AndrÃ© Schultz

List of Publications by Year
in descending order

73	987	516710	526287
papers	citations	16 h-index	27 g-index

74	74	74	1374
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Airway surface liquid pH is not acidic in children with cystic fibrosis. Nature Communications, 2017, 8, 1409.	12.8	84
2	Assessment of bronchodilator responsiveness in preschool children using forced oscillations. Thorax, 2007, 62, 814-819.	5.6	82
3	Changing Prevalence of Lower Airway Infections in Young Children with Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 590-599.	5.6	49
4	Aspergillus Infections and Progression of Structural Lung Disease in Children with Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 688-696.	5.6	42
5	Episodic Viral Wheeze and Multiple Trigger Wheeze in preschool children: A useful distinction for clinicians?. Paediatric Respiratory Reviews, 2011, 12, 160-164.	1.8	40
6	Efficacy of oral amoxicillinâ€“clavulanate or azithromycin for non-severe respiratory exacerbations in children with bronchiectasis (BEST-1): a multicentre, three-arm, double-blind, randomised placebo-controlled trial. Lancet Respiratory Medicine,the, 2019, 7, 791-801.	10.7	37
7	Childhood interstitial lung diseases in immunocompetent children in Australia and New Zealand: a decadeâ€™s experience. Orphanet Journal of Rare Diseases, 2017, 12, 133.	2.7	35
8	Induced sputum to detect lung pathogens in young children with cystic fibrosis. Pediatric Pulmonology, 2017, 52, 182-189.	2.0	33
9	Aerosol Inhalation From Spacers and Valved Holding Chambers Requires Few Tidal Breaths for Children. Pediatrics, 2010, 126, e1493-e1498.	2.1	32
10	Investigating selfâ€efficacy, disease knowledge and adherence to treatment in adolescents with cystic fibrosis. Journal of Paediatrics and Child Health, 2017, 53, 488-493.	0.8	26
11	The clinical significance of oropharyngeal cultures in young children with cystic fibrosis. European Respiratory Journal, 2018, 51, 1800238.	6.7	25
12	Early pulmonary inflammation and lung damage in children with cystic fibrosis. Respirology, 2015, 20, 569-578.	2.3	21
13	Chronic wet cough in Aboriginal children: It's not just a cough. Journal of Paediatrics and Child Health, 2019, 55, 833-843.	0.8	20
14	Biomarkers in Paediatric Cystic Fibrosis Lung Disease. Paediatric Respiratory Reviews, 2015, 16, 213-218.	1.8	19
15	An introduction to clinical trial design. Paediatric Respiratory Reviews, 2019, 32, 30-35.	1.8	19
16	Duration of amoxicillin-clavulanate for protracted bacterial bronchitis in children (DACS): a multi-centre, double blind, randomised controlled trial. Lancet Respiratory Medicine,the, 2021, 9, 1121-1129.	10.7	19
17	We won't find what we don't look for: Identifying barriers and enablers of chronic wet cough in Aboriginal children. Respirology, 2020, 25, 383-392.	2.3	18
18	Recognition and Management of Protracted Bacterial Bronchitis in Australian Aboriginal Children. Chest, 2021, 159, 249-258.	0.8	17

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19	CF derived scoring systems do not fully describe the range of structural changes seen on CT scans in PCD. Pediatric Pulmonology, 2019, 54, 471-477.	2.0	17
20	Usefulness of parental response to questions about adherence to prescribed inhaled corticosteroids in young children. Archives of Disease in Childhood, 2012, 97, 1092-1096.	1.9	16
21	BAL Inflammatory Markers Can Predict Pulmonary Exacerbations in Children With Cystic Fibrosis. Chest, 2020, 158, 2314-2322.	0.8	16
22	Letters to the Editor. Journal of Paediatrics and Child Health, 2008, 44, 604-606.	0.8	15
23	Developing a smartphone application to support social connectedness and wellbeing in young people with cystic fibrosis. Journal of Cystic Fibrosis, 2020, 19, 277-283.	0.7	15
24	Surfactant protein disorders in childhood interstitial lung disease. European Journal of Pediatrics, 2021, 180, 2711-2721.	2.7	15
25	Prevalence of chronic wet cough and protracted bacterial bronchitis in Aboriginal children. ERJ Open Research, 2019, 5, 00248-2019.	2.6	14
26	Ivacaftor and Airway Inflammation in Preschool Children with Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2021, 204, 605-608.	5.6	14
27	Pressurised metered dose inhaler-spacer technique in young children improves with video instruction. European Journal of Pediatrics, 2016, 175, 1007-1012.	2.7	13
28	Side effects of medications used to treat childhood interstitial lung disease. Paediatric Respiratory Reviews, 2018, 28, 68-79.	1.8	13
29	The impact of chest computed tomography and chest radiography on clinical management of cystic fibrosis lung disease. Journal of Cystic Fibrosis, 2020, 19, 641-646.	0.7	13
30	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
31	Conducting decolonizing research and practice with Australian First Nations to close the health gap. Health Research Policy and Systems, 2021, 19, 127.	2.8	13
32	Incentive device improves spacer technique but not clinical outcome in preschool children with asthma. Journal of Paediatrics and Child Health, 2012, 48, 52-56.	0.8	12
33	Humidified high-flow nasal cannula oxygen for bronchiolitis: should we go with the flow?. Archives of Disease in Childhood, 2018, 103, 303-303.	1.9	12
34	Identifying pediatric lung disease: A comparison of forced oscillation technique outcomes. Pediatric Pulmonology, 2019, 54, 751-758.	2.0	12
35	Validation of Methodology for Recording Breathing and Simulating Drug Delivery Through Spacers and Valved Holding Chambers. Journal of Aerosol Medicine and Pulmonary Drug Delivery, 2010, 23, 311-322.	1.4	11
36	ABCA3 lung disease in an ex 27 week preterm infant responsive to systemic glucocorticosteroids. Pediatric Pulmonology, 2016, 51, E1-E3.	2.0	11

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37	Current options in aerosolised drug therapy for children receiving respiratory support. Anaesthesia, 2017, 72, 1388-1397.	3.8	11
38	Multiâ€centre ethics and research governance review can impede nonâ€interventional clinical research. Internal Medicine Journal, 2019, 49, 722-728.	0.8	11
39	Phenotypeâ€directed treatment of preâ€schoolâ€aged children with recurrent wheeze. Journal of Paediatrics and Child Health, 2012, 48, E73-8.	0.8	9
40	Individual pharmacokinetic variation leads to underdosing of ciprofloxacin in some cystic fibrosis patients. Pediatric Pulmonology, 2017, 52, 319-323.	2.0	8
41	Does machine learning have a role in the prediction of asthma in children?. Paediatric Respiratory Reviews, 2022, 41, 51-60.	1.8	8
42	Outpatient Management of Asthma in Children. Clinical Medicine Insights Pediatrics, 2013, 7, CMPed.S7867.	1.4	7
43	Trial Refresh: A Case for an Adaptive Platform Trial for Pulmonary Exacerbations of Cystic Fibrosis. Frontiers in Pharmacology, 2019, 10, 301.	3.5	7
44	Frequency of protracted bacterial bronchitis and management <scp>preâ€respiratory</scp> referral. Journal of Paediatrics and Child Health, 2022, 58, 97-103.	0.8	7
45	CrossTalk opposing view: mucosal acidification does not drive early progressive lung disease in cystic fibrosis. Journal of Physiology, 2018, 596, 3439-3441.	2.9	6
46	Discrete choice experiment to evaluate preferences of patients with cystic fibrosis among alternative treatment-related health outcomes: a protocol. BMJ Open, 2019, 9, e030348.	1.9	5
47	Danger of using an unreliable classification system for preschool wheeze. European Respiratory Journal, 2009, 33, 944-945.	6.7	4
48	Assessment of different techniques for the administration of inhaled salbutamol in children breathing spontaneously via tracheal tubes, supraglottic airway devices, and tracheostomies. Paediatric Anaesthesia, 2020, 30, 1363-1377.	1.1	4
49	Paediatric headbox as aerosol and droplet barrier. Archives of Disease in Childhood, 2022, 107, 65-67.	1.9	4
50	Variation in treatment preferences of pulmonary exacerbations among Australian and New Zealand cystic fibrosis physicians. BMJ Open Respiratory Research, 2021, 8, e000956.	3.0	4
51	Preferred health outcome states following treatment for pulmonary exacerbations of cystic fibrosis. Journal of Cystic Fibrosis, 2022, 21, 581-587.	0.7	4
52	To track or not to track: wheeze phenotypes in preschool children. European Respiratory Journal, 2018, 51, 1800042.	6.7	3
53	Cough swabs less useful but induced sputum very useful in symptomatic older children with cystic fibrosis. Lancet Respiratory Medicine,the, 2018, 6, 410-411.	10.7	3
54	Redesign of the Australian Cystic Fibrosis Data Registry: A multidisciplinary collaboration. Paediatric Respiratory Reviews, 2021, 37, 37-43.	1.8	3

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55	Respiratory follow-up to improve outcomes for Aboriginal children: twelve key steps. The Lancet Regional Health - Western Pacific, 2021, 15, 100239.	2.9	3
56	Genomic testing for children with interstitial and diffuse lung disease (chILD): parent satisfaction, understanding and health-related quality of life. BMJ Open Respiratory Research, 2022, 9, e001139.	3.0	2
57	Lung abscess: 14â€‰years of experience in a tertiary paediatric hospital. ANZ Journal of Surgery, 0, , .	0.7	2
58	Vascular ring: Unmasked. Journal of Paediatrics and Child Health, 2017, 53, 503-506.	0.8	1
59	Cystic Fibrosis Survival Gap Closing between the United States and Canada. Donâ€™t Leave Anyone Behind!. American Journal of Respiratory and Critical Care Medicine, 2018, 197, 701-703.	5.6	1
60	Rebuttal from Stephen M. Stick and AndrÃ© Schultz. Journal of Physiology, 2018, 596, 3445-3446.	2.9	1
61	Reply to Turnbull et al. and to Hulme et al.. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 750-752.	5.6	1
62	Acute haemoptysis, fever and abdominal pain in an adolescent from northern Australia. Thorax, 2021, 76, 951-953.	5.6	1
63	Tuberculosis in Australia's Top End First Nations highlights health and life expectancy gaps: a call to arms. The Lancet Regional Health - Western Pacific, 2021, 15, 100253.	2.9	1
64	Adherence to CF treatment can be improved with the right approach!. Thorax, 2022, 77, 428-428.	5.6	1
65	Fissure adjacent partial lobe atelectasis in primary ciliary dyskinesia. Journal of Paediatrics and Child Health, 2022, 58, 683-686.	0.8	1
66	A pilot study of disease related education and psychotherapeutic support for unresolved grief in parents of children with CF. Scientific Reports, 2022, 12, 5746.	3.3	1
67	Preservation of Lung Function in Cystic Fibrosis: Are Macrolides the Answer?. American Journal of Respiratory and Critical Care Medicine, 2018, 198, 1114-1116.	5.6	0
68	Is it time to move on from episodic viral wheeze and multiple trigger wheeze?. Pediatric Pulmonology, 2019, 54, 1499-1450.	2.0	0
69	Research Note: Adaptive trials. Journal of Physiotherapy, 2019, 65, 113-116.	1.7	0
70	Suboptimal asthma care: Lessons from Australia and a way forward. Respirology, 2020, 25, 45-46.	2.3	0
71	Primary Nasal Epithelial Cells as a Surrogate Cell Culture Model for Type-II Alveolar Cells to Study ABCA-3 Deficiency. Frontiers in Medicine, 2022, 9, 827416.	2.6	0
72	Implementation of a strategy to facilitate effective medical follow-up for Australian First Nations children hospitalised with lower respiratory tract infections: study protocol. BMC Pulmonary Medicine, 2022, 22, 92.	2.0	0

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73	Reducing exacerbations in children and adults with primary ciliary dyskinesia using erdosteine and/or azithromycin therapy (REPEAT trial): study protocol for a multicentre, double-blind, double-dummy, 2Ã—2 partial factorial, randomised controlled trial. BMJ Open Respiratory Research, 2022, 9, e001236.	3.0	0