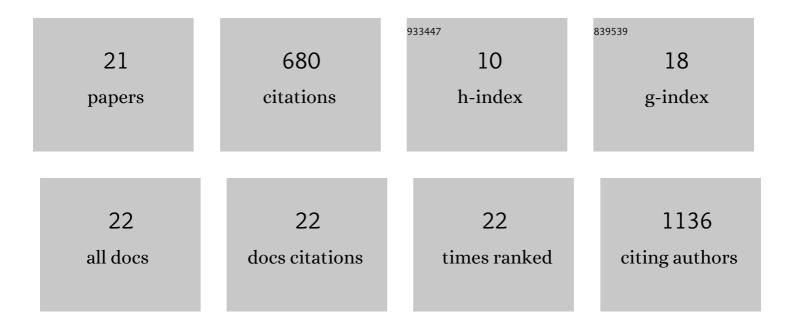
Andrew P Prayle

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3120441/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	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
2	Association Between Treatments and Short-Term Biochemical Improvements and Clinical Outcomes in Post-Severe Acute Respiratory Syndrome Coronavirus-2 Inflammatory Syndrome. Pediatric Critical Care Medicine, 2021, 22, e285-e293.	0.5	20
3	Gastrointestinal complications of cystic fibrosis. Paediatrics and Child Health (United Kingdom), 2020, 30, 345-349.	0.4	4
4	Assessing the impact of posture on diaphragm morphology and function using an open upright MRI system—A pilot study. European Journal of Radiology, 2020, 130, 109196.	2.6	1
5	Observational Study of Pulse Transit Time in Children With Sleep Disordered Breathing. Frontiers in Neurology, 2020, 11, 316.	2.4	2
6	Looking under the bonnet of bronchopulmonary dysplasia with MRI. Thorax, 2020, 75, 100-100.	5.6	3
7	What's hot that the other lot got. Thorax, 2019, 74, 923-924.	5.6	0
8	Key paediatric messages from the 2018ÂEuropean Respiratory Society International Congress. ERJ Open Research, 2019, 5, 00241-2018.	2.6	1
9	β ₂ -agonists do not work in children under 2â€years of age: myth or maxim?. Breathe, 2019, 15, 273-276.	1.3	5
10	Gaps in the evidence for treatment decisions in cystic fibrosis: a systematic review. Thorax, 2019, 74, 229-236.	5.6	12
11	Do guidelines for treating chest disease in children use Cochrane Reviews effectively? A systematic review. Thorax, 2018, 73, 670-673.	5.6	3
12	Percutaneous lines for delivering intravenous antibiotics in people with cystic fibrosis. The Cochrane Library, 2017, 2017, CD008243.	2.8	1
13	Growth and nutrition in children with ataxia telangiectasia. Archives of Disease in Childhood, 2016, 101, 1137-1141.	1.9	15
14	Rate of improvement of CF life expectancy exceeds that of general population—Observational death registration study. Journal of Cystic Fibrosis, 2014, 13, 410-415.	0.7	66
15	From pipeline to patient: new developments in cystic fibrosis therapeutics. Expert Opinion on Pharmacotherapy, 2013, 14, 323-329.	1.8	3
16	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
17	Pneumonia in the developed world. Paediatric Respiratory Reviews, 2011, 12, 60-69.	1.8	41
18	Karyomegalic-like nephropathy, Ewing's sarcoma and ifosfamide therapy. Pediatric Nephrology, 2011, 26, 1163-1166	1.7	25

#	Article	IF	CITATIONS
19	Aminoglycoside use in cystic fibrosis: therapeutic strategies and toxicity. Current Opinion in Pulmonary Medicine, 2010, 16, 604-610.	2.6	94
20	Side effects of aminoglycosides on the kidney, ear and balance in cystic fibrosis. Thorax, 2010, 65, 654-658.	5.6	119
21	Homeostasis and the Respiratory System. , 0, , 11-15.		0