.	2.9	26

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127	Aerosolized antibiotic therapy for chronic cystic fibrosis airway infections: continuous or intermittent?. Respiratory Medicine, 2011, 105, S9-S17.	2.9	21
128	The Management of Pre-School Wheeze. Paediatric Respiratory Reviews, 2011, 12, 70-77.	1.8	27
129	Pneumonia in the developed world. Paediatric Respiratory Reviews, 2011, 12, 60-69.	1.8	41
130	Evaluating the effectiveness of a schools-based programme to promote exercise self-efficacy in children and young people with risk factors for obesity: Steps to active kids (STAK). BMC Public Health, 2011, 11, 830.	2.9	13
131	A Glycopeptide Dendrimer Inhibitor of the Galactoseâ€Specific Lectin LecA and of Pseudomonas aeruginosa Biofilms. Angewandte Chemie - International Edition, 2011, 50, 10631-10635.	13.8	149
132	Risk Factors for Chronic Kidney Disease in Adults with Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2011, 184, 1147-1152.	5.6	72
133	Association between socioeconomic status, sex, and age at death from cystic fibrosis in England and Wales (1959 to 2008): cross sectional study. BMJ: British Medical Journal, 2011, 343, d4662-d4662.	2.3	79
134	Treatment massive haemoptysis in cystic fibrosis with tranexamic acid. Journal of the Royal Society of Medicine, 2011, 104, 49-52.	2.0	17
135	Antibiotic adjuvant therapy for pulmonary infection in cystic fibrosis. , 2010, , CD008037.		6
136	Aminoglycoside use in cystic fibrosis: therapeutic strategies and toxicity. Current Opinion in Pulmonary Medicine, 2010, 16, 604-610.	2.6	94
137	Garlic as an inhibitor of Pseudomonas aeruginosa quorum sensing in cystic fibrosisâ€"a pilot randomized controlled trial. Pediatric Pulmonology, 2010, 45, 356-362.	2.0	116
138	Minimizing the toxicity of aminoglycosides in cystic fibrosis. Journal of the Royal Society of Medicine, 2010, 103, 3-5.	2.0	7
139	Pseudomonas eradication in cystic fibrosis: who will join the ELITE?. Thorax, 2010, 65, 281-282.	5.6	4
140	Side effects of aminoglycosides on the kidney, ear and balance in cystic fibrosis. Thorax, 2010, 65, 654-658.	5.6	119
141	Oral versus i.v. antibiotics for community-acquired pneumonia in children: a cost-minimisation analysis. European Respiratory Journal, 2010, 35, 858-864.	6.7	25
142	Once-daily versus multiple-daily dosing with intravenous aminoglycosides for cystic fibrosis. , 2010, , CD002009.		17
143	Respiratory medicines for children: current evidence, unlicensed use and research priorities. European Respiratory Journal, 2010, 35, 247-265.	6.7	39
144	European best practice guidelines for cystic fibrosis neonatal screening. Journal of Cystic Fibrosis, 2009, 8, 153-173.	0.7	196

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145	Oral Prednisolone for Preschool Children with Acute Virus-Induced Wheezing. New England Journal of Medicine, 2009, 360, 329-338.	27.0	296
146	Antibiotic strategies for eradicating Pseudomonas aeruginosa in people with cystic fibrosis. , 2009, , CD004197.		63
147	Treatment strategies for cystic fibrosis: what's in the pipeline?. Expert Opinion on Pharmacotherapy, 2009, 10, 1191-1202.	1.8	14
148	Bronchial asthma on Mount Kilimanjaro is not a disadvantage. Thorax, 2008, 63, 936-937.	5.6	14
149	Case-control study of acute renal failure in patients with cystic fibrosis in the UK. Thorax, 2008, 63, 532-535.	5.6	100
150	Exacerbations in cystic fibrosis: 3 {middle dot} Management. Thorax, 2007, 63, 180-184.	5.6	68
151	Survey of acute renal failure in patients with cystic fibrosis in the UK. Thorax, 2007, 62, 541-545.	5.6	78
152	Asthma as a Barrier to Children's Physical Activity: In Reply. Pediatrics, 2007, 119, 1248-1249.	2.1	45
153	Effective treatment strategies for paediatric community-acquired pneumonia. Expert Opinion on Pharmacotherapy, 2007, 8, 1091-1101.	1.8	22
154	Comparison of oral amoxicillin and intravenous benzyl penicillin for community acquired pneumonia in children (PIVOT trial): a multicentre pragmatic randomised controlled equivalence trial. Thorax, 2007, 62, 1102-1106.	5.6	87
155	Population pharmacokinetics of tobramycin administered thrice daily and once daily in children and adults with cystic fibrosis. Journal of Cystic Fibrosis, 2007, 6, 327-333.	0.7	61
156	Clinical trials in cystic fibrosis. Journal of Cystic Fibrosis, 2007, 6, 85-99.	0.7	41
157	Antibiotic strategies for eradicating Pseudomonas aeruginosa in people with cystic fibrosis. , 2006, , CD004197.		33
158	Once-daily versus multiple-daily dosing with intravenous aminoglycosides for cystic fibrosis. , 2006, , CD002009.		29
159	Update on treatment of pulmonary exacerbations in cystic fibrosis. Current Opinion in Pulmonary Medicine, 2006, 12, 440-444.	2.6	36
160	Absence of Cochleotoxicity Measured by Standard and High-Frequency Pure Tone Audiometry in a Trial of Once- versus Three-Times-Daily Tobramycin in Cystic Fibrosis Patients. Antimicrobial Agents and Chemotherapy, 2006, 50, 2293-2299.	3.2	36
161	Asthma as a Barrier to Children's Physical Activity: Implications for Body Mass Index and Mental Health. Pediatrics, 2006, 118, 2443-2449.	2.1	152
162	A Randomized, Controlled Trial of an Interactive Educational Computer Package for Children With Asthma. Pediatrics, 2006, 117, 1046-1054.	2.1	88

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163	Prophylactic Antibiotics in Cystic Fibrosis: A Conviction without Evidence?. Pediatric Pulmonology, 2005, 40, 471-476.	2.0	35
164	Once versus three-times daily regimens of tobramycin treatment for pulmonary exacerbations of cystic fibrosisâ€"the TOPIC study: a randomised controlled trial. Lancet, The, 2005, 365, 573-578.	13.7	176
165	Tobramycin dosing in cystic fibrosis. Lancet, The, 2005, 365, 1767-1768.	13.7	8
166	Multiresistant pulmonary infection in cystic fibrosisâ€"prevention is better than cure. Lancet, The, 2005, 366, 433-435.	13.7	8
167	Educational interventions â€“ computers for delivering education to children with respiratory illness and to their parents. Paediatric Respiratory Reviews, 2005, 6, 215-226.	1.8	19
168	Prophylactic anti-staphylococcal antibiotics for cystic fibrosis. , 2003, , CD001912.		61
169	Aminoglycoside Prescribing and Surveillance in Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2003, 167, 819-823.	5.6	61
170	Parental attitudes: antenatal diagnosis of cystic fibrosis. Archives of Disease in Childhood, 2002, 87, 284-286.	1.9	13
171	Pneumonia due to viral and atypical organisms and their sequelae. British Medical Bulletin, 2002, 61, 247-262.	6.9	20
172	The Asthma Files: Evaluation of a multimedia package for childrenâ€™s asthma education. Paediatric Nursing, 2002, 14, 32-35.	0.1	19
173	Once-daily tobramycin monotherapy in cystic fibrosis. Pediatric Pulmonology, 2002, 33, 406-406.	2.0	2
174	Lack of concordance in the use and monitoring of intravenous aminoglycosides in UK cystic fibrosis centers. Pediatric Pulmonology, 2002, 33, 165-165.	2.0	21
175	Antibiotics and Acute Renal Failure in Children with Cystic Fibrosis. Paediatric and Perinatal Drug Therapy, 2002, 5, 65-67.	0.5	12
176	Drug Development for Children. Paediatric and Perinatal Drug Therapy, 2002, 5, 2-3.	0.5	0
177	Double click for health: the role of multimedia in asthma education. Archives of Disease in Childhood, 2001, 85, 447-449.	1.9	20
178	Trends in passive smoking in cystic fibrosis, 1993-1998. Pediatric Pulmonology, 2001, 31, 133-137.	2.0	10
179	Acute rib fracture pain in CF. Thorax, 2001, 56, 819-819.	5.6	10
180	Antibiotic therapy against Pseudomonas aeruginosa in cystic fibrosis: a European consensus. European Respiratory Journal, 2000, 16, 749.	6.7	556

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181	Rationalised prescribing for community acquired pneumonia: a closed loop audit. Archives of Disease in Childhood, 2000, 83, 320-324.	1.9	58
182	Systematic review of antistaphylococcal antibiotic therapy in cystic fibrosis. Thorax, 2000, 55, 251-251.	5.6	1
183	A study of a single high potency multivitamin preparation in the management of cystic fibrosis. Journal of Human Nutrition and Dietetics, 1998, 11, 493-500.	2.5	2
184	Impact of a case management protocol for childhood pneumonia in a rural Zambian hospital. Annals of Tropical Paediatrics, 1998, 18, 155-160.	1.0	6
185	Effect of respiratory virus infections including rhinovirus on clinical status in cystic fibrosis.. Archives of Disease in Childhood, 1995, 73, 117-120.	1.9	163
186	Strictures of ascending colon in cystic fibrosis and high-strength pancreatic enzymes. Lancet, The, 1994, 343, 85-86.	13.7	309
187	Transmission of Pseudomonas cepacia by social contact in cystic fibrosis. Lancet, The, 1993, 342, 434-435.	13.7	9
188	Receiver operating characteristic curves for comparison of serial neutrophil band forms and C reactive protein in neonates at risk of infection.. Archives of Disease in Childhood, 1992, 67, 808-812.	1.9	44
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