Sanja Stanojevic

List of Publications by Citations

151 papers	9,693 citations	38 h-index	97 g-index
170 ext. papers	12,881 ext. citations	6.5 avg, IF	6.04 L-index

#	Paper	IF	Citations
151	Multi-ethnic reference values for spirometry for the 3-95-yr age range: the global lung function 2012 equations. *European Respiratory Journal*, **2012**, 40, 1324-43	13.6	2784
150	Global asthma prevalence in adults: findings from the cross-sectional world health survey. *BMC Public Health*, **2012**, 12, 204	4.1	819
149	Standardization of Spirometry 2019 Update. An Official American Thoracic Society and European Respiratory Society Technical Statement. *American Journal of Respiratory and Critical Care Medicine*, **2019**, 200, e70-e88	10.2	719
148	Reference ranges for spirometry across all ages: a new approach. *American Journal of Respiratory and Critical Care Medicine*, **2008**, 177, 253-60	10.2	513
147	Recommendations for a Standardized Pulmonary Function Report. An Official American Thoracic Society Technical Statement. *American Journal of Respiratory and Critical Care Medicine*, **2017**, 196, 1463-1472	10.2	278
146	Comparison of urokinase and video-assisted thoracoscopic surgery for treatment of childhood empyema. *American Journal of Respiratory and Critical Care Medicine*, **2006**, 174, 221-7	10.2	257
145	Official ERS technical standards: Global Lung Function Initiative reference values for the carbon monoxide transfer factor for Caucasians. *European Respiratory Journal*, **2017**, 50,	13.6	194
144	Efficacy and safety of lumacaftor and ivacaftor in patients aged 6-11 years with cystic fibrosis homozygous for F508del-CFTR: a randomised, placebo-controlled phase 3 trial. *Lancet Respiratory Medicine,the*, **2017**, 5, 557-567	35.1	176
143	Lung clearance index at 4 years predicts subsequent lung function in children with cystic fibrosis. *American Journal of Respiratory and Critical Care Medicine*, **2011**, 183, 752-8	10.2	170
142	Reference values for lung function: past, present and future. *European Respiratory Journal*, **2010**, 36, 12-9	13.6	158
141	Effect of pulmonary exacerbations on long-term lung function decline in cystic fibrosis. *European Respiratory Journal*, **2012**, 40, 61-6	13.6	146
140	Lung Clearance Index and HRCT are complementary markers of lung abnormalities in young children with CF. *Thorax*, **2011**, 66, 481-8	7.3	133
139	Spirometry centile charts for young Caucasian children: the Asthma UK Collaborative Initiative. *American Journal of Respiratory and Critical Care Medicine*, **2009**, 180, 547-52	10.2	133
138	Lung clearance index as an outcome measure for clinical trials in young children with cystic fibrosis. A pilot study using inhaled hypertonic saline. *American Journal of Respiratory and Critical Care Medicine*, **2013**, 188, 456-60	10.2	118
137	Age- and height-based prediction bias in spirometry reference equations. *European Respiratory Journal*, **2012**, 40, 190-7	13.6	115
136	A contemporary survival analysis of individuals with cystic fibrosis: a cohort study. *European Respiratory Journal*, **2015**, 45, 670-9	13.6	111
135	Influence of secular trends and sample size on reference equations for lung function tests. *European Respiratory Journal*, **2011**, 37, 658-64	13.6	111

134	Lung function from infancy to the preschool years after clinical diagnosis of cystic fibrosis. *American Journal of Respiratory and Critical Care Medicine*, **2008**, 178, 42-9	10.2	111
133	Survival Comparison of Patients With Cystic Fibrosis in Canada and the United States: A Population-Based Cohort Study. *Annals of Internal Medicine*, **2017**, 166, 537-546	8	107
132	Age- and size-related reference ranges: a case study of spirometry through childhood and adulthood. *Statistics in Medicine*, **2009**, 28, 880-98	2.3	98
131	Age and height dependence of lung clearance index and functional residual capacity. *European Respiratory Journal*, **2013**, 41, 1371-7	13.6	89
130	Progression of Lung Disease in Preschool Patients with Cystic Fibrosis. *American Journal of Respiratory and Critical Care Medicine*, **2017**, 195, 1216-1225	10.2	88
129	Longitudinal trends in nutritional status and the relation between lung function and BMI in cystic fibrosis: a population-based cohort study. *American Journal of Clinical Nutrition*, **2013**, 97, 872-7	7	87
128	Trade-offs in relative limb length among Peruvian children: extending the thrifty phenotype hypothesis to limb proportions. *PLoS ONE*, **2012**, 7, e51795	3.7	76
127	Multiple breath nitrogen washout: a feasible alternative to mass spectrometry. *PLoS ONE*, **2013**, 8, e56868	3.7	72
126	Clinical and demographic factors associated with post-lung transplantation survival in individuals with cystic fibrosis. *Journal of Heart and Lung Transplantation*, **2015**, 34, 1139-45	5.8	70
125	The Global Lung Initiative 2012 reference values reflect contemporary Australasian spirometry. *Respirology*, **2012**, 17, 1150-1	3.6	68
124	The Global Lung Function Initiative (GLI) Network: bringing the world's respiratory reference values together. *Breathe*, **2017**, 13, e56-e64	1.8	59
123	Preschool Multiple-Breath Washout Testing. An Official American Thoracic Society Technical Statement. *American Journal of Respiratory and Critical Care Medicine*, **2018**, 197, e1-e19	10.2	56
122	Use of FEV in cystic fibrosis epidemiologic studies and clinical trials: A statistical perspective for the clinical researcher. *Journal of Cystic Fibrosis*, **2017**, 16, 318-326	4.1	54
121	Physiologic endpoints for clinical studies for cystic fibrosis. *Journal of Cystic Fibrosis*, **2016**, 15, 416-23	4.1	48
120	Factors associated with response to treatment of pulmonary exacerbations in cystic fibrosis patients. *Journal of Cystic Fibrosis*, **2015**, 14, 755-62	4.1	47
119	Change of Outcomes in Pediatric Intestinal Failure: Use of Time-Series Analysis to Assess the Evolution of an Intestinal Rehabilitation Program. *Journal of the American College of Surgeons*, **2016**, 222, 1180-1188.e3	4.4	46
118	Lung clearance index in cystic fibrosis subjects treated for pulmonary exacerbations. *European Respiratory Journal*, **2015**, 46, 1055-64	13.6	41
117	Normative data for multiple breath washout outcomes in school-aged Caucasian children. *European Respiratory Journal*, **2020**, 55,	13.6	40

116	Data Resource Profile: The UK Cystic Fibrosis Registry. *International Journal of Epidemiology*, **2018**, 47, 9-10e	7.8	40
115	The changing epidemiology and demography of cystic fibrosis. *Presse Medicale*, **2017**, 46, e87-e95	2.2	38
114	Effect of pulmonary exacerbations treated with oral antibiotics on clinical outcomes in cystic fibrosis. *Thorax*, **2017**, 72, 327-332	7.3	38
113	Randomized controlled trial of biofilm antimicrobial susceptibility testing in cystic fibrosis patients. *Journal of Cystic Fibrosis*, **2015**, 14, 262-6	4.1	36
112	The impact of switching to the new global lung function initiative equations on spirometry results in the UK CF registry. *Journal of Cystic Fibrosis*, **2014**, 13, 319-27	4.1	36
111	A Systematic Approach to Multiple Breath Nitrogen Washout Test Quality. *PLoS ONE*, **2016**, 11, e0157523	3.7	35
110	Inhaled hypertonic saline in preschool children with cystic fibrosis (SHIP): a multicentre, randomised, double-blind, placebo-controlled trial. *Lancet Respiratory Medicine,the*, **2019**, 7, 802-809	35.1	34
109	Effectiveness of a stepwise Pseudomonas aeruginosa eradication protocol in children with cystic fibrosis. *Journal of Cystic Fibrosis*, **2017**, 16, 395-400	4.1	33
108	Inter-test reproducibility of the lung clearance index measured by multiple breath washout. *European Respiratory Journal*, **2017**, 50,	13.6	33
107	Oral non-steroidal anti-inflammatory drug therapy for cystic fibrosis. *Cochrane Database of Systematic Reviews*, **2007**, CD001505		32
106	Restoring Pulmonary and Sleep Services as the COVID-19 Pandemic Lessens. From an Association of Pulmonary, Critical Care, and Sleep Division Directors and American Thoracic Society-coordinated Task Force. *Annals of the American Thoracic Society*, **2020**, 17, 1343-1351	4.7	32
105	Global Lung Function Initiative 2012 reference equations for spirometry in the Norwegian population. *European Respiratory Journal*, **2016**, 48, 1602-1611	13.6	32
104	Prolongation of antibiotic treatment for cystic fibrosis pulmonary exacerbations. *Journal of Cystic Fibrosis*, **2015**, 14, 770-6	4.1	31
103	The association between socioeconomic status and obesity in Peruvian women. *Obesity*, **2012**, 20, 2283-8		30
102	A standardized approach to estimating survival statistics for population-based cystic fibrosis registry cohorts. *Journal of Clinical Epidemiology*, **2016**, 70, 206-13	5.7	29
101	Alternative outcomes for the multiple breath washout in children with CF. *Journal of Cystic Fibrosis*, **2015**, 14, 490-6	4.1	29
100	The Global Lung Function Initiative: dispelling some myths of lung function test interpretation. *Breathe*, **2013**, 9, 462-474	1.8	29
99	The all-age spirometry reference ranges reflect contemporary Australasian spirometry. *Respirology*, **2011**, 16, 912-7	3.6	29

98	Reference equations for pulmonary function tests in preschool children: a review. *Pediatric Pulmonology*, **2007**, 42, 962-72	3.5	28
97	Official ERS technical standard: Global Lung Function Initiative reference values for static lung volumes in individuals of European ancestry. *European Respiratory Journal*, **2021**, 57,	13.6	28
96	The CF Canada-Sick Kids Program in individual CF therapy: A resource for the advancement of personalized medicine in CF. *Journal of Cystic Fibrosis*, **2019**, 18, 35-43	4.1	28
95	Lung clearance index to monitor treatment response in pulmonary exacerbations in preschool children with cystic fibrosis. *Thorax*, **2018**, 73, 451-458	7.3	27
94	Stunting, adiposity, and the individual-level "dual burden" among urban lowland and rural highland Peruvian children. *American Journal of Human Biology*, **2014**, 26, 481-90	2.7	27
93	Secular changes in relative leg length confound height-based spirometric reference values. *Chest*, **2015**, 147, 792-797	5.3	27
92	The effect of rural-to-urban migration on social capital and common mental disorders: PERU MIGRANT study. *Social Psychiatry and Psychiatric Epidemiology*, **2012**, 47, 967-73	4.5	27
91	A randomized controlled trial to evaluate the lung clearance index as an outcome measure for early phase studies in patients with cystic fibrosis. *Respiratory Medicine*, **2016**, 112, 59-64	4.6	25
90	Effectiveness of inhaled tobramycin in eradicating Pseudomonas aeruginosa in children with cystic fibrosis. *Journal of Cystic Fibrosis*, **2014**, 13, 172-8	4.1	24
89	Factors influencing the acquisition of Stenotrophomonas maltophilia infection in cystic fibrosis patients. *Journal of Cystic Fibrosis*, **2013**, 12, 575-83	4.1	24
88	Oral non-steroidal anti-inflammatory drug therapy for lung disease in cystic fibrosis. *Cochrane Database of Systematic Reviews*, **2013**, CD001505		23
87	Comparison of Nutrition and Lung Function Outcomes in Patients with Cystic Fibrosis Living in Canada and the United States. *American Journal of Respiratory and Critical Care Medicine*, **2018**, 197, 768-775	10.2	23
86	Clinical Outcomes Associated with Achromobacter Species Infection in Patients with Cystic Fibrosis. *Annals of the American Thoracic Society*, **2017**, 14, 1412-1418	4.7	22
85	Effect of equipment dead space on multiple breath washout measures. *Respirology*, **2015**, 20, 459-66	3.6	22
84	Oral non-steroidal anti-inflammatory drug therapy for lung disease in cystic fibrosis. *The Cochrane Library*, **2016**, 4, CD001505	5.2	20
83	Oral non-steroidal anti-inflammatory drug therapy for lung disease in cystic fibrosis. *The Cochrane Library*, **2019**, 9, CD001505	5.2	20
82	ERS/ATS technical standard on interpretive strategies for routine lung function tests.. *European Respiratory Journal*, **2021**,	13.6	19
81	Linking COPD epidemiology with pediatric asthma care: Implications for the patient and the physician. *Pediatric Allergy and Immunology*, **2019**, 30, 589-597	4.2	18

80	Global Lung Function Initiative equations improve interpretation of FEV1 decline among patients with cystic fibrosis. *European Respiratory Journal*, **2015**, 46, 262-4	13.6	18
79	Is asthma a vanishing disease? A study to forecast the burden of asthma in 2022. *BMC Public Health*, **2013**, 13, 254	4.1	18
78	Correcting for tissue nitrogen excretion in multiple breath washout measurements. *PLoS ONE*, **2017**, 12, e0185553	3.7	18
77	Integrating the multiple breath washout test into international multicentre trials. *Journal of Cystic Fibrosis*, **2020**, 19, 602-607	4.1	17
76	A guide to interpreting estimated median age of survival in cystic fibrosis patient registry reports. *Journal of Cystic Fibrosis*, **2018**, 17, 213-217	4.1	15
75	Associations between arterial oxygen saturation, body size and limb measurements among high-altitude Andean children. *American Journal of Human Biology*, **2013**, 25, 629-36	2.7	15
74	Do the Global Lung Function Initiative 2012 equations fit my population?. *European Respiratory Journal*, **2016**, 48, 1782-1785	13.6	15
73	Can thrombophilia predict recurrent catheter-related deep vein thrombosis in children?. *Blood*, **2018**, 131, 2712-2719	2.2	14
72	Bronchodilator responsiveness using spirometry in healthy and asthmatic preschool children. *Archives of Disease in Childhood*, **2013**, 98, 112-7	2.2	14
71	Bronchodilator responsiveness in wheezy infants and toddlers is not associated with asthma risk factors. *Pediatric Pulmonology*, **2012**, 47, 421-8	3.5	12
70	Longitudinal study of Stenotrophomonas maltophilia antibody levels and outcomes in cystic fibrosis patients. *Journal of Cystic Fibrosis*, **2017**, 16, 58-63	4.1	11
69	Surname-inferred Andean ancestry is associated with child stature and limb lengths at high altitude in Peru, but not at sea level. *American Journal of Human Biology*, **2015**, 27, 798-806	2.7	11
68	Birth month associations with height, head circumference, and limb lengths among Peruvian children. *American Journal of Physical Anthropology*, **2014**, 154, 115-24	2.5	11
67	Lung clearance index response in patients with CF with class III CFTR mutations. *Thorax*, **2016**, 71, 476-7	7.3	11
66	Projecting the impact of delayed access to elexacaftor/tezacaftor/ivacaftor for people with Cystic Fibrosis. *Journal of Cystic Fibrosis*, **2021**, 20, 243-249	4.1	11
65	Some types of exercise are more effective than others in people with chronic low back pain: a network meta-analysis. *Journal of Physiotherapy*, **2021**, 67, 252-262	2.9	11
64	Performance of oscillometric blood pressure devices in children in resource-poor settings. *European Journal of Cardiovascular Prevention and Rehabilitation*, **2008**, 15, 362-4		10
63	Standardisation of lung function test interpretation: Global Lung Function Initiative. *Lancet Respiratory Medicine,the*, **2018**, 6, 10-12	35.1	10

62	Validation of multiple breath washout devices. *Journal of Cystic Fibrosis*, **2017**, 16, e22-e23	4.1	9
61	Deep-learning algorithm helps to standardise ATS/ERS spirometric acceptability and usability criteria. *European Respiratory Journal*, **2020**, 56,	13.6	9
60	Secular and seasonal trends in obesity in Chilean preschool children, 1996-2004. *Journal of Pediatric Gastroenterology and Nutrition*, **2008**, 47, 339-43	2.8	9
59	Lung Clearance Index to Track Acute Respiratory Events in School-Age Children with Cystic Fibrosis. *American Journal of Respiratory and Critical Care Medicine*, **2021**, 203, 977-986	10.2	9
58	Cluster and CART analyses identify large subgroups of adults with cystic fibrosis at low risk of 10-year death. *European Respiratory Journal*, **2019**, 53,	13.6	8
57	Early detection using qPCR of Pseudomonas aeruginosa infection in children with cystic fibrosis undergoing eradication treatment. *Journal of Cystic Fibrosis*, **2018**, 17, 723-728	4.1	8
56	The association between changes in height and obesity in Chilean preschool children: 1996-2004. *Obesity*, **2007**, 15, 1012-22	8	8
55	Epidemiology of Clonal Pseudomonas aeruginosa Infection in a Canadian Cystic Fibrosis Population. *Annals of the American Thoracic Society*, **2018**, 15, 827-836	4.7	8
54	Oral Azithromycin and Response to Pulmonary Exacerbations Treated with Intravenous Tobramycin in Children with Cystic Fibrosis. *Annals of the American Thoracic Society*, **2019**, 16, 861-867	4.7	7
53	Clinical characteristics of cystic fibrosis patients prior to lung transplantation: An international comparison between Canada and the United States. *Clinical Transplantation*, **2018**, 32, e13188	3.8	7
52	Effect of changes in tidal volume on multiple breath washout outcomes. *PLoS ONE*, **2019**, 14, e0219309	3.7	7
51	A1C as a diagnostic criteria for diabetes in low- and middle-income settings: evidence from Peru. *PLoS ONE*, **2011**, 6, e18069	3.7	7
50	Interpretation of Spirometry in Saskatchewan First Nations Adults. *Annals of the American Thoracic Society*, **2018**, 15, 1237-1239	4.7	7
49	Comparison of facemask and mouthpiece interfaces for multiple breath washout measurements. *Journal of Cystic Fibrosis*, **2018**, 17, 511-517	4.1	6
48	An observational study of the lung clearance index throughout childhood in cystic fibrosis: early years matter. *European Respiratory Journal*, **2020**, 56,	13.6	6
47	A clinical tool to calculate post-transplant survival using pre-transplant clinical characteristics in adults with cystic fibrosis. *Clinical Transplantation*, **2017**, 31, e12950	3.8	5
46	Development and external validation of 1- and 2-year mortality prediction models in cystic fibrosis. *European Respiratory Journal*, **2019**, 54,	13.6	5
45	Pulmonary Function Reference Equations: A Brief History to Explain All the Confusion. *Respiratory Care*, **2020**, 65, 1030-1038	2.1	5

44	The utility of moment ratios and abbreviated endpoints of the multiple breath washout test in preschool children with cystic fibrosis. *Pediatric Pulmonology*, **2020**, 55, 649-653	3.5	5
43	The prevalence of asthma in Canadian children of South Asian descent. *Pediatric Pulmonology*, **2014**, 49, 43-8	3.5	5
42	Reference values for spirometry: the way forward for our patients. *Respirology*, **2011**, 16, 869	3.6	5
41	Ethnically specific norms for ventilatory function. *International Journal of Epidemiology*, **2012**, 41, 1490; author reply 1491-2	7.8	5
40	Resumption of pulmonary function testing during the post-peak phase of the COVID-19 pandemic. *Canadian Journal of Respiratory, Critical Care, and Sleep Medicine*, **2020**, 4, 156-159	0.6	5
39	Special considerations for the treatment of pulmonary exacerbations in children with cystic fibrosis. *Expert Review of Respiratory Medicine*, **2016**, 10, 1221-1228	3.8	5
38	Incidence and risk factors of paediatric cystic fibrosis-related diabetes. *Journal of Cystic Fibrosis*, **2019**, 18, 874-878	4.1	4
37	The burden of asthma among the South Asian and Chinese population residing in Ontario. *Canadian Respiratory Journal*, **2014**, 21, 346-350	2.1	4
36	Spirometric thresholds and biased interpretation of test results. *Thorax*, **2014**, 69, 1146	7.3	4
35	Paediatric reproducibility limits for the forced expiratory volume in 1 s. *Thorax*, **2020**, 75, 891-896	7.3	4
34	Real-world use of ivacaftor in Canada: A retrospective analysis using the Canadian Cystic Fibrosis Registry. *Journal of Cystic Fibrosis*, **2021**, 20, 1040-1045	4.1	4
33	Low lung function in early adulthood: morbidity and death. *Lancet Respiratory Medicine,the*, **2017**, 5, 910-911	35.11	3
32	Determinants of lung disease progression measured by lung clearance index in children with cystic fibrosis. *European Respiratory Journal*, **2021**, 58,	13.6	3
31	Choosing the Better Global Lung Initiative 2012 Equation in South African Population Groups. *American Journal of Respiratory and Critical Care Medicine*, **2020**, 202, 1724-1727	10.2	3
30	Bridging the survival gap in cystic fibrosis: An investigation of lung transplant outcomes in Canada and the United States. *Journal of Heart and Lung Transplantation*, **2021**, 40, 201-209	5.8	3
29	Evaluation of a multiple breath nitrogen washout system in children. *Pediatric Pulmonology*, **2020**, 55, 2108-2114	3.5	2
28	Changes in the parent cystic fibrosis questionnaire-revised (CFQ-R) with respiratory symptoms in preschool children with cystic fibrosis. *Journal of Cystic Fibrosis*, **2020**, 19, 492-498	4.1	2
27	Recommendations for epidemiological studies on COPD. *European Respiratory Journal*, **2012**, 39, 1277-8; author reply 1278-9	13.6	2

26	Exposure to emergency contraception in an undergraduate medical curriculum. *Journal of Obstetrics and Gynaecology Canada*, **2003**, 25, 391-5	1.3	2
25	Development of a symptom menu to facilitate Goal Attainment Scaling in adults with Down syndrome-associated Alzheimer's disease: a qualitative study to identify meaningful symptoms. *Journal of Patient-Reported Outcomes*, **2021**, 5, 5	2.6	2
24	Survival and Lung Transplant Outcomes for Individuals With Advanced Cystic Fibrosis Lung Disease Living in the United States and Canada: An Analysis of National Registries. *Chest*, **2021**, 160, 843-853	5.3	2
23	Validation of short- and long-term demographic forecasts using the Canadian Cystic Fibrosis Registry. *European Respiratory Journal*, **2020**, 55,	13.6	1
22	Disentangling the discordance between epidemiological associations and physiological mechanisms. *Thorax*, **2014**, 69, 869	7.3	1
21	Patterns of Symptom Tracking by Caregivers and Patients With Dementia and Mild Cognitive Impairment: Cross-sectional Study.. *Journal of Medical Internet Research*, **2022**, 24, e29219	7.6	1
20	International consensus on lung function testing during the COVID-19 pandemic and beyond.. *ERJ Open Research*, **2022**, 8,	3.5	1
19	Ethnically Diverse Normative Data for Diffusing Capacity and Lung Volumes: Another Research Priority. *Annals of the American Thoracic Society*, **2020**, 17, 128	4.7	1
18	How Local SARS-CoV-2 Prevalence Shapes Pulmonary Function Testing Laboratory Protocols and Practices During the COVID-19 Pandemic. *Chest*, **2021**, 160, 1241-1244	5.3	1
17	Clinimetric Properties of the Lung Clearance Index in Adults and Children With Cystic Fibrosis. *Chest* , **2016**, 150, 1412-1413	5.3	1
16	Lung compartment analysis assessed from N multiple-breath washout in children with cystic fibrosis. *Pediatric Pulmonology*, **2020**, 55, 1671-1680	3.5	1
15	Unsupervised phenotypic clustering for determining clinical status in children with cystic fibrosis. *European Respiratory Journal*, **2021**, 58,	13.6	1
14	CFTR-function and ventilation inhomogeneity in individuals with cystic fibrosis. *Journal of Cystic Fibrosis*, **2021**, 20, 641-647	4.1	1
13	Impact of cross-sensitivity error correction on representative nitrogen-based multiple breath washout data from clinical trials. *Journal of Cystic Fibrosis*, **2021**,	4.1	1
12	Multiple breath washout: measuring early manifestations of lung pathology.. *Breathe*, **2021**, 17, 210016	1.8	0
11	Author reply: Do Tunisians have a European ancestry?. *European Respiratory Journal*, **2021**, 58,	13.6	0
10	Reply to: When adopting Global Lung Function Initiative reference values, can we also adapt them to a local context as needed?. *European Respiratory Journal*, **2021**, 58,	13.6	0
9	Letter to the Editor, International Journal of COPD [Letter]. *International Journal of COPD*, **2020**, 15, 2307-2308	3	

8	FEV1:FVC Thresholds for Defining Chronic Obstructive Pulmonary Disease. *JAMA - Journal of the American Medical Association*, **2019**, 322, 1609-1610	27.4
7	Markers of early lung disease: the search continues *Pediatric Pulmonology*, **2014**, 49, 1253-4	3.5
6	Resumption of pulmonary function testing during the COVID-19 pandemic. *Canadian Journal of Respiratory, Critical Care, and Sleep Medicine*,1-4	0.6
5	Selection and Appropriate Use of Spirometric Reference Equations for the Pediatric Population. *Respiratory Medicine*, **2015**, 181-193	0.2
4	Transparency and open access in CF research. *Journal of Cystic Fibrosis*, **2020**, 19, e13	4.1
3	Respiratory physiology. *Canadian Journal of Respiratory, Critical Care, and Sleep Medicine*, **2021**, 5, 114-116	0.6
2	Clinical Characteristics Associated With Lung Function Decline in Individuals With Adult-Diagnosed Cystic Fibrosis: Contemporary Analysis of the Canadian CF Registry. *Chest*, **2021**, 160, 65-73	5.3
1	Reference Equations for Pulmonary Function Tests. *Respiratory Medicine*, **2018**, 271-289	0.2